LIFE DOESN’T COME WITH INSTRUCTIONS, BUT LIVING WITH LUNG CANCER NOW HAS THIS SURVIVOR’S GUIDEBOOK.

INSIDE YOU WILL GET THE MOST UP-TO-DATE NAVIGATION TOOLS ON:

- The Diagnosis Process
- Lung Cancer Staging
- Treatment Options
- Clinical Trials
- Living with Lung Cancer
- Financing Your Care
- Hope for Surviving Lung Cancer
- And More

In my house we call it “the lung bible.” It has been an invaluable resource for me, my family and (sadly) a newly diagnosed friend.

—Diane Broderick

I love this handbook! It has great information that I have shared with my family and friends. I often pick it up and re-read. Each time I learn something new, or it triggers something that I need to talk to my doctor(s) about.

—Kimberly Buchmeier

When my husband was first diagnosed with stage IV Adenocarcinoma we were paralyzed with fear and found ourselves starved for good information about the fight we were beginning. The University of Colorado/Anschutz hospital advised us to contact the foundation and we were sent this wonderful handbook that answered all of our questions and enabled us to feel knowledgeable when choosing an oncologist and when facing surgery and treatments. It is an invaluable resource to us.

—Peter & Donna Blum

This is the most comprehensive manual I’ve ever seen written...focused for the lung cancer patient.

—Roy S. Herbst, MD, PhD
Ensign Professor of Medicine (Oncology), Professor of Pharmacology, Chief of Medical Oncology, Associate Director for Translational Research, Director—Thoracic Oncology Research Program, Yale Comprehensive Cancer Center, Yale School of Medicine

WWW.GO2FOUNDATION.ORG

NAVIGATING LUNG CANCER
360° OF HOPE
FIFTH EDITION – FALL 2020

GO2 FOUNDATION FOR LUNG CANCER

BONNIE J. ADDARIO
SURVIVOR SINCE 2003
NAVIGATING LUNG CANCER
360° OF HOPE
5TH EDITION - FALL 2020

GO2 FOUNDATION FOR LUNG CANCER
Empower Everyone. Ignore No One.
In April 2019, the Addario Lung Cancer Foundation (ALCF), formerly the Bonnie J. Addario Lung Cancer Foundation, and the Lung Cancer Alliance in Washington, D.C., announced joining forces to become the GO2 Foundation for Lung Cancer.

“We are thrilled to blend our talent, resources, commitment, and compassion to create this powerhouse foundation,” said GO2 Foundation co-founder and board chair Bonnie J. Addario. “The GO2 Foundation will transform what it means to live with lung cancer and increase the patient survival rate of the world’s number one cancer killer annually. Together we will continue to break barriers and save lives.”

www.GO2foundation.org
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For Additional Copies: To order additional copies go to www.go2foundation.org, call us at 1-800-298-2436 or email info@go2foundation.org.

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ACKNOWLEDGMENTS

The GO2 Foundation for Lung Cancer is proud to publish the 5th Edition of *Navigating Lung Cancer 360° of Hope* with the most recent advancements and updates to help patients in real time. We are eternally grateful to Dr. Shane Dormady for his leadership and expertise editing this guidebook and to the leading lung cancer clinicians who provided insight and direction on the original content.

It is only through the generosity of our supporters that we are able to publish and re-publish this guidebook and offer it free of charge to the lung cancer community, lung cancer patients and their families. For this 5th edition, special thanks goes to Amgen, AstraZeneca, Bristol-Myers Squibb, Daiichi-Sankyo, EMD Serono, Foundation Medicine, Genentech, Janssen Oncology, Jazz Pharmaceuticals, Eli Lilly and Company, Merck, Novartis, Regeneron I Sanofi Genzyme, and Takeda.

If you or your company would like information on supporting future editions, please contact Andrea Beltramo Parks: andrea@go2foundation.org.
Dear Patients,

I have been with the Bonnie J. Addario Lung Cancer Foundation (ALCF), and now the GO2 Foundation for Lung Cancer, from its beginning in 2006 and I am honored to help them lead patients to the best possible outcomes.

If you are holding this handbook or reading it online, you’ve taken the first best step to connecting yourself with the most experienced clinicians worldwide and with a foundation that provides the information you need on your cancer journey.

One of the most important resources that we provide is the Living Room. On the third Tuesday of every month, you can tune into the https://go2foundation.org/resources-and-support/lung-cancer-living-room/ to meet other patients who are either exactly where you are or years ahead and LIVING with lung cancer—patients all over the world. I have spoken many times over the years at the Living Room, am a co-author of this handbook, and am part of a team of leaders in lung cancer who support this foundation and its patients. This foundation is a lifeline for many of my patients and we are all here to help you and your family in every way we can.

It’s important to know that there are many, many kinds of lung cancer. It’s complex. The way we approach lung cancer today, as opposed to 11 years ago, is through personalized medicine and individualized care, because each of you is different.

The pace of advancement and learning new things in lung cancer has never been faster than it is today. More and more, lung cancer is a poster child for all the other cancers where we can take the information from the laboratory and translate it into how we take care of patients. All of us need to have our running shoes on if we’re going to keep up. Today, when something is found in the laboratory it takes less than a year to make its way to the clinic and right to the patient.

From the start, getting information must be part of your lung cancer journey. Most of my patients are informed before they come to see me. Sometimes they pick up bad information, but by and large they know a lot. Having knowledgeable patients make decisions together with their doctors, as a partnership, is very positive approach for both patients and physicians. It is important for you to know that often, there are options, not always right or wrong answers.

You are a patient who now holds in your hands a resource that is your best first step toward understanding and living with lung cancer. In this book and through the lung cancer Living Room you will become educated, informed and most important…you will find answers, directions, options and HOPE.

What we want all our patients to do is live their lives every day to the fullest.

Warmly,

David R. Gandara, MD
Professor of Medicine at the University of California, Davis School of Medicine and Associate Director of Clinical Research, and Director of Thoracic Oncology
UC Davis Comprehensive Cancer Center
29 QUESTIONS TO ASK YOUR ONCOLOGIST AND HEALTHCARE TEAM

1. What type of lung cancer do I have?
2. How does the type of cancer I have affect my treatment options?
3. What stage is my cancer?
4. If my cancer has spread, where else is it in my body?
5. How does the stage affect my treatment options?
6. Do I need more tests before we talk about my treatment plan will be?
7. Has my biopsy tissue been sent out for comprehensive biomarker testing?
8. Which testing was my tissue sent out for and how is that decided?
9. If a biomarkers are found, how will that impact my treatment plan?
10. How can I learn more about my treatment options? ie: Chemo, Surgery, Radiation, immunotherapy, and targeted therapy.
11. If the best treatment for me is not covered by my insurance, what resources are available to help with access/payment?
12. Are there any clinical trials I should consider?
13. What cancer centers or universities are specializing in my type of cancer?
14. Can I get a second opinion at one of these centers and still be treated here locally by you?
15. How long will I be on treatment before I know it is working?
16. How often are my follow up scans?
17. When should we re-biopsy and is liquid biopsy an option for me?
18. What are the side effects of my treatment?
19. How are these side effects managed?
20. I want children in my future, should I consider fertility preservation before starting treatment?
21. Will my treatment affect my daily routine?
22. Can I still work during treatment?
23. Can I travel during treatment?
24. Will I need oxygen for certain activities or traveling by plane?
25. What resources are provided for people with lung cancer?
26. Who is my contact person here for any questions I may have?
27. Are there support groups available for me and my loved ones?
28. Where can I get treatment related to help such as wigs or transportation to appointments?
29. Is there a social worker or patient navigator that can help me?
TO EVERYONE TOUCHED BY LUNG CANCER,

I was diagnosed with lung cancer at the age of 56. I was a wife, a mother, a grandmother, business woman, and one of millions of Americans diagnosed with lung cancer. Faced with a 16% survival rate¹ and following a 14-hour surgery, radiation and chemotherapy treatments that invaded my formerly predictable world, I became a lung cancer survivor with a new purpose in life.

Despite losing three family members to lung cancer, when the doctor said, “you have lung cancer,” I realized I knew very little about the disease. So, I searched for information. I was surprised by how difficult it was to find credible information on lung cancer, treatment options, and how to live with cancer. Everyone kept saying that “cancer is a journey” but no one could provide me with a roadmap. I was lost and I was only just diagnosed.

In 2006, the Addario Lung Cancer Foundation (ALCF) was founded to empower those diagnosed with lung cancer through education and to fund novel research efforts that directly impact patients, today. Our innovative patient education programs are designed and led by lung cancer experts with the goal of supporting you and your family throughout your diagnosis. We support promising research projects through our grants program and the formation of the Addario Lung Cancer Medical Institute (ALCMI). In 2009, we merged with the Lung Cancer Alliance to become the GO2 Foundation for Lung Cancer. Together, we have raised almost $110M to help patients navigate through their lung cancer journeys.

The 5th Edition of this guidebook is the culmination of years of research, conversations with lung cancer experts and patients, my personal experience, and the experience of many patients in both foundations. It is designed to be a resource throughout your cancer journey whether you are newly diagnosed, facing a relapse, or a loved one of someone living with lung cancer. You will find questions to pose to your doctor, detailed explanations of complex treatment options, and access to additional resources in the cancer community.

Lung cancer research is advancing rapidly. In recent years, we have seen new drugs brought to market, many clinical trials started across the nation, the advancement of biomarker testing, and even better side effect management. All of which is needed to improve lung cancer survivability. To this end, we are committed to keeping this guidebook updated with the latest information available.

It is my greatest hope that this guidebook is helpful to you throughout your cancer journey and that we have a positive impact on your life. If I can leave you with one message, it is that you are not alone. Visit our website, join one of our support groups or fundraisers, or simply call us—we are here to help you throughout your journey.

With love,

Bonnie J. Addario, Lung Cancer Survivor
GO2 Foundation for Lung Cancer co-founder and Board Chair
“The most comprehensive and accessible resource available for lung cancer patients.”

—Arlene, Survivor
This guidebook is dedicated to
all lung cancer patients
and their families and friends.

As vital information becomes available,
new print editions of this guidebook will be released
with updated PDFs available on our website.

Check our website (www.go2foundation.org) or
Amazon.com to make sure you have
the most current edition.
# TABLE OF CONTENTS

<table>
<thead>
<tr>
<th>Section</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>DISEASE OVERVIEW</strong></td>
<td>09</td>
</tr>
<tr>
<td><strong>DIAGNOSIS PROCESS</strong></td>
<td>19</td>
</tr>
<tr>
<td>Radiographic Tests</td>
<td>19</td>
</tr>
<tr>
<td>Biopsy Procedures</td>
<td>21</td>
</tr>
<tr>
<td>Biomarker testing</td>
<td>29</td>
</tr>
<tr>
<td>Other Diagnostic Tests</td>
<td>36</td>
</tr>
<tr>
<td>Diagnosis Timeline</td>
<td>38</td>
</tr>
<tr>
<td>Multidisciplinary Healthcare Team</td>
<td>42</td>
</tr>
<tr>
<td><strong>LUNG CANCER STAGING</strong></td>
<td>45</td>
</tr>
<tr>
<td><strong>NON-SMALL CELL LUNG CANCER - TREATMENTS</strong></td>
<td>59</td>
</tr>
<tr>
<td>Overview</td>
<td>61</td>
</tr>
<tr>
<td>Surgery</td>
<td>62</td>
</tr>
<tr>
<td>Chemotherapy/Immunotherapy/Targeted Therapy</td>
<td>69</td>
</tr>
<tr>
<td>Radiation Therapy</td>
<td>76</td>
</tr>
<tr>
<td>Proton Therapy</td>
<td>82</td>
</tr>
<tr>
<td>Pulmonary Therapy</td>
<td>86</td>
</tr>
<tr>
<td>Other Treatment Options</td>
<td>90</td>
</tr>
<tr>
<td>Summary of Treatment Options by Disease Stage</td>
<td>93</td>
</tr>
<tr>
<td><strong>TARGETED THERAPY</strong></td>
<td>100</td>
</tr>
<tr>
<td><strong>IMMUNOTHERAPY</strong></td>
<td>109</td>
</tr>
<tr>
<td>Topic</td>
<td>Page</td>
</tr>
<tr>
<td>------------------------------------------------------------</td>
<td>------</td>
</tr>
<tr>
<td>SMALL CELL LUNG CANCER - TREATMENTS</td>
<td>119</td>
</tr>
<tr>
<td>CLINICAL TRIALS</td>
<td>126</td>
</tr>
<tr>
<td>LIVING WITH LUNG CANCER</td>
<td>139</td>
</tr>
<tr>
<td>Transitional Care Planning</td>
<td>141</td>
</tr>
<tr>
<td>Nutrition</td>
<td>148</td>
</tr>
<tr>
<td>Traveling</td>
<td>149</td>
</tr>
<tr>
<td>Alternative / Complementary Care</td>
<td>150</td>
</tr>
<tr>
<td>FINANCING YOUR CANCER CARE</td>
<td>151</td>
</tr>
<tr>
<td>END-OF-LIFE PLANNING</td>
<td>159</td>
</tr>
<tr>
<td>OUR GENEROUS SUPPORTERS</td>
<td>169</td>
</tr>
<tr>
<td>GLOSSARY</td>
<td>173</td>
</tr>
<tr>
<td>REFERENCES</td>
<td>179</td>
</tr>
<tr>
<td>ADDITIONAL REFERENCES</td>
<td>184</td>
</tr>
<tr>
<td>INDEX</td>
<td>187</td>
</tr>
<tr>
<td>GO2 TEAM</td>
<td>192</td>
</tr>
<tr>
<td>ADDARIO CENTERS OF EXCELLENCE</td>
<td>194</td>
</tr>
</tbody>
</table>
YOUR DATA COULD OPEN THE DOOR TO A CURE
We believe patients hold the keys to discovery.

MORE DATA. MORE SURVIVORS.

Together patients and caregivers can build a knowledge base to enable researchers and clinicians to better identify and understand this disease. Knowledge will power breakthroughs in diagnosis and treatment and drive improvements in patient care and outcomes. Thank you for joining us in the fight against lung cancer.

LUNGCANCERREGISTRY.ORG
DISEASE OVERVIEW

We believe patients hold the keys to discovery. Together patients and caregivers can build a knowledge base to enable researchers and clinicians to better identify and understand this disease. Knowledge will power breakthroughs in diagnosis and treatment and drive improvements in patient care and outcomes. Thank you for joining us in the fight against lung cancer.

LUNGCANCER.ORG

MORE DATA. MORE SURVIVORS.
My cancer center gave me information regarding the Patient Education Handbook as soon as I was diagnosed with lung cancer. As the diagnosis process continued to find out type, stage and genomic mutation, I would reference the handbook throughout my appointments and procedures.

I got hope after meeting several stage IV lung cancer survivors and speaking with Bonnie at the World Conference on Lung Cancer Pancake Walk. The event just so happened to be a drive away in Denver, just weeks after my lung cancer diagnosis. I framed photos from that day. When I look at them it brings back that initial feeling of hope.

—Lisa Moran, survivor
DISEASE OVERVIEW

After you receive a diagnosis of lung cancer, it is normal to feel scared and alone. We want to help you understand your disease, what you can do to help take care of yourself, and what we can do to help you. This guidebook will help you know what to expect during this process. We know that having information when you need it is critical; however, this guidebook will not replace your interactions with your healthcare team.

What is lung cancer?
In a healthy body, normal cells grow, mature, and eventually die and are replaced by other healthy cells. Occasionally, abnormal cells in the body begin to develop and grow. If your body recognizes these cells as “abnormal,” the body’s defense mechanisms may kick into action to destroy the abnormal cells much the same as when bacteria are destroyed by white blood cells. In the case of cancer, your body sees these abnormal cells as part of your body, so it does not attack them and as a result, the cells begin to grow out of control.

DNA, which stands for deoxyribonucleic acid, is the molecule in every cell that controls how that cell grows and functions. In a cancer cell, the DNA is damaged and is reproduced in other abnormal cells. In most types of cancer, these abnormal cells begin to stick together and form tumors. Tumors are usually classified as benign (non-cancerous) or malignant (cancerous).

When we talk about lung cancer, we are talking about this out-of-control, malignant growth that starts in the lung tissue. As the cancer cells grow and multiply, the normal cells in the lung are replaced by the malignant cells.
Cancer cells can develop in any part of the body and then spread to other parts of the body through the blood and lymph systems. When this happens, the cancer is said to have metastasized and the resulting tumors are called metastatic tumors or metastases. Lung cancer that starts in the lung is called primary lung cancer; if the cancer started in another part of the body and metastasizes to the lung, it is called secondary lung cancer.

The lymphatic system (or lymph system for short) is a system much like the blood system in the body. The lymph system is responsible for carrying nutrients to the cells and waste away from the cells. The lymph nodes are special parts of the lymph system that are responsible for filtering the wastes out of the liquid that passes through. When waste collects in the lymph node, it swells and becomes painful. These lymph nodes are in many different places in your body. That is why your doctor and nurses will feel around your neck, in your armpits, in your groin and other areas. They are looking for these swollen glands.

What causes lung cancer?
Primary lung cancer is caused by the out-of-control growth of cells that do not die as in the normal cell pattern. The cause of lung cancer may not always be known.

Carcinogens are those things that can cause cancer. Normal cells in the lung can be affected by carcinogens in the environment, genetic factors, or a combination of those factors. Exposure to carcinogens may form molecules in the body called free radicals which damage cells and alter the DNA of the cell. This damage may cause cancer.

Environmental factors include things such as smoking, secondhand smoke, radon gas, air pollutants, asbestos, heavy metals, and chronic dust exposure. Genetic factors may include an inherited (passed from parent to child) or a genomic mutation. A genomic
mutation is damage to the gene that increases the chances of developing a particular kind of cancer.

What are the signs and symptoms of lung cancer?
It is important to recognize the signs and symptoms of lung cancer in order to ensure a reliable diagnosis. A sign is something that can be seen by someone else; for example, a rash is a clinical sign. A symptom is something that cannot be seen by someone else but must be described by the person; for example, a headache is a symptom. Early in the disease, lung cancer may not produce any signs or symptoms. However, as the disease progresses, certain key signs and symptoms may develop.
Possible signs and symptoms of lung cancer may include:

- A cough that does not seem to be related to a specific illness, a change in a chronic cough, or a cough that does not go away
- Shortness of breath particularly if it is not related to physical activity or if the shortness of breath seems worse than it should be for the amount of activity (“I walk to the corner and have to sit down and catch my breath before I can walk back”)
- New wheezing that is unrelated to a specific illness (“When I breathe, it sounds like I’m whistling”)
- Coughing up blood (hemoptysis)
- Chest pain
- A hoarse voice or a marked change in voice
- Chronic fatigue (“I just can’t seem to get enough rest; I’m always tired”)
- Weight loss with no known cause
- Headaches
- Painful lumps in the neck, armpits, or groin caused by inflammation of the lymph glands as the cancer spreads through the lymphatic system
All of these signs and symptoms can be caused by other diseases and conditions and may not indicate a diagnosis of lung cancer. However, when several of these symptoms exist, particularly if they do not get better in a short period of time, you should visit your healthcare provider for diagnosis and treatment.

**What should I ask my healthcare provider?**

We understand that this is a scary time for you and your family and we want you to know that we are here to help.

Before your first appointment with your doctor, and at every appointment after that, be prepared with a written list of questions. Between appointments, keep a pad of paper and pencil with you so that you are always ready to jot down a question that comes to mind. At every appointment, ask all of your questions and ask for clarification when the healthcare provider gives an answer you don’t understand. Write down the answer to each question. Read the answers back to your provider to make sure you have recorded the information correctly. If possible, take a friend or spouse with you to each appointment. Two sets of ears and two brains are more likely to hear and remember all the information. If your healthcare provider agrees, it might be helpful to take a tape recorder to the appointment and record the discussion.

**Throughout this guidebook, you will find suggested questions or points to discuss with your healthcare team in boxes like this one.**

**Are there different types of lung cancer?**

Five types of lung cancer have been identified: Non-small cell lung carcinoma (NSCLC), small cell lung carcinoma (SCLC), mesothelioma, carcinoid, and sarcoma. NSCLC and SCLC represent about 96% of all lung cancers. These two types of lung cancer are identified by the size of abnormal cells and the way the cancer spreads in the body.
Treatments for these two types of cancer are different so it is critical that the type of cancer is correctly identified.

Non-Small Cell Lung Cancer (NSCLC)

NSCLC represents about 85 to 90% of all lung cancers and can be further described as:

- Adenocarcinoma
- Epidermoid or squamous cell carcinoma
- Pancoast or pulmonary sulcus tumor
- Large cell undifferentiated carcinoma

**Adenocarcinoma**: Adenocarcinoma is the most common type of lung cancer accounting for 40% of all cases. Typically, this type of lung cancer starts growing in tissue on the outside surface of the lung. The tumor in a lung adenocarcinoma is made up of cells that tend to line up in small masses. These tumors vary in size and how fast they grow.

**Pancoast Tumor**: A Pancoast tumor is sometimes called a pulmonary or superior sulcus tumor. Typically, this type of lung cancer is found at the top of the lung and has a tendency to spread to ribs and bones of the spine. Since the Pancoast tumor usually grows on the top of the lung, it is very close to nerves and the spine; these facts make surgery on these tumors very difficult. Pancoast tumors account for fewer than 5% of all primary lung cancers.
Small Cell Lung Cancer (SCLC)
SCLC represents about 10 to 15% of all lung cancers. These lung cancers typically grow rapidly and are aggressive forms of lung cancer. SCLC can be further defined as small cell carcinoma (oat cell cancer) or combined small cell carcinoma. In addition, SCLC is usually described as limited or extensive.

SCLC tumors may also cause paraneoplastic syndromes. A paraneoplastic syndrome is a collection of symptoms that develops as a result of cancer but is not directly related to the cancer cells. Usually, these symptoms are caused when the SCLC tumor produces hormones or other specialized proteins that cause an inflammatory response in the body. The body's immune system reacts to these substances and can begin attacking normal nervous system cells causing problems in the nervous system.

Lung Mesothelioma
Malignant lung mesothelioma is diagnosed in 2,000 to 3,000 people in the United States each year. The mesothelium is the lining that covers the body's internal organs and cavities. This rare form of cancer is most commonly found in the pleura, or outer lining, of the lungs and internal lining of the chest wall thus the name “lung mesothelioma.” Pleural mesothelioma accounts for approximately 70% of all mesothelioma cases. For more information on this disease, visit the National Cancer Institute at http://www.cancer.gov/cancertopics/pdq/treatment/malignantmesothelioma/patient.
Carcinoid
Carcinoid tumors in the lungs are extremely rare representing about 1% of all lung cancer cases. Carcinoid tumors grow slowly in the lining of the lungs. Because the carcinoid tumors are composed of endocrine cells and secrete hormones, they are often considered endocrine tumors. These very slow-growing carcinoid tumors can often be treated with radiation, surgery, chemotherapy, and immunotherapy. People with certain genetic disorders (multiple endocrine neoplasia type 1 and neurofibromatosis type 1) may be at a higher risk of developing carcinoid tumors.

Sarcoma
Sarcoma is another extremely rare cancer that is seen in about 1% of all lung cancers. Typically, a sarcoma is found in bone or other soft tissues. The sarcoma is different from other tumors because of the cells in which it grows. For more information on sarcoma, visit the Sarcoma Foundation of America at http://www.curesarcoma.org/index.php/patient_resources/.
THE LIVING ROOM

A LUNG CANCER COMMUNITY OF COURAGE

BONNIE J. ADDARIO
WITH JON LAND

AVAILABLE ON AMAZON.COM
DIAGNOSIS PROCESS
This handbook is an invaluable guide to the world of lung cancer. It creates a sense of knowledge and understanding when it comes to the brash new reality you and loved ones are now living. It’s a comprehensive guide that gives you the security of knowledge in an overwhelming situation—a great place to begin your path to recovery and wellness. Find easy to comprehend definitions that lay a foundation of understanding to help construct a plan that is best-suited for you in this journey.

The foundation is a warm embrace when you feel like you’re in free fall. With the compassion of a saint, the tenacity of a bulldog, and the strength of an army, every single staff member is sincerely here for you and wants the very best for you and your loved ones. I’m so blessed to have the resources of the GO2 Foundation for Lung Cancer. They are a wonderful support system and definitely someone you want on your team!

—Bekah, survivor
My doctor found a spot on my lung –
What happens next?
First, take a deep breath and know that this spot may
not be lung cancer; it may be something such as a
benign (non-cancerous) nodule, infection, or many
other things. The next steps in the process will help
your doctor determine, or diagnose, the problem.

Your doctor will talk to you about what tests will
be done to determine if the spot is cancer. Usually,
your plan will include some sort of radiographic,
or x-ray, tests. Your doctor may also want to do a
biopsy of the spot. A biopsy involves taking a
tissue sample from the area on or around the
lung and examining it under a microscope.

**RADIOGRAPHIC TESTS**

The radiographic, or x-ray, tests described here are not painful. The most painful thing
you will experience during these tests is a needle stick for those tests that require an
injection of a radioactive liquid.

**Computed Tomography (CAT or CT Scan):** A CAT or CT scan is done using a
special x-ray machine that gives a more detailed picture inside your body than
a normal chest x-ray. CT scans can find very small tumors in the lung and can
help to determine if the cancer has spread to the lymph nodes around
the lungs. This scan will help your doctor know what size the tumor
is and the exact location of the tumor.

Questions to ask your physician
during the diagnosis process:
• What tests will I need to
determine if I have lung cancer?
• Should I watch and wait?
• If I decide to watch and wait, how
  long before you check the spot or
  nodule again?
• What is active surveillance?
• What are the changes the spot or
  nodule is benign (non-cancerous)?
• Do I need x-ray tests?
• Do I need a biopsy?
• How long will it take to get my
  biopsy results?
• What will each test show?
Positron Emission Tomography Scan (PET Scan): A PET scan is done by another very specialized machine that rotates around your body giving a three dimensional view of your body and allowing your doctors to see the differences between malignant and benign areas. Before the PET scan, a member of your healthcare team will inject a small amount of sugar water with radioactive isotopes into a vein. A radioactive isotope is an atom that emits radiation that can be “seen” by the radiological equipment. As the PET scanner rotates, it shows a picture of where the isotopes are deposited in the cells. Malignant tumors show up brighter in the scan because the cancer cells are more active and are using more of the sugar water mixture than normal cells do.

Magnetic Resonance Imaging (MRI): The MRI uses huge magnets, magnetic fields, and radio waves to create clear images of many different areas of the body such as the brain, muscles, joints, and blood vessels. Before this test, the x-ray technician will ask you to remove all metal (rings, glasses, bracelets, etc.) that may be attracted to the magnets.

Bone Scan: A bone scan is a very specific test that may be used to determine if cancer has spread to the bones. Again, with this test, the radiology technician will inject a small amount of sugar water with a radioactive isotope solution into your vein. This fluid begins to accumulate in areas of abnormal bone growth where a radiation scanner can measure the radioactivity levels and record them on x-ray film providing a clear picture of areas that might have cancerous tumors.
BIOPSY PROCEDURES

You will want to discuss the biopsy procedures listed here with your physician to understand which procedures are necessary in your unique situation. We have highlighted several points to discuss with your doctor to help you get the information you need to make informed decisions about your care.

**Fine Needle Aspiration (FNA)** is usually performed by an interventional radiologist (a doctor who specializes in doing procedures using radiology) or pulmonologist (a doctor who specializes in lung disease). In this procedure, the doctor will insert a needle through the chest wall into the tumor. Cells from the tumor are pulled into the syringe and are then examined by a pathologist under a microscope. The pathologist is the doctor on your team who specializes in diagnosing disease by examining tissue and body fluids. The fine needle aspiration procedure is done with the help of a CT scanner, fluoroscopy (live x-ray images done using a fluoroscope) or MRI to guide the needle to the exact location of the tumor. Before this procedure, the biopsy site (area that will be stuck by the needle) is numbed so the procedure should not hurt.

It is important to obtain enough tumor tissue for diagnosis and biomarker testing. Ask your doctor if an FNA biopsy will collect enough tissue for both diagnosis and biomarker testing.

**Core Needle Biopsy** is usually performed by an interventional radiologist or pulmonologist. This procedure is similar to the FNA, but the doctor can usually get a larger piece of tissue with core needle biopsy. Using this method, the pathologist will have enough tissue to determine the type of lung cancer and for biomarker testing. The core needle biopsy is usually done with the aid of...
some sort of x-ray equipment to guide the needle to the exact location of the tumor. Again, before this procedure, the biopsy site (area that will be stuck by the needle) is numbed so the procedure should cause minimal discomfort.

**Traditional Bronchoscopy** is typically performed by a pulmonologist. In this procedure, a flexible tube called a bronchoscope is passed down the nose or mouth into the trachea, bronchi and larger tubes in the lungs. A bronchoscopy allows the physician to actually see the central regions of the lungs and take a tissue sample for the pathologist to examine. Usually done under local anesthesia with sedation, your healthcare team may do this procedure as an outpatient, so you will not have to spend the night in the hospital. A quick procedure, a bronchoscopy usually takes less than an hour. You may spend several hours “recovering” from the procedure. During this time, the healthcare team will make sure you are awake and not having any problems before you are sent home with your family.

**Robotic Bronchoscopy** is a robotically-assisted procedure performed by a pulmonologist or thoracic surgeon to biopsy suspicious nodules in the lung. The procedure is as safe as a traditional bronchoscopy and provides the physician with advanced control that comes from robotic technology.

Robotic bronchoscopy may be used to reach small tumors that are in the periphery of the lung. As the physician moves the scope deeper into the lung, the tubes get more windy and smaller. It can be very challenging for physicians to reach these small tumors using traditional, manual bronchoscopes.

During the procedure, the physician guides the robotic bronchoscope through the mouth into the lungs using a controller that looks similar to a video game controller.
This controller allows the physician to make very precise movements required to maneuver through the small tubes in the lung.

The robotic bronchoscopy combines traditional camera views into the lung with computer-assisted navigation based on 3-D models of the patient’s own lung anatomy. Now, physicians have the benefit of seeing inside the lung while getting additional navigational guidance to reach the precise location of the tumor for biopsy.

**Electromagnetic Navigation Bronchoscopy™ Procedure:** Also known as ENB™ procedures, Electromagnetic Navigation Bronchoscopy™ procedures are performed by a pulmonologist or thoracic surgeon. ENB™ procedures provide a minimally invasive approach to accessing difficult-to-reach areas of the lung aiding in the diagnosis of lung disease. Using tiny instruments, your physician will take a sample of the nodule for testing. In some cases, small markers may be placed near the lung nodule to help guide a physician delivering follow-up treatment or therapy.
Thoracentesis is performed by an interventional radiologist or pulmonologist. If any of the x-ray procedures show that fluid is present in the chest cavity outside of the lungs, your doctor may insert a thin needle into the chest between the ribs to pull out a sample of the fluid. If you are having trouble breathing because of the amount of fluid in the chest, the doctor may remove more of the fluid to help your breathing. The pathologist will examine the fluid that is removed from the chest.

Lymph Node Biopsy is performed by an interventional radiologist or pulmonologist. A lymph node biopsy is done after the initial diagnosis of lung cancer to see if cancer has spread from the lung to the lymph nodes. Lymph node biopsy is an important step in determining the stage of the lung cancer. This procedure can be done in one of three ways: by inserting a needle directly into the lymph node, by using a needle during a bronchoscopy or mediastinoscopy, or by complete removal of the lymph node with surgery. Any of these methods will typically be done on an outpatient basis with local anesthesia. The type of anesthesia and recovery will differ by type of procedure.

Mediastinoscopy is performed by a thoracic or general surgeon. For this procedure, you will go to the operating room where you will be given general anesthesia so that you are asleep during the procedure. Your surgeon will place a tube called a mediastinoscope through a small incision in your neck. With the bronchoscopy, the surgeon is able to see inside your lungs; during the mediastinoscopy, the surgeon will be examining the mediastinum (the area between and in front of your lungs). During this procedure, the surgeon can biopsy any lymph nodes or masses seen outside of the lungs. The mediastinoscopy may be done at the same time as a bronchoscopy; if so, both procedures will take less than two hours. If no other procedures are done, a mediastinoscopy usually takes about 45 minutes and may be done on an outpatient basis.
Video-Assisted Thoracoscopic Surgery (VATS) is performed by a thoracic surgeon. For the VATS procedure, you will be taken to the operating room and will be given general anesthesia so that you are asleep during the procedure. A thoracoscope is placed into the chest through an incision in the chest wall. A thoracoscope is a camera on the end of flexible tubing that allows your doctor to look into your chest. Your surgeon can then look at the surface of the lung and the chest wall. Your doctor may use the VATS technique to remove some lung cancer tumors. This procedure is less invasive than a thoracotomy and has a shorter recovery time.

Thoracotomy is performed by a thoracic surgeon. A thoracotomy is something like a VATS procedure; however, instead of inserting a scope through a small incision, your surgeon will make a larger incision into the chest in order to see the lung directly. In a thoracotomy, a tumor, lung tissue or lymph nodes may be removed. This procedure is done under general anesthesia and you will probably be in the hospital for 3 to 5 days. Your surgeon may elect to do the VATS procedure instead of a thoracotomy.

What happens to my biopsy and what does it tell my physician?
When your doctor completes the biopsy, he or she will send the biopsy tissue to the lab where the pathologist takes a very small slice of the tissue to look at under a microscope. Each type of cell looks very different under a microscope so the pathologist will be able to tell the type of tumor you have and whether it is benign or malignant (cancerous or not).

If the piece of biopsy tissue is large enough, the pathologist may also be able to “grade” the tumor. When the pathologist grades the tumor, he is comparing the tumor cells to normal cells. The tumor grade describes how much the cells from the biopsy
tissue resemble normal lung cells. Tumor grades are different for different kinds of cancer, but typically, a lower grade is better. Based on what he sees under the microscope, the pathologist will also determine how fast the tumor may grow and spread.

When your pathologist grades your tumor, he will also send your tissue for biomarker testing. Because we know that different types of lung cancer are caused by new or acquired mutations that have different genomic forms, we can use biomarker testing to identify the specific genomic makeup of the tumor. Knowing the specific genetic form of your tumor may help your team to create a treatment plan that is specific to your tumor.

On tissue biopsies that are larger, the pathologist will also look at the lung tissue around the tumor to see if there are cancer cells outside of the tumor and in the lung tissue. The pathologist prepares a pathology report that includes all of the findings and sends that report to the rest of your healthcare team.

Your healthcare team will use the tumor grade and other findings to begin to develop a treatment plan designed specifically for you. Your doctor will help you understand exactly what the grade of your tumor means and how the grade will help guide your treatment.
Why does the doctor need to repeat the biopsy?

When you are initially diagnosed the pathologist’s first priority is to use your tissue biopsy to determine which specific type of lung cancer you have. This often requires looking at multiple slices of the tumor tissue and applying special stains to the tissue to help with the right diagnosis. The result is often that the tissue is used up and there is nothing left for genomic testing. This happens with 30% to 50% of lung cancer biopsies, especially when they are smaller needle biopsies (1–3).

The more common reason for repeating genomic testing is because the initial tissue sample was only partly genotyped or undergenotyped. Sadly, this happens the majority of the time and typically occurs when you are only tested for EGFR and ALK, and nothing is found. This could be because tests less sensitive than NGS were used for EGFR or ALK which may miss a significant number of mutations (4–8). Another common reason for undergenotyping is because local non-NGS testing for EGFR and ALK depletes or exhausts your tissue sample and so they cannot test for additional genes even though they may be targetable.

Cancer cells evolve over time, especially when the cancer has been treated. Your cancer one year from diagnosis for example, isn’t necessarily the same cancer originally seen under the microscope when you were first diagnosed. The only way to know what the cancer has potentially evolved into is to re-biopsy the tissue and examine it for changes.

Fortunately, there are “liquid biopsy tests” available that use various technologies so that you can get complete genomic testing with simple blood test. The GeneStrat® blood-based genomic test from Biodesix provides actionable mutations that can direct the use of targeted therapy in 72 hours. Another method, used by the Guardant360 test, is likely to find a mutation that drives your cancer in lung adenocarcinoma in all but about ¼ of patients. When you are initially diagnosed there is rarely more than one driver mutation found. Some are targetable and many are not, but if any driving mutation is found then there is no need to repeat an invasive biopsy.

If your cancer has been treated with a targeted therapy but has progressed, a liquid biopsy may be used initially instead of repeating an invasive biopsy. The results typically come back in 3-14 days, depending on the test utilized, and if the blood test does not detect the new mutations driving your cancer, then your doctor may recommend a repeat invasive tissue biopsy.
We’re at a pivotal point in the fight against cancer. Researchers have a greater understanding of its molecular causes than ever before. Dr. Tony Mok so eloquently stated at a recent World Conference on Lung Cancer, it’s time for the research establishment, pharma and academia, to remember the reason for the research is the patient.

Patients are at the center of the fight.

— Bonnie J. Addario
BIOMARKER TESTING

One of the goals of your treatment is to determine whether your tumor will respond to a particular drug or treatment. Historically, lung cancer was treated based only on type and stage with cytotoxic chemotherapy. Cytotoxic means cell-killing and typically this chemotherapy kills rapidly dividing cells. Chemotherapy may be very effective in some people, but because the cancer cells are not the only rapidly dividing cells, there are often side effects when blood cell producing cells or hair cells (which also rapidly divide) are killed. We are learning that different types of lung cancer have different genomic forms that we can identify through biomarker testing. Identifying the specific genomic makeup of the tumor may allow your team to tailor your treatment plan to the specific tumor.

What is biomarker testing?
Biomarker testing, also called assays or profiles, can help your treatment team identify specific biomarkers that are in your tumor. Molecules contain biomarkers that are specific genes or proteins that are mutated within the cancer cells and determine how your cancer will respond to treatment. A biomarker (or biological marker) is a very distinctive substance that indicates if a particular subtype disease is present. Biomarkers can be proteins, genes, or other biological substances. Generally, a tissue biopsy is performed to obtain an ideal amount of your tumor tissue for testing. When the tumor is biopsied, your oncologist and pathologist should look for certain biomarkers that have been associated with lung cancer. A “liquid biopsy,” or blood-based test, can also be used to conduct biomarker testing non-invasively.

The results of these tests determine your distinct ‘molecular fingerprint.’ Just as no two fingerprints are alike, neither are molecular fingerprints. The information contained in your unique molecular fingerprint gives

The role of biomarker testing in lung cancer has grown in the past year. Ask your doctor if biomarker testing is available to you. If not, contact us at 1-800-298-2436 to learn how to have your lung cancer tested.
your oncologist or treating physician insights into how to personalize your lung cancer treatment plan.

Each time you have a biopsy, your doctor may order biomarker testing on the tissue.

When the pathologist identifies specific biomarkers, this may indicate a genetic mutation and/or fusion. A genetic mutation is anything that changes the structure of a gene. A genetic fusion is a gene that is formed when the genetic material from two previously separate genes are mixed or fused together to form a new cancer-causing gene. We are learning that there are certain genes that can work to produce or suppress lung cancers. When there are changes in the structure (or mutations) of these genes, lung cancer can be the result.

What specific information is obtained from my biomarker testing and how does it determine my personalized treatment?

Different molecular tests are done depending on the laboratory where the tissue is sent. Many major cancer centers at teaching hospitals can perform molecular testing. However, most labs will test the DNA from your tumor in tissue or blood for only EGFR mutations and ALK fusions, and omit the other five genomic alterations which can be treated with targeted therapies instead of chemotherapy. The seven genes recommended for testing in the national guidelines are found over one third of patients with non-squamous non-small cell lung cancer. Many patients tissue biopsy is used up or exhausted with one-at-a time testing for EGFR and ALK, so that the other five genes (BRAF, MET, ERBB2 (HER2), RET, ROS1, and KRAS) go untested. Yet 20% or one in five patients with lung adenocarcinoma have one of these other alterations.
This is one reason guidelines recommend that everyone insist on comprehensive genotyping of their tumor with tissue or blood, as with NGS of tissue or blood all seven genomic targets can be tested. More and more drug therapies that are targeted or matched to cancers with these and other specific mutations are being developed all the time so we expect new clinical trials and treatment options for lung cancer patients who get NGS testing of their tissue or blood.

- **EGFR**: The *EGFR* gene produces a protein called epidermal growth factor receptor. In 10% of patients with non-small cell lung cancer (NSCLC), the *EGFR* gene is mutated. Nearly 50% of the lung cancers resulting from *EGFR* mutation happen in people who have never smoked.
• **KRAS:** The KRAS gene is mutated in about 25% of people with NSCLC. There are three drugs that are commonly used to treat lung cancer – gefitinib (Iressa™) and erlotinib (Tarceva®) and afatinib (Gilotrif®). The KRAS gene is mutated in about 25% of people with NSCLC. Ongoing clinical trials are showing promise for lung cancers with certain KRAS mutations. Other options for patients with KRAS mutations may be chemotherapy or immunotherapy.

• **ALK fusions:** In addition to alterations in the **EGFR** or **KRAS** genes, another abnormality called an ALK fusion may be the genomic alteration driving your lung cancer. This mutation occurs when two genes (such as **EML4** and **ALK**) become fused into a form that increases the activity of the **ALK** oncogene. The **EML4-ALK** and other ALK fusions are found in nearly 5% of patients with non-small cell lung tumors and is highly responsive to a targeted therapy. It is also present in about 10 to 15% of people with non-small cell lung cancer who have never smoked.

• **BRAF:** The fourth identified mutation, **BRAF**, occurs in about 3% of people with lung cancer. Like **KRAS**, this mutation seems to happen most often in patients who are either current smokers or who smoked in the past.

The **BRAF** mutation produces a protein that transmits signals within a cell to its interior. In a cancer tumor, this signal can cause cells to divide and cancer to grow. About half of the mutations in **BRAF** are now targetable.

• **MET:** There are several different alterations in the **MET** gene which are responsive to matched therapy. These include point mutations, a deletion of a part of the gene known as exon 14, or amplification where the tumor is driven by extra copies of the **MET** gene even though it is not mutated. All three of these types of alterations are not routinely measured unless comprehensive genomic testing with tissue or blood with NGS is used. This is another reason we recommend NGS testing, because all three of these alterations in **MET** may respond to targeted therapy. About 6% of patients with lung adenocarcinoma have mutations in the **MET** gene, and it occurs with higher frequencies in current or past smokers than in non-smokers.
• **ROS1**: ROS1 fusions, like ALK fusions, are formed when the ROS1 gene and a second gene break apart and fuse together with the result of activating the ROS1 oncogene. These targetable alterations occur in about 2% of lung adenocarcinoma and are responsive to targeted therapies.

• **RET**: RET fusions also occur in about 1% of lung adenocarcinomas. Recently these have been shown to be targetable with available matched therapy drugs and exciting new RET inhibitors now have approved targeted therapies. See the chapter on Targeted Therapies for more information on the role of biomarker testing and treatment decisions.

**Blood-based proteomic testing**

If genetic sequencing is similar to the “script” of biology, then proteomics (the study of proteins) is the live video capturing biology in action. One of the advantages of proteomics is that cancer patient subgroups have been identified based on specific protein signatures expressed by the tumor cells or a patient’s immune response to the tumor. These protein signatures can be detected in the patient’s blood (no tissue biopsy needed) and can be used to help inform the patient’s treatment plan.
Next-Generation Sequencing
Genomic testing or profiling identifies the underlying DNA alterations that are driving the tumor’s growth. This information may help physicians understand which targeted treatment options are available for a patient based on their tumor’s unique genomic profile. A new technology called next-generation sequencing (NGS) is often referenced in relation to molecular or genomic testing for lung cancer. NGS is a tool for sequencing large amounts of DNA accurately in a short period of time, but it can be applied in many different ways.

Standard genomic testing examines only one or a limited set of cancer-related genes and does not provide a complete picture. Some tests may use NGS to look for a few types of alterations in predetermined “hotspot” regions within genes where alterations more commonly occur. However, tumors often contain multiple alterations that would be missed by these more narrowly focused genomic tests, limiting potential treatment options.

In fact, national guidelines for non-small cell lung cancer recommend comprehensive NGS testing so that no potentially treatable genomic alterations are overlooked. For example, hotspot testing for the most common mutations in the EGFR gene would still miss 1/6 of the EGFR mutations that occur.²

A comprehensive genomic profiling test uses NGS to look at all of the cancer-related genes in a single sample of tumor tissue and detects all types of alterations. This approach provides the information your physician needs in one single test to help guide a tailored treatment approach using targeted therapies. You and your doctor can use the results from a comprehensive genomic profile to discuss possible treatment options, including FDA-approved targeted therapies or novel targeted treatments under development in clinical trials.
If you would like comprehensive genomic profiling performed on your lung cancer tumor, you can find more information and a discussion guide for you and your physician at www.dontguesstestlungcancer.com or call 1-800-298-2436.

How is my tumor tissue for biomarker testing obtained?
One of your doctors will do a biopsy to get a tissue sample from your tumor. Cancer diagnosis always begins with a tissue biopsy so that a pathologist can determine which type of cancer you have. Biopsies can be performed in a number of ways. It is important to get a big enough piece of tumor tissue to do biomarker testing. Fine needle aspiration (FNA) biopsy may not provide enough tissue for biomarker testing, so your oncologist may recommend one of the following methods for biopsy.

- Core needle biopsy performed by an interventional radiologist
- Bronchoscopy performed by a pulmonologist
- Lymph node biopsy performed by an interventional radiologist or pulmonologist
- Mediastinoscopy performed by a thoracic or general surgeon
- Computed tomography (CT), fluoroscopy, ultrasound, or MRI-guided core needle biopsy performed by an interventional radiologist or pulmonologist
- Video-assisted thoracoscopic surgery performed by a thoracic surgeon
- Electromagnetic Navigation Bronchoscopy™ procedure performed by a pulmonologist or thoracic surgeon

What if the biopsy does not produce enough tissue to test for all the known genetic mutations?
There are seven genes whose alterations cause non-small cell lung cancer (NSCLC), that currently have an FDA approved drug associated with them. Patients with these alterations comprise almost 85% of lung cancers (small cell lung cancer incidence has gradually declined to < 15 of lung cancers). (ref Govindan R: Changing Epidemiology of Small-Cell Lung Cancer in the United States Over the Last 30 Years: Analysis of the Surveillance, Epidemiologic, and End Results Database. J Clin Oncol 24:4539–4544, 2006 ) Recent National Comprehensive Cancer Network guidelines (ref www.nccn.org) strongly recommend broad molecular profiling to cover all seven genomic targets in NSCLC: (EGFR, ALK, ROS1, BRAF, MET, RET and HER2) as these have matched therapy
treatments which produce responses 2-3 times higher than chemotherapy alone. If the
doctor does not get enough tissue to test for all genetic markers, NCCN guidelines
recommend blood (plasma) testing (a “liquid biopsy”) when a repeat invasive tissue
biopsy is infeasible (please note these blood biopsies although on the brink of being
FDA approved, they are currently not). (ref NCCN).

Where will my tissue be tested?
A Clinical Laboratory Improvement Amendments certified laboratory, most commonly
referred to as a CLIA certified laboratory, will usually test the tumor tissue. If your
hospital does not have a lab that offers the biomarker testing, your oncologist will ask
that the tissue be sent to another lab.

How long does it take to get my results?
Your oncologist will get the results of the biomarker testing within 3 to 10 business
days. Your oncologist may call you with the results or discuss them with you at your
next appointment. Either way, your oncologist will discuss the results and treatment
options that may be appropriate for you based on those results.

OTHER DIAGNOSTIC TESTS

Pulmonary Function Test (PFT)
The PFT is a breathing test to determine how well your lungs are working. This
non-painful test may be performed in your pulmonologist’s office or in the hospital
on an outpatient basis.

Pulse Oximetry (Pulse Ox)
One of the common symptoms you might experience with lung cancer is shortness
of breath. Your team may use a device called a pulse oximeter to measure the level of
oxygen in your blood. The pulse oximeter is placed on your fingertip for a minute. A low
level of oxygen in your blood may prompt your doctor to order extra oxygen for you
during your illness.
High Altitude Simulation Test (HAST)
The High Altitude Simulation Test (HAST) is a test your doctor may use to find out if you will need oxygen when you fly or travel to a high altitude city or country. You may also hear HAST called a “hypoxia altitude simulation test.” When you fly or are at a high altitude, you may be at risk for cardiopulmonary (heart or lung) problems due to the lower oxygen available. During the HAST exam, your doctor will take your blood pressure, pulse and respiratory rate before the test begins and while you are breathing your normal air mixture. The doctor may also connect you to a cardiac monitor that will allow the team to monitor your heart rhythm. After those initial measurements, you will breathe air that contains a lower percentage of oxygen than you may be used to. During the 20 to 30 minutes of the test, your doctor will monitor you for any significant symptoms you have while breathing the air with lower oxygen. If you have symptoms during the test, the doctor will test you again while giving you oxygen to ensure that the additional oxygen will prevent the symptoms. The doctor doing the test will send results of the HAST to your oncologist and pulmonologist.

Complete Blood Count (CBC)
Chemotherapy and radiation therapy can temporarily affect the cells of the bone marrow that produce normal blood cells, so your healthcare team will want to keep track of this important test before and during treatment. A CBC can also alert your doctor to certain abnormalities in the blood that may indicate problems with the function of your kidneys or liver. Your doctor will order a CBC on a regular basis to determine if your blood has the correct number and types of cells.
Sputum Cytology
For the sputum cytology test, a member of your healthcare team will ask you to cough up a sample of mucus (sputum) from as deep in your lungs as possible. Lung cancer cells can shed into the airway and mix with mucus there. When you produce a sputum sample, the or pathologist will examine the sputum looking for normal or abnormal cells. Cytology is the study of cells and a cytologist is a scientist who studies the identification of cancer cells.

DIAGNOSIS TIMELINE

How long should I expect to wait for results and, ultimately, a diagnosis?
The timeline for the diagnosis of lung cancer may vary greatly based on who your physician is, the institution where you are being treated, your treatment plan and possible other diagnostic tests that may need to be performed. Following is a timeline that the GO2 Foundation would be pleased to see as the standard of care for the diagnosis of lung cancer.

Following an x-ray that shows a suspicious spot on the lung, you should get a CT scan. If the CT scan shows a spot, your doctor will schedule a biopsy. Following the biopsy,
depending on the type of lung cancer found in your tumor, your doctor might want to perform biomarker testing. If your oncologist recommends infusion (in the vein) chemotherapy and you decide to have an intravenous port inserted, your team will schedule and perform this procedure. Depending on the treatment recommended by your oncologist, access to the chemotherapy drugs may take time. In the ideal circumstances, the time from identification of a possible tumor to treatment may take up to two weeks. This timeline may vary depending on the availability of services in your area — but this timeline should be the goal for your treatment team.

My doctor says I have lung cancer. What happens next?
Your family doctor or the doctor who helped diagnose the lung cancer will refer you to an oncologist who will work with you to design your treatment plan. You may also have a radiation oncologist if you will be receiving radiation treatments for the cancer.

Your healthcare team may also consist of many other people from many different specialties whose job it is to help you understand your disease and make your treatments as comfortable as possible.

Can I get a second opinion?
Before starting treatment, you may want a second opinion about your diagnosis and treatment plan. Many insurance companies cover the cost of a second opinion if you or your doctor requests it. There are many ways to find a doctor for a second opinion. The best way is to have your doctor refer you to someone he or she trusts. If your doctor refers you, the time to get an appointment may be much shorter. You can also get names of physicians and medical centers specializing in lung cancer by contacting 1-800-298-2436, calling or writing a local or state medical society, talking to social workers at your local hospital, or asking at a nearby medical school for names of specialists they recommend. Your nearest cancer center or cancer support group may also be excellent sources of names for second opinions.
Before going to see another doctor for a second opinion, be sure to collect all your medical records, including x-rays and pathology reports, to take with you. In some cases, you may be able to have the hospital or your doctor send your records directly to the doctor you will be seeing. Be patient since this process is not always smooth. Ask your doctor if a delay for a second opinion will have a negative impact on your health. In most cases, a delay of less than two weeks will make very little difference in the effectiveness of your treatment plan. Be sure to check with your insurance company about expenses they will cover if you must travel to another city for a second opinion. Some companies will cover all or part of this expense.

What is the difference between a community cancer center and an academic medical center?
Depending on the treatment prescribed by your oncologist and the options available in your community, you may receive your treatment in a number of settings. First, you may have appointments in your oncologist’s office located in a medical office building or community cancer center. The oncologist’s office may have a laboratory in the office; this will mean that most lab tests can be done without having to go to another office. Your oncologist may also have an infusion center in his or her office where you will be able to receive chemotherapy treatments.

Second, you may have access to a community cancer center where you can receive most of the care you will need during your treatment. In 2007, the National Cancer Institute (NCI) began the Community Cancer Center Program providing funding to community cancer centers around the country. It is likely that there is a cancer center close to you where the goal is to provide high quality care while advancing cancer research efforts. Many cancer centers are associated with hospitals where you can easily receive your laboratory tests, diagnostic tests and procedures, radiation and chemotherapy treatments and surgical procedures. In addition, most cancer centers
also have extensive social service, financial counseling, and other support services you may need during your treatment course.

Finally, you may live close to an academic medical center that is associated with a variety of healthcare schools. If you do have access to an academic center, you may be able to receive more specialized treatment using technologies that are more advanced. Often, an academic medical center will have available innovative treatments that may not be available in a community hospital. Be aware that since these academic centers are associated with medical, nursing and other healthcare schools, your treatment team will likely include students and researchers learning new skills and conducting clinical trials. An academic medical center will also have extensive social service, financial counseling, and other support services you may need during your treatment course.

Depending on the type of facilities near you, the resources available may be very different. It is important to find a cancer center that has the resources you need to support you during your cancer treatment. Where you receive your care is as important as finding an oncologist who specializes in treating lung cancer. We are here to help—contact the GO2 Foundation for Lung Cancer at 1-800-298-2436 for a referral to a cancer center.
MULTIDISCIPLINARY HEALTHCARE TEAM

A multidisciplinary team is ideal! The following is a list of medical professionals you may have on your healthcare team. Some of these people may have different titles, and the same person may fill some of these roles, but you should have access to these services:

**Medical Oncologist:** Physician who treats cancer using medications and chemotherapy drugs.

**Radiation Oncologist:** Physician who specializes in treating cancer using various forms of radiation therapy.

**Radiologist:** Physician who uses x-rays and special radiology procedures in cancer diagnosis and treatment. This includes CT scans, MRI, and PET scans.

**Thoracic Surgeon:** Physician who specializes in the surgical treatment of cancer and other diseases of the chest.

**Pulmonologist:** Physician who specializes in the evaluation and treatment of lung problems.

**Pathologist:** Physician who analyzes tumor tissues removed by biopsy or surgery in order to diagnose and stage cancer and other diseases.

**RN Navigator:** Registered nurse who will help you and your family by offering education, support and coordination of services in the process of diagnosis and treatment of cancer.

**Chemotherapy Nurse:** Registered nurse who specializes in the delivery of chemotherapy and other cancer treatments, helping you deal with any side effects and placing IVs for infusions.
Research Nurse: Registered nurse who administers and provides nursing care if you are involved in clinical trials.

Symptom Management Care Coordinator: Registered nurse or physician who will help you manage symptoms associated with disease and treatment of cancer.

Radiation Technician: Licensed professional who will guide you through radiation treatments, inject dyes or contrast for radiation tests, and care for you during radiation treatments.

Social Worker: Licensed professional who will be available to assist you and your family with supportive counseling and community resources.

Registered Dietitian: Licensed professional who will help you develop a nutritional plan based on your specific needs.

GO2 Foundation for Lung Cancer: One of the largest patient-founded, patient-focused, and patient-driven philanthropies devoted exclusively to eradicating lung cancer through research, early detection, education, and treatment. The Foundation is available to assist you along your cancer journey. Simply call us at 1-800-298-2436.

What should I do if a multi-disciplinary team is not readily available?
If you live in an area far from a cancer center or major medical center, you may want to travel to have a second opinion or to access more resources. A medical oncologist at a major medical center may be willing to work with your local oncologist to ensure that you get the most advanced care as possible at your local hospital or clinic. If you are unable to travel to a cancer center or major medical center, ask your local oncologist for help identifying the resources you need to manage your treatment course. We are here to help; feel free to contact us at 1-800-298-2436 for information on local resources.
“Treatments can not only extend people’s lives, but also improve the quality of their lives.”

– Jacob Sands, MD,
Scientific Leadership Board
The Patient Education Handbook was and still is my go to for information. It’s been five years since I was diagnosed and I still refer back to the handbook for information.

—Kimberly Buchmeier, survivor
LUNG CANCER STAGING

In addition to grading the tumor, your healthcare team will also stage your lung cancer.

What does the stage of my lung cancer mean?
The stage of your lung cancer tells your oncologist the size of the primary tumor, the number of lymph nodes with cancer cells in them, and if the cancer has spread to other organs. Knowing the stage of the lung cancer is critical because the stage of lung cancer will help you and your oncologist determine the types of treatment that will be most effective for you.

You may be familiar with the traditional staging of lung cancer in which your oncologist may describe the stages as Stage I, II, III or IV. In this staging, the higher number indicates lung cancer that is more extensive. Oncologists also use the TNM system as a way to determine the stage of the lung cancer.

T, N, M Lung Cancer Staging
The TNM staging system was developed by the American Joint Committee on Cancer (AJCC) and the International Union Against Cancer (UICC). Since the development of this system, it has become one of the most commonly used staging systems for cancer. Your oncologist will use the TNM classification system to stage your cancer based on standard criteria.

Questions for your physician about staging your lung cancer:
• What stage is my lung cancer and what does that mean for me?
• Has the cancer spread from my lung to other parts of my body?
• Will I need more tests before deciding which treatments to take?
According to the 2010 definitions of staging, the letters T, N and M, identify three major pieces of information used for tumor staging:

- **T** = describes the size of the primary tumor
- **N** = describes the number of lymph nodes with cancer cells in them
- **M** = describes the presence of metastatic tumors in distant organs

If your team uses this staging, your doctor might describe the stage of your lung cancer, for example, as T1, N1, M0. This designation would mean that the primary tumor has been identified but is relatively small (T1). There are lymph nodes involved (N1), but the cancer has not spread to other organs (M0).

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**Stage I, II, III, IV Lung Cancer Staging**

**Stage 0 or Carcinoma in Situ:** If the oncologist says that you have Stage 0, this means that your doctor found abnormal cells in your airway, often with a sputum cytology test. These cells may grow and invade the lung.

**Stage I:** If your oncologist says your lung cancer is Stage I, it means that a tumor has been found in one lung only and there is no cancer found in the lymph nodes.
Staging chart for Stage I:

<table>
<thead>
<tr>
<th>Stages 0 to I of lung cancer</th>
<th>TNM (Tumor, Nodes, Metastasis)</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Occult carcinoma</td>
<td>TX, N0, M0</td>
<td>TX = Primary tumour cannot be assessed, or tumour proven by the presence of malignant cells in sputum or bronchial washings but not visualized by imaging or bronchoscopy. N0 = No regional lymph node metastasis M0 = No distant metastasis</td>
</tr>
<tr>
<td>0</td>
<td>Tis, N0, M0</td>
<td>Tis = Carcinoma in situ N0 = No regional lymph node metastasis M0 = No distant metastasis</td>
</tr>
<tr>
<td>IA</td>
<td>T1a, N0, M0</td>
<td>T1a = Tumour ≤2 cm in greatest dimension T1b = Tumour &gt;2 cm but ≤3 in greatest dimension N0 = No regional lymph node metastasis M0 = No distant metastasis</td>
</tr>
<tr>
<td>IB</td>
<td>T2a, N0, M0</td>
<td>T2a = Tumour &gt;3 cm but ≤5 cm in greatest dimension N0 = No regional lymph node metastasis M0 = No distant metastasis</td>
</tr>
</tbody>
</table>

Provided courtesy of the International Association for the Study of Lung Cancer.\(^1\)
Stage II: Stage II lung cancer means that your doctor has found cancer in one lung only and there may be lymph node involvement on the same side as the lung cancer. In Stage II cancer, the cancer is not found in the lymph nodes in the mediastinum. The mediastinum is the area between the lungs from the breastbone and the spinal column.

Stage chart for Stage II:

<table>
<thead>
<tr>
<th>Stage II of lung cancer</th>
<th>TNM (Tumor, Nodes, metastasis)</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>IIA</td>
<td>T1a, N1, M0</td>
<td>T1a = Tumour ≤ 2 cm in greatest dimension</td>
</tr>
<tr>
<td></td>
<td>T1b, N1, M0</td>
<td>T1b = Tumour &gt; 2 cm but ≤ 3 cm in greatest dimension</td>
</tr>
<tr>
<td></td>
<td>T2a, N1, M0</td>
<td>T2a = Tumour &gt; 3 cm but ≤ 5 cm in greatest dimension</td>
</tr>
<tr>
<td></td>
<td>T2b, N0, M0</td>
<td>N1 = metastasis in ipsilateral peribronchial and/or ipsilateral hilar lymph nodes and intrapulmonary nodes, including involvement by direct extension</td>
</tr>
<tr>
<td></td>
<td></td>
<td>M0 = No distant metastasis</td>
</tr>
</tbody>
</table>

Provided courtesy of the International Association for the Study of Lung Cancer.¹¹
**Stage II of lung cancer**

<table>
<thead>
<tr>
<th></th>
<th>TNM (Tumor, Nodes, metastasis)</th>
<th>Definition</th>
</tr>
</thead>
</table>
| IIB | T2b, N1, M0 | T2b = Tumour >5 cm but ≤7 cm in greatest dimension  
N1 = metastasis in ipsilateral peribronchial and/or ipsilateral hilar lymph nodes and intrapulmonary nodes, including involvement by direct extension  
M0 = No distant metastasis |
|  | T3, N0, M0 | T3 = Tumour >7 cm or one that directly invades any of the following: chest wall (including superior sulcus tumours), diaphragm, phrenic nerve, mediastinal pleura, parietal pericardium; or tumour in the main bronchus less than 2 cm distal to the carina but without involvement of the carina; or associated atelectasis or obstructive pneumonitis of the entire lung or separate tumour nodule(s) in the same lobe as the primary.  
N0 = No regional lymph node metastasis  
M0 = No distant metastasis |
Stage IIIA: Stage IIIA lung cancer means that there may be one or more tumors in the same lobe of the lung. In this stage, the cancer has also spread to the lymph nodes on the same side of the lung as the cancer or to where the trachea joins the bronchus, the chest wall, or the lining around the lung.

Staging chart for Stage IIIA:

<table>
<thead>
<tr>
<th>Stage IIIA of lung cancer</th>
<th>TNM (Tumor, Nodes, Metastasis)</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>IIIA</td>
<td>T1a, N2, M0</td>
<td>T1a = Tumour ≤2 cm in greatest dimension</td>
</tr>
<tr>
<td></td>
<td>T1b, N2, M0</td>
<td>T1b = Tumour &gt;2 cm but ≤3 cm in greatest dimension</td>
</tr>
<tr>
<td></td>
<td>T2a, N2, M0</td>
<td>T2a = Tumour &gt;3 cm but ≤5 cm in greatest dimension</td>
</tr>
<tr>
<td></td>
<td>T2b, N2, M0</td>
<td>T2b = Tumour &gt;5 cm but ≤7 cm in greatest dimension</td>
</tr>
<tr>
<td></td>
<td></td>
<td>N2 = Metastasis in ipsilateral mediastinal and/or subcarinal lymph node(s)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>M0 = No distant metastasis</td>
</tr>
<tr>
<td>Stage IIIA of lung cancer</td>
<td>TNM (Tumor, Nodes, Metastasis)</td>
<td>Definition</td>
</tr>
<tr>
<td>---------------------------</td>
<td>-------------------------------</td>
<td>------------</td>
</tr>
<tr>
<td></td>
<td>T3, N1, M0</td>
<td>T3 = Tumour &gt;7 cm or one that directly invades any of the following: chest wall (including superior sulcus tumours), diaphragm, phrenic nerve, mediastinal pleura, parietal pericardium; or tumour in the main bronchus less than 2 cm distal to the carina but without involvement of the carina; or associated atelectasis or obstructive pneumonitis of the entire lung or separate tumour nodule(s) in the same lobe as the primary. N1 = Metastasis in ipsilateral peribronchial and/or ipsilateral hilar lymph nodes and intrapulmonary nodes, including involvement by direct extension. M0 = No distant metastasis.</td>
</tr>
<tr>
<td></td>
<td>T3, N2, M0</td>
<td>T3 = Tumour &gt;7 cm or one that directly invades any of the following: chest wall (including superior sulcus tumours), diaphragm, phrenic nerve, mediastinal pleura, parietal pericardium; or tumour in the main bronchus less than 2 cm distal to the carina but without involvement of the carina; or associated atelectasis or obstructive pneumonitis of the entire lung or separate tumour nodule(s) in the same lobe as the primary. N2 = Metastasis in ipsilateral mediastinal and/or subcarinal lymph node(s). M0 = No distant metastasis.</td>
</tr>
<tr>
<td></td>
<td>T4, N0, M0</td>
<td>T4 = Tumour of any size that invades any of the following: mediastinum, heart, great vessels, trachea, recurrent laryngeal nerve, oesophagus, vertebral body, carina; separate tumour nodule(s) in a different ipsilateral lobe to that of the primary. N0 = No regional lymph node metastasis. M0 = No distant metastasis.</td>
</tr>
</tbody>
</table>
## LUNG CANCER STAGING

<table>
<thead>
<tr>
<th>Stage IIIA of lung cancer</th>
<th>TNM (Tumor, Nodes, metastasis)</th>
<th>Definition</th>
</tr>
</thead>
</table>
| IIIA                     | T4, N1, M0                    | T4 = Tumour of any size that invades any of the following: mediastinum, heart, great vessels, trachea, recurrent laryngeal nerve, oesophagus, vertebral body, carina; separate tumour nodule(s) in a different ipsilateral lobe to that of the primary. 
N1 = Metastasis in ipsilateral peribronchial and/or ipsilateral hilar lymph nodes and intrapulmonary nodes, including involvement by direct extension 
M0 = No distant metastasis |
**Stage IIIB:** In Stage IIIB lung cancer, there may be separate tumors in any of the lobes of the lung and the cancer may have spread to the chest wall, diaphragm, lining of the lung or chest wall, lining of the heart or the heart, major blood vessels that lead to or from the heart, esophagus, sternum, or spine.

**Staging chart for Stage IIIB:**

<table>
<thead>
<tr>
<th>Stage IIIB of lung cancer</th>
<th>TNM (Tumor, Nodes, Metastasis)</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>IIIB</td>
<td>T4, N2, M0</td>
<td>TX = Primary tumour cannot be assessed, or tumour proven by the presence of malignant cells in sputum or bronchial washings but not visualized by imaging or bronchoscopy. N0 = No regional lymph node metastasis M0 = No distant metastasis Any T N3 = Metastasis in contralateral mediastinal, contralateral hilar, ipsilateral or contralateral scalene, or supraclavicular lymph node(s) M0 = No distant metastasis</td>
</tr>
</tbody>
</table>

Provided courtesy of the International Association for the Study of Lung Cancer.¹¹
Stage IIIC: A new stage grouping has also been created for the most advanced local disease categories, T3 and T4 associated with N3 disease but category M0. Now classified as stage IIIC. The separation is justified by the different treatment approaches used in such cases.

Staging chart for Stage IIIC:

<table>
<thead>
<tr>
<th>Stage IIIC of lung cancer</th>
<th>TNM (Tumor, Nodes, Metastasis)</th>
<th>Definition</th>
</tr>
</thead>
</table>
| IIIC                     | T3, N3, M0                    | T3 = Tumour more than 5 cm but not more than 7 cm in greatest dimension, or directly invades any of the following structures: chest wall (including parietal pleura and superior sulcus tumours), phrenic nerve, parietal pericardium; or associated with separate tumour nodule(s) in the same lobe as the primary.  
N3 = Metastasis in contralateral mediastinal, contralateral hilar, ipsilateral or contralateral scalene, or supraclavicular lymph node(s).  
M0 = No distant metastasis.  
T4 = Tumour more than 7 cm in greatest dimension, or invades any of the following structures: diaphragm, mediastinum, heart, great vessels, trachea, recurrent laryngeal nerve, oesophagus, vertebral body, carina; or associated with separate tumour nodule(s) in a different ipsilateral lobe to that of the primary. |
| T4, N3, M0               |                               |            |

Provided courtesy of the International Association for the Study of Lung Cancer.
**Stage IV:** In Stage IV lung cancer, there are one or more tumors in both lungs and cancer may be found in the fluid around the lung. The cancer may have spread through to other organs of the body, often the brain, liver, or the bones.

Staging chart for Stage IV:

<table>
<thead>
<tr>
<th>Stage IV of lung cancer</th>
<th>TNM (Tumor, Nodes, Metastasis)</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>IV</td>
<td>Any T, Any N, M1</td>
<td>Any T</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Any N</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• NX = Regional lymph nodes cannot be assessed</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• N0 = No regional lymph node metastasis</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• N1 = Metastasis in ipsilateral peribronchial and/or ipsilateral hilar lymph nodes and intrapulmonary nodes, including involvement by direct extension</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• N2 = Metastasis in ipsilateral mediastinal and/or subcarinal lymph node(s)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• N3 = Metastasis in contralateral mediastinal, contralateral hilar, ipsilateral or contralateral scalene, or supraclavicular lymph node(s)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>M1 = Distant Metastasis</td>
</tr>
</tbody>
</table>

Provided courtesy of the International Association for the Study of Lung Cancer. 

While staging is a useful classification, it is important to remember that you must discuss these stages with your healthcare team who will help you understand what the stage means in the context of your specific diagnosis and treatment plan.
I love this organization. They are making the world a better place. Everyone that works there is talented, dedicated and empathetic. It’s a phenomenal team. I’m so grateful for them.

— Lisa Goldman
NON-SMALL CELL LUNG CANCER TREATMENTS
We’re improving the standard of care by bringing new and improved treatment options to lung cancer patients. The future of cancer care lies in the concept of personalized medicine—a model that focuses on the individual, not just the disease.

—Bruce Gellman, Board Member
OVERVIEW
After you are diagnosed with lung cancer, your next question is sure to be “What can be done to treat the cancer?” Your individual treatment plan will depend on the type of lung cancer, its stage and your overall health. As you begin to plan your treatment with your healthcare team, it is important that you keep a list of questions you have. This can be a confusing time so remember to write everything down as you discuss your treatment plan with the team.

Possible treatments for lung cancer include surgery, chemotherapy, radiation therapy, targeted therapy, immunotherapy or a combination of these.

Lung cancer treatments fall into three categories:

- Local therapy: Surgery and radiation therapy are local therapies. They remove or destroy

Questions for your physician regarding lung cancer treatments:

- Has my cancer been tested for multiple biomarkers?
- What are my treatment choices?
- What is the goal of my treatment (curative, stable disease, palliative or symptom management)?
- Will I have more than one type of treatment?
- What are the expected benefits of each type of treatment?
- How will treatment affect my normal activities and daily life?
- What can we do to control the side effects?
- Are there other treatment options available to me?
- Are there any clinical trials available to me?
- What can I do to prepare for treatment?
- Will I need to be hospitalized? If so, for how long?
- What is the cost of treatment? Will my insurance cover the cost?
- Do I have time to seek a 2nd opinion or to think about my treatment options? If so, how long before I need to start treatment?
cancer tumors in the lungs. If lung cancer has spread to other parts of the body, such as other organs or bones, your doctor may use one of these local therapies to control the disease in those specific areas as well.

- **Systemic therapy:** Chemotherapy and targeted therapies are **systemic therapies**. These drugs enter the bloodstream to destroy or control cancer everywhere in the body. Systemic therapy is taken by mouth or given through a vein in the arm or a port that is inserted in your chest (intravenous).

- **Immunotherapy:** **Immunotherapies** are types of treatments that use substances to stimulate or suppress the immune system to help the body fight cancer. Some types of immunotherapy only target certain types of cells in the immune system.

### SURGERY FOR THE TREATMENT OF NSCLC

Surgery may be effective for treatment of stages I - III in non-small cell lung cancer.

**When is surgery used to treat lung cancer?**
If your NSCLC tumor has not spread to other tissues outside of the lungs, your oncologist may recommend surgery to remove the tumor. Surgery may be the first type of treatment you receive, or your oncologist may recommend other non-surgical treatments first. In some cases, chemotherapy, immunotherapy or radiation will be used first to shrink the cancer tumor before surgery. The specific order of treatment depends on the size of the cancer tumor and whether the cancer has spread outside of the lungs.

**Questions for your physician about surgery:**
- What type of surgery do you recommend?
- How long will I be in the hospital?
- What side effects should I expect?
- Will I feel pain? If so, how will it be controlled?
- When can I get back to my normal activities?
The tissue that is removed from the lung (specimen) is sent to the pathologist who will look at the edges (or margins) of the specimen to see if the tumor has been completely removed. If there are tumor cells at the margin, it may mean that the entire tumor was not removed. These results will determine treatment following surgery.

**What types of surgery might be used to treat me?**

The following are types of surgery for lung cancer:

**To remove a small portion of the lung:**

*Wedge Resection*: In wedge resection, your surgeon will remove a small portion of the lung that contains the tumor. For stage I and II tumors, your surgeon may elect to use VATS, thoracotomy, or the *da Vinci* surgical system.

*Segmental Resection or Segmentectomy*: In a segemental resection, your surgeon will remove a slightly larger portion of tissue than with a wedge resection but not the whole lobe.

**To remove a lobe of the lung:**

*Sleeve Resection*: In a sleeve resection, the surgeon will try to keep as much of the lung as possible by removing only the lobe (part of lung) with a cancerous tumor. In this surgery, the surgeon will remove the lobe with the cancerous tumor and part of the bronchus (air passage); the lobe of the lung that is left is connected to the remaining bronchus.

*Lobectomy*: In a lobectomy, your surgeon will remove an entire lobe of the lung.

Ask your oncologist for a referral to a thoracic surgeon to assist with decisions involving surgery and to perform major surgical procedures on your lung.
To remove the entire lung:

**Pneumonectomy**: In a pneumonectomy, your surgeon will remove the entire lung.

Sleeve resections and pneumonectomy are used when the lung cancer tumor is larger and closer to the middle of the chest. A lobectomy is used when the lung cancer tumor is more peripherally located (away from the center of the chest).

To remove lymph nodes:

**Lymph Node Dissection or Lymphadenectomy**: During a lymph node dissection, your surgeon will remove several lymph nodes around your tumor to determine if any cancer cells are outside of your lungs. This will help your oncologist determine the stage of the lung cancer and the most appropriate treatment plan. If the pathologist finds cancer cells in the lymph nodes, you may receive chemotherapy after surgery to kill those cells.

**Treatment of Pneumothorax or Recurrent Pleural Effusion**

A pneumothorax is a collection of air in the space that separates your lung from your chest wall. When this happens, part of your lung may collapse making it difficult for you to breathe. A pleural effusion happens when fluid collects in the pleural layers that surround the lungs. This condition can also cause difficulty breathing. A pleurodesis is a chemical or surgical procedure that may be done to prevent these conditions from recurring (happening another time).

Your oncologist may do a chemical pleurodesis by injecting a drug into the pleural space around the lungs through a drain or tube in your chest. The drug acts as an irritant that closes the pleural space and prevents fluid from entering the space.
You will be given a local anesthetic that will numb the area on your chest where the tube enters. You may also be given a drug to relax you before the procedure starts.

A surgical pleurodesis is done by making an incision into your chest and rubbing the pleural layers with a rough pad to irritate the pleural lining. Your surgeon may also remove some of the parietal tissue. Either of these surgical procedures will be done under anesthesia.

Advantages of surgical removal of a lung tumor:

- If the margins of the tumor and lymph nodes outside of the lung do not contain cancer cells, surgery can be a cure for lung cancer.
- Because the surgeon removes all or most of the tumor, the size of the tumor tissue will be large enough for biomarker testing and to stage the tumor. The combination of accurate staging and biomarker testing will enable your oncologist to develop an individualized treatment plan specific to your type of lung cancer.
- Your surgeon can do a pleurodesis to prevent fluid from collecting between the lung and its lining.

Disadvantages of surgical removal of a lung tumor:

- Long recovery time
- Not all the cancer may be removed
- Risks associated with invasive surgery

What to expect during and after surgery:

- The surgeon will do the surgery procedure in the operating room.
- An anesthesiologist will use general anesthesia to put you to sleep for the procedure.
• You will remain in the hospital for about one week for recovery.
• Your doctor may order an epidural for pain control and other drugs to control pain.
• Your surgeon may insert a chest tube to drain any fluid that might collect after surgery.
• The respiratory therapist will teach you some breathing and strengthening exercises to help you recover more quickly after surgery.
• Your doctor may prescribe an inhaler filled with medicine to help you if you have trouble breathing.

Possible side effects of a surgical procedure:

• You may have pain from the surgery or chest tube incision; be sure to ask for pain medicine before your pain is severe. Controlling pain will be a large part of your recovery.
• You may experience some neuralgia (numbness) on the side of your chest where you had surgery.
• If fluid builds up around the lungs, you might develop a condition called a pleural effusion. This condition may cause you to have trouble breathing. Call your doctor if you notice shortness of breath that does not go away when you rest.
• The anesthesiologist will insert a tube into your throat during surgery to help you breathe during the procedure. This tube may injure one or both of your vocal cords resulting in hoarseness or difficulty speaking.
• Depending on the extent of the surgery, you may have weeks of recovery time following surgery.
• Women may want to avoid wearing a bra for a week or two after surgery because of pain and discomfort around the ribs.
• There are other possible side effects. Your surgeon and the surgical team will discuss the benefits and risks of surgery and anesthesia prior to surgery. Be sure to ask questions.

Tips for recovering from surgery – A patient’s perspective

After any surgery for lung cancer, you may have side effects because of the surgery. I know because I have been there. Your healthcare team will be able to tell you many things you can do to recover after surgery, but there are some things only another patient can tell you. A few things I have learned in my own journey that may help you as you recover from a surgical procedure include:

• Be sure you talk to your team before the surgical procedure so you know exactly what to expect after the surgery.
• After lung surgery, the incision area may be sore. A cold pack for 20 minutes at a time may help relieve the swelling at the site. Talk to your surgeon to make sure this is something you can do.
• It may help if you sleep with your head and shoulders raised. This may help your lungs to expand more fully and allow you to breathe better.
• Unless your surgeon says you should stay in bed, be sure to get up in a chair several times a day and walk a little more each day. Unless your condition requires that you stay in bed, you will recover faster if you get up and moving as soon as possible after surgery.
• The first day or two after surgery, you should take pain medicine regularly in order to make it easier to move; however, the sooner you stop taking pain medicine the more energy you will have.
• Eat small meals often. These small meals will allow you to have energy throughout the day without having a full stomach that may interfere with your breathing. As you eat more frequent but smaller meals, be sure you are drinking lots of fluid.
• Keep all appointments with your healthcare team and report any symptoms that you think are not normal following your surgery!

Information presented in Navigating Lung Cancer 360° of Hope is not intended as a substitute for the advice given by your health care provider. We recommend you follow the instructions provided to you by your healthcare team. Contact your physician with any questions or concerns.
Chemotherapy can be used to treat lung cancer stages I - IV NSCLC and SCLC, limited and extensive.

When is chemotherapy used to treat lung cancer?
Your oncologist may use chemotherapy to destroy or control growth of cancer in the body. Chemotherapy is a cancer treatment that uses drugs in pill form or intravenously (IV inserted into a vein or delivered through a port in the chest wall) to stop the growth of cancer cells, either by killing the cells or by stopping them from dividing. Your oncologist may also refer to chemotherapy as systemic therapy because it circulates throughout the body. If you receive chemotherapy, your oncologist may prescribe only one of these drugs. Most of the time, your oncologist will prescribe chemotherapy drugs in some combination of drugs. When you receive several different chemotherapy drugs, this combination of drugs is called a chemotherapy regimen.

Questions for your physician regarding chemotherapy:
- Will I have one drug or a combination of drugs?
- What are the benefits of chemotherapy?
- When will treatment start and how long will it last?
- How often will I receive chemotherapy?
- Where do I go for treatment?
- Will I need someone to help me get home after my chemotherapy?
- How will we know the treatment is working?
- What side effects should I tell you about?
- Can I prevent or treat any of the side effects?
- Will I have side effects after the treatment is completed?
- Can I take vitamins while I am on chemotherapy?
- Do I have to eat certain foods or avoid certain foods?
Your oncologist will determine the dose and schedule of your chemotherapy regimen based on the type, stage and molecular profile of your tumor. Usually, you will receive your chemotherapy treatment in cycles, with each period of treatment followed by a recovery period. You will receive a first chemotherapy regimen called a first-line treatment. If the first-line treatment is not effective, you may receive another combination of chemotherapy drugs called a second-line treatment. The Food and Drug Administration (FDA) classifies different chemotherapy drugs as first- or second-line treatments. This means each chemotherapy drug has been determined to be effective as either a first- or second-line treatment.

In NSCLC, chemotherapy drugs can be used as neoadjuvant therapy, which is treatment before surgery. Your oncologist may prescribe neoadjuvant therapy to shrink the tumor so that surgery will be easier or more effective. Neoadjuvant chemotherapy treatments are usually used in stage IIIA cancer. Your doctor will use these drugs before surgery to kill cancer cells in lymph nodes in the chest before surgery. After the chemotherapy, surgery will be done and more chemotherapy will probably be done. Sometimes, your oncologist may also prescribe radiation after the surgery and chemotherapy treatments.

Chemotherapy drugs may be used as adjuvant therapy. Adjuvant therapy is any therapy that is started after surgery. Your oncologist may prescribe adjuvant therapy to kill cancer cells that may not have been removed during surgery or which may have spread from the primary tumor.

"Numerous clinical trials have shown statistical improvement in outcomes for using at least two drugs (doublets) for adjuvant treatment in stage IB, stage II and stage III disease as well as in first-line treatment for stage IV disease." —Shane Dormady, MD, PhD
Your surgeon may not be able to remove late stage non-small cell lung cancers by surgery. In this case, your oncologist will probably prescribe chemotherapy to try to destroy cancer cells or control the growth of the tumor. A number of chemotherapy regimens can be used to treat non-small cell lung cancer. These are usually used for stages III - IV for NSCLC. For information on SCLC treatments, see the chapter titled the Small Cell Lung Cancer—Treatments.

If you have a good response after the first-line treatment, your oncologist may prescribe maintenance therapy. There are two types of maintenance therapy: continuation maintenance and switch maintenance. **Continuation maintenance therapy** means that your oncologist will continue using at least one of the chemotherapy drugs you received during your first-line treatment. **Switch maintenance** means that your oncologist will prescribe a different chemotherapy drug – one that was not part of your first-line treatment.

**What kinds of chemotherapy might be used to treat me?**
Your oncologist may prescribe one or more chemotherapy drugs that you will receive in your vein (IV or intravenous) or by pill. If your treatment involves drugs in the vein, you will receive these drugs in the hospital or in your cancer center’s infusion center. If your treatment involves drugs in pill form, you will be able to take these at home.

**What specific FDA-approved drugs, or chemotherapies, are approved for NSCLC?**
We understand the following list may be overwhelming. It is important to understand that for NSCLC, a platinum-based chemotherapy is the backbone of the “recipe” and with first-line therapy your doctor will add another drug to the platinum-drug. As you move to second-, third- and fourth-line therapy, some of the drugs are just used one at a time.
## Non-Small Cell Lung Cancer - Treatments

### Chemotherapies

<table>
<thead>
<tr>
<th>Brand Name</th>
<th>Generic Name</th>
<th>Cancer Type</th>
</tr>
</thead>
<tbody>
<tr>
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### Immunotherapies

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### Targeted Therapies

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This Foundation is tenacious, innovative, collaborative and determined that lung cancer patients will no longer be left behind.

—Jaimi Julian Thompson
Advantages of chemotherapy treatments:

- May cure the cancer
- Can slow the cancer’s growth
- May keep the cancer from spreading
- Can kill cancer cells that may have spread to other parts of the body from the original tumor
- Can shrink the tumor prior to surgery
- Can destroy any cancer cells that are still present after surgery and/or radiation
- Can relieve symptoms caused by the cancer

Disadvantages of chemotherapy treatments:

- May need more than one chemotherapy regimen
- Side effects of chemotherapy drugs

What to expect with your chemotherapy regimen:

- If your treatment plan consists of many treatments with chemotherapy drugs in your veins, your oncologist may suggest that you have a permanent IV site, or port, put under your skin near your collarbone. This port allows easy access to your bloodstream and protects the veins in your arms.
- Unless you develop complications that require that you be in the hospital, your healthcare team will probably give you your IV treatments on an outpatient basis at the hospital or cancer center.

If your chemotherapy drug is in pill form, your oncologist will tell you how and when to take the pill. You will be able to take the pills at home.
Possible side effects of chemotherapy:

Side effects of chemotherapy will depend on the type and length of your treatment and your body’s own reaction to the chemotherapy drugs. Although not an exhaustive list, you may experience some of the following:

- Fatigue
- Feeling weak, loss of strength
- Nausea and vomiting
- Hair loss
- Drop in white-blood cells, increasing chance of infection
- Drop in red-blood cells, increasing chance of anemia
- Skin and nail changes
- Peripheral neuropathy (tingling, burning, weakness or numbness in your hands/feet)

Possible long-term effects of chemotherapy:

- Menopause
- Infertility (if you are of child bearing age, consider talking with your physician about fertility preservation options)
- Damage to your heart or lungs
- Bone disease (brittle/necrotic)

RADIATION THERAPY FOR THE TREATMENT OF LUNG CANCER

When is radiation therapy used to treat lung cancer?

Your oncologist may prescribe traditional radiation therapy as part of your treatment plan. Radiation therapy treats lung cancer by using high-energy x-ray beams to destroy cancerous tumors. Because radiation is focused directly on the tumor, you may hear it referred to as local therapy as opposed to chemotherapy that goes through your whole body and is called a systemic therapy. Cancer experts classify radiation therapy as a local therapy because it is aimed directly at the tumor.
Radiation treatments are sometimes given along with chemotherapy. This is known as combination or combined modality therapy. Combination therapy may have more side effects than radiation or chemotherapy alone, but can be more effective at destroying the cancer cells.

Questions to ask your physician regarding radiation therapies:

- What is the probability that radiation therapy will work for me? If it works, what are the chances that the cancer will come back in the same place or other places?
- What are the chances that the cancer will spread if I do not have radiation therapy?
- How will the radiation therapy be given?
- How many treatments will I receive per week and for how long?
- What side effects should I expect and how do I manage them?
- Will I also need other treatments, such as chemotherapy, surgery, or hormone therapy? If so, when will I receive them and in what order?
- Will I need a special diet during or after my radiation treatment?
- Can I drive myself to and from the treatment facility? Do you recommend I bring a friend or family member?
- Will I be able to continue my normal activities during treatment? If not, how soon after treatment will I be able to resume them such as work, aerobic exercise and sexual activity?
- How can I expect to feel during treatment and in the weeks following radiation therapy?
- What symptoms/problems after radiation should I tell you about?
- After my treatment is completed, how often will I need to return for checkups?

Are there different types of radiation?

External Beam Radiation
The most common kind of radiation treatment to treat lung cancer is “external beam radiation.” This treatment uses a machine called a linear accelerator to treat your lung cancer with high-energy photons, or “x-rays.” The high-energy x-rays are aimed at the tumor and destroy the DNA of the cancer cells. External beam radiation can be
used to treat both small cell and non-small cell lung cancers of all stages. Your radiation oncologist will talk to you about the type of treatment that is recommended in your particular case. Your external beam radiation treatment will usually last for 6 to 8 weeks. Types of external beam radiation include:

• **3-D Conformal Radiation Therapy**
  One of the most common types of external beam radiation therapy is three-dimensional conformal radiotherapy (3DCRT). This type of radiation therapy is a complicated treatment process that begins with x-rays that your radiation oncologist will use to create 3-D images of the tumor and the normal tissues around it. Your healthcare team will use these 3-D pictures to plan your individual treatment that will deliver radiation directly to your tumor and the area at risk around it. With 3DCRT, your radiation oncologist will be able to use multiple beams to deliver radiation to the tumor while limiting the amount of radiation to the healthy tissue around it.

• **Intensity-Modulated Radiation Therapy (IMRT)**
  Intensity-modulated radiation therapy, most commonly called IMRT, is an advanced form of 3DCRT. Your radiation oncologist will use specialized software and hardware to focus small “beamlets” of radiation to treat only the tumor while limiting dose to the healthy tissue around the tumor. This allows your doctor to treat tumors that might have been untreatable in the past because they were too close to healthy organs.

In some cases, there may be fewer side effects than with conventional radiation treatments. Treatment times for each IMRT treatment may be longer than with other techniques because daily set-up is very precise and requires multiple measurements.
• **Image Guidance Radiation Therapy (IGRT)**
  One of the problems with treating lung cancer with radiation treatment is that the tumor moves as you breathe. With Image Guidance Radiation Therapy (IGRT) tracking, the radiation beam turns on *only* when the tumor is in the path of the beam. IGRT is another radiation treatment that targets the tumor only, limiting the exposure of normal tissue around the tumor.

• **Volumetric Arc Therapy (VMAT)**
  Volumetric Arc Therapy (VMAT) is the most advanced form of IMRT. It allows for treatment of the tumor as the radiation machine is moving. This means your treatment may be faster.

**Radiosurgery**

Another way to treat lung cancer is with radiosurgery, also called “stereotactic radiotherapy.” Radiosurgery does not involve actual surgery with a knife, but instead uses many pinpoint radiation beams that focus on one small area and treat it with a very high dose of radiation.

When radiosurgery is used for cancers in the lung or elsewhere in the body (outside of the head), it is called “stereotactic body radiation therapy” or SBRT. SBRT can be given with a traditional radiation device or with a machine especially designed for radiosurgery. SBRT can be used in place of traditional surgery in some patients with early stage disease who either cannot or choose not to have surgery. New studies released in 2011 showed that radiosurgery has results that are as good as or better than traditional surgery in some stage I patients.\(^\text{12,13}\)

A typical radiosurgery course of treatment takes 1 to 5 treatments, as opposed to 6 or 7 weeks with other types of external beam radiation. Each SBRT treatment may last several hours.
What to expect with Radiation Therapy

External beam radiation is usually given once per day, Monday through Friday, for 6 to 8 weeks. During treatment, you will be lying on a table and the machine will move around you. You will hear some noises from the machine, but the treatment itself is painless, much like having a chest x-ray or dental x-ray. In order to make sure you are in the proper position, your radiology technician may place small dots in the form of a tattoo in order to aim the radiation beam at exactly the same spot each time you receive radiation therapy. You will be asked to lie still for 15 to 30 minutes depending on the length of the treatment. Although each individual treatment is painless, a small dose of radiation is given each day and you should be aware of the possible side effects that may develop during the course of your treatment:

- Your skin may look and feel as if it has been sunburned. You will be given skin creams and instructions on how to manage this, and it will go away within a few weeks after your treatment is complete.
- Mild or moderate fatigue may begin after the first few weeks of treatment. This fatigue will peak by the end of treatment. Approximately 4 to 8 weeks following the end of treatment, your fatigue should be much better. Fatigue may be worse if you are receiving a combination of chemotherapy and radiation.
- Your esophagus is often exposed to radiation during lung cancer treatment.
- This exposure to the radiation can result in temporary sore throat and pain with swallowing that you may first notice 3 to 4 weeks into your treatment. You may find that soft foods or liquids are easier to swallow during this time, and your doctor may prescribe medication to help with discomfort. Your sore throat and difficulty swallowing should be better within 2 to 3 weeks after your treatments are completed.
- You may develop a temporary cough or change in breathing during radiation. This is usually managed with cough medications and sometimes a short course of steroids.
Radiation pneumonitis is a radiation pneumonia caused by treatment. This complication occurs in 5 to 15% of patients and typically happens 2 to 6 months after your treatment is complete. This is a particularly important side effect because, if it is not treated, pneumonitis can be very serious. If you develop shortness of breath, chest pain when you breathe, cough or a low-grade fever after finishing radiation, be sure to report the symptoms to your oncologist. Pneumonitis is usually diagnosed with a chest x-ray and it is treated with steroids. With appropriate treatment, you will probably not have any lasting problems.

Radiation fibrosis is scarring of the lung that develops after radiation treatment. The amount of scarring depends on how much of your normal lung had to be treated, and the dose of radiation to that lung. Depending on the severity of the fibrosis, it can cause shortness of breath and coughing. Your oncologist may want you to be on oxygen if scarring develops.

Advantages of Radiation Therapy

- May cure the cancer
- May be used to shrink tumors to relieve pain or make surgery possible
- May be used as a targeted therapy to decrease the amount of time needed to give radiation, and leave healthy tissue unharmed.

Disadvantages of Radiation Therapy

- Side effects such as those listed above
- Unless you receive radiosurgery, you may spend time at appointments every day for several weeks.
PROTON THERAPY

Proton therapy (also called proton beam therapy) is a type of radiation treatment that uses protons rather than x-rays to treat cancer. A proton is a positively charged particle that is part of an atom, the basic unit of all chemical elements, such as hydrogen or oxygen. At high energy, protons can destroy cancer cells.

Like standard x-ray radiation, proton therapy is a type of external-beam radiation therapy. It painlessly delivers radiation through the skin from a machine outside the body. Protons, however, can target the tumor with lower radiation doses to surrounding normal tissues—approximately 60% lower, depending on the location of the tumor.

Traditional radiation treatment can damage the tissue around the tumor. However, with proton therapy, the protons’ energy hits the tumor site, delivering a smaller dose to surrounding healthy tissue. With standard treatment, doctors may need to reduce the radiation dose to limit side effects, resulting from damage to healthy tissue. With treatment using protons, on the other hand, doctors can select an appropriate dose, knowing that there will likely be fewer early and late side effects of radiation on the healthy tissue.

Compared with standard radiation treatment, proton therapy has several benefits. It reduces the risk of radiation damage to healthy tissues; may allow a higher radiation dose to be directed at some types of tumors, which may keep the tumor from growing or spreading; and may result in fewer and less severe side effects (such as low blood counts, fatigue, and nausea) during and after treatment.

(Source: Cancer.net)
GET THE RIGHT TREATMENT AT THE RIGHT TIME

Know your options
New ways to understand and treat lung cancer are being tested and approved more quickly than ever before. Knowing your treatment options makes you an active and empowered member of your team.

For treatment navigation, call our treatment specialists at 1.800.298.2436 or visit lungmatch.org.

Lung MATCH
Brain metastases
Radiation therapy is commonly used to treat brain metastases from lung cancer. In some cases, radiation is used to try to prevent brain metastases in people who are at high risk for developing them.

**External beam radiation** is used for treatment of the entire brain. Standard whole brain radiation is used for tumors that can be seen as well as for abnormal cells that can only be seen through a microscope. Your radiation oncologist will prescribe treatments lasting 2 to 4 weeks. Whole brain radiation may result in memory and cognition deficits unless hippocampal avoidance whole brain radiation is used instead of standard whole brain radiation. Avoiding radiation dose to the hippocampus minimizes memory and cognition deficits.

**Radiosurgery** is focused treatment that targets only the visible tumors. Typically, tumors up to 3 to 4 cm can be treated. There are various ways to treat brain metastases with commercially available radiosurgery devices including linear accelerator-based treatment, Gamma Knife®, Cyberknife®, Novalis and TrueBeam™. All of these treatments use pinpoint x-ray beams to target tumors with a high dose of radiation, and there are no clinical outcome data favoring one over another.

At times, radiosurgery is used in combination with whole brain radiation therapy for brain metastases. This combination can work well because whole brain radiation therapy treats the microscopic disease with a low dose of radiation and radiosurgery can deliver a high dose directly to the visible tumors.

Side effects from radiation to the brain vary depending on the type of treatment, but can include fatigue, weakness, hair loss, and neurologic effects including memory loss and speech problems.
New and experimental types of radiation treatment for lung cancer

Brachytherapy
Brachytherapy is the delivery of radiation treatment using radioactive seeds. These seeds may be placed in the target area and left for a specific length of time or may be left in the area permanently. Unlike external beam radiation, brachytherapy delivers radiation from inside the body.

Endoluminal high dose rate (HDR) brachytherapy
In HDR brachytherapy treatment, your pulmonologist places a high dose radioactive seed into the lung tumor using a small catheter through a bronchoscope. The seed is left in temporarily and then removed.

Mesh brachytherapy
This type of brachytherapy uses permanent radiation seeds (called a mesh) that are placed on the area where the tumor had been after a lung cancer is removed surgically. Mesh brachytherapy delivers a precise dose of radiation and reduces the risk of recurrence.

NanoKnife Electroporation
The NanoKnife® Irreversible Electroporation (IRE) System is a treatment that uses electrical energy to destroy soft tissue tumors. Probes are placed in the tumor and then brief electrical pulses are sent through the probes.
PULMONARY THERAPY

Interventional pulmonology is a specialty within pulmonary medicine that focuses on treating lung cancer and other airway diseases performed by a pulmonary medicine doctor who has additional advanced training in minimally invasive techniques and will be involved with your team in diagnosis, staging and treatment of lung cancer. A pulmonologist will usually do any number of procedures that will assist your oncologist, thoracic surgeon or radiation oncologist from biopsy procedures to treatment or symptom management.

The advantages of interventional pulmonary therapy are (1) less invasive procedure, (2) more precise biopsies and delivery of treatment, and (3) decreased recovery time. Your pulmonologist will describe the benefits and risks of each procedure.

Biopsy Procedures

Electromagnetic Navigation Bronchoscopy™ Procedure: Also known as ENB™ procedures, Electromagnetic Navigation Bronchoscopy™ procedures are performed by a pulmonologist or thoracic surgeon. ENB™ procedures provide a minimally invasive approach to accessing difficult-to-reach areas of the lung aiding in the diagnosis of lung disease.

Unlike a traditional bronchoscopy, the devices used in an ENB procedure create a real-time, image-guided map for doctors to access the deepest regions of your lungs. This allows them to see tumors and nodes that can’t be seen or accessed by
a traditional bronchoscope; making the need for more invasive techniques and exploratory surgeries unnecessary. This technology may be suitable for you if:

- You cannot undergo more aggressive procedures
- You have multiple tumors
- You want a diagnosis and/or staging before undergoing surgery
- You may be a candidate for stereotactic radiosurgery for fiducial marker placement at the time of biopsy
- You want to obtain additional lung tissue for genetic testing

**Endobronchial Ultrasound (EBUS) and Radial Probe Ultrasound (REBUS)**

With EBUS or REBUS, your pulmonologist uses a special ultrasound equipped bronchoscope. Your doctor may use this technique to do biopsies on multiple lesions. The procedure is much more accurate and the risk of puncturing a blood vessel is minimal, because the pulmonologist can see the needle as it is placed inside the tumor. Your doctor may use this technique to biopsy lymph nodes in the middle of the chest (EBUS) or in peripheral lung areas (REBUS).
Narrow band imaging
Narrow band imaging uses a specialized light at specific wavelengths that can detect abnormal blood vessels in the airway. These abnormal vessels may indicate tumor growth. Using these abnormal vessels, the pulmonologist may be able to guide a biopsy during bronchoscopy. Although not fully validated by scientific data, this technique may be used as a supplemental tool in some programs.

Treatment & Symptom Management Procedures

Argon Plasma Coagulation (APC)
For an APC procedure, your pulmonologist will use this technology to destroy tumors or stop bleeding. Using APC, your pulmonologist will use an argon gas jet to apply heat to specific areas without having to make direct contact with the area. As a result, the pulmonologist can treat a larger area, which will often shorten the procedure time.

Cryosurgery, laser
Using a bronchoscope, your pulmonologist may use cryotherapy to destroy tumors in your airways by freezing the tissue. The pulmonologist will apply a super-cooled probe over the entire surface of the tumor. This procedure is often used with argon plasma coagulation to open any airways that are blocked by a tumor or by scar tissue that forms as part of the healing process.
Fiducial placement for Stereotactic Body Radiation (SBRT)

Some tumors cannot be treated through traditional surgery, but may respond very well to stereotactic radiosurgery. To make sure that SBRT is delivered to the exact location of the tumor, Covidien’s SuperLock™ fiducial markers are placed in or near the tumor and can be placed at the same time as your ENB™ procedure for biopsy using Covidien’s superDimension™ system. A fiducial marker is simply a small gold seed or platinum coil that is placed around a tumor to act as a radiologic landmark.

High dose rate (HDR) brachytherapy, also called Image-Guided Brachytherapy (IGBT)

Using some sort of radiology tool or x-ray, your pulmonologist will place a catheter into the lung tumor and deliver high dose radiation by passing radioactive seeds through the catheter. This technique minimizes damage to lung tissue and delivers higher dose of radiation to the tumor; as a result, more cancer cells are destroyed.

Airway Stenting

Airway stents are small, expandable tubes that your pulmonologist may use to open bronchial tubes (airways) that are occluded or narrowed due to the lung tumor or scar tissue. Some covered stents can also be used to prevent the cancer from growing back into the breathing tube and compromising lung function.
Pleuroscopy
When a laparoscopy is performed in the chest, it is called pleuroscopy or medical thoracoscopy. A small instrument with a camera is inserted into the chest cavity through a very small incision, enabling the pulmonologist to perform diagnostic and therapeutic procedures inside the chest.

Balloon bronchoplasty
A balloon bronchoplasty is a technique that your pulmonologist may use to open a narrow airway using a balloon. It is very similar to how coronary arteries are opened during a heart angioplasty. Bronchoplasty is particularly useful when an airway is narrowed because of scarring after a tracheotomy, for example. Depending on the location of the narrowing, the dilation (or widening) of the airway can be performed using a flexible or rigid bronchoscope. It can also be done prior to stent placement.

OTHER TREATMENT OPTIONS

Photodynamic therapy (PDT)
Photodynamic therapy is a cancer treatment that uses a drug called a photosensitizer or porfimer sodium (brand name Photofrin®) and a certain type of light to kill cancer cells. After it is injected into a vein, the photosensitizer drug is exposed to certain wavelengths of light and becomes active. This activation of the photosensitizer produces a certain kind of oxygen that kills the tumor and nearby cells or blood vessels that are feeding the tumor. The photosensitizer may also activate your immune system to destroy the tumor cells.

PDT is usually only used on small tumors since the light that is used cannot pass through bigger tumors. Your doctor may use PDT to relieve the symptoms of a non-small lung cancer that is blocking your airway. To do this, the doctor will use
a bronchoscope to shine the light on the tumor. Your doctor may use Photodynamic therapy along with other therapies like chemotherapy and/or radiation.

PDT is performed at only select academic centers in the United States. Generally, a pulmonologist or interventional radiologist performs the procedure though, occasionally, a surgeon will be certified to do so.

The Food and Drug Administration (FDA) has approved PDT for the treatment of non-small cell lung cancer when the tumor cannot be treated using other treatment options. The FDA also approved PDT to relieve symptoms caused by these tumors when they block airways in the lung.

**Advantages of PDT:**

- Causes little damage to healthy tissue
- Is less invasive than surgery to remove tumors
- Can be done on an outpatient basis
- Provides targeted therapy directly to the tumor

**Disadvantages of PDT:**

- Cannot treat very large tumors or tumors in body cavities because the light used in PDT can only pass through a small amount of tissue
- Generally, PDT cannot be used for tumors that have metastasized or spread
- to other areas
What to expect with PDT:

- PDT is usually performed during an outpatient visit or short hospital stay
- A member of your healthcare team will inject the photosensitizer drug 24 to 72 hours before the procedure
- The drug will be absorbed by all cells but stays in cancer cells longer than normal cells
- After the photosensitizer has left most of the normal cells, the tumor is exposed to the special light to activate the drug and kill the tumor cells

Side effects of PDT:

- Porfimer sodium may make your skin and eyes sensitive to light for about 6 weeks after injection. You should avoid direct sunlight during the time of your treatments.
- The treatment may cause burns, swelling, pain, and scarring in otherwise healthy tissue.
- PDT may cause temporary side effects, including coughing, trouble swallowing, stomach pain, painful breathing, or shortness of breath.

To learn more about Photodynamic therapy for cancer visit the National Cancer Institute website at http://www.cancer.gov/cancertopics/factsheet/Therapy/photodynamic.

Vaccine therapy
Research and clinical trials are being conducted in the United States on the use of a “lung cancer vaccine.” In this treatment, the vaccine is used to stimulate antibody production. The antibodies produced by your immune system target the cancer cells and destroy them. See the Clinical Trials chapter for more information on how to find clinical trials in your area.
SUMMARY OF TREATMENT OPTIONS FOR NSCLC:
STAGES I, II, III, AND IV

The following is a summary of treatment options for individuals diagnosed with NSCLC.

**Stage 0:** Stage 0 lung cancer is lung cancer found only in the lining of your air passages. Most lung cancer is diagnosed at later stages; stage 0 lung cancer patients are usually discovered through sputum cytology. If you are diagnosed with stage 0 lung cancer, it is probably because you participated in a lung screening trial or because you are considered to be at high risk. Stage 0 lung cancer is also known as carcinoma in situ.

*Carcinoma in situ* are tumors that are present in only a few layers of cells. These tumors have not grown (or metastasized) outside the lining of your air passages however, they may progress to invasive cancer. Standard treatments may include surgical resection usually by segmentectomy or wedge resection. The goal of the treatment is to remove as little normal tissue as possible. Occasionally, if the tumor is more centrally located, your surgeon may have to do a lobectomy.

**Stage I:** If you are diagnosed with stage I lung cancer, this means that the cancer is located in one lung and has not spread to your lymph nodes or outside of the chest. At this early stage, surgery is usually the treatment of choice. Be aware that your oncologist may recommend a multi-treatment approach in which two or more types of treatment are combined. Your team will discuss with you the type of surgery they recommend and whether the addition of chemotherapy or radiation is appropriate. Talk to your oncologist about the potential risks and benefits for each treatment option.

Surgical removal of the cancer may be accomplished through various techniques including segmentectomy (removal of a small segment of the lung), lobectomy.
(removal of a lobe of the lung) or pneumonectomy (removal of the entire lung). When determining treatment, your oncologist will take into account your age and general health, as well as where the cancer is located. Your oncologist and surgeon will try to remove as little of the lung as possible in order to preserve as much lung function as possible. Your oncologist will be able to tell you if you are not a surgical candidate based on your age or on concurrent health conditions that might make surgery too risky. If you are not a good candidate for surgery, your oncologist will talk to you about using newer imaging techniques such as positron emission tomography (PET) scan that can more accurately stage your cancer so that radiation can be used.

The five-year survival rate for stage I NSCLC is approximately 60 to 80% with surgery. However, even in early stages of lung cancer, cancer cells may have spread outside the lung and may not be found. Therefore, your oncologist may recommend chemotherapy, radiation therapy, or immunotherapy before or after surgery.

**Stage II:** About 30% of lung cancer are diagnosed at this stage.

Stage II tumor is one that has been found in one lung and may be present in lymph nodes on the same side of the chest but not in the lymph nodes of the mediastinum. Your oncologist will probably identify surgery as the best first-line treatment if your age and general health are good. However, if you are diagnosed with stage II NSCLC, you may require more than one treatment type to increase the effectiveness of treatment and prevent recurrence.

Surgical options are usually the same for stage II as stage I. Surgery is the treatment of choice for patients with stage II NSCLC. A lobectomy, pneumonectomy, or segmental resection, wedge resection, or sleeve resection may be performed as appropriate. Your oncologist will do a careful assessment of your overall health to determine the risks
and benefits of surgery. For stage II NSCLC tumors, surgical removal results in 20 to 30% of patients being alive without return of the cancer within 5 years of surgery.\(^{16}\)

If it is determined that surgery may not have removed all cancer cells, your oncologist may recommend chemotherapy and/or radiation as further treatment. If your oncologist determines that you are not a good candidate for surgery, he may recommend stereotactic body radiation therapy (SBRT) to kill any remaining cancer cells.

**Stage III:** About 30% of lung cancer is diagnosed at stage IIIA or IIIB.\(^{17}\)

**Stage IIIA:** A stage IIIA tumor has extended into lymph nodes in the tracheal area outside the lung. These lymph nodes may be around the diaphragm or chest wall and will be on the same side of the body on which the cancer started. Some stage IIIA NSCLC tumors can be treated with surgery and others cannot.

If the stage IIIA tumor can be treated with surgery, your oncologist may recommend some combination of surgery, chemotherapy, radiation or a clinical trial of new treatments. Because all tumors are different, your oncologist and treatment team will decide what treatments should be done and in what order they will be most effective.

If the stage IIIA tumor cannot be treated with surgery, your oncologist may recommend some combination of chemotherapy, external radiation, internal radiation or a clinical trial of new treatments. Because all tumors are different, your oncologist and treatment team will decide what treatments should be done and in what order they will be most effective.
Stage IIIB: A stage IIIB tumor is a cancer that has extended into lymph nodes in the neck or in the opposite lung from where the cancer started. It is very common for your oncologist to provide more than one type of treatment if your tumor is a stage IIIB NSCLC. Some combination of chemotherapy, internal or external radiation therapy, surgery, immunotherapy or clinical trials may be prescribed as part of your treatment plan. Timing of each treatment will be based on your age and general health.

Stage IIIC: A stage IIIC tumor is a cancer that has extended into distant lymph nodes. In general, the tumor size is 5-7 cm or the tumor has spread to nearby organs.” Source: Reference: Rami-Porta R, Asamura H, Travis WD, Rusch VW. Lung Cancer — Major Changes in the American Joint Committee on Cancer Eighth Edition Cancer Staging Manual. CA Cancer J Clin. 2017;67:138–155.

Stage IV: About 40% of NSCLC is diagnosed at stage IV.18

If you are diagnosed with a stage IV NSCLC tumor, the cancer has spread into both lungs or more distant parts of the body. A diagnosis of a stage IV tumor must include one or more of the following:

- There is at least one tumor in each lung;
- Cancer cells are found in fluid around the lungs or the heart;
- Cancer has spread to other parts of the body.

Again, your individual treatment plan will be developed based on your age and general health. Treatment options for stage IV NSCLC may include radiation therapy, chemotherapy, immunotherapy, and targeted therapy. Radiation therapy is used mainly for pain control rather than an intent to cure. Treatment options may include
combinations of chemotherapy, immunotherapy, or targeted therapy, (see Biomarker testing section), external beam radiation therapy for local tumor growth (see Radiation Therapy section), or brachytherapy if you have tumors that obstruct your airway (see Radiation Therapy section). New drugs and combinations of treatments are being studied and a clinical trial may be available to you.

Stage IV disease complications

Bone metastases

Palliative radiation
Often stage IV patients have tumors in their bones, or bone metastases. Many times these bone metastases result in pain, decreased ability to move, anemia, bone fractures and in some cases, if near the spine, paralysis. Treatment for these tumors is usually radiation therapy for several days to relieve pain and shrink the tumor. Chemotherapy can also shrink bone metastases.

Brittle bones
Chemotherapy and complications from lung cancer can lead to brittle bones, or osteoporosis. Your oncologist may prescribe one of a number of drugs to decrease your chance of developing brittle bones. Ask your oncologist if one of these medicines might be right for you.

Monthly infusions of zoledronic acid (Zometa®) or subcutaneous injections of denosumab (Xgeva®) are used in patients with bone metastases to prevent new bone lesions from forming and to help heal existing bone lesions.

Note: Before taking one of these medicines, your oncologist may recommend supplements to improve your levels of calcium and vitamin D. Talk to your
oncologist about having dental work done before starting any of the medications for brittle bones since the drugs normally given may cause breakdown of the jaw bone resulting in loose teeth, swelling and infection of the jaw and gums, and loss of gum tissue. Be sure your dentist knows that you are taking (or will be taking) one of the drugs for brittle bones.

Wasting syndrome, or cachexia
Wasting syndrome is the loss of body mass that cannot be reversed by eating correctly. This syndrome may cause weight loss, muscle wasting, also known as atrophy, extreme fatigue and weakness, and loss of appetite. If you develop wasting syndrome, you may not be able to tolerate treatments as well so it is important that your team treat this syndrome aggressively. If you develop wasting syndrome, your oncologist may prescribe steroids. A few drugs in clinical development and available via clinical trials may prevent wasting syndrome when given with first-line chemotherapy. Ask your physician if participating in a clinical trial for wasting syndrome might be right for you.

Need for oxygen
For many different reasons, a lung cancer patient may experience the need for oxygen: flying, traveling to a high altitude location, symptoms from fluid build-up in the lung, removal of the lobe of a lung or the entire lung itself or other complications. You may have your physician order an oxygen tank for your use at home and when you travel.

Pneumonia
Lung cancer can, for many reasons, weaken your immune system, putting you at risk for pneumonia. Pneumonia is an infection of the lung. It is important to go to your doctor to be checked for pneumonia if you experience a continuing or worsening cough, chest pain, difficulty breathing or fever. You may be required to stay in the hospital to
receive intravenous (in the vein) antibiotics, or you may be able to be treated at home with oral antibiotics. Pneumonia needs to be treated to avoid more serious issues with breathing and circulatory problems.

**Fluid in or around the lungs, or pleural effusion**

This fluid build-up often contains cancer cells. It causes coughing and can cause severe shortness of breath. It may require a surgical procedure, called *pleurodesis*, to essentially ‘glue’ the lung to its lining to keep fluid build-up from happening. This procedure involves inserting a chest tube to insert chemicals to induce a scar, thus ‘gluing’ the lung to its lining. The chest tube must remain in for a few days at least until the fluid has completely drained out of the lung. Another option is to have a drainage catheter (tube) inserted into the lung for about 30 days. Each day the patient or a caregiver connects the catheter to a simple vacuum tube that drains the collecting fluid. When not in use there is a cap placed on the tube. This is a good option for patients with pleural effusions to be at home and to continue receiving chemotherapy, if indicated.

**Embolism**

Cancer can make your blood thicker than usual and this can lead to blood clots. When a blood clot travels through the blood stream into your lung, it is called a pulmonary embolism. This is similar to a blood clot that becomes lodged in your leg resulting in a deep vein thrombosis.

Symptoms of a pulmonary embolus include sudden shortness of breath, chest pain and coughing up blood. Symptoms of deep vein thrombosis include swelling or severe pain in your leg. Both of these conditions can be treated once they are identified, so if you experience any of these symptoms, you should contact your oncologist immediately so that you can be evaluated and treatment can be started.
When you are diagnosed with cancer, the most important thing that you can arm yourself with is knowledge and education. The GO2 Foundation for Lung Cancer provides the tools to arm yourself, through the Patient Education Handbook, the website, in person support from staff. The foundation has been a game changer for me.

—Jim Brown, survivor
TARGETED THERAPY
I was diagnosed at Stage IV when I was 28-years old in 2012. I am now NED (No Evidence of Disease). This Foundation was integral in the success of my cancer journey, and I am so grateful to them. During treatment, I promised myself that I would beat this, and then get to work helping patients the way the Foundation had helped me. I was also able to look beyond illness to my dream raising a family. I am now the mother of two baby girls.

—Emily Bennett Taylor, survivor
TARGETED THERAPIES

Targeted therapy and personalized medicine
Your oncologist may prescribe “targeted therapy” if your lung cancer is diagnosed as stages III or IV NSCLC. You may also receive these therapies after surgery as maintenance therapy.

What are targeted therapies?
Targeted therapy is a term you might hear that describes a type of lung cancer treatment that uses drugs to identify and attack cancer cells specifically, while sparing normal cells. Cancer cells can create “mutant proteins” and other genomic abnormalities, such as fusions that cause two unrelated genes to ‘fuse’. These mutant proteins and fused genes are what cause cancer cells to grow, divide and spread, and are therefore good targets for drugs that act like “guided missiles” to attack only these “mutant” or abnormal proteins and genes. Although targeted therapies have side effects, they are generally better tolerated than chemotherapy.

Why are targeted therapies important?
Because the genomic alterations driving a cancer, such as protein mutations and gene fusions, in each tumor are different, the treatments for each tumor will be different. These personalized treatments recognize that what works for one type of lung cancer may not work for another. Targeted therapies are a relatively new line of research and treatment for lung cancer. If your oncologist is not familiar with biomarker testing and targeted therapies, it is acceptable and advisable to get a second opinion about your treatment options.

What targeted therapies are available?
Although many gene mutations have been identified in lung cancer tumors, many of the mutations do not have effective targeted therapies identified yet (research is ongoing to find treatments for all of these genomic
alterations, with many promising new experimental drugs in clinical trials). Currently, four known mutations have effective targeted therapies available, and approved by the U.S. Food and Drug Administration (FDA). These alterations are $EGFR$ mutations, $EGFR$ T790M, ALK fusions, and ROS1 fusions. If your tumor tests positive for alterations outside of these four, ask your doctor for clinical trials as a treatment option.

- **Epidermal Growth Factor Receptor (EGFR):**
  About 15% of patients diagnosed with non-small cell lung cancer have mutations in the $EGFR$ gene. Biomarker testing for alteration in $EGFR$ may tell whether certain types of targeted drugs called tyrosine kinase inhibitors (TKIs) would be beneficial in treating your lung cancer.

If your tumor is $EGFR$ mutation negative or “wildtype” (that is, an $EGFR$ mutation is not found in the tumor), your oncologist may still prescribe a TKI since the drug may slow cancer growth. Typically, in the case of $EGFR$ mutation negative tumors, TKI’s will be used as a second-line treatment after chemotherapy.

- **$EGFR$ T790M:** T790M is a point mutation in the $EGFR$ gene that is associated with resistance to epidermal growth factor receptor ($EGFR$)-directed targeted therapies such as erlotinib. If you have an $EGFR$ positive lung cancer and have become resistant to the drugs that target the $EGFR$ mutation, you may have the T790M “resistance” mutation. Approximately 60% of all patients that stop responding to $EGFR$-directed targeted therapies over time, become resistant to therapies like erlotinib because their cancers have evolved the T790M mutation to bypass resistance.
treatment and continue to grow. Both tissue and blood tests have been approved as diagnostic tests to look for EGFR T790M. If you test positive for EGFR T790M, a drug called Tagrisso™ (osimertinib) has been approved to target this mutation.

• **ROS1 rearrangements:** About 1% to 2% of individuals with non-small cell lung cancer have an abnormality in which the ROS1 gene is fused to part of another gene. These are called, “ROS1 translocations” or “ROS1 fusions.” The ROS1 gene makes a protein called ROS, which is found within the membrane of human cells. There are several approved therapies for targeting ROS1 fusion-positive lung cancer.

• **Anaplastic Lymphoma Kinase (ALK):** About 5% of non-small cell lung cancers are driven by fusions of the ALK gene. The ALK fusion or re-arrangements produce an abnormal ALK protein that causes the cells to spread and grow. Targeted drug therapies have been approved in treating ALK positive lung cancer.

A great deal of research is being done about mutations of proteins and genes that might cause lung cancer.

The following list was created by GO2 Foundation for Lung Cancer with assistance from our contributing authors:
TABLE OF CONTENTS

TARGETED THERAPIES

DRUGS THAT TARGET

<table>
<thead>
<tr>
<th>E G F R</th>
<th>GILOTRIF (afatinib)</th>
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<td>IRESSA (gefitinib)</td>
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<td>TAGRISSO (osimertinib)</td>
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<td>PORTRAZZA (necitumab)</td>
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Some of the most common changes in NSCLC are changes in the EGFR gene. Gilotrif, Iressa, Tagrisso, Tarceva and Vizimpro are oral drugs that can all be prescribed as first treatment. They are used to target many different types of EGFR changes.

You should talk with your treatment team about which one of these drugs is best for you. All of them target the most common EGFR changes, but some of the drugs can also target less common changes in the EGFR gene. For example, Gilotrif is approved for the EGFR S768I, L861Q and G719X changes and Tagrisso can target EGFR with the T790M change.

Portrazza targets EGFR and is given by vein. It is approved for a type of NSCLC called squamous cell. It is given with chemotherapy.

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<th>A L K</th>
<th>ALECENSA (alectinib)</th>
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<td>XALKORI (crizotinib)</td>
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<td>ZYKADIA (ceritinib)</td>
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Changes involving the gene ALK can also be found in NSCLC. These oral drugs are approved to treat such “ALK positive” lung cancer when it has spread. Alecensa, Alunbrig, Xalkori and Zykadia are all approved as first treatments, but can also be given as later treatments.

Lorbrena is approved for use after the cancer has stopped responding to at least one prior ALK treatment.

Importantly, some of these drugs work to treat cancer in the brain and spinal cord better. Some are better if your cancer has certain changes in ALK. Some may also have fewer side effects. It is important to talk with your healthcare team about which therapy is best for you.

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<th>R O S 1</th>
<th>XALKORI (crizotinib)</th>
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<td>ROZLYTREK (entrectinib)</td>
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Changes in the ROS1 gene also occur in NSCLC. Xalkori and Rozlytrek also target changes in this gene and are approved for treatment only when the cancer has spread.

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<th>B R A F</th>
<th>TAFLINAR (dabrafenib)</th>
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<td>WITH MEKINIST (trametinib)</td>
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BRAF is another gene that can change in NSCLC. These two oral drugs are approved for NSCLC with a change in BRAF called V600E. The drugs are approved only for cancer that has spread.
<table>
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<th>DRUGS THAT TARGET cont.</th>
<th>Description</th>
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| **RET**                | RETEVMO (selpercatinib)  
Changes in the RET gene can be found in 1-2% of all NSCLCs. Retevmo targets changes in this gene and has been approved for cancers that have spread with changes in RET. |
| **MET**                | TABRECTA (capmatinib)  
Changes in the MET gene can be found in 1-3% of all NSCLCs. Tabrecta targets changes in this gene and has been approved for cancers that have spread with changes in MET. In addition to Tabrecta being approved, there are several drugs currently in clinical trials for cancers with changes in MET. |
| **NTRK**               | VITRAKVI (larotrectinib)  
Changes involving the NTRK genes, known as NTRK gene fusions, can be found in less than 1% of all NSCLCs.  
NTRK gene fusions can produce abnormal TRK proteins which can cause cancer to develop and grow in sites throughout the body. There can be gene fusions of NTRK1, NRTK2, or NTRK3.  
Vitrakvi and Rozlytrek are oral ‘TRK inhibitors’ that work by blocking signals that the abnormal TRK proteins send to cells to grow and divide uncontrollably. By blocking this signal, Vitrakvi stops the growth and spread of the cancer cells. |
| Other targeted therapies | AVASTIN (bevacizumab)  
Some targeted therapies work in a different way. These drugs target proteins that cause blood vessel growth. The goal is to “starve” the cancer by cutting off the blood supply that feeds it. Unlike other targeted therapies, these therapies do not need biomarker testing.  
The VEGF protein family causes blood vessel growth around cancers. Avastin targets VEGF-A. It can be taken in combination with chemotherapy and as a first treatment. It is given by vein.  
Mvasi is nearly the same therapy as Avastin. It is called a biosimilar.  
Cyramza targets VEGFR2. It can be used after the cancer has grown during prior treatment. It is given together with chemotherapy or Tarceva (for patients with changes in the EGFR gene). It is given by vein. |
| MVASI (bevacizumab-awwb) |
| CYRAMZA (ramucirumab)  |
TARGETED THERAPIES

Biomarker testing will help your oncologist determine if the lung tumor is one of the seven well-known genetic alterations that have an approved targeted drug (EGFR, ROS1, ALK, BRAF, RET, MET, NTRK). If the testing is negative on all seven genes, your oncologist may be able to enroll you in a clinical trial for another targeted therapy or may elect to treat you with other more traditional therapies. Much research is being done on other genetic mutations that may be treatable someday.

The use of Next Generation Sequencing (NGS) also called Comprehensive Biomarker Testing (CBT) is a deep dive look into the molecular makeup of your cancer regardless of cancer type and may help your doctor determine which clinical trial might be best for you. By the time you are reading this guidebook, researchers may have identified additional mutations. Please contact GO2 Foundation for Lung Cancer at 1.800.298.2436 or visit lungmatch.org for more information and an updated list of molecular mutations.
IMMUNOTHERAPY
IMMUNOTHERAPY FOR THE TREATMENT OF SMALL CELL AND NON SMALL CELL LUNG CANCER

What is the immune system and how does it work?

- The immune system is a collection of unique cells and substances they produce, that act as the body’s defense mechanism against infections and anything ‘foreign.’
- Immune cells travel through the body and keep track of all cells and substances normally found in the body. These cells are trained to recognize pathogens like bacteria, viruses, etc. and abnormal cells in the body as ‘foreign’ and eliminate them.
- This process of recognition and elimination is based on the presence of molecules (such as proteins) on the surface of all cells that the immune cells use to distinguish between ‘self’ and foreign.

What is Cancer Immunotherapy?

- Immunotherapy is a treatment modality that employs several different tricks to stimulate the patient’s own immune system to fight their cancer.
- Cancer cells have devised unique ways to evade surveillance and elimination by the immune system by cloaking themselves to appear as normal cells.
- Immunotherapy aims to either specifically un-‘cloak’ these cancer cells and expose them to the immune system, OR, train the immune system to fight harder and smarter in a general, non-specific fashion.
- Immunotherapy holds great potential for treating cancer, as no other therapy can compare with the elaborate network of cellular interactions and pathways employed by the human body to rid itself of foreign entities.
Immunotherapy has several advantages over chemotherapy and targeted therapy:

- Immunotherapy has demonstrated a low toxicity profile relative to chemotherapy and targeted therapy.
- Because the biological system is sensitive to even very minor alterations, the immune system can detect relatively low numbers of cancer cells and mount a response to eliminate them.
- The immune system has a strong ‘memory,’ in that it remembers the foreign cells it was exposed to and each time it encounters those cells again, it gets activated and works to eliminate them. This immune memory bestows longer lasting tumor control, as against chemotherapy and targeted therapy that need to be constantly replenished in the body. Since immune responses stimulated by immunotherapy, once generated, are always remembered
by the body and activated each time cancer recurs, this therapeutic modality produces *durable, sustained tumor response*.

**What are the various types of immunotherapy effective for lung cancer?**

There are currently three types of immunotherapies that are being evaluated to treat cancer, listed below.

1. Immune modulators such as Immune Checkpoint Inhibitors
2. Cancer Vaccines
3. Adoptive T cell Transfer

Five drugs belonging to the first category of Immune Checkpoint Inhibitors are now FDA-approved for use by patients with unresectable Stage III and advanced, metastatic non-small cell lung cancer patients. These drugs are Durvalumab (IMFINZI, manufactured by AstraZeneca), Nivolumab (Opdivo, manufactured by Bristol Myers Squibb), Pembrolizumab (Keytruda, manufactured by Merck) and Atezolizumab (Tecentriq, manufactured by Genentech), Yervoy (Ipilimumab, manufactured by Bristol Myers Squibb).

**What are Immune Checkpoint Inhibitors?**

- The main role of the immune system is to keep track of what is ‘self’ and, identify and eliminate anything that is ‘foreign.’
- In order to prevent the immune system from attacking its own normal ‘self’ cells, the body has evolved several checks and balances that keep the immune system under control.
- These checks and balances are like the brakes in a car that prevent the car from going into overdrive, and are designed to prevent or abort actions that could be self-destructive.
• A breach in these systems results in the immune system recognizing normal cells as ‘non-self’ and eliminating them, resulting in auto-immune diseases like lupus and arthritis.
• Cancer cells have evolved ways and means to overtake these normal checkpoints, locally block the immune response in the vicinity of the tumor, and effectively escape detection and elimination by the immune system.
• Two immune checkpoints that have been targeted recently to lift the brakes from the immune system so that it goes full force to attack cancer cells are CTLA4 and PD-1/PDL1.
• Checkpoint inhibitors basically undo the local blockade of the immune response evoked by cancer cells and allow the immune system to resume normal function and attack the tumor.

How do PD-1/PDL1 checkpoint inhibitors work?

• PD-1 stands for Programmed Death receptor-1.
• It is a protein expressed on the surface of immune cells, specifically T cells, a kind of white blood cells that fight infection and other foreign bodies.
• PD-1 interacts with a protein on the surface of normal cells of the body, PDL1 (Programmed Death Ligand 1).
• This PD1-PDL1 interaction is an immune checkpoint, i.e. a signal to the immune system to not mount an attack on the body’s own cells.
• Cancer cells usurp this mechanism and express PDL1 on their surface to fool the immune system into believing that they are normal cells.
• Therefore, blocking the PD-1/ PDL1 interaction is the target for anti-cancer immunotherapy, as PD-1 or PDL1 inhibitors allow the immune system to then recognize the cancer cells as foreign and eliminate them.
• It is encouraging that lung cancer cells express PDL1 on their surface and are therefore amenable to PD-1 and PDL1 blockade.
• PD-1/ PDL1 checkpoint inhibitors are molecules that bind to either PD-1 (expressed on immune cells) or PDL1 (expressed on cancer cells) and block the surface of these proteins, preventing them from interacting with each other.
• The tolerability of these immune checkpoint inhibitors has been generally good, with few dose-limiting toxicities reported. The most common irAEs or immune related Adverse Events reported are: dermatologic (rash, pruritus, and vitiligo), gastrointestinal (diarrhea and colitis), endocrine (hypothyroidism and hyperthyroidism), and hepatic (hepatitis and increased liver function enzymes) events, as well as pneumonitis, uveitis, infusion-related events, and fatigue.
• A Phase III clinical trial assessing the efficacy of an antibody that blocks PD-L1 found that the treatment improved progression-free survival by more than 11 months compared to a placebo in patients with unresectable Stage III NSCLC.
• PD-1 vs. PDL1: Although the response rates are similar for the inhibitors that bind to PD-1 on immune cells and those that block PD-L1 on tumor cells, early data suggest that there may be a slight safety advantage in targeting PD-L1. A phase I trial of a PD-1 inhibitor reported a 3% incidence of drug-related pneumonitis (inflammation of the lung tissue) but this side effect has so far been less severe or absent with the PD-L1 inhibitors.
• Effects in smokers vs. non-smokers: Early results suggest that both kinds of inhibitors-anti-PD1 and anti-PDL1- seem to benefit smokers more than non-smokers. Results of a phase I trial of a PD-L1 inhibitor presented at the 2013 European Cancer Congress indicated that 26% of smokers responded to the drug, but only 10% of never-smokers responded. Researchers speculate that this is probably due to the greater number of mutations present in smokers’ tumors, an abundance that would probably present the newly awakened immune response with a far greater array of tumor antigens to respond to, and mount a response against.
Which immunotherapeutic drugs are currently approved for the treatment of lung cancer?

Currently, five immunotherapy drugs are FDA-approved for the treatment of patients with advanced, metastatic non-small cell lung cancer. For more information, contact our GO2 Foundation treatment specialists at 1.800.298.2436 or visit lungmatch.org.

What are the side effects of treatment with Immunotherapeutic drugs?

Most common side effects observed in the clinical trials that evaluated these four immunotherapeutic agents for lung cancer patients were fatigue, decreased appetite, dyspnea, cough and pneumonitis. However, it is important to note that immunotherapeutic agents may be associated with immune-mediated side effects in the lungs, colon and hormone-producing glands. Immune-mediated adverse reactions observed with these drugs in clinical trials have included pneumonitis, colitis, hepatitis, hypophysitis, hyperthyroidism, hypothyroidism, type 1 diabetes mellitus, and nephritis. Based on the severity of the adverse reaction, these immunotherapeutic drugs should either be withheld or discontinued and corticosteroids should be administered.
What is CTLA-4?

CTLA-4 (Cytotoxic T-Lymphocyte Associated protein 4) is one well known immune checkpoint protein. Certain drugs are currently approved for lung cancer and may work particularly well when used with PD-L1 checkpoint inhibitors.

What is Combination Therapy?

- Combination therapy is combining one or more different therapeutics for increased efficacy and tumor shrinkage, such that the effects of the combination are greater than the effects produced by sum of the parts.
- Combining two different therapeutics may be sequential (one after the other) or concurrent (both therapies administered together).
- Studies are underway to understand if and how immunotherapy may be combined with chemotherapy and/or radiotherapy. These studies are based on the hypotheses that the antigens released from dying cancer cells upon effective chemotherapy, may serve to stimulate the immune system, mount a tumor-specific immune response, and thereby enhance the efficacy of the immunotherapeutic.
- A combination of the immune checkpoint inhibitor Ipilimumab with chemotherapy has shown encouraging results in both small cell and non-small cell lung cancer.
- Combinations of anti-PD1/PD-L1 and CTLA4 immunotherapeutic agents have been approved for lung cancer patients.
What are cancer vaccines?

- A vaccine is typically a biological agent used to stimulate and train the immune system to recognize this agent as ‘foreign’, mount a response to eliminate it from the body, and create ‘memory’ such that if the agent is encountered again, the body readily clears it from the system.
- Vaccines may be either prophylactic (they prevent future infection by the agent) or therapeutic (they treat current infections).
- Cancer vaccines are therapeutic. These vaccines use proteins expressed on the surface of cancer cells to train the immune system to recognize tumors and destroy them.
- So far, there is only one FDA-approved vaccine for cancer: Provenge approved for the treatment of advanced prostate cancer in April 2010.
- There is excitement around the potential use of cancer vaccines for lung cancer as lung tumors over-express specific proteins such as MAGE-3 (overexpressed in 42% of all lung cancers, 35% early stage and 55% late stage NSCLCs), NY-ESO-1 (overexpressed in 30% of all lung cancers), p53 (overexpressed in 50% of lung cancers), survivin, MUC-1, etc., which can serve as agents to train the immune system to recognize these proteins on cancer cells and specifically kill those cells.
What is Adoptive T Cell Transfer?

- The third major type of immunotherapy currently being evaluated for lung cancer is adoptive T Cell transfer which is a process that involves 1) removing a patient’s immune cells, specifically the T cells from their body, 2) treating these cells with various chemicals and other biological factors in a lab dish such that they recognize antigens on tumors and mount a strong immune response, and 3) re-inject these activated immune cells back into the patient’s body.

Patient response to cancer immunotherapy

- One of the challenges of immunotherapy for lung cancer is the variability in patient response: while some patients see very durable and lasting responses, others only get a partial response to the therapy and progress, while others see no response at all.
- Studies are underway to understand the underlying reasons for these differences in response to immunotherapy in lung cancer patients. These studies will hopefully uncover biomarkers of response to these therapies that can be used to better select patients who are most likely to respond, while sparing toxicity and side effects in those unlikely to respond, thereby allow tailoring therapy to patients based on the specifics of their cancer.
- Since immunotherapies are designed to stimulate the immune system, these agents are not suited for patients that have a history of autoimmune disorders or previous immunosuppressive therapy.
SMALL CELL LUNG CANCER TREATMENTS
Educated and empowered patients do much better.

—Bonnie J. Addario, survivor
Overview

Small cell lung cancer (SCLC) is the other type of lung cancer. Although it is less common than non-small cell lung cancer (NSCLC), SCLC grows and spreads through the body early in the disease – sometimes before any symptoms are noticed. Of all lung cancers, only about 10 to 15% are SCLC and almost all of these cases are found in people who currently smoke or have smoked cigarettes.² Because of the connection with cigarette smoking, SCLC is a little more common in men than in women. In addition, SCLC is usually described as limited or extensive.

SCLC is called “limited stage” when the total area involved by the disease can be targeted with one radiation field. This means that small cell lung cancer can still be limited stage if it moves to lymph nodes in the middle of the chest known as the mediastinum. Most importantly, having limited stage disease means you can be treated with curative intent. “Extensive stage” SCLC is cancer that has spread outside of one radiation field and usually means the disease cannot be cured, only controlled for a period of time.

SCLC typically starts in the bronchi (large breathing tubes) located behind the breastbone in the middle of the chest. True to the name, the cells in SCLC are smaller than the cells in NSCLC; however, because these cells grow very quickly, the tumors they create can be larger than NSCLC tumors. This type of lung cancer also tends to metastasize rapidly, or spread to other areas of the body, such as the brain, liver, or bones faster than NSCLC in most cases.
How is SCLC treated?

Chemotherapy is the main treatment for small cell lung cancers. Since SCLC may spread before you notice any symptoms, removing the lung tumor by surgery rarely cures the cancer. Even when surgery is used to treat SCLC, it is never the only treatment you will receive. Laser treatments and experimental treatments available in a clinical trial may also be used to treat SCLC.

**Surgery**
Surgery is rarely used to treat SCLC and if it is, it is rarely the only treatment since the cancer has typically spread before it is diagnosed. Your thoracic surgeon may also use one of the surgical techniques, previously described, to obtain tissue to determine the type of cancer and how far it has spread.

**Chemotherapy**
Since SCLC tends to travel outside of the lung, chemotherapy treatments are designed to kill cancer cells that have metastasized into other areas of the body. Taken by mouth or injected into a vein, there are many different types of chemotherapy that your oncologist may prescribe for you. Most often, a platinum-based drug such as cisplatin or carboplatin is coupled with etoposide, which has been found to be most effective treatment for limited or extensive stage small cell lung cancer.

Additional chemotherapy drugs that have been approved for treatment of small cell lung cancer are included in the following table:
## Small Cell Lung Cancer - Treatments

### Chemotherapies

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<tr>
<th>Brand Name</th>
<th>Generic Name</th>
<th>Cancer Type</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abitrexate, Folez, Folex PFS, Methotrex-ate LPF, Mexate, Mexate-AQ</td>
<td>methotrexate</td>
<td>SCLC</td>
</tr>
<tr>
<td>Adriamycin</td>
<td>doxorubicin</td>
<td>SCLC</td>
</tr>
<tr>
<td>Camptosar, CPT-11</td>
<td>irinotecan</td>
<td>SCLC</td>
</tr>
<tr>
<td>Hycamtin</td>
<td>topotecan hydrochloride</td>
<td>SCLC</td>
</tr>
<tr>
<td>Paraplat, Paraplatin</td>
<td>carboplatin</td>
<td>SCLC</td>
</tr>
<tr>
<td>Platinol, Platinol AQ</td>
<td>cisplatin</td>
<td>SCLC</td>
</tr>
<tr>
<td>Toposar, VePesid</td>
<td>etoposide</td>
<td>SCLC</td>
</tr>
<tr>
<td>Zepsyre</td>
<td>lurbinectedin</td>
<td>SCLC</td>
</tr>
<tr>
<td>Afinitor</td>
<td>everolimus</td>
<td>NET</td>
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### Immunotherapies

<table>
<thead>
<tr>
<th>Brand Name</th>
<th>Generic Name</th>
<th>Cancer Type</th>
</tr>
</thead>
<tbody>
<tr>
<td>Imfinzi</td>
<td>durvalumab</td>
<td>PD-L1 SCLC</td>
</tr>
<tr>
<td>Keytruda</td>
<td>pembrolizumab</td>
<td>PD-1 SCLC</td>
</tr>
<tr>
<td>Opdivo</td>
<td>nivolumab</td>
<td>PD-1 SCLC</td>
</tr>
<tr>
<td>Tecentriq</td>
<td>atezolizumab</td>
<td>PD-L1 SCLC</td>
</tr>
</tbody>
</table>

### Targeted Therapies

Not currently available
Radiation Therapy
For SCLC, your oncologist may prescribe radiation treatments. Treatment with radiation may also help to relieve symptoms such as breathing problems. Your team may use many different types of radiation therapies to treat your SCLC. Radiation treatments are usually used in a treatment plan with chemotherapy.

Treatment for limited SCLC
If you are diagnosed with limited SCLC, the first option might be surgery if the tumor is small. However, it is more likely that you will be started on a combination of chemotherapy and radiation therapy.

About 50% of people with SCLC will develop metastases to the brain during their cancer journey. Your oncologist may also prescribe prophylactic cranial irradiation (PCI) to prevent spread of the cancer to your brain. PCI is a kind of radiation treatment that may be used to kill cancer cells in the brain that may not be visible on x-rays or scans.

Treatment for extensive SCLC
If you are diagnosed with extensive SCLC, chemotherapy will usually be the first-line of treatment prescribed by your oncologist. If the tumor shrinks, your doctor will usually prescribe prophylactic cranial irradiation (PCI) treatments.

Monthly infusions of zoledronic acid (Zometa®) or subcutaneous injections of denosumab (Xgeva®) are used in patients with bone metastases to prevent new bone lesions from forming and to help heal existing bone lesions.
to prevent metastasis to the brain. PCI is a kind of radiation treatment that may be used to kill cancer cells in the brain that may not be visible on x-rays or scans. Your oncologist may recommend a clinical trial as a course of treatment. Clinical trials are studies that have shown enough promise that they are now being done on humans. See the Clinical Trials chapter for more information on how to find clinical trials in your area.

**Treatment for recurrent SCLC**

Even with aggressive treatment, small cell lung cancer may come back or recur. It is a type of cancer that responds extremely well to radiation and chemotherapy in most cases. The problem is that the responses generally do not last very long and are not “durable.”

When the diagnosis of SCLC is made, you should discuss with your healthcare team the treatment plan you prefer such as chemotherapy and/or radiation. The plan may include treatment of the disease or symptom management (see the Transitional Care Planning section in the Living with Lung Cancer chapter for further discussion of treatment plans).
We’re very fortunate to have the Foundation and it’s been our saving grace. It really has been. It is giving us hope and they keep us going every day for our mom.

—Yvette, caregiver
I really appreciate the GO2 Foundation for Lung Cancer, all the work they’ve done and the guidance they’ve given me.

—Robert, survivor
C L I N I C A L  T R I A L S

What are Clinical Trials?
“A clinical trial provides the means by which your doctors can evaluate an important scientific question relating to your cancer. In most cases, the question of interest is whether a new drug or novel treatment approach is better than an existing treatment or at least worthy of further evaluation.” Paul Hesketh, MD, Lahey Clinical Medical Center. A clinical trial is a research study that has progressed from a scientific question through laboratory testing and is now ready for human volunteers. Clinical trials are critical to the development of new lung cancer treatments, ways to ease the symptoms of lung cancer treatments, and collect tumor or blood samples for research. These new treatments may include drugs, surgical procedures, and new ways to manage side effects. The clinical trials process is overseen by the Food and Drug Administration (FDA), a local institutional review board (also known as an ethics committee), and a physician specifically trained to manage clinical trials.

A clinical trial may be referred to as a “research study,” “study,” or “trial.” The team that manages the clinical trial is often referred to as the “clinical trials team,”

Questions to ask your oncologist and healthcare team about the clinical trial you are considering:

- What do you hope to learn from this clinical trial?
- Has the experimental treatment/procedure been studied before?
- What phase is this clinical trial?
- Who will be in charge of my care during the trial?
- Will my care change based upon my response to the treatment during the trial?
- What are the risks and benefits?
- How long will the trial last?
- Who pays for the trial?
- Will my insurance cover the treatment?
- Will I be paid?
- Can I be forced or asked to leave the trial?
- Can I learn the results of the trial?
“research staff,” or “study staff.” Do not let the names confuse you as they all mean the same thing.

**What types of clinical trials might be available?**

There are several types of clinical trials for which you might be eligible. Your eligibility for any trial will be based on very specific requirements, so it is important you discuss these requirements with your oncologist and the study staff. Clinical trials may be classified as:

- **Prevention trials** – Prevention trials explore factors that may increase or decrease your risk of developing lung cancer.
- **Screening trials** – Screening trials develop new and better ways to detect cancer.
- **Diagnostic trials** – Diagnostic trials develop better tests or procedures for diagnosing cancer.
- **Treatment trials** – When most people think of clinical trials, treatment trials are the ones that most often come to mind. Treatment trials evaluate specific medications, radiation treatments, and new surgical techniques to treat cancer.
- **Supportive care trials** – Supportive care trials, or quality-of-life trials, evaluate medications, radiation treatments, and new surgical techniques to decrease symptoms of cancer or the side effects of cancer treatments.

**What are Clinical Trial phases?**

In order for a new drug to be approved by the Food and Drug Administration (FDA) for use in humans, the drug must pass through a rigorous testing process. This testing process is called a clinical trial and is composed of four different phases usually referred to as phases I through IV.

Phase I trials are the first level in which the researchers evaluate safety, determine
safe amount of drug and identify side effects that might occur with the treatment. Before this phase, the treatment has already been researched at length in the lab and on animals and the drug has been determined to be ready for use in humans. The research team will adjust the amount of the treatment you receive at different intervals in the trial while monitoring the treatment’s side effects. Typically, there may be only 20 to 80 people selected to participate in a phase I clinical trial.

Phase II trials begin after a treatment has been found to be safe in phase I trials. During phase II, the research team will use a specific treatment, or combination of treatments, to determine the effectiveness for a specific type of cancer. A phase II clinical trial may include 100 to 300 people.

Phase III trials will be done when a treatment is found to be effective in phase II trials. During this phase, the treatment will be tested on a large number of patients comparing standard treatments (treatment you receive outside of the clinical trial) with the new treatment. If you participate in a phase III clinical trial, you may be randomly assigned to a control or test group. If you are assigned to the control group, you will receive the standard treatment for your specific type and stage of lung cancer. If you are assigned to the test group, you will receive the new treatment. Results from the two groups will be closely monitored by the research team to determine which treatment is most effective and the side effects of the treatment. Phase III trials include up to 3,000 patients.

Phase IV trials begin after the treatment has been approved by the FDA. In phase IV clinical trials, the treatment will be given to a much larger group of patients. In this phase, additional information will be gathered about effectiveness, side effects that might not have been previously identified, and safety issues that can only be identified in a larger group of participants.
How can I learn about the purpose, risk and benefits of a clinical trial?
Informed consent is the process of learning the facts about the clinical trial before deciding whether you want to participate. To help you decide whether to participate, the doctors and nurses involved in the trial, called the study staff, will explain the details of the trial. The study staff will provide you with an informed consent document that includes details about the trial: its purpose, length of time the clinical trial will be open, any required procedures, and the key contacts. The study staff should provide updated information to you throughout the trial.

What are the potential benefits of clinical trials?
Participating in a clinical trial may have several potential benefits for you. By participating in the trial, you will:

- Play an active role in determining the direction of your health care
- Have access to new treatments before they are widely available
- Receive expert medical care at leading health care facilities
- Help others by contributing to medical research
What are the risks of clinical trials?
Before you agree to participate in a clinical trial, you should talk to your oncologist and the doctor in charge of the trial to make sure you understand the possible risks. You should understand that the treatment being used may not be better and side effects may be worse than the standard treatment. Because the treatment is new, your healthcare team may not know all of the side effects that you will experience. A clinical trial may require more time and attention from your healthcare team and from you than a non-clinical trial treatment regimen. This extra time may include trips to the cancer center, more treatments, hospital stays and complex dosage requirements.

When do I ask my healthcare team about participating in a clinical trial?
In a study done in 1999, the American Society of Clinical Oncologists found that only 3% of adults with cancer participate in clinical trials. This low level of participation in clinical trials means that advances in cancer care do not happen as quickly as they might. Your participation in clinical trials can help to develop new cancer treatments for all cancer patients.

Any time you are facing a treatment decision, you should ask about clinical trials that might be appropriate for you. Clinical trials are not just for advanced stage lung cancer – clinical trials are available for all stages of lung cancer. Ideally, your entire healthcare team will be available to talk to you about new treatments that may be available. For example, your oncologist, radiologist, and surgeon may each have access to information about different clinical trials. Once you know about clinical trials that might be appropriate, you should discuss the options with your entire team who can help you understand the benefits and risks based on your specific lung cancer and health status.

Who takes care of me while I am in a clinical trial?
When you participate in a clinical trial, your healthcare needs and treatments will be
managed by the clinical trial doctor (who may or may not be your oncologist) and the study staff (research nurse, research coordinator, laboratory personnel). This team will manage your care throughout your participation in the clinical trial.

The clinical trial and study staff is overseen by the Institutional Review Board (IRB) at the hospital, research facility, or cancer center. The role of the IRB is to make sure the trial is safe and is being managed properly. Typically, you will find that you will receive a very high quality of care while participating in a clinical trial because the study staff will closely monitor your condition while you participate in the trial.

**How long does a clinical trial last?**
The length of clinical trials will vary based upon the research being studied. Some trials such as a tissue or blood collection trial may only involve a single visit. Other trials may last several years such as might occur in the case of a treatment trial. The informed consent form will detail the length of the clinical trial and should include how often you will be required to go to doctor visits, treatments, and follow up procedures.

Participating in a clinical trial is a commitment on your part. That said, you have the right to stop participating in a clinical trial at any time. Your clinical trial doctor may end your participation as well if the treatment is found to be unsafe, ineffective, if the clinical trial closes (research is complete) or for any other reason they deem appropriate. Be sure to understand your responsibilities in the clinical trial before you agree to participate.

**What does it cost to participate in a clinical trial?**
Clinical trials are a critical part of cancer care. Most of the time, if you enroll in a clinical trial, the cost of tests, procedures, drugs, extra doctor visits, and any research related
to the trial will be covered by the agency or company that sponsors the clinical trial. The sponsor may be a government agency, a college or university, a medical center, a non-profit organization, a drug company, or another private company.

Your health insurance plan may say that your participation in a clinical trial is “experimental” or “investigational.” In this case, your insurance may not cover the costs of routine care including doctor visits, hospital stays, and tests or treatments that you would have normally received. Many states have laws in place regarding insurance coverage for clinical trials. Ask your study staff and your insurance company about the costs before you participate.

How can I find clinical trials?
There are over 2,500 clinical trials in the US available to the lung cancer community at the time this guidebook was printed. However, not all clinical trials will be available in your area. Clinical trials may be open at only one cancer center; others may be open in hundreds of cancer centers across the country. The number of participating centers depends on the disease being studied, the phase of the clinical trial, and the complexity of the clinical trial.

If you are interested in participating in a clinical trial, there are many sources of information. The two best sources of information are:

- Your healthcare team (e.g. oncologist, radiologist, pulmonologist, etc.) – Ask your healthcare team if a clinical trial is appropriate for you at this time and what clinical trials are available at your center. If no trials are available at your center, ask your oncologist which investigation drugs or procedures might be right for you. With this information, you can search the government database for clinical trials in your area.
• U.S. National Institutes of Health (NIH) website of clinical trials located at http://ClinicalTrials.gov. There are many other internet sites with information on clinical trials, but these sites are generally built on information from the NIH website. This website lists both federally funded and privately supported clinical trials.

The NIH clinical trial list includes over 136,000 clinical trials available worldwide not just in the US. When you access the site, search for a clinical trial using the most specific information you have. For example, if your diagnosis is small cell lung cancer, search for “SCLC in the US.” A list will open showing all of the studies that are in the database. In the listing, you will be able to tell the status of the clinical trial (Completed, Recruiting, Not yet Recruiting, Active, etc.). The list will include what conditions are being targeted in the trial and what treatments are actually being tested (drug, radiation therapy, etc.). Clicking on the name of the study will open a new window that shows extensive information about the specific study including how long the trial is expected to last, eligibility requirements, how outcomes will be measured and contacts for the trial. If you find clinical trials that may be applicable to you, it is critical that you discuss them with your healthcare team.21

Searching for clinical trials may be very confusing since the resulting list may contain hundreds of possibilities. We are here to help — contact us for assistance with identifying clinical trials in your area that may be of interest to you.
HELPLINE 1.800.298.2436

Help is here...
For more information about lung cancer and current treatments, to discuss support options or for referral to other resources, please contact us: 1.800.298.2436 or support@go2foundation.org

GO2 FOUNDATION FOR LUNG CANCER
Empower Everyone. Ignore No One.
THE LUNG CANCER LIVING ROOM

BRING HOPE HOME
EVERY THIRD TUESDAY

JOIN US

Every month our Lung Cancer Living Room – Bring Hope Home – A Patient Education and Support Series group sessions BRING HOPE HOME to patients and their families. Through presentations by Lung Cancer specialists, physicians and researchers, this unrestricted forum covers all topics: early detection, treatment options, molecular and genetic testing, clinical trials, drug discoveries, personalized medicine, nutrition, surgical equipment and procedures, up-to-date news about advancements and more.

We share our personal stories, get advice and support from others, critical information from caring doctors and access to researchers who, like us, are fighting Lung Cancer 24/7.

LOCATION

Your Living Room via Facebook Live (https://www.facebook.com/GO2Foundation/) or YouTube Live (https://www.youtube.com/go2foundationforlungcancer)
LIVING WITH LUNG CANCER

Navigating Lung Cancer: 360º of Hope

THE LUNG CANCER LIVING ROOM

Partners

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LOCATION

Your Living Room via Facebook Live (https://www.facebook.com/GO2Foundation/) or YouTube Live (https://www.youtube.com/go2foundationforlungcancer)
When you are overwhelmed or don’t know where to turn, this foundation becomes a ‘home’ — a safe haven of caring people who will share their amazing knowledge and give you the courage and strength (along with faith in God) to face the journey knowing you are making the best possible choices for YOU.”

—Janice Lalley, survivor
LIVING WITH LUNG CANCER

When you are diagnosed with lung cancer and start receiving treatments, you begin to realize the many changes that are happening in your life. As your health status and treatment plans change, the care you receive will also change. During these periods of change, you may have problems moving from one phase into the next. Your healthcare team should help you move between phases by working with you to create a Transitional Care Plan.

Transitional Care Planning
Transitional Care Planning will help you find a healthy balance between your disease and the rest of your life. While you have treatments, doctor’s appointments, and days when you are not feeling your best, your family, finances, and job situation will continue to move on. You may become depressed or anxious about these issues that you just simply cannot manage right now. Transitional Care Planning can help you identify and manage these problems to minimize the impact on your treatment and healing process. Your healthcare team will help educate and support you and your family by providing support and providing referrals to resources that you may need during your care.

As your lung cancer gets better or worse, your treatment goals will change. During active treatment, you may be receiving chemotherapy, radiation, surgery, some combination therapy, or a new, experimental treatment. You will also be receiving supportive care to treat symptoms of the lung cancer and side effects from your treatments. Palliative therapy will be given to improve the quality of your life at any time in your cancer journey or to make you comfortable at the end of your life. Because each of these types of care are different, your Transitional Care Plan can help you and your family adjust by helping with the day-to-day issues, medical problems and emotional issues that will arise in each phase. And because you are a unique person, your Transitional Care
Plan will also be unique. Your healthcare team will do many assessments to determine what care is needed to make the changes you experience go smoothly.

**Transitional Care Planning Assessments**

Some kind of assessment will be done at every contact with your healthcare team. Your oncologist will examine you, the nurse will ask about side effects, general health status, appetite, and any issues you may be having in your life. The social worker and/or financial counselor will help you with any issues you might be having in your employment or financial life. As your treatments change, your team will help to identify any new needs or stresses you or your family might have. Specifically, your healthcare team will do physical, care setting, support system, spiritual and mental health, and legal assessments. These assessments may not be done at every visit, but will certainly be done when there are changes in your health status or treatment plan.

**Physical Assessment**

Throughout your treatment, but especially when you are diagnosed with lung cancer, when you are receiving treatments, or when there is a change in your treatment, you will receive regular physical assessments by various members of your healthcare team. In general, your doctors and nurses will be primarily responsible for your physical assessment. In addition to asking you questions about your symptoms and quality of life, the healthcare team will usually do a hands-on exam. This exam might include:

- Measurement of your temperature, pulse, respiratory rate, weight and blood pressure
- General examination to look for signs of infection
- Listening to your heart and lungs
- Feeling your armpits, neck, groin and other areas of your body to check for swollen lymph nodes
• Drawing blood
• Doing x-rays or other radiological procedures
• Doing pulmonary (lung) tests to determine how your lungs are functioning

Care Setting Assessment
During your treatments, you may receive treatment in many different care settings. Some of your care may occur in the hospital, but you may also receive care in an outpatient cancer center, home, a nursing home or a rehabilitation center. As your care moves from one setting to another, your healthcare team will help you plan for this change in care setting. New team members may become involved in your care; this will depend on the type of care you need at any given point in your cancer journey.

As you move from one care setting to another, your team will assess your needs and the physical arrangements in the new setting. If the team determines that you need medical equipment or assistive devices to help you move around, they will help you find those services.

Support Systems Assessment
Your healthcare team will do a full assessment of your support systems—those people and groups around you who are willing to help you during your illness. This assessment will also include a review of those people for whom you might be responsible. If you have young children or elderly parents that you care for, your team will help you determine how roles and relationships might change during your treatments.

It is sometimes very hard to ask for help. However, this is one time when you will need people around you who care about you and are willing to help. We know that...
your loved ones and friends will feel honored when you ask them to help you during this journey. At the same time, you might find that you have many friends who want to help – and you may want one person to act as the gatekeeper. Your gatekeeper can be responsible for taking phone calls, answering emails, helping schedule visits and setting up a schedule for your friends who want to be involved with you during your treatment.

What things can your support team do for you? Of course, you are the one who is in control of the help you need, but your loved ones can help with specific tasks when you just don’t feel up to doing daily chores. Duties that you might consider sharing with your team include things such as:

- Cooking meals for you and your family. Meals that can be frozen and defrosted for use at any time are particularly good.
- Babysitting. If you have young children, your friends with children may be very willing to take yours for a “play date.” This may be useful when you are having treatments and have to be out of the house for several hours.
- Driving you to appointments. Many of the treatments you might receive can make you tired; having someone drive you to and from appointments will be critical. There may be other options available for transportation to and from appointments:
  - American Cancer Society’s Road to Recovery: to find out if Road to Recovery is available in your area visit their website at http://www.cancer.org/Treatment/SupportProgramsServices/road-to-recovery or call 1-800-227-2345.
  - Cancer Care: provides free, professional support for anyone affected by cancer. To learn more about their programs visit www.cancercare.org or call 1-800-813-HOPE (4673).
—Speak with your social worker. They may be able to direct you to a local transportation program.
—Local religious organizations may have people who would be happy to help you with transportation.

• Light housekeeping. Again, treatments are likely to make you feel tired. A friend will likely be thrilled if you ask for help with vacuuming or light dusting. If you need additional help with housekeeping activities, contact Cleaning For A Reason – the non-profit organization that connects maid services with women enduring cancer treatments. To learn more call 1-877-337-3348 or visit www.cleaningforareason.org.

• Lodging while a loved one is receiving treatment away from home. If you have to travel and stay overnight for your treatments, you and your family may need a place to stay. Often, your personal support system may have a place for you to stay. If not, there may be other resources available to you:
  —American Cancer Society’s Hope Lodge: For more information on Hope Lodge visit their website at http://www.cancer.org/Treatment/SupportProgramsServices/HopeLodge/index
  —Ask your healthcare team if they know of discounted housing for out of town patients.

• Talking! You will probably want to talk to people throughout your treatment. Some of your support team will feel comfortable listening to you talk about your lung cancer; others will be great to sit and gossip with you. Both of these groups are important to you.

Anything else where you just need a friend to be with you. Remember, that your loved ones are likely to feel helpless, but they really want to help you. They will be honored that you asked them to help.
Spiritual and Mental Health Assessment

When you receive a diagnosis of lung cancer, you will very likely experience a broad range of emotions. It is very likely that your first reaction may be one of disbelief and denial – “There has to be a mistake. I can’t have lung cancer.” As you begin to accept that the diagnosis is real, you may feel angry – “This isn’t fair. What did I do to deserve this?” It is also very common to get depressed and have feelings of hopelessness during your treatment. This is particularly true when you are not feeling well and find that you cannot do those things you can usually do. Throughout your treatment, you may very well be scared. This fear may be related to the diagnosis itself, the treatments you are going through, or simply the fear of the unknown. All of these reactions are normal.

During a spiritual or mental health assessment, your healthcare team will ask you questions about how you and your family feel about your treatments and treatment plan. They may ask you about what things are most important to you since those things will affect your plan. The team may ask questions about how you and your family usually deal with stress. Are there things you have done in the past that you can use during this time? Are there other stressors in the home that will interfere with treatment and your ability to concentrate on healing?

You may look to your church, religion or spiritual beliefs to help you cope with your diagnosis and treatments. There are studies that show that spirituality may help you adjust to your diagnosis and treatments in a way that will help you cope with the new stressors in your life. Your spirituality may be expressed as an organized religion, yoga, the arts, or any other outlet that allows you express your feelings about life. If you are a member of a religious or spiritual community or church, the other members of the community can be an excellent source of support to you and to your family.
The diagnosis of lung cancer has a profound effect on you, but it also has a huge impact on your loved ones. There are many support groups available to you and your family and friends. The GO2 Foundation Lung Cancer Living Room® is a monthly in person and online support group that welcomes patients, families and friends. Our hope is that during these monthly meetings, we can share stories, talk about problems you are having, share ideas and practices that have helped, raise awareness of lung cancer, and offer any kind of support to patients, survivors and families. There are many other support groups. To find these groups, search “lung cancer support groups” online. Your local treatment center or hospital may also have a support group. Your social worker or case manager should be able to give you contact information for those groups.

As you progress through treatment, dealing with stress and depression will be critical. If you find yourself having problems coping, ask your oncologist for a referral to a mental health professional. Depending on the kind of help you need, your doctor may refer you to one of a number of different professionals: psychiatrists, psychologists and psychiatric clinical nurse specialist. A psychiatrist is a medical doctor who provides counseling, medication and other treatments for mental and emotional disorders. A clinical psychologist is a professional with advanced training in psychology. This professional provides counseling for individuals with mental or emotional needs. A psychiatric clinical nurse specialist is a master’s prepared nurse with advanced training in mental health nursing. This nurse may provide counseling or teaching
for patients and families with mental health needs. Be sure that the mental health professional you work with has experience working with cancer patients. A mental health professional without experience in cancer care may not understand the physical and emotional issues and stressors with which you are faced. Your oncologist will be able to direct you to an appropriate mental health specialist. Remember, there is no shame in asking for this support!

**Legal Assessment**

Your financial counselor should help by doing an assessment of your legal and financial status. Your team will ask about several legal documents that will help your doctors and family make decisions about your care. Specifically, your team will assess your insurance program, access to patient assistance programs as well as whether or not you have Advance Directives, Health Care Proxy, and a Durable Power of Attorney.

**Nutrition**

Nutrition is a critical piece of your cancer journey. During treatment, you may have side effects that cause you to lose your appetite. A nutritionist or dietician experienced in cancer care can help you identify a diet that will taste good to you and provide the nutrition you need. There are foods that may interfere with your treatments or help boost your immune system; a qualified nutritionist will help you identify those foods. There are many cookbooks available with easy recipes for cancer patients.

Be sure your healthcare team includes a nutritionist or dietician who can help develop a menu that is best for you while going through treatments. Ask your oncologist to help find the right nutritionist for you.
In addition to meals provided by your immediate support group, your nutritionist or dietician can help you find other sources for delivery of meals when you need them. Many communities have a Meals on Wheels program. To find out if there is a program in your area, visit their website at http://www.mealsonwheelsandmore.org/programs/. Check with your church or other local religious organizations that might have meal programs. Your social worker or nutritionist should be able to find contacts for you.

Traveling

If you travel during cancer treatment, be sure to take a copy of your medical records and a list of all of your medications including brand and generic names, dose and frequency. Also, be sure you have contact information for your oncologist. If you have to get medical care while you are on the road, the information you can provide will be valuable to those caregivers who don’t know you.

Traveling with oxygen

Some airlines provide oxygen for therapeutic or medical purposes (often at an additional cost). There are also portable oxygen concentrators for flying, traveling or simply to do things outside the home setting. Check with your airlines to see if you can purchase inflight medical oxygen or bring your own. Either way, you will need a signed order from your physician. Make sure to plan ahead and check with each airlines for your options and arrangements.

If traveling by bus or train, two weeks’ notice is often needed for booking travel with a portable oxygen concentrator.

Before you fly or visit a high altitude location, ask your doctor to perform a High Altitude Simulation Test (HAST) to determine if you will need oxygen when you are traveling.

Inogen is a provider of portable oxygen systems. Visit www.inogen.com for more information.
Alternative / Complementary Therapies

If you do research about lung cancer, you will find a lot of information about alternative or complementary therapies. Complementary therapy is any treatment that is used along with standard treatment. These treatments may enhance the treatments prescribed by your oncologist. Alternative therapies are treatments that are used as a substitute for the standard treatment prescribed by your oncologist. These treatments are used instead of the standard treatments.

According to the MD Anderson Cancer Center, these therapies may or may not be useful “…to promote wellness, manage symptoms associated with cancer and its treatment, or treat cancer. When properly combined with standard cancer treatments, some complementary therapies can enhance wellness and quality of life.” However, others may be harmful or actually interfere with your medical treatment. It is imperative to discuss any of these alternative treatments with your team as your complementary or alternative therapy may interfere with your standard cancer care.

Before using any complementary or alternative therapy, be sure to discuss the therapy with your oncologist and healthcare team.
FINANCING YOUR CANCER CARE
The Foundation recommended I go talk to another lung cancer specialist up in Colorado—Dr. Camidge. They opened the doors to the right treatment, but they also opened their doors to their hearts. That’s the best thing.

—Henry Randall “Hank” Baskett, Jr., survivor
Health and Disability Insurance
Your healthcare team should include a certified benefits counselor or a social worker who can help guide you through the process of applying for benefits should you become disabled by lung cancer. These professionals have been specifically trained to help you determine if you are eligible for financial help through your healthcare insurance or Social Security. You may also qualify for long or short-term Social Security disability benefits through the Department of Labor. The Medicare prescription Part D section may be available to you. If you are not eligible for Medicare, there are prescription assistance programs that might be helpful to you. Retirement or veterans benefits may help those who are eligible for them. State and community programs may exist including home-based programs.

When you are preparing to speak with a social worker or benefits counselor, be sure to have the following information:

- Recent statements from your insurance company
- Bank account information
- Medications that you are currently taking (for Medicare Rx or other prescription benefit programs)
- Veterans benefits and separation documents
- Retirement statements concerning benefits you are already receiving
- Social Security statements and card (if available)
- Disability benefits you are currently receiving
Medicare
Medicare is the nationally sponsored program that guarantees that elderly and disabled Americans have access to health insurance and health care. You may qualify for Medicare benefits if you are aged 65 or older or if you have certain disabilities or end stage renal disease (ESRD). Your certified benefits coordinator or professional social worker can help you determine if you might be eligible for benefits. Your certified benefits coordinator or social work can also help you through the application process if you are eligible for benefits.

Medicaid
Medicaid is the national and state sponsored program that guarantees that certain low-income families and people with certain disabilities have access to health care. As with Medicare, the process for qualifying and applying for Medicaid benefits as a lung cancer patient is extremely complicated. Your certified benefits coordinator or social work can also help you through the application process if you are eligible for benefits.

The Consolidated Omnibus Budget Reconciliation Act (COBRA)
The Consolidated Omnibus Budget Reconciliation Act (COBRA) gives you the right to choose to continue your health benefits when you are no longer able to work. This coverage is the same as that provided by your group health plan and is available to you for limited periods. You may qualify under certain circumstances such as voluntary or involuntary job loss, reduction in the number of hours worked, and other life events. You may be required to pay the entire premium for coverage up to 102% of the cost to the plan.

COBRA outlines how you may elect to continue coverage. It also requires your employer to provide notice. For more information, go to the US Department of Labor website at http://www.dol.gov/dol/topic/health-plans/cobra.htm.
Social Security Disability Insurance (SSDI)
The Social Security Administration (SSA) has a specific medical listing for cancer of the lung and a five-step evaluation process they will use to evaluate your claim. To find out if you are eligible for SSDI benefits visit the Social Security Disability program website at http://ssa.gov/disability/ or call 1-800-772-1213.

High-risk medical insurance
Many states offer high-risk medical plans for lung cancer patients with pre-existing conditions. For a list of states offering these plans and how the new Patient Protection and Affordable Care Act affects you, visit the healthinsurance.org website at http://www.healthinsurance.org.

Special rates for the uninsured or for creating a payment plan
Many hospitals will work with you and your family to create a payment plan that suits your budget. To find out more, call your hospital's financial services office. You may also apply for a reduced rate for services such as diagnostic tests, treatments and other bills related to your lung cancer treatment.

Patient Advocate Foundation (PAF)
PAF's Co-Pay Relief Program provides direct financial relief for insurance co-pays for drugs associated with the treatment of NSCLC. For more information visit their website at http://www.copays.org/resources/lung.php. On this site, you may also find helpful information for solving insurance and healthcare access problems.

Note: If one of these organizations is not enrolling at the time you call or if you do not qualify for benefits, ask the organization you contact which organizations are enrolling new patients. Not all of these organizations are open to enroll lung cancer patients year-round.
Patient Access Network (PAN)
The Patient Access Network provides direct financial relief for insurance co-pays for drugs associated with the treatment of NSCLC. You can sign up on their website: http://www.panfoundation.org/ or call toll-free to 1-866-316-7263.

Healthwell Foundation
The Healthwell Foundation may be able to help you cover coinsurance, copayments, healthcare premiums and other costs for some treatments. The Foundation supports a limited number of diseases at any one time and the list changes frequently. For more information on the diseases covered and the funding process visit their website at http://healthwellfoundation.org/or call 1-800-675-8416.

Chronic Disease Fund
The Chronic Disease Fund assists eligible individuals with paying for drugs, co-pay assistance, and travel assistance. For more information visit their website at http://www.cdfund.org/ or call 1-877-968-7233.

Cancercare
Cancercare provides limited financial assistance, counseling with certified oncology social workers, support groups for patients and caregivers and community programs in Connecticut, New Jersey, and New York. If you live in one of those states, visit their website at http://www.cancercare.org/diagnosis/lung_cancer for more information.

For assistance paying for drug treatments search online using the keywords “Prescription Assistance” and “[your state].”
Pharmaceutical companies

Pharmaceutical companies may provide financial assistance to pay for drugs provided by the company if you meet certain financial requirements.

If you are having trouble paying for your treatment, check with your pharmaceutical company, local pharmacist or your oncologist for information on financial assistance programs. It is virtually always necessary to provide one’s tax return for this process, so be sure to have a copy handy for the application.
I see firsthand how Bonnie and this Foundation reach out and touch so many lives ... lung cancer patients, survivors, and caretakers.

—Sally Samuels, survivor
END-OF-LIFE PLANNING
It takes a family to support lung cancer patients, and the GO2 Foundation for Lung Cancer family is doing just that.

—Adeeti Ullal
At some point in your cancer journey, you will be asked about the plans you have made for end-of-life care. You may have already created an end-of-life plan such as a will and Advance Directives. If not, we understand having these discussions now may be very difficult for you, your family and even for your healthcare team. These discussions may become even more difficult as you become more ill. Having these discussions and making these decisions early in your cancer journey can help you and your family feel less stress should your treatment plan change.

End-of-life plans include directions on how to manage pain, where you want treatment (e.g. hospice, home, hospital), legal documents such as Advance Directives and Health Care Proxy, as well as preplanning funeral services. Many of these end-of-life plans may be guided by your philosophical or religious beliefs and your spiritual advisor may be very helpful as you think about these issues. If your belief system requires or prohibits certain actions or treatments, your family and healthcare team must know about these limitations before the time when the decision must be made. If you have not made these plans before being diagnosed with lung cancer, it is important that you begin to think and talk about them and document your plans.

Although these discussions are difficult, your support system must understand what you want in order to provide the treatment you would choose for yourself. It is also important that these discussions continue throughout the course of your treatment; decisions you make at the time of diagnosis may change over time as your disease and treatments change. As your feelings about treatment change, you need to be sure your family and healthcare team know about these changes. By having your plans documented, you can relax knowing that your family will not have to make decisions when they may be upset…and that the decisions they make will be those you want.
You will also be able to focus all your attention on your treatment plan.

There are a series of legal documents to complete in order to capture your healthcare wishes. In order to complete the documents described below, you will want to talk to your family and healthcare team about what treatments and medications you want to receive – and at what point in your treatment you will not want to receive them. You may also want to speak with an attorney to help complete the documents.

Healthcare Pre-Planning
It is important for everyone to plan for end-of-life – this is even more important when you are diagnosed with a serious illness such as cancer. End-of-life planning will allow you to concentrate on taking care of your health knowing that the rest of your team understands exactly what you want. End-of-life planning will also relieve the stress your family may feel because they will know exactly how you wish to be treated. As you review your end-of-life planning, the discussions may be uncomfortable at the beginning so it may be helpful to include your healthcare team, legal advisor, spiritual advisor, and your family.

A cancer diagnosis may carry with it a variety of legal issues, including insurance coverage, employment and taking time off work, access to health care and government benefits, and estate planning. These issues can be overwhelming to you. If you do not deal with these legal issues, you may find that even though you have made it through treatment, you have lost your job, home, or insurance.

Online Resources
There are online resources that can provide great information as you begin planning. One very good resource is the Cancer Legal Resource Center (CLRC) that is sponsored
by the Loyola Law School of Los Angeles and the Disability Rights Legal Center. This center offers free information for you, your family, and your healthcare team. In addition to the online resources available at https://thedrlc.org/cancer/, the center also offers support on a toll-free assistance line (1-866-THE-CLRC). When you call this number, you will be connected to an appropriate person (attorney, accountant, or insurance professional) to help you with your specific question.

The National Cancer Institute (NCI) at the National Institutes of Health is another good resource for end-of-life planning. This site will give you ideas about planning your care and managing symptoms at the end of life. For more information, visit the NCI website at http://www.cancer.gov/cancertopics/pdq/supportivecare/lasthours/patient.

Important Documents
As you do pre-planning, you will want to prepare several documents. Although these documents do not guarantee your wishes will be followed, they will provide guidance for your family and healthcare team if you are unable to make decisions for yourself.

- An Advance Directive (AD) is a generic term that your healthcare team will use to describe a document in which you describe what medical treatment(s) you want to receive if you are unable to tell your oncologist what you want. For example, you may want to receive all treatments that are available to you – or you may not want any. The document known as a Living Will is a certain type of advance directive that may or may not be a legal document in your state. Each state has a specific format for the Living Will or Advance Directive. Your attorney will be able to help you determine the specific format that is legal in your state.
The AD document will also typically describe whether you wish to be resuscitated in the event your heart stops. A Do Not Resuscitate (DNR) order means that you do not want CPR if your heart stops. You can also revise this document at any time before, during or after treatment. Be sure your healthcare team has a copy of your current AD document. It is also critical that you discuss your wishes with your friends and family members. Let them know what your wishes are and how you want to be treated. It is extremely important to discuss your wishes with the person you name as your Health Care Proxy (see more information below).

There are online websites where you can quickly and inexpensively create a Living Will/Advance Directive that will be legal in your state. When you have completed the advance directive document, be sure to share copies with your family, healthcare team, hospital and health care proxy.

- The Aging with Dignity Five Wishes Online (www.agingwithdignity.org) allows you to complete the form online or print a blank copy to complete by hand.
- The Do Your Own Will site (www.doyourownwill.com/living-will/states.html) allows you to download the Living Will specific to your state for completion off line. This site is also a good resource for general information about wills and estate planning.
- Caring Connections is an organization that offers resources including a free Advance Directives document specific to your state (http://www.caringinfo.org/i4a/pages/index.cfm?pageid=3289).

- Your Health Care Proxy (HCP) document will identify the person you want to make medical decisions for you if you are unable to make your own. This person, or proxy, may also be known as your durable power of attorney for healthcare.
• The HCP is different from the Durable Power of Attorney. A Durable Power of Attorney (DPOA) names a person who has the power to make legal decisions for you. Your HCP and DPOA may be the same person, if you so choose.

In addition to the legal documents, be sure your health care proxy or family members or the person you most trust have access to information about all your will, living will/advance directives, credit cards, bank accounts, phone numbers, email accounts, investment accounts, and any other documents they may need in the event you cannot make decisions for yourself. We recommend you keep a file in a safe place that includes all these important documents.

Legacy planning is a wonderful way to leave a meaningful mark on your community and the world. As you go through this journey, you can discover many ways to make a difference in the world by leaving a gift to benefit those causes in which you believe. Giving a gift to our foundation can allow you and your family to leave a legacy that will touch others who are diagnosed with lung cancer in the future.

Our staff at the GO₂ Foundation for Lung Cancer would be honored to discuss appropriate opportunities for legacy giving and recognition. Please call 1-800-298-2436.

Funeral or Memorial Service Pre-planning
While it is difficult to comprehend your own mortality as you are fighting to cure your lung cancer, some people find it helpful to themselves and their families to pre-plan a funeral or memorial services. Planning your service will help your family because you will make all of the decisions, sparing your loved ones these difficult decisions when you are gone.
The LIVESTRONG Foundation is a great resource for managing the pre-planning process. This organization provides many suggestions and resources that will walk you through pre-planning for your service. LIVESTRONG provides step-by-step instructions to begin the process and some things you should think about as you work through the plan. This site will give you information about funeral costs and different options for paying for the funeral. To access the information on this site, visit http://www.livestrong.org/.

Palliative Care vs. Hospice

**Palliative Care**

Palliative care teams are a relatively new formal concept in health care although the concept of providing comfort care is not new at all. In palliative care, the goal of the team is to prevent and/or relieve pain and suffering. This suffering might be physical, mental or emotional. The desired outcome is always that your quality of life will be improved.

Some people are confused about the differences between palliative care and hospice. Palliative care can be delivered at any point in your treatment including at the end of life; hospice care is typically given when the illness cannot be cured. Whereas hospice care is usually delivered in the home or a hospice facility, palliative care may be delivered in any environment.

**Hospice Care**

While many people see hospice care as a last resort, we encourage you and your family to consider hospice as a caring support system. According to the Hospice Foundation, “Hospice is the ‘something more’ that can be done for the patient
and the family when the illness cannot be cured. It is a concept based on comfort-oriented care. Referral into hospice is a movement into another mode of therapy, which may be more appropriate for terminal care.”23 Visit the Hospice Foundation site at http://www.hospicefoundation.org/ to learn more about how to find hospice care in your area.

Grief
Grief is a natural reaction to a diagnosis of lung cancer. Grief is the emotional suffering you feel because your health and life have been changed. The process of grieving is unique to you. Your grief will be influenced by your personality, individual coping style, diagnosis and overall physical health. Ignoring the emotional pain you feel will not make it go away. You might find that it is helpful to talk to a counselor or close friend about what you are feeling. Ask your physician to refer you to a social worker or counselor that specializes in cancer care.

The five stages of grief include:

- Denial – “The diagnosis is not correct” – This stage of grief is characterized by shock and disbelief.
- Anger – “What did I do to deserve this?” – This stage is characterized by feelings of resentment.
- Bargaining – Usually expressed as trying to make a deal with some higher power – “If you make this not happen, I will become a better person” – This stage is characterized by feelings of fear and guilt.
- Depression – “I am so sad/upset/down I cannot get up in the morning” – This stage of grief may be characterized by physical symptoms including fatigue, insomnia, nausea and vomiting.
- Acceptance – “I can deal with this no matter what happens” – This stage is characterized by feelings of relief and peace.
It is common to go back and forth between these stages. One day you may be angry and the next you may be depressed. Finding ways to cope with grief is important. First, find a strong support system that will allow you to share your feelings regardless of what they are. Second, take care of yourself. Eat right and stop and rest when you get tired.

Finally, do not be ashamed to get professional help if your grief becomes overwhelming. We are here to help. Do not hesitate to contact us at 1-800-298-2436.
OUR GENEROUS SUPPORTERS
As vital information becomes available, new print editions of this guidebook will be released with updated PDFs available on our website. Check our website (www.go2foundation.org) or Amazon.com to make sure you have the most current edition.
In 2003, my mom was diagnosed with lung cancer. Her life changed from one filled with business and family obligations to a life defined by doctor’s visits, chemo and radiation therapy, and surgery. When my mother was diagnosed with lung cancer, my world changed as well. I was a wife, a mother, and an entrepreneur. Suddenly, I was the daughter of a cancer patient trying to support my mother on a daily basis while trying to find answers to complex healthcare issues. This guidebook is the culmination of years of conversations with patients, doctors, researchers … just about anyone with any information relating to lung cancer. I am grateful to our generous supporters without whom this guidebook would not exist. Thanks to their willingness to support us and the lung cancer community at large, we are getting closer to our goal of making lung cancer a survivable disease.

Throughout the guidebook we encourage you to call us with any questions. I want you to know that I understand the journey you are on and I am willing to help. Please feel free to call me with any questions.

Sincerely,

Danielle Hicks
Chief Patient Officer & Daughter of a Lung Cancer Survivor
Without the CT test that you recommended, I would never have suspected I had lung cancer.

—Evy Schiffman, survivor
GLOSSARY
I belong to this foundation and host a run every year in Gainesville, Florida, because I want to help all patients have a fighting chance to be warriors in this battle.

—Caren Gorenberg, survivor
Adjuvant therapy: any therapy that is started after surgery.

Benign: non cancerous

Biomarkers: biomarker or biological marker is a very distinctive substance that indicates a particular disease is present.

Biomarker testing: also called assays or profiles, can help your treatment team identify specific biomarkers that are in a tumor.

Bronchi: the trachea (windpipe) divides into two main bronchi which is a passage of airway that allows air into the lungs.

Carcinogens: substances that can cause cancer.

Chemotherapy regimen: a combination of chemotherapy drugs.

donPCR: Digital Droplet™ polymerase chain reaction is a method for performing digital PCR based on emulsion droplet technology that is known for its highly sensitivity and quick turn-around time. Blood-based ddPCR measures circulating nucleic acids isolated from plasma, and can be optimized for rare mutation detection from blood.

DNA (deoxyribonucleic acid): the molecule in every cell that controls how that cell grows and functions.

Electromagnetic Navigation Bronchoscopy™ procedure: also known as an ENB™ procedure, this is a minimally invasive approach to accessing difficult-to-reach areas of the lung using the superDimension™ navigation system to aid in the diagnosis of lung disease.

ENB™ procedure: see Electromagnetic Navigation Bronchoscopy™ procedure.
Fiducial marker: a small gold seed or platinum coil that is placed around a tumor to act as a radiologic landmark.

Free radicals: exposure to carcinogens may form molecules in the body called free radicals which damage cells and alter the DNA of the cell.

Genetic fusion: a gene that is formed when the genetic material from two previously separate genes are mixed.

Genetic mutation: a change in the structure of a gene.

Hemoptysis: coughing up of blood or of blood-stained sputum.

Lymph nodes: part of the lymph system that are responsible for filtering the wastes out of the liquid that passes through.

Lymphatic system: responsible for carrying nutrients to the body’s cells and waste away from the cells.

Malignant: cancerous

Mesothelium: the lining that covers the body’s internal organs and cavities.

Metastases, Metastasized: cancer that moves from its site of origination to another part of the body.

Neoadjuvant therapy: any therapy (chemotherapy or radiation) that is started before surgery.

Next generation sequencing: a technique or method of sequencing large amounts of DNA accurately in a short period of time.
**Pleura**: outer lining of the lungs.

**Pleurodesis**: a procedure that involves inserting a chest tube to insert chemicals to induce a scar, thus ‘gluing’ the lung to its lining.

**Primary lung cancer**: lung cancer that starts in the lung.

**Prophylactic cranial irradiation (PCI)**: a kind of radiation treatment that may be used to kill cancer cells in the brain that may not be visible on x-rays or scans.

**Radioactive isotope**: an atom that emits radiation that can be seen by the radiological equipment.

**Secondary lung cancer**: cancer formed in another part of the body and travels to the lung.

**Thorascopy**: a camera on the end of flexible tubing that allows your doctor to look into your chest.

**Trachea**: also known as the “windpipe,” is a tube that connects the pharynx or larynx to the lungs, allowing the passage of air.

**Tumor**: a group of cells that stick together. Can be benign (non-cancerous) or malignant (cancerous).
EVENTS

GET INVOLVED
AND FIGHT LUNG CANCER AT
NATIONWIDE EVENTS

JOIN THE FIGHT

Join our community of people across the country as we come together and raise funds and awareness for lung cancer!

Our GO2 Foundation National Walk/Run Series raises much-needed funds for the GO2 Foundation. Each event provides fun, family-friendly activities, including music, a walk/run, event awards and more. Come together with lung cancer survivors, caregivers, friends and loved ones as we unite to help those impacted by lung cancer in your community.

If you are interested in learning more about the program, please contact us at walk@go2foundation.org.

www.go2foundation.org/events/
REFERENCES
I want to find a cure.

—Ellis Cox


INDEX
INDEX

#
3-D Conformal Radiation Therapy (3DCRT) 78

A
Adenocarcinoma 13, 27, 30, 32-33
Adjuvant therapy 175
Airway Stenting 89
ALK 27, 30, 32-33, 35, 73, 104-106, 108
Alternative Therapies 150
Anaplastic Lymphoma Kinase (ALK) 105
APC 88
Argon Plasma Coagulation 88

B
Balloon bronchoplasty 90
Benign 9, 19, 20, 25, 223, 177
Biomarker(s) 21, 26, 29-38, 39, 65, 97, 103, 107-108, 118, 127, 160, 175
Biomarker testing 21, 29-38, 65, 97, 103, 107
Biopsy 19, 21, 29-30, 33, 35, 36, 38, 42, 86-89, 104-105, 119
Blood-based proteomic testing 33
Bone metastases 97, 124
Brachytherapy 85, 89, 97
BRAF 30, 32, 35, 73, 106, 108
Brain metastases 84-85
Brigatinib (Alunbrig™) 73, 106
Bronchoscopy 22-24, 35, 49, 55, 86, 88, 175

C
Cachexia 98
Cancercare 144, 156
Carcinogens 10, 175, 6
Care Setting Assessment 143
CAT Scan/CT Scan 19, 21, 38, 42
CBC 37
Chemotherapy 15, 29, 30, 32, 36-37, 39, 40, 42, 61-62, 64, 69-71, 75-77, 103-104, 122, 124, 125, 141, 175-176
Chemotherapy Nurse 42
Chronic Disease Fund 156
Clinical Trial Phases 130
Clinical Trials 31-32, 34, 41, 43, 61, 70, 92, 98, 115, 125, 127-136
COBRA 200
Complementary Therapy 196
Complete Blood Count (CBC) 37
Core Needle Biopsy 21, 35
Cryosurgery 86
CTLA4 113, 116
CT Scan 19, 21, 38, 42
Cytologist 38
Cytology 38, 48, 93
Cytotoxic T Lymphocyte Antigen 116

D
ddPCR Blood-based Testing 34
Diagnosis Process 19-43
DNA 9, 10, 30, 34, 77, 175-176
Durvalumab (Imfinzi™) 72, 112, 115, 123

E
EBUS 87
EGFR 27, 30, 31-32, 34-35, 73, 104-108
EGFR T790M 104, 105
Electromagnetic Navigation Bronchoscopy™ procedure 23, 35, 86, 175
Embolism 99
EML4-ALK 32
ENB™ 23, 35, 86, 175
Endobronchial Ultrasound 87
End-of-Life Planning 161-168
Endoluminal high dose rate (HDR) brachytherapy 85
Epidermoid or squamous cell carcinoma 13
<table>
<thead>
<tr>
<th>INDEX</th>
</tr>
</thead>
</table>

**ERBB2** 30
Erlotinib (Tarceva®) 32, 73, 104-105, 106
Extensive Stage 121-122
External Beam Radiation 77-80, 84-85, 97

**F**
Fiducial marker/placement 87, 89, 176
Fine Needle Aspiration 21, 35
FNA 21, 35

**G**
Gefitinib (Iressa™) 32, 73, 106
Genetic mutation 30, 35, 108, 176
Grief 167-168

**H**
HAST 37, 149
Health and Disability Insurance 153
Healthwell Foundation 156
HER2 30, 35
High Altitude Simulation Test 37, 149
High dose rate (HDR) brachytherapy 85, 89
Hospice Care 166-167

**IGRT** 79
Image Guidance Radiation Therapy 79
Imfinzi™ 72, 112, 115, 123
Immune System 14, 62, 90, 92, 98, 110-113, 116-117
Immunotherapy 15, 32, 61-62, 94, 96-97, 109-118
Important Documents 163-165
IMRT 78-79
Intensity-Modulated Radiation Therapy 78-79
Iressa™ 32, 73, 106

**K**
Keytruda® 72, 112, 115, 123
KRAS 30-32

<table>
<thead>
<tr>
<th>L</th>
</tr>
</thead>
</table>
Large cell undifferentiated carcinoma 13
Legal 142, 148, 161-165
Limited Stage 121
Lobectomy 63-64, 93-94
Local therapy 61, 76
Lung Cancer Staging 45-57
Lymphadenectomy 64
Lymphatic system 10-11, 176
Lymph Node Biopsy 24, 35
Lymph Node Dissection 64
Lymph nodes 10, 35, 64-65, 47-54, 87, 93-96,

<table>
<thead>
<tr>
<th>M</th>
</tr>
</thead>
</table>
Maintenance therapy 71, 103
Malignant 9, 14, 20, 25, 49, 55, 176-177
Mediastinoscopy 24, 35
Medical Oncologist 42, 43
MET 30, 32, 35, 105
Mesh brachytherapy 85
Mesothelioma 12, 14
Metastases 10, 20, 84, 97, 106, 124
Molecular fingerprint 29
MRI 20-21, 35, 42,
Mucus 38
Multidisciplinary Healthcare Team 42

<table>
<thead>
<tr>
<th>N</th>
</tr>
</thead>
</table>
NanoKnife Electroporation 85
Narrow band imaging 88
Neoadjuvant (or Neo-adjuvant) therapy 70, 176
Next-Generation Sequencing 34-36
NIH clinical trial list 136
Non-small cell lung carcinoma 12
NSCLC 12, 13, 31-35, 62, 69, 70-73, 93-96, 103-107, 114-115, 117, 121, 155, 156
Opdivo® 72, 112, 115, 123

Pain 10-11, 62, 66-67, 161, 166-167, 80-81, 92, 96-99
Palliative Care 166
Palliative radiation 97
PAN 156
Pancoast or pulmonary sulcus tumor 13
Pancoast Tumor 13
Paraneoplastic syndromes 14
Pathologist 21-22, 24-27, 30, 35, 38, 42, 63-64
Patient Access Network 156
Patient Advocate Foundation 155
PD-1 113-115, 72, 123
PD-1/PDL1 checkpoint inhibitors 113-114
PDL1 113-114
PDT 90-92
PET Scan 20, 22, 42
PFT 36
Photodynamic therapy 90-92
Pleural Effusion 64, 66, 99
Pleurodesis 64-65, 99, 177
Pleuroscopy 90
Pneumonectomy 64, 94
Pneumonia 81, 98-99
Pneumothorax 64
Porfimer sodium 90, 92
Port 62, 69, 75
Positron Emission Tomography Scan 20, 94
Pulmonary Function Test 36
Pulmonologist 35-37, 42, 85-89, 90, 135
Pulse Ox 36
Pulse Oximetry 36

Radial Probe Ultrasound (REBUS) 87
Radiation Oncologist 39, 42, 78, 84, 86
Radiation Technician 43
Radiographic Tests 19
Radiosurgery 79, 81, 84, 87, 89
REBUS 87
Recurrent SCLC 125
Registered Dietitian 43
Research Nurse 43, 134
RET 30, 33, 35, 108
RN Navigator 42
Robotic Bronchoscopy 22-23
ROS 30, 33, 35, 73, 104-105, 106, 108
Sarcoma 12, 15
SCLC 12, 14, 69, 71, 115, 121-125, 136
Segmental Resection 63
Segmentectomy 63, 93
Side effects 29, 41-42, 75-76, 78, 80-82, 84, 92, 103,
106, 115, 118, 129, 130-131, 133, 141-142, 148
Signs and symptoms 11-12
Sleeve Resection 63-64, 94
Small Cell Lung Cancer 14, 35, 110, 121-125, 136
Small cell lung carcinoma 12
Social Worker 39, 43, 142, 145, 147, 149, 153-154,
156, 167
Spiritual and Mental Health Assessment 142, 146
Sputum Cytology 38, 48, 93
Stage 0 48, 93
Stage I 47-49, 63, 79, 93, 94
Stage II 50-51, 94-95, 70
Stage III 70, 95-96, 112, 114, 52-56
Stage IIIA 52-54, 70, 95
Stage IIIB 55, 96
Stage IIIC 56, 94
Stage IV 20, 57, 70, 96, 97
Stereotactictic radiotherapy 79
Support Systems 143
Symptom Management Care Coordinator 43
Systemic Therapy 62, 69, 76
T
Tarceva® 32, 73, 105, 105-107
Targeted Therapies 30, 33, 34, 62, 73, 103-108, 123
Tecentriq® 72, 112, 115, 123
Thoracentesis 24
Thoracic Surgeon 22, 25, 35, 42, 86, 122
Thoracotomy 25, 63
Three-dimensional conformal radiotherapy (3DCRT) 78
TNM staging system 47-57
Transitional Care Planning 187-191
Tumor 225
Tumor grade 25-26

V
VATS 25, 63
Video-Assisted Thoracoscopic Surgery (VATS) 25, 63
VMAT 79
Volumetric Arc Therapy 79

W
Wedge Resection 63, 93-94

Y
Yervoy® 72, 112, 115
The GO2 Foundation for Lung Cancer is a shining light to those diagnosed with lung cancer. They are knowledgeable, helpful, dedicated, compassionate, energetic people—from the person who answers the phone (thanks Kim!) to every single person who is part of the foundation! They go the extra mile and are making a difference for those diagnosed with lung cancer. As a lung cancer survivor, I am proud and so grateful to be on their team.

—Jane Millman, Survivor

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Multiple resources exist across the nation to provide patients and families with medical help, social services, financial guidance, clinical trial information and advice about living with Lung Cancer (diet, exercise, etc). Lung Cancer resources, both local and national, are valuable for anyone looking for help in understanding and coping with this disease.

1-800-298-2436 or support@go2foundation.org

Team members are available Monday – Friday, 9:00 a.m. to 5:00 p.m. (ET).
I have been associated with the foundation since the beginning. Here I am an 23-year survivor of Stage IV Lung Cancer who has really tried to understand the basics of lung cancer and its therapy options thoroughly. ALCMI and the projects we fund—any one of them—may lead to an innovative new protocol for early detection, a new therapy drug, or a new basic understanding of what lung cancer really is. This is an exciting time to be on the front lines of a science that can and will change very fast—and I, and we at the foundation will be there.

—Wells Whitney, survivor
Born out of the vision of the world’s leading lung cancer experts, the Addario Lung Cancer Medical Institute (ALCMI) was founded in 2008 as a premier virtual lung cancer institute. Today, ALCMI partners with key opinion leaders, scientists, patients and our sister organization, the GO2 Foundation for Lung Cancer, to advance cutting-edge research that is transforming what it means to live with lung cancer.

**BY THE NUMBERS**

<table>
<thead>
<tr>
<th>Category</th>
<th>Number</th>
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</thead>
<tbody>
<tr>
<td>Centers of Excellence and Screening Centers</td>
<td>700+ (and growing)</td>
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<tr>
<td>Studies that are open, planned, and complete</td>
<td>15</td>
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<tr>
<td>Specimens in biorepository</td>
<td>2500</td>
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<tr>
<td>Patients registrants in the LUNG CANCER Registry</td>
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<tr>
<td>Institutions</td>
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<tr>
<td>International locations</td>
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<td>United Kingdom</td>
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**PARTNERSHIPS**

- **Biotechnology**
- **Pharmaceutical Industry**
- **GO2 Foundation for Lung Cancer**
- **Diagnostic Companies**

[www.ALCMI.net](http://www.ALCMI.net)
A critical part of this approach is the Community Hospital Center of Excellence Program, which is designed to accelerate lung cancer detection and patient treatment by utilizing the most advanced technology available to compassionate lung cancer specialists. Because 80% of patients receive treatment at their local Community Hospital, these are the centers where the greatest good can be done for the greatest number and we can improve the overall lung cancer patient survival rate.

Spearheaded by a pilot program led by renowned oncologist Dr. Shane Dormady at El Camino Hospital in Silicon Valley, GO2 is working with an elite team of specialists to create an unsurpassed paradigm for lung cancer treatment worldwide — a patient-centric, collaborative model to provide all patients, regardless of where they live, access to the newest and most effective diagnostic and therapeutic techniques.

This new “standard of care” established at Community Hospitals will be accompanied by a formal seal of excellence awarded by GO2 and will ensure that no lung cancer patient is left behind.
LIFE DOESN’T COME WITH INSTRUCTIONS, BUT LIVING WITH LUNG CANCER NOW HAS THIS SURVIVOR’S GUIDEBOOK.

INSIDE YOU WILL GET THE MOST UP-TO-DATE NAVIGATION TOOLS ON:

• THE DIAGNOSIS PROCESS
• LUNG CANCER STAGING
• TREATMENT OPTIONS
• CLINICAL TRIALS
• LIVING WITH LUNG CANCER
• FINANCING YOUR CARE
• HOPE FOR SURVIVING LUNG CANCER
• AND MORE

In my house we call it “the lung bible.” It has been an invaluable resource for me, my family and (sadly) a newly diagnosed friend.

—Diane Broderick

I love this handbook! It has great information that I have shared with my family and friends. I often pick it up and re-read. Each time I learn something new, or it triggers something that I need to talk to my doctor(s) about.

—Kimberly Buchmeier

When my husband was first diagnosed with stage IV Adenocarcinoma we were paralyzed with fear and found ourselves starved for good information about the fight we were beginning. The University of Colorado/Anchutz hospital advised us to contact the foundation and we were sent this wonderful handbook that answered all of our questions and enabled us to feel knowledgeable when choosing an oncologist and when facing surgery and treatments. It is an invaluable resource to us.

—Peter & Donna Blum

This is the most comprehensive manual I’ve ever seen written...focused for the lung cancer patient.

—Roy S. Herbst, MD, PhD
Ensign Professor of Medicine (Oncology), Professor of Pharmacology, Chief of Medical Oncology, Associate Director for Translational Research, Director—Thoracic Oncology Research Program, Yale Comprehensive Cancer Center, Yale School of Medicine

www.GO2FOUNDATION.ORG