Treating Osteosarcoma

If you or your child has been diagnosed with osteosarcoma, the cancer care team will discuss treatment options with you. It’s important to weigh the benefits of each option against the possible risks and side effects.

How is osteosarcoma treated?

Three main types of treatment are used for osteosarcoma:

- Surgery for Osteosarcoma
- Chemotherapy and Other Drugs for Osteosarcoma
- Radiation Therapy for Osteosarcoma

Common treatment approaches

Most often, chemotherapy is given both before and after surgery. It can help lower the risk that the cancer will come back after treatment. It might also allow the surgeon to do a less extensive operation to remove the cancer. Radiation therapy is used less often.

- Treatment Based on the Extent of the Osteosarcoma

Who treats osteosarcoma?

Because osteosarcoma is rare, only doctors in major cancer centers have a lot of experience treating these cancers.

A team approach is recommended when treating osteosarcoma. For children and teens, this team includes the child’s pediatrician as well as children’s cancer specialists. Treatment is best done at a children’s cancer center. For adults with osteosarcoma, the treatment team typically includes the patient’s primary care doctor, as well as specialists...
at a major cancer center. Doctors on the treatment team might include:

- An **orthopedic surgeon** (a surgeon who specializes in muscles and bones) who is experienced in treating bone tumors
- A **medical or pediatric oncologist** (a doctor who treats cancer with chemotherapy and other drugs)
- A **radiation oncologist** (a doctor who treats cancer with radiation therapy)
- A **physiatrist** (a doctor specializing in rehabilitation and physical therapy)

For both adults and children, the team might also include other doctors, physician assistants (PAs), nurse practitioners (NPs), nurses, psychologists, social workers, rehabilitation specialists, and other health professionals.

**Making treatment decisions**

Treatment for osteosarcoma is often effective, but it can also cause serious side effects. It’s important to discuss all treatment options as well as their possible side effects with the cancer care team so you can make an informed decision. It’s also very important to ask questions if you’re not sure about anything.

If time allows, getting a second opinion from another doctor experienced in treating osteosarcoma is often a good idea. This can give you more information and help you feel more confident about the treatment plan you choose. If you aren’t sure where to go for a second opinion, ask your doctor for help.

Once treatment starts, members of the treatment team can help you deal with side effects, stress, and financial and other issues related to treatment.

For cancer in children and teens, many of these issues can be more complex. As a parent, taking care of a child with cancer can be a very big job. It’s important to remember that you will have a lot of help. Many people will be involved in your child’s overall care. It’s also important for you to know that the health professionals who treat children with osteosarcoma are using the experience and knowledge gained from many decades of studying the treatment of this disease.

- **Questions to Ask About Osteosarcoma**
How to Talk to Your Child’s Cancer Care Team

Seeking a Second Opinion

Thinking about taking part in a clinical trial

Today, most children and teens with cancer are treated at specialized children’s cancer centers. These centers offer the most up-to-date treatment by conducting clinical trials (studies of promising new therapies). Children’s cancer centers often conduct many clinical trials at any one time, and in fact most children treated at these centers take part in a clinical trial as part of their treatment.

Adults with cancer also typically have the option to participate in clinical trials as a way to get state-of-the-art cancer treatment. In some cases they may be the only way to get access to newer treatments. They are also the best way for doctors to learn better methods to treat cancer. Still, they might not be right for everyone. Talk to your treatment team if you’re interested in learning more about clinical trials, and ask about the pros and cons of enrolling in one of them.

If you would like to learn more about clinical trials, start by asking the treatment team if your clinic or hospital conducts clinical trials.

Clinical Trials

Considering complementary and alternative methods

You may hear about alternative or complementary methods that the doctor hasn’t mentioned. These methods can include vitamins, herbs, and special diets, or other methods such as acupuncture or massage, to name a few.

Complementary methods refer to treatments that are used along with your regular medical care. Alternative treatments are used instead of standard medical treatment. Although some of these methods might be helpful in relieving symptoms or helping people feel better, many have not been proven to work. Some might even be harmful.

Be sure to talk to your cancer care team about any method you are thinking about using. They can help you learn what is known (or not known) about the method, which can help you make an informed decision.

Complementary and Alternative Medicine
Preparing for treatment

Before treatment, the doctors and other members of the team will help you understand the tests that will need to be done. The team’s social worker will also counsel you about some of the issues that might come up during and after treatment, and might be able to help you find housing and financial aid if needed.

- [When Your Child Has Cancer](#)

Help getting through cancer treatment

Your cancer care team will be your first source of information and support, but there are other resources for help when you need it. Hospital- or clinic-based support services can also be an important part of your care. These might include nursing or social work services, financial aid, nutritional advice, rehab, or spiritual help. For children and teens with cancer and their families, other specialists can be an important part of care as well.

The American Cancer Society also has programs and services – including rides to treatment, lodging, and more – to help you get through treatment. Call our National Cancer Information Center at 1-800-227-2345 and speak with one of our trained specialists.

- [Finding Help and Support When Your Child Has Cancer](#)
- [Find Support Programs and Services in Your Area](#)

*The treatment information given here is not official policy of the American Cancer Society and is not intended as medical advice to replace the expertise and judgment of your cancer care team. It is intended to help you and your family make informed decisions, together with your doctor. Your doctor may have reasons for suggesting a treatment plan different from these general treatment options. Don’t hesitate to ask him or her questions about your treatment options.*

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**Surgery for Osteosarcoma**

Surgery is an important part of treatment for almost all osteosarcomas. It includes:
The biopsy to diagnose the cancer
• The removal of the tumor(s)

Whenever possible, it’s very important that the biopsy and the surgery to remove the tumor be planned together, and that an experienced orthopedic surgeon does both the biopsy and the surgery to remove the tumor. The biopsy should be done in a certain way to give the best chance that less extensive surgery will be needed later on.

The main goal of surgery is to remove all of the cancer. If even a small amount of cancer is left behind, it might continue to grow and make a new tumor, and might even spread to other parts of the body. To lower the risk of this happening, surgeons remove the tumor plus some of the normal tissue that surrounds it. This is known as a wide excision.

A doctor called a pathologist will look at the removed tissue under a microscope to see if there are cancer cells at the margins (outer edges).

• If cancer cells are seen at the edges of the tissue, the margins are called positive. Positive margins can mean that some cancer was left behind.
• When no cancer cells are seen at the edges of the tissue, the margins are said to be negative, clean, or clear. A wide excision with clean margins helps limit the risk that the cancer will come back in the place where it started.

The type of surgery done depends mainly on the location and size of the tumor. Although all operations to remove osteosarcomas are complex, tumors in the limbs (arms or legs) are generally not as hard to remove as those in the jaw bone, at the base of the skull, in the spine, or in the pelvic (hip) bone.

**Tumors in the arms or legs**

Tumors in the arms or legs might be treated with either:

- **Limb-salvage (limb-sparing) surgery**: removing the cancer and some surrounding normal tissue but leaving the limb basically intact
- **Amputation**: removing the cancer and all or part of an arm or leg

**Limb-salvage surgery**
Most patients with tumors in the arms or legs can have limb-sparing surgery, but this depends on where the tumor is, how big it is, and whether it has grown into nearby structures.

Limb-salvage surgery is a very complex operation. The surgeons who do this type of operation must have special skills and experience. The challenge is to remove the entire tumor while still saving the nearby tendons, nerves, and blood vessels to keep as much of the limb’s function and appearance as possible. If the cancer has grown into these structures, they will need to be removed along with the tumor. In such cases, amputation may sometimes be the best option.

The section of bone that is removed along with the osteosarcoma is replaced with a piece of bone from another part of the body or from another person (a bone graft) or with a man-made device made of metal and other materials that replaces part or all of a bone (an internal prosthesis). Some newer devices combine a graft and a prosthesis.

Complications of limb-salvage surgery can include infections and grafts or rods that become loose or broken. Patients who have limb-salvage surgery might need more surgery in the following years, and some might still eventually need an amputation.

Using an internal prosthesis in growing children is especially challenging. In the past, it required occasional operations to replace the prosthesis with a longer one as the child grew. Newer prostheses have become very sophisticated and often can be made longer without any extra surgery. They have tiny devices in them that can lengthen the prosthesis when needed to make room for a child’s growth. But even these prostheses may need to be replaced with a stronger adult prosthesis once the child’s body stops growing.

It takes about a year, on average, for patients to learn to walk after limb-salvage surgery on a leg. Physical rehabilitation after limb-salvage surgery is more intense than after amputation, and it’s extremely important. If the patient doesn’t actively take part in the rehabilitation program, the salvaged arm or leg might become useless.

**Amputation**

For some patients, amputation may be the best option. For example, if the tumor is very large or if it extends into the nerves and/or the blood vessels, it might not be possible to save the limb.

The surgeon determines how much of the arm or leg needs to be amputated based on the results of MRI scans and an examination of removed tissue by the pathologist during the surgery.
Surgery is usually planned so that muscles and the skin will form a cuff around the remaining bone. This cuff will fit into the end of a prosthetic (artificial) limb. Another option might be to implant a prosthesis into the remaining bone, with the end of the prosthesis remaining outside the skin. This can then be attached to an external prosthesis.

**Reconstructive surgery** can help some patients who lose a limb to function as well as possible. For example, if the leg must be amputated mid-thigh (including the knee joint), the lower leg and foot can be rotated and attached to the thigh bone, so that the ankle functions as a new knee joint. This surgery is called *rotationplasty*. Of course, the patient would still need a prosthetic limb to replace the lower part of the leg.

With proper physical therapy, a person is often able to walk on his/her own 3 to 6 months after a leg amputation.

If the osteosarcoma is in the shoulder or upper arm and amputation is needed, in some cases the area with the tumor can be removed and the lower arm reattached so that the patient has a functional, but much shorter, arm.

**Rehabilitation after surgery**

This may be the hardest part of treatment, and it cannot be described here completely. Patients and parents should meet with a rehabilitation specialist before surgery to learn about their options and what might be required after surgery.

If a limb is amputated, the patient will need to learn to live with and use a prosthetic limb. This can be particularly hard for growing children if the prosthetic limb needs to be changed to keep up with their growth.

When only the tumor and part of the bone is removed in a limb-sparing operation, the situation can sometimes be even more complicated, especially in growing children. More surgery might be needed in the future to replace an internal prosthesis with one more suited to their growing body size.

Both amputation and limb-sparing surgery can have pros and cons. For example, limb-sparing surgery, although often more acceptable than amputation, tends to lead to more complications because of its complexity. Growing children who have limb-sparing surgery are also more likely to need further surgery later.

When researchers have looked at the results of the different surgeries in terms of quality of life, there has been little difference between them. Perhaps the biggest problem can be for teens, who may worry about the social effects of their operation.
Emotional issues can be very important, and support and encouragement are needed for all patients. (See Living as an Osteosarcoma Survivor.)

Tumors that start in other areas

Tumors in the pelvic (hip) bones can often be hard to remove completely with surgery. But if the tumor responds well to chemotherapy first, surgery (sometimes followed by radiation therapy) may get rid of all of the cancer. Pelvic bones can sometimes be reconstructed after surgery, but in some cases pelvic bones and the leg they are attached to might need to be removed.

For tumors in the lower jaw bone, the entire lower half of the jaw may be removed and later replaced with bones from other parts of the body. If the surgeon can’t remove all of the tumor, radiation therapy may be used as well.

For tumors in areas like the spine or the skull, it may not be possible to remove all of the tumor safely. Cancers in these bones may require a combination of treatments such as chemotherapy, surgery, and radiation.

Joint fusion (arthrodesis): Sometimes, after the removal of a tumor that involves a joint (an area where two bones come together), it might not be possible to reconstruct the joint. In this case, surgery might be done to fuse the two bones together. This is most often used for tumors in the spine, but it might also be used in other parts of the body, such as a shoulder or hip. While it can help stabilize the joint, the person will have to learn to compensate for the resulting loss of motion.

Surgical treatment of metastases

If the osteosarcoma has spread to other parts of the body, these tumors need to be removed to have a chance at curing the cancer.

Osteosarcoma most often spreads to the lungs. If surgery can be done to remove these metastases, it must be planned very carefully. Things to be considered before the operation include:

- The number of tumors
- The location of the tumor(s) (one lung or both lungs)
- The size of the tumor(s)
- How well the tumor(s) responded to chemotherapy
- The person’s overall health
Since the chest CT scan done before surgery might not show all of the lung tumors, the surgeon will have a treatment plan in case more tumors are found during the operation.

Patients who have tumors in both lungs and respond well to chemotherapy can have surgery on one lung at a time. Removing tumors from both lungs at the same time may be another option.

Some lung metastases may not be able to be removed because they are too big or are too close to important structures in the chest (such as large blood vessels). Patients whose overall health isn't good (for example, because of heart, liver, or kidney problems) might not be able to withstand the stress of anesthesia and surgery to remove the metastases.

A small number of osteosarcomas spread to other bones or to organs like the kidneys, liver, or brain. Whether these tumors can be removed with surgery depends on their size, location, and other factors.

**Side effects of surgery**

**Short-term risks and side effects:** Surgery to remove an osteosarcoma is often a long and complex operation. Serious short-term side effects are not common, but they can include reactions to anesthesia, excess bleeding, blood clots, and infections. Pain is common after the operation and might require strong pain medicines for a while after surgery as the site heals.

**Long-term side effects:** The long-term side effects of surgery depend mainly on where the tumor is and what type of operation is done. Most osteosarcomas occur in bones of the arms or legs, and some of the long-term issues from surgery on these tumors are described above.

Complications of limb-sparing surgery can include bone grafts or prostheses that might become loose or broken. This is more likely than with bone surgery done for other reasons because the chemotherapy used before and after surgery can increase the risk of infection and affect wound healing. Infections are also a concern in people who have had amputations, especially of part of a leg, because the pressure placed on the skin at the site of the amputation can cause the skin to break down over time.

As mentioned before, physical therapy and rehabilitation are very important after surgery for osteosarcoma. Following the recommended rehab program offers the best chance for good long-term limb function. Even with rehab, people might still have to
adjust to long-term issues such as changes in how they walk or do other tasks, and changes in appearance. Physical, occupational, and other therapies can often help people adjust and cope with these challenges.

For more general information on surgery as a treatment for cancer, see Cancer Surgery\textsuperscript{12}.

**Hyperlinks**


**References**

Chemotherapy and Other Drugs for Osteosarcoma

Chemotherapy (chemo) is the use of drugs to treat cancer. The drugs are usually given into a vein and can reach and destroy cancer cells throughout the body.

Chemo is an important part of the treatment for most people with osteosarcoma (although some patients with low-grade osteosarcoma might not need it). Most osteosarcomas don’t appear to have spread beyond the main tumor when they are first found. But in the past, when these cancers were treated with surgery alone, the cancer would often come back in other parts of the body, where it would be very hard to control. Giving chemo along with surgery helps lower the risk of these cancers coming back.


Last Revised: October 8, 2020
Most osteosarcomas are treated with chemo before surgery (known as neoadjuvant chemotherapy) for about 10 weeks. In some people with osteosarcoma in an arm or leg bone, this can shrink the tumor, which might help make surgery easier. Chemo is then given again after surgery (known as adjuvant chemotherapy) for up to a year.

Chemo in is given in cycles, with each period of treatment followed by a rest period to give the body time to recover. Each cycle typically lasts for a few weeks.

**Chemo drugs used to treat osteosarcoma**

The drugs used most often to treat osteosarcoma include:

- **Methotrexate** (given in high doses, along with leucovorin to help limit side effects)
- **Doxorubicin (Adriamycin)**
- **Cisplatin or carboplatin**
- **Ifosfamide**
- **Cyclophosphamide**
- **Etoposide**
- **Gemcitabine**

Usually, 2 or more drugs are given together. Some common combinations of drugs include:

- High-dose methotrexate, doxorubicin, and cisplatin (known as the MAP regimen), sometimes with ifosfamide. This is used more often in children, teens, and young adults.
- Doxorubicin and cisplatin. This is used more often in older adults.
- Ifosfamide and etoposide
- Ifosfamide, cisplatin (or carboplatin), and epirubicin

Many experts recommend that the drugs be given in very high doses when possible.

Before starting chemo, the doctor might advise putting a catheter (a thin, soft tube) into a large vein in the chest. This is sometimes called a venous access device (VAD) or central venous catheter (CVC). The catheter is inserted surgically while the patient is sedated (sleepy) or under general anesthesia (in a deep sleep). One end of the catheter stays in the vein, while the other end lies just under or outside the skin. This lets the health care team give chemo and other drugs and draw blood samples without having to stick needles into the veins each time. The catheter usually remains in place for
several months and can make having chemo less painful. If such a device is used, the health care team will teach you how to care for it to reduce the risk of problems such as infections.

**Side effects of chemo**

Chemo drugs can cause side effects³. Children tend to have less severe side effects from chemo than adults and often recover from side effects more quickly. Because of this, doctors can give them higher doses of chemo to try to kill the tumor.

The side effects of chemo drugs depend on the type, dose, and the length of time they are taken.

**General side effects of chemo**

- Nausea and vomiting
- Loss of appetite
- Diarrhea
- Hair loss
- Mouth sores

Chemo can damage the bone marrow, where new blood cells are made. This can lead to low blood cell counts, which can result in:

- Increased chance of infection (from a shortage of white blood cells)
- Bleeding or bruising after minor cuts or injuries (from a shortage of platelets)
- Fatigue or shortness of breath (from low red blood cell counts)

A major concern with chemo used to treat osteosarcoma is that it can lead to dangerously low white blood cell levels and an increased risk of serious infections. Because of this, drugs called growth factors (such as filgrastim, also known as G-CSF) may be given along with the chemo to help the body make new white blood cells as quickly as possible.

Most of the side effects above tend to go away after treatment is finished. Often there are ways to make these side effects less severe. For example, drugs can be given to help prevent or reduce nausea and vomiting, or to help get blood counts back to normal levels. Be sure to discuss any questions you have about side effects with the cancer care team, and tell them about any side effects so that they can be controlled.
Side effects of certain chemo drugs

Some side effects are specific to certain drugs. Many of these side effects are uncommon, but they are possible. Before treatment, ask your cancer care team about the possible side effects of the drugs you or your child will be getting.

- **Ifosfamide** and cyclophosphamide can damage the lining of the bladder, which can cause blood in the urine. The chance of this happening can be lowered by giving a drug called mesna during chemotherapy, along with plenty of fluids.
- **Cisplatin** and carboplatin can cause nerve damage (called neuropathy) leading to numbness, tingling, or pain in the hands and feet. This often goes away or gets better once treatment is stopped, but it might last a long time in some people. These drugs can sometimes affect hearing, especially of high-pitched sounds. Kidney damage can also occur after treatment. Giving lots of fluid before and after the drug is infused can help prevent this.
- **Etoposide** can also cause nerve damage. It can also increase the risk of later developing acute myeloid leukemia (AML), a cancer of white blood cells. Fortunately, this is not common.
- **High-dose methotrexate** can damage the white matter of the brain (called leukoencephalopathy) and can also affect the liver and kidneys. Before starting high-dose methotrexate, medicines are given to help protect the kidneys. Methotrexate blood levels may be checked to see how much leucovorin (also called folinic acid) should be given to help limit any damage to normal tissues.
- **Doxorubicin (Adriamycin)** can damage the heart muscle. The risk of this goes up with the total amount of the drug that is given, so doctors are careful to limit the total dose. Your (child’s) doctor may order a heart function test (such as an echocardiogram) before and during treatment to see if this drug is affecting the heart. A drug called dexrazoxane may be given along with the chemo to help lessen the possible damage.

Some chemo drugs can affect your (child’s) ability to have children (fertility) later in life. Ask the cancer care team about the possible effects of treatment on fertility, and ask if there are options for preserving fertility, such as sperm banking or egg preservation.

The doctors and nurses will watch closely for side effects. Don’t hesitate to ask the cancer care team any questions about side effects.

For more information on the possible late or long-term side effects of chemo, including
infertility and second cancers, see Late Effects of Childhood Cancer Treatment\(^7\).

**Tests to check for side effects of chemo:** Before each treatment, lab test results will be checked to be sure the liver, kidneys, and bone marrow are working well. Other tests might be done during and after treatment as well.

- The **complete blood count (CBC)** checks levels of white blood cells, red blood cells, and blood platelets. These will be watched closely during and after chemo. The white blood cells and platelets usually reach their lowest point about 2 weeks after chemo is given, though this can occur earlier with high-dose regimens.
- **Blood chemistry tests** measure certain blood chemicals that tell doctors how well the liver and kidneys are working. Some chemo drugs can damage these organs.
- An **audiogram** might be done to check hearing, which can be affected by certain chemo drugs.
- If doxorubicin is to be given, tests such as an **echocardiogram** (an ultrasound of the heart) may be done before and during treatment to check heart function.

For more information, see Chemotherapy\(^8\).

**Other drugs to treat osteosarcoma**

If chemo drugs are no longer helpful, other types of drugs might be an option in some situations.

**Targeted therapy drugs**

Doctors are now studying newer drugs that target specific parts of tumor cells (or nearby cells) as a way to treat osteosarcoma. These targeted drugs work differently from standard chemo drugs. They might be helpful sometimes when chemo drugs are no longer working.

For example, **regorafenib (Stivarga)**, **sorafenib (Nexavar)**, and **cabozantinib (Cabometyx)** are drugs that affect a tumor’s ability to develop new blood vessels, which it needs to grow. These drugs have been shown to help some people with osteosarcoma in early studies. Although these drugs are not yet FDA-approved to treat osteosarcoma, they are approved to treat other types of cancer, and trying one of them might be an option if standard chemo drugs are no longer helpful.

Common side effects of these drugs can include fatigue, loss of appetite, hand-foot syndrome (redness and irritation of the hands and feet), high blood pressure, weight
loss, diarrhea, and abdominal (belly) pain. Less common but more serious side effects can include problems with blood flow to the heart, bleeding, abnormal thyroid test results, and perforations (holes) in the stomach or intestines.

For more on this type of treatment, see Targeted Therapy⁹.

**Radioactive drugs (radiopharmaceuticals)**

Drugs that include a radioactive element can sometimes be helpful in people with advanced osteosarcoma. These drugs are injected into the blood and travel to the bones, where they give off small amounts of radiation that can help slow tumor growth and treat symptoms such as pain. For more information, see Radiation Therapy for Osteosarcoma.

**Hyperlinks**


**References**

Radiation Therapy for Osteosarcoma

Radiation therapy uses high-energy rays or particles to kill cancer cells.

Osteosarcoma cells are not easily killed by radiation, so radiation therapy doesn’t play a major role in treating this type of cancer. But sometimes radiation can be useful if the tumor can’t be removed completely by surgery. For example, osteosarcoma can start in hip bones or in the bones of the face, particularly the jaw. In these situations, often it’s not possible to remove all of the cancer. After as much of the tumor is removed as possible, radiation is given to try to kill the remaining cancer cells. Chemotherapy is then often given after the radiation.

Radiation can also be used to help slow tumor growth and control symptoms like pain.
and swelling if surgery is not possible, or if the cancer has come back\textsuperscript{1}.

**External beam radiation therapy**

This is the type of radiation therapy most often used to treat osteosarcoma. A machine outside the body focuses high-energy beams on the tumor to kill the cancer cells.

Before treatments start, the radiation team takes careful measurements with imaging tests such as MRI scans to determine the correct angles for aiming the radiation beams and the proper dose of radiation. This planning session is called **simulation**.

Most often, radiation treatments are given 5 days a week for several weeks. Each treatment is much like getting an x-ray, although the dose of radiation is much higher. The treatment is not painful. For each session, you (or your child) will lie on a special table while a machine delivers the radiation from precise angles.

Each treatment lasts only a few minutes, although the setup time – getting into place for treatment – usually takes longer. Young children may be given medicine to make them sleep so they will not move during the treatment.

Newer techniques, such as **intensity modulated radiation therapy (IMRT)**, **conformal proton beam therapy**, and **stereotactic radiosurgery (SRS)**, let doctors aim the radiation at the tumor more precisely while reducing how much nearby healthy tissues get. This may offer a better chance of increasing the success rate and reducing side effects. Many doctors now recommend using these approaches when they are available. (See [What’s New in Osteosarcoma Research?\textsuperscript{2}])

**Possible side effects of radiation therapy**

The side effects of external radiation therapy depend on the dose of radiation and where it is aimed.

**Short-term problems** can include effects on skin areas that receive radiation, which can range from mild sunburn-like changes and hair loss to more severe skin reactions. Radiation to the abdomen or pelvis can cause nausea\textsuperscript{3}, diarrhea, and urinary problems. Talk with your (child’s) health care team about the possible side effects because there may be ways to relieve some of them.

In children, radiation therapy can slow bone growth. For example, radiation to the bones in one leg might result in it being shorter than the other. Radiation to the facial bones may cause uneven growth, which might affect how a child looks. But if a child is
fully or almost fully grown, this is less likely to be an issue.

Depending on where the radiation is aimed, it can also damage other organs:

- Radiation to the **chest wall or lungs** can affect lung and heart function.
- Radiation to the **jaw area** might affect the salivary glands, which could lead to dry mouth and tooth problems.
- Radiation therapy to the **spine or skull** might affect the nerves in the spinal cord or brain. This could lead to nerve damage, headaches, and trouble thinking, which usually become most serious 1 or 2 years after treatment. Radiation to the spine might cause numbness or weakness in part of the body.
- Radiation to the **pelvis** can damage the bladder or intestines, which can lead to problems with urination or bowel movements. It can also damage reproductive organs, which could affect a child’s fertility later in life, so doctors do their best to protect these organs by shielding them from the radiation or moving them out of the way whenever possible.

Another major concern with radiation therapy is that it might cause a new cancer to form in the part of the body that was treated with the radiation. The higher the dose of radiation, the more likely this is to occur, but the overall risk is small and should not keep children who need radiation from getting it.

To lower the risk of serious long-term effects from radiation, doctors try to use the lowest dose of radiation therapy that is still effective. Still, it’s important to continue follow-up visits with your (child’s) doctor so that if problems come up they can be found and treated as early as possible.

**Radioactive drugs (radiopharmaceuticals)**

Bone-seeking radioactive drugs, such as **samarium-153-EDTMP (Quadramet)** or **radium-233 (Xofigo)**, are sometimes used to slow tumor growth and treat symptoms such as pain in people with advanced osteosarcoma. These drugs are injected into a vein and collect in the bones. Once there, the radiation they give off kills the cancer cells.

These drugs are especially helpful when cancer has spread to many bones, since external beam radiation would need to be aimed at each affected bone. In some cases, these drugs are used together with external beam radiation aimed at the most painful **bone metastases**.
The major side effect of these drugs is a lowering of blood cell counts, which could increase the risk for infections or bleeding, especially if the blood counts are already low.

For more detailed information on radiation therapy, see Radiation Therapy.

Hyperlinks


References


Treatment Based on the Extent of the Osteosarcoma

Treatment for osteosarcoma depends on several factors, including the location, extent, and grade of the cancer, if doctors think it can be removed (resected) completely, and on a person’s age and overall health.

Localized, resectable osteosarcoma

These cancers have not been found to have spread to other parts of the body, and all of the visible tumor can be removed (resected) by surgery.

High grade: Most osteosarcomas are high grade, meaning they will probably grow and spread quickly if not treated. The usual treatment for these cancers is as follows:

- Biopsy to establish the diagnosis
- Chemotherapy (chemo), usually for about 10 weeks
- Surgery to remove the tumor, preferably by the same surgeon who did the biopsy. If cancer is found at the edge of the surgery specimen (meaning some cancer might have been left behind), a second surgery might be done to try to remove any remaining cancer. Radiation therapy might be given to the area as well.
- More chemo (for up to a year). If the initial chemo killed most of the cancer cells, the same drugs are often given again after surgery. If the initial chemo didn’t work well, different drugs might be tried (although not all doctors agree that switching drugs is needed).

Chemo is a very important part of treatment for these cancers. Even when imaging tests don’t show that the cancer has spread to distant parts of the body, there are still likely to
be very small areas of cancer spread that can’t be detected with tests. If chemo isn’t given, the cancer is more likely to come back after surgery.

In rare cases, surgery might be the first treatment (before chemo), especially for people who are elderly.

**Low grade:** A small number of osteosarcomas are low grade, meaning they are likely to grow slowly. Patients with low-grade, resectable osteosarcomas can often be cured with surgery alone (without chemo). However, if the tumor removed by surgery is found to be high grade on lab tests, chemo might then be recommended.

**Localized, non-resectable osteosarcoma**

These cancers have not been found to have spread to other parts of the body, but they can’t be removed (resected) completely by surgery. For example, they may be too large or too close to vital structures in the body to be resected completely.

As with other osteosarcomas, a biopsy is needed first to establish the diagnosis.

**Chemotherapy** is usually the first treatment for these cancers. If the tumor shrinks enough to become resectable, it is then removed with surgery. This is followed by more chemotherapy for up to a year.

If the tumor still can’t be removed completely after chemotherapy, **radiation therapy** can often be used to try to keep the tumor under control and to help relieve symptoms. More chemo might be another option, either instead of or after radiation therapy. If the first chemo regimen didn’t work very well, different chemo drugs might be tried.

Because these tumors can be hard to treat, a [clinical trial](#) of newer treatments may be a good option in many cases.

**Metastatic osteosarcoma**

These cancers have already spread to distant parts of the body when they are diagnosed. Most often they have spread to the **lungs**. As with other osteosarcomas, a biopsy is needed first to establish the diagnosis.

**Chemotherapy** is usually the first treatment for these cancers. If all of the tumors are thought to be resectable after chemotherapy, they are removed with surgery, sometimes in more than one operation. This is followed by more chemo for up to a year.
If some of the tumors remain unresectable after chemo, radiation therapy can often be used to try to keep them under control and to help relieve symptoms. More chemo might be another option, either instead of or after radiation therapy. If the first chemo regimen didn’t work very well, different chemo drugs might be tried.

Newer targeted therapy drugs such as regorafenib (Stivarga), sorafenib (Nexavar), or cabozantinib (Cabometyx) might also be an option at some point as well, although more research will be needed to see how effective these drugs are (see What’s New in Osteosarcoma Research?4).

Because these tumors can be hard to treat, clinical trials6 of newer treatments may be a good option in many cases.

**Recurrent osteosarcoma**

Recurrent cancer means that the cancer has come back after treatment. It may come back locally (near where the first tumor was) or in other parts of the body. Most of the time, if osteosarcoma recurs it will be in the lungs.

If possible, surgery to remove the tumor(s) is an important part of treatment, as it offers the best chance for long-term survival. If the cancer recurs at the original site on an arm or leg after limb-sparing surgery, amputation of the limb may be recommended.

Chemotherapy is often part of the treatment for recurrent cancers as well. If the cancer is not resectable, chemo might be used to try to shrink the tumor(s), which might then allow surgery to be done. If the cancer is resectable, chemo might be given after surgery. For more advanced cancers, chemo might be used to try to help relieve symptoms.

Radiation therapy might be part of treatment as well. It can sometimes help keep tumor growth in check and help relieve symptoms.

If the cancer is still growing, newer targeted therapy drugs such as regorafenib (Stivarga), sorafenib (Nexavar), or cabozantinib (Cabometyx) might be an option at some point as well, although more research is needed to see how effective these drugs are (see What’s New in Osteosarcoma Research?6).

Because these tumors can be hard to treat, clinical trials7 of newer treatments may be a good option.
The treatment information given here is not official policy of the American Cancer Society and is not intended as medical advice to replace the expertise and judgment of your cancer care team. It is intended to help you and your family make informed decisions, together with your doctor. Your doctor may have reasons for suggesting a treatment plan different from these general treatment options. Don’t hesitate to ask him or her questions about your treatment options.

Hyperlinks


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Last Revised: October 8, 2020

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