About Your Total Pelvic Exenteration

This guide will help you get ready for your total pelvic exenteration (eg-ZEN-teh-RAY-shun) at Memorial Sloan Kettering (MSK). It will also help you understand what to expect during your recovery.

Use this guide as a source of information in the days leading up to your surgery. Bring it with you on the day of your surgery. You and your healthcare team will refer to it as you learn more about your recovery.

Your surgery

Surgery date: ___________________________________________________

Surgery location: Presurgical Center (PSC) on the 6th floor
1275 York Avenue (between East 67th and East 68th Streets)
New York, NY 10065
Take the B elevator to the 6th floor.

Expected discharge date: ________________________________________

It’s best to plan your ride home from the hospital ahead of time. This will help you keep from waiting for your ride after you’re discharged from (leave) the hospital. For more information, read the section “Leaving the hospital.”

Visit www.mskcc.org/pe/pelvic_exenteration to view this guide online.
# Table of contents

## About Your Surgery
- About your urinary system ......................................................... 3
- About your gastrointestinal system ........................................ 4
- About your gynecologic system .................................................. 5
- Meeting with your healthcare team ........................................... 6

## Before Your Surgery
- Getting ready for your surgery .................................................. 8
- Within 30 days of your surgery .................................................. 10
- 7 days before your surgery .......................................................... 12
- 2 days before your surgery ........................................................... 13
- 1 day before your surgery ............................................................. 13
- The morning of your surgery ....................................................... 16

## After Your Surgery
- In the Post-Anesthesia Care Unit (PACU) ...................................... 22
- In your hospital room ................................................................. 24
- At home ....................................................................................... 27
- When to call your healthcare provider ........................................ 31

## Support Services
- Gynecologic surgery support services ..................................... 34
- MSK support services ................................................................. 34
- External support services ............................................................ 37

## Educational Resources
- Caring for Your Ileostomy or Colostomy ..................................... 39
- Common Medications Containing Aspirin, Other Nonsteroidal Anti-inflammatory Drugs (NSAIDs), or Vitamin E
- Herbal Remedies and Cancer Treatment
- How to Use Your Incentive Spirometer
About Your Surgery

A total pelvic exenteration is a surgery to remove organs from your urinary, gastrointestinal, and gynecologic systems. You may be having this surgery because you have cancer in your cervix or another organ in your urinary, gastrointestinal, or gynecologic systems.

About your urinary system

Your urinary system is made up of organs that make urine (pee) and get it out of your body. It includes your:

- Kidneys. These filter toxins out of your blood and make urine.
- Ureters. These carry urine from your kidneys to your bladder.
- Bladder. This stores your urine until you feel like you need to urinate.
- Urethra. This carries urine from your bladder out of your body. In females, it’s located in front of the vagina.

Figure 1 shows the parts of your urinary system.

During your surgery, your surgeon will remove your bladder and urethra. This means you’ll need a new place for urine to leave your body. Your surgeon will connect your ureters to a new urinary collection system called a urinary diversion or urostomy.
Types of urinary diversions

There are 2 types of urinary diversions: an ileal conduit and a urinary pouch. Each type opens onto your abdomen (belly). The opening is called a stoma.

- With an ileal conduit, your urine will drain into a bag that you put around your stoma. You'll drain the bag as needed.
- With a urinary pouch, you'll put a drainage catheter (thin, flexible tube) into the stoma several times a day to drain your urine.

Your healthcare provider will talk with you about which urinary diversion you'll have. Your nurse will give you a resource that describes the type of diversion you'll have. They'll also teach you how to care for it.

About your gastrointestinal system

Your gastrointestinal system includes your:

- Mouth
- Esophagus (food pipe)
- Small intestine
- Colon (large intestine)
- Rectum (bottom end of your colon)
- Anus

Figure 2 shows the parts of your gastrointestinal system.

During your surgery, your surgeon will remove part of your colon. They'll also remove your rectum and anus. This means you'll need a new place for stool (poop) to leave your body.

Your surgeon will bring the end of your remaining colon to the outside of your abdomen (belly). This creates a colostomy, which is a new place for stool to leave your body. The opening itself is called a stoma. A pouch will cover the stoma to collect your stool.
Your healthcare team will teach you how to care for your stoma as you recover. Read the resource *Caring for Your Ileostomy or Colostomy* for more information about living with a colostomy. You can find it in the “Educational Resources” section of this guide.

**About your gynecologic system**

Your gynecologic system includes your:

- Ovaries
- Fallopian tubes
- Uterus
- Cervix
- Vagina

Figure 3 shows the parts of your gynecologic system.

During your surgery, your surgeon will remove your ovaries, fallopian tubes, uterus, and cervix. This means you won’t be able to have children naturally. If you’d like information about fertility (ability to get pregnant) options and your cancer treatment, read the resource *Fertility Preservation: Options for Females Starting Cancer Treatment*. You can ask your healthcare provider for a copy of find it online at www.mskcc.org/pe/fertility_starting_treatment

Your surgeon may also remove all or part of your vagina during your surgery. They’ll talk with you about this if it’s needed.

For some people, the vagina can be reconstructed. Ask your surgeon if this is an option for you. If it is, you’ll meet with a plastic surgeon to discuss this surgery. The reconstruction would be done at the end of your surgery. Your plastic surgeon will create a new vaginal canal from the muscles and skin of other areas of your body. Your nurse will tell you about how to care for your new vagina. If you don’t want your vagina reconstructed, the area can be closed or covered with a flap of skin.

If your vagina isn’t removed, you should be able to have vaginal intercourse (sex). If you have vaginal reconstruction, you should be able to have vaginal intercourse again when the area heals. If you don’t have vaginal reconstruction, remember that sex can include
other forms of intimacy. Ask your surgeon if your clitoris will be removed and how much feeling you’ll still have in your vaginal area.

Your healthcare providers will tell you what to expect after surgery. For example, sensations in your vaginal area may not be the same as they were before your surgery. Your surgeon can refer you to our Female Sexual Medicine and Women’s Health Program so you can meet with a therapist and discuss your concerns.

**Meeting with your healthcare team**

You’ll see many different healthcare providers before your surgery. Each one will describe their role in your care and answer your questions. You may see team members such as:

- Surgeons from other departments who will work with your surgeon to complete your surgery.
- A psychologist who works with people having this surgery.
- A wound, ostomy, and continence (WOC) nurse who will talk with you about the stoma you’ll have.

It may be helpful for you to write down questions as you think of them. Bring your list of questions to your appointments.

You may also be able to speak with someone who has had the surgery. Your social worker can help plan this for you.
Before Your Surgery

The information in this section will help you get ready for your surgery. Read this section when your surgery is scheduled and refer to it as your surgery date gets closer. It has important information about what you need to do before your surgery.

As you read through this section, you can use the space below to write down any questions you want to ask your healthcare provider.

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Getting ready for your surgery

You and your healthcare team will work together to get ready for your surgery.

Help us keep you safe during your surgery by telling us if any of the following statements apply to you, even if you aren’t sure.

☐ I take a blood thinner, such as:
  - Aspirin
  - Heparin
  - Warfarin (Jantoven® or Coumadin®)
  - Clopidogrel (Plavix®)
  - Enoxaparin (Lovenox®)
  - Dabigatran (Pradaxa®)
  - Apixaban (Eliquis®)
  - Rivaroxaban (Xarelto®)

There are others, so be sure your healthcare provider knows all the medications you’re taking.

☐ I take prescription medications (medications my healthcare provider prescribes), including patches and creams.

☐ I take over-the-counter medications (medications I buy without a prescription), including patches and creams.

☐ I take dietary supplements, such as herbs, vitamins, minerals, or natural or home remedies.

☐ I have a pacemaker, automatic implantable cardioverter-defibrillator (AICD), or other heart device.

☐ I have sleep apnea.

☐ I’ve had a problem with anesthesia (medication to make me sleep during surgery) in the past.

☐ I’m allergic to certain medication(s) or materials, including latex.

☐ I’m not willing to receive a blood transfusion.

☐ I drink alcohol.

☐ I smoke or use an electronic smoking device (such as a vape pen, e-cigarette, or Juul®).

☐ I use recreational drugs.

About drinking alcohol

The amount of alcohol you drink can affect you during and after your surgery. It’s important to talk with your healthcare providers about how much alcohol you drink. This will help us plan your care.

- If you stop drinking alcohol suddenly, it can cause seizures, delirium, and death. If we know you’re at risk for these complications, we can prescribe medications to help keep them from happening.
• If you drink alcohol regularly, you may be at risk for other complications during and after your surgery. These include bleeding, infections, heart problems, and a longer hospital stay.

Here are things you can do before your surgery to keep from having problems:

• Be honest with your healthcare providers about how much alcohol you drink.
• Try to stop drinking alcohol once your surgery is planned. If you develop a headache, nausea (feeling like you’re going to throw up), increased anxiety, or can’t sleep after you stop drinking, tell your healthcare provider right away. These are early signs of alcohol withdrawal and can be treated.
• Tell your healthcare provider if you can’t stop drinking.
• Ask your healthcare provider questions about drinking and surgery. As always, all your medical information will be kept confidential.

About smoking
If you smoke, you can have breathing problems when you have surgery. Stopping even for a few days before surgery can help. If you smoke, your healthcare provider will refer you to our Tobacco Treatment Program. You can also reach the program by calling 212-610-0507.

About sleep apnea
Sleep apnea is a common breathing disorder that causes you to stop breathing for short periods of time while sleeping. The most common type is obstructive sleep apnea (OSA). With OSA, your airway becomes completely blocked during sleep. OSA can cause serious problems during and after surgery.

Please tell us if you have sleep apnea or if you think you might have it. If you use a breathing device (such as a CPAP device) for sleep apnea, bring it with you the day of your surgery.

Using MyMSK
MyMSK (my.mskcc.org) is your MSK patient portal. You can use MyMSK to send and receive messages from your healthcare team, view your test results, see your appointment dates and times, and more. You can also invite your caregiver to create their own account so they can see information about your care.

If you don’t have a MyMSK account, you can visit my.mskcc.org, call 646-227-2593, or call your doctor’s office for an enrollment ID to sign up. You can also watch our video How to Enroll in MyMSK: Memorial Sloan Kettering’s Patient Portal
Within 30 days of your surgery

Presurgical Testing (PST)

You’ll have a PST appointment before your surgery. The date, time, and location will be printed on the appointment reminder from your surgeon’s office. You can eat and take your usual medications the day of your appointment.

It’s helpful to bring the following things to your PST appointment:

- A list of all the medications you’re taking, including prescription and over-the-counter medications, patches, and creams.
- Results of any tests done outside of MSK, such as a cardiac stress test, echocardiogram, or carotid doppler study.
- The name(s) and telephone number(s) of your healthcare provider(s).

During your PST appointment, you’ll meet with a nurse practitioner (NP). They work closely with anesthesiology staff (specialized healthcare providers who will give you anesthesia during your surgery). Your NP will review your medical and surgical history with you. You may have tests, such as an electrocardiogram (EKG) to check your heart rhythm, a chest x-ray, blood tests, and any other tests needed to plan your care. Your NP may also recommend that you see other healthcare providers.

Your NP will talk with you about which medications you should take the morning of your surgery.

Identify your caregiver

Your caregiver plays an important role in your care. Before your surgery, you and your caregiver will learn about your surgery from your healthcare providers. After your surgery, your caregiver will take you home when you’re discharged from the hospital. They’ll also help you care for yourself at home.
For caregivers
Resources and support are available to help manage the responsibilities that come with caring for a person going through cancer treatment.

For information, visit www.mskcc.org/caregivers or read A Guide for Caregivers. You can ask your healthcare provider for a copy or find it online at www.mskcc.org/pe/guide_caregivers

Complete a Health Care Proxy form
If you haven’t already completed a Health Care Proxy form, we recommend you complete one now. If you’ve already completed one or have any other advance directives, bring them to your next appointment.

A health care proxy is a legal document that identifies the person who will speak for you if you can’t communicate for yourself. The person you identify is called your health care agent.

Talk with your healthcare provider if you’d like to complete a health care proxy. You can also read the resources Advance Care Planning and How to Be a Health Care Agent for information about health care proxies, other advance directives, and being a health care agent. You can find them online at www.mskcc.org/pe/advance_care_planning and www.mskcc.org/pe/health_care_agent or ask your healthcare provider for a copy.

Do breathing and coughing exercises
Practice taking deep breaths and coughing before your surgery. Your healthcare provider will give you an incentive spirometer to help expand your lungs. For more information, read the resource How to Use Your Incentive Spirometer. You can find it in the “Educational Resources” section of this guide.

Buy a 4% chlorhexidine gluconate (CHG) solution antiseptic skin cleanser (such as Hibiclens®)
4% CHG solution is a skin cleanser that kills germs for 24 hours after you use it. Showering with it before your surgery will help lower your risk of infection after surgery. You can buy a 4% CHG solution antiseptic skin cleanser at your local pharmacy without a prescription.
Buy bowel preparation supplies, if needed

Your healthcare provider may tell you to do a bowel preparation (clear the stool from your body) before your surgery. If they do, they’ll tell you what to do. You’ll need to buy the following supplies:

- 1 (238-gram) bottle of polyethylene glycol (MiraLAX®). You can buy this at your local pharmacy. You don’t need a prescription.
- 1 (64-ounce) bottle of a clear liquid. For examples of clear liquids, read the section “Follow a clear liquid diet, if needed.”
- Extra clear liquids to drink while you’re following a clear liquid diet, if needed.

Your healthcare provider will give you a prescription for antibiotics to take as part of your bowel preparation. Be sure to also pick these up.

7 days before your surgery

Follow your healthcare provider’s instructions for taking aspirin

If you take aspirin or a medication that contains aspirin, you may need to change your dose or stop taking it 7 days before your surgery. Aspirin can cause bleeding.

Follow your healthcare provider’s instructions. **Don’t stop taking aspirin unless they tell you to.** For more information, read the resource *Common Medications Containing Aspirin, Other Nonsteroidal Anti-inflammatory Drugs (NSAIDs), or Vitamin E*. You can find it in the “Educational Resources” section of this guide.

Stop taking vitamin E, multivitamins, herbal remedies, and other dietary supplements

Stop taking vitamin E, multivitamins, herbal remedies, and other dietary supplements 7 days before your surgery. These things can cause bleeding. For more information, read the resource *Herbal Remedies and Cancer Treatment*. You can find it in the “Educational Resources” section of this guide.
2 days before your surgery

Stop taking nonsteroidal anti-inflammatory drugs (NSAIDs)

Stop taking NSAIDs, such as ibuprofen (Advil® or Motrin®) and naproxen (Aleve®), 2 days before your surgery. These medications can cause bleeding. For more information, read the resource Common Medications Containing Aspirin, Other Nonsteroidal Anti-inflammatory Drugs (NSAIDs), or Vitamin E. You can find it in the “Educational Resources” section of this guide.

1 day before your surgery

Follow a clear liquid diet, if needed

Your healthcare provider will tell you if you need to follow a clear liquid diet the day before your surgery. A clear liquid diet includes only liquids you can see through. Examples are listed in the “Clear liquid diet” table. While you’re following this diet:

- Don’t eat any solid foods.
- Try to drink at least 1 (8-ounce) glass of clear liquid every hour while you’re awake.
- Drink different types of clear liquids. Don’t just drink water, coffee, and tea.
- Don’t drink sugar-free liquids unless you have diabetes and a member of your healthcare team tells you to.

For people with diabetes

If you have diabetes, ask the healthcare provider who manages your diabetes what you should do while you’re following a clear liquid diet.

- If you take insulin or another medication for diabetes, ask if you need to change the dose.
- Ask if you should drink sugar-free clear liquids.
- Make sure to check your blood sugar level often while you’re following a clear liquid diet. If you have any questions, talk with your healthcare provider.
Clear liquid diet

<table>
<thead>
<tr>
<th>Drink</th>
<th>Do Not Drink</th>
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<tr>
<td><strong>Soups</strong></td>
<td>• Clear broth, bouillon, or consommé</td>
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| **Sweets** | • Gelatin (such as Jell-O®)  
• Flavored ices  
• Hard candies (such as Life Savers®) | • All others |
| **Drinks** | • Clear fruit juices (such as lemonade, apple, cranberry, and grape juices)  
• Soda (such as ginger ale, 7-Up®, Sprite®, and seltzer)  
• Sports drinks (such as Gatorade®)  
• Black coffee  
• Tea  
• Water | • Juices with pulp  
• Nectars  
• Milk or cream  
• Alcoholic drinks |

Start your bowel preparation, if needed

Your healthcare provider will tell you if you need to do a bowel preparation the day before your surgery. If you do, follow the instructions below.

The morning of the day before your surgery, mix all 238 grams of MiraLAX with 64 ounces of clear liquid until the MiraLAX powder dissolves. Once the powder is dissolved, you can put the mixture in the refrigerator if you want to.

At 5:00 PM on the day before your surgery, start drinking the MiraLAX mixture. It will cause frequent bowel movements, so make sure you’re near a bathroom.

- Drink 1 (8-ounce) glass of the mixture every 15 minutes until it’s gone.
- When you finish the MiraLAX mixture, drink 4 to 6 glasses of clear liquids. You can keep drinking clear liquids until midnight, but you don’t have to.
- Apply zinc oxide ointment or Desitin® to the skin around your anus after every bowel movement. This helps prevent irritation.

At 7:00 PM on the day before your surgery, take your antibiotics as instructed.

At 11:00 PM on the day before your surgery, take your antibiotics as instructed.
Note the time of your surgery

A staff member from the Admitting Office will call you after 2:00 PM the day before your surgery. If your surgery is scheduled for a Monday, they’ll call you on the Friday before. If you don’t get a call by 7:00 PM, call 212-639-5014.

The staff member will tell you what time to arrive at the hospital for your surgery. They’ll also remind you where to go. This will be the following location:

- Presurgical Center (PSC) on the 6th floor
- 1275 York Avenue (between East 67th and East 68th Streets)
- New York, NY 10065
- Take the B elevator to the 6th floor.

Shower with a 4% CHG solution antiseptic skin cleanser (such as Hibiclens)

The night before your surgery, shower using a 4% CHG solution antiseptic skin cleanser.

1. Use your normal shampoo to wash your hair. Rinse your head well.
2. Use your normal soap to wash your face and genital area. Rinse your body well with warm water.
3. Open the 4% CHG solution bottle. Pour some into your hand or a clean washcloth.
4. Move away from the shower stream. Rub the 4% CHG solution gently over your body from your neck to your feet. Don’t put it on your face or genital area.
5. Move back into the shower stream to rinse off the 4% CHG solution. Use warm water.
6. Dry yourself off with a clean towel after your shower.
7. Don’t put on any lotion, cream, deodorant, makeup, powder, perfume, or cologne after your shower.

Instructions for eating before your surgery

Do not eat anything after midnight the night before your surgery. This includes hard candy and gum.
The morning of your surgery

Remember, don’t eat anything after midnight the night before your surgery.

Instructions for drinking before your surgery

If your healthcare provider gave you a CF(Preop)® drink, finish it 2 hours before your scheduled arrival time. **Do not drink anything else after midnight the night before your surgery, including water.**

If your healthcare provider didn’t give you a CF(Preop) drink, you can drink a total of 12 ounces of water between midnight and 2 hours before your scheduled arrival time. **Do not drink anything else.**

Do not drink anything starting 2 hours before your scheduled arrival time. This includes water.

Take your medications as instructed

If your healthcare provider told you to take certain medications the morning of your surgery, take only those medications with a sip of water. Depending on what medications you take, this may be all, some, or none of your usual morning medications.

Shower with a 4% CHG solution antiseptic skin cleanser (such as Hibiclens)

Shower with a 4% CHG solution antiseptic skin cleanser before you leave for the hospital. Use it the same way you did the night before.

Don’t put on any lotion, cream, deodorant, makeup, powder, perfume, or cologne after your shower.
Things to remember

- Wear something comfortable and loose-fitting.
- If you wear contact lenses, wear your glasses instead. Wearing contact lenses during surgery can damage your eyes.
- Don’t wear any metal objects. Remove all jewelry, including body piercings. The equipment used during your surgery can cause burns if it touches metal.
- Leave valuable items at home.
- If you’re menstruating (have your monthly period), use a sanitary pad, not a tampon. You’ll get disposable underwear, as well as a pad if needed.

What to bring

- Your breathing device for sleep apnea (such as your CPAP device), if you have one.
- Your incentive spirometer, if you have one.
- Your Health Care Proxy form and other advance directives, if you completed them.
- Your cell phone and charger.
- Only the money you may want for small purchases (such as a newspaper).
- A case for your personal items (such as your eyeglasses, hearing aids, dentures, prosthetic devices, wig, and religious articles), if you have any.
- This guide. Your healthcare team will use it to teach you how to care for yourself after surgery.

Where to park

MSK’s parking garage is on East 66th Street between York and First Avenues. If you have questions about prices, call 212-639-2338.

To reach the garage, turn onto East 66th Street from York Avenue. The garage is about a quarter of a block in from York Avenue, on the right-hand (north) side of the street. There’s a tunnel you can walk through that connects the garage to the hospital.

There are also other garages on East 69th Street between First and Second Avenues, East 67th Street between York and First Avenues, and East 65th Street between First and Second Avenues.
Once you’re in the hospital
When you get to the hospital, take the B elevator to the 6th floor. Check in at the desk in the PSC waiting room.

Many staff members will ask you to say and spell your name and birth date. This is for your safety. People with the same or a similar name may be having surgery on the same day.

When it’s time to change for surgery, you’ll get a hospital gown, robe, and nonskid socks to wear.

Meet with a nurse
You’ll meet with a nurse before surgery. Tell them the dose of any medications you took after midnight (including prescription and over-the-counter medications, patches, and creams) and the time you took them.

Your nurse may place an intravenous (IV) line in one of your veins, usually in your arm or hand. If your nurse doesn’t place the IV, your anesthesiologist will do it in the operating room.

Meet with an anesthesiologist
You’ll also meet with an anesthesiologist before surgery. They will:

- Review your medical history with you.
- Ask if you’ve had any problems with anesthesia in the past, including nausea or pain.
- Talk with you about your comfort and safety during your surgery.
- Talk with you about the kind of anesthesia you’ll get.
- Answer your questions about your anesthesia.

Your surgeon or anesthesiologist may also talk with you about placing an epidural catheter (thin, flexible tube) in your spine (back). An epidural catheter is another way to give you pain medication after your surgery.

Get ready for your surgery
When it’s time for your surgery, you’ll need to remove your hearing aids, dentures, prosthetic device(s), wig, and religious articles, if you have them.

You’ll either walk into the operating room or a staff member will bring you there on a stretcher. A member of the operating room team will help you onto the operating bed.
and place compression boots on your lower legs. These gently inflate and deflate to help blood flow in your legs.

Once you're comfortable, your anesthesiologist will give you anesthesia through your IV line and you'll fall asleep. You’ll also get fluids through your IV line during and after your surgery.

During your surgery

After you're fully asleep, a breathing tube will be placed through your mouth and into your windpipe to help you breathe. A urinary (Foley) catheter will also be placed to drain urine (pee) from your bladder.

Once your surgery is finished, your surgeon will close your incisions with staples or sutures (stitches). They may also place Steri-Strips™ (thin pieces of surgical tape) or Dermabond® (surgical glue) over them and cover them with a bandage.

Your breathing tube is usually taken out while you’re still in the operating room.

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After Your Surgery

The information in this section will tell you what to expect after your surgery, both during your hospital stay and after you leave the hospital. You’ll learn how to safely recover from your surgery.

As you read through this section, you can use the space below to write down any questions you want to ask your healthcare provider.

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In the Post-Anesthesia Care Unit (PACU)

When you wake up after your surgery, you’ll be in the PACU. A nurse will be keeping track of your body temperature, pulse, blood pressure, and oxygen levels. You may be getting oxygen through a thin tube that rests below your nose or a mask that covers your nose and mouth. You’ll also have compression boots on your lower legs.

Pain medication

You’ll get epidural or IV pain medication while you’re in the PACU.

- If you’re getting epidural pain medication, it will be put into your epidural space (the space in your spine just outside your spinal cord) through your epidural catheter.
- If you’re getting IV pain medication, it will be put into your bloodstream through your IV line.

You’ll be able to control your pain medication using a button called a patient-controlled analgesia (PCA) device. For more information, read the resource Patient-Controlled Analgesia (PCA). You can find it online at www.mskcc.org/pe/PCA or ask one of your healthcare providers for a copy.

Tubes and drains

You’ll have many tubes, drains, pouches, and bandages on your abdomen (see Figure 4). They may include:

- Jackson-Pratt (JP) drains to drain fluid from around your incisions. They’ll be removed when your incisions stop draining.
- A catheter to drain urine from your urinary pouch.
- A drainage tube at the incision site. This serves as a safety catheter. It’s placed in case the catheter in your urinary pouch gets blocked or comes out. It will be clamped shut.

Figure 4. Drains, pouches, and bandages after your surgery
• A colostomy pouch on your colostomy stoma to collect stool.
• A urostomy pouch on your ileal conduit stoma to collect urine.

You’ll also have:
• Bandages and drains on your upper inside thighs if you had your vagina reconstructed.
• An IV line to give you fluids.
• Compression boots on your calves to help your blood flow. You’ll wear these whenever you’re in bed.

Your abdomen and colostomy stoma will become more swollen over the next few days, and then the swelling will go down.

Most of the drains and tubes will be removed over the next days to weeks. In about 3 to 6 weeks, most people will have either 1 stoma for a urinary pouch or a pouch for the ileal conduit. Your colostomy stoma will also be covered with a pouch to collect the stool (see Figure 5).

Your WOC nurse and other nursing staff will teach you how to care for your stomas and pouches as you recover.

Moving to your hospital room

Most people stay in the PACU overnight. After your stay in the PACU, a staff member will take you to your hospital room.
In your hospital room

The length of time you’re in the hospital after your surgery depends on your recovery. Most people stay in the hospital for about 4 to 7 days.

When you’re taken to your hospital room, you’ll meet one of the nurses who will care for you while you’re in the hospital. While you’re in the hospital, your healthcare providers will teach you how to care for yourself while you’re recovering from your surgery.

You won’t be able to sit for 6 to 8 weeks after your surgery. You can lie on your back or side or you can stand. Your healthcare providers will help you with this.

Managing your pain

You may have pain at your incision sites and your abdomen.

- If you have an epidural catheter, you’ll get pain medication into your epidural space. You’ll be able to control your medication using a PCA device.
- If you don’t have an epidural catheter, you’ll get pain medication in your IV line.
- Once you’re able to eat normal food, you’ll get oral pain medication (medication you swallow).

Your healthcare providers will ask you about your pain often and give you medication as needed. If your pain isn’t relieved, tell one of your healthcare providers. It’s important to control your pain so you can use your incentive spirometer and move around. Controlling your pain will help you recover better.

You’ll get a prescription for pain medication before you leave the hospital. Talk with your healthcare provider about possible side effects and when to start switching to over-the-counter pain medications.

Moving around and walking

Moving around and walking will help lower your risk for blood clots and pneumonia (lung infection). It will also help you start passing gas and having stool in your colostomy pouch. Your nurse, physical therapist, or occupational therapist will help you move around, if needed.

Read the resource Call! Don’t Fall! to learn what you can do to stay safe and keep from falling while you’re in the hospital. You can ask your healthcare provider for a copy or find it online at www.mskcc.org/pe/call_dont_fall
Exercising your lungs

It's important to exercise your lungs so they expand fully. This helps prevent pneumonia.

- Use your incentive spirometer 10 times every hour you’re awake. For more information, read the resource *How to Use Your Incentive Spirometer*. You can find it in the “Educational Resources” section of this guide.
- Do coughing and deep breathing exercises. A member of your care team will teach you how.

Eating and drinking

You won’t be able to eat for 1 to 2 days after your surgery. You’ll get fluids through an IV line during this time. Then, you’ll follow a clear liquid diet. After that, you can start eating normal foods again, as tolerated.

When you’re able to eat normal foods again, it’s important to eat a balanced diet high in protein to help you heal after surgery. Your diet should include a healthy protein source at each meal, as well as fruits, vegetables, and whole grains.

For more tips on increasing the calories and protein in your diet, read the resource *Eating Well During Your Cancer Treatment*. You can ask your healthcare provider for a copy or find it online at www.mskcc.org/pe/eating_cancer_treatment

If you have questions about your diet, ask to see a clinical dietitian nutritionist.

Caring for your drains and incisions

You'll likely have some bleeding from most of your incisions or around your drains. You may also have some discharge and tenderness around your incisions and drains for a few days. Your nurses will show you and tell you what’s normal and expected as they care for you. You’ll also start to learn how to care for your stoma(s) and incisions.

Some of the tubes and drains may be removed before you’re discharged from the hospital. If you go home with the drains, your nurses will teach you how to care for them.

If you have an incision that was closed with staples, the staples are usually taken out before you leave the hospital. Some people will have them removed at their first follow-up visit.
Showering

You can shower while you’re in the hospital. Your healthcare provider will tell you when it’s OK to start. Wash your incisions with a 4% CHG solution antiseptic skin cleanser such as Hibiclens until your staples are removed.

Planning for discharge

Before you go home, one of your healthcare providers will give you the supplies you need to care for yourself for the first month. After that, you can order your supplies from an outside source.

Your discharge nurse case manager will arrange to have a visiting nurse see you at home. The visiting nurse will help you care for your drains and stoma.

You may still have drains in when you go home. Ask your healthcare provider when they’ll be removed. Your visiting nurse will help you while you’re learning to care for your incisions and urinary diversion.

You’ll get a prescription for pain medication. Take your medication as instructed by your healthcare provider and as you need it. Don’t drive or drink alcohol while you’re taking prescription pain medication.

Read the resource *Gynecology Service: What You Should Know About Going Home* for more information about getting ready for your discharge. You can find it online at www.mskcc.org/pe/gyn_going_home or ask your healthcare provider for a copy.

Leaving the hospital

By the time you’re ready to leave the hospital, your incisions will have started to heal. Before you leave, look at your incisions with one of your healthcare providers. Knowing what they look like will help you notice any changes later.

On the day of your discharge, plan to leave the hospital around 11:00 AM. Before you leave, your healthcare provider will write your discharge order and prescriptions. You’ll also get written discharge instructions. One of your healthcare providers will review them with you before you leave.

If your ride isn’t at the hospital when you’re ready to be discharged, you may be able to wait in the Patient Transition Lounge. A member of your healthcare team will give you more information.
At home

Read the resource *What You Can Do to Avoid Falling* to learn what you can do to stay safe and keep from falling at home and during your appointments at MSK. You can find it online at www.mskcc.org/pe/avoid_falling or ask your healthcare provider for a copy.

Filling out your Recovery Tracker

We want to know how you’re feeling after you leave the hospital. To help us continue caring for you, we’ll send questions to your MyMSK account every day for 10 days after you leave the hospital. These questions are known as your Recovery Tracker.

Fill out your Recovery Tracker every day before midnight (12:00 AM). It only takes 2 to 3 minutes to complete. Your answers to these questions will help us understand how you’re feeling and what you need.

Based on your answers, we may reach out to you for more information or ask you to call your surgeon’s office. You can always contact your surgeon’s office if you have any questions. For more information, read *About Your Recovery Tracker*. You can find it online at www.mskcc.org/pe/recovery_tracker or ask your healthcare provider for a copy.

Managing your pain

People have pain or discomfort for different lengths of time. You may still have some pain when you go home and will probably be taking pain medication. Some people have soreness, tightness, or muscle aches around their incision for 6 months or longer. This doesn’t mean something is wrong.

Follow the guidelines below to help manage your pain at home.

- Take your medications as directed and as needed.
- Call your healthcare provider if the medication prescribed for you doesn’t ease your pain.
- Don’t drive or drink alcohol while you’re taking prescription pain medication. Some prescription pain medications can make you drowsy. Alcohol can make the drowsiness worse.
- As your incision heals, you’ll have less pain and need less pain medication. An over-the-counter pain reliever such as acetaminophen (Tylenol®) or ibuprofen (Advil® or Motrin®) will ease aches and discomfort.
  - Follow your healthcare provider’s instructions for stopping your prescription pain medication.
  - Don’t take more of any medication than the amount directed on the label or as instructed by your healthcare provider.
Read the labels on all the medications you’re taking, especially if you’re taking acetaminophen. Acetaminophen is an ingredient in many over-the-counter and prescription medications. Taking too much can harm your liver. Don’t take more than 1 medication that contains acetaminophen without talking with a member of your healthcare team.

- Pain medication should help you resume your normal activities. Take enough medication to do your activities and exercises comfortably. It’s normal for your pain to increase a little as you start to be more active.
- Keep track of when you take your pain medication. It works best 30 to 45 minutes after you take it. Taking it when you first have pain is better than waiting for the pain to get worse.

Some prescription pain medications (such as opioids) may cause constipation (having fewer bowel movements than usual). Talk with your healthcare provider about how to prevent and manage constipation.

Caring for your incision

Take a shower every day to clean your incision. Follow the instructions in the “Showering” section below.

It’s normal for the skin below your incision to feel numb. This happens because some of your nerves were cut during your surgery. The numbness will go away over time.

Call your healthcare provider’s office if:

- The skin around your incision is very red.
- The skin around your incision is getting redder.
- You see drainage that looks like pus (thick and milky).
- Your incision smells bad.

If you go home with staples in your incision, your healthcare provider will take them out during your first appointment after surgery. It’s OK to get them wet.

Caring for your colostomy and urinary diversion

Your nurse will teach you how to care for your colostomy and urinary diversion before you leave the hospital. Read the resource About Your Ileostomy or Colostomy for more information about caring for your colostomy. You can find it in the “Educational Resources” section of this guide. Your WOC nurse will give you information about caring for your urinary diversion.
Showering

**Take a shower every day to clean your incision.** If you have staples in your incision, it’s OK to get them wet.

Take your bandage(s) off before you shower. Use soap during your shower, but don’t put it directly on your incision. Don’t rub the area around your incision.

After you shower, pat the area dry with a clean towel. Leave your incision uncovered or cover it with a small bandage if your clothing (such as the waistline of your pants) may rub it.

Don’t take a bath for the first 4 weeks after your surgery.

Eating and drinking

You can eat all the foods you did before your surgery unless your healthcare provider gives you other instructions. Eating a balanced diet with lots of calories and protein will help you heal after surgery. Try to eat a good protein source (such as meat, fish, or eggs) at each meal. You should also try to eat fruits, vegetables, and whole grains.

It’s also important to drink plenty of liquids. Choose liquids without alcohol or caffeine. Try to drink 8 to 10 (8-ounce) glasses of liquids every day. For more tips on eating and drinking after surgery, read the resource *Eating Well During Your Cancer Treatment*. You can find it online at www.mskcc.org/pe/eating_cancer_treatment or ask your healthcare provider for a copy.

If you have questions about your diet, ask to see a clinical dietitian nutritionist.

Physical activity and exercise

When you leave the hospital, your incision will look like it’s healed on the outside, but it won’t be healed on the inside. For the first 6 to 8 weeks after your surgery:

- Don’t lift anything heavier than 10 pounds (4.5 kilograms).
- Don’t do any strenuous activities (such as jogging and tennis).
- Don’t play any contact sports (such as football).

Talk with your healthcare provider before you start doing strenuous exercise or lifting heavy objects.

Doing aerobic exercise, such as walking and stair climbing, will help you gain strength and feel better. Walk at least 2 to 3 times a day for 20 to 30 minutes. You can walk outside or indoors at your local mall or shopping center.
It’s normal to have less energy than usual after your surgery. Recovery time is different for each person. Increase your activities each day as much as you can. Always balance activity periods with rest periods. Rest is an important part of your recovery.

Sexual activity

If you have a vagina or a reconstructed vagina, don’t put anything inside it or have vaginal intercourse until your first appointment after surgery. At this appointment, your healthcare provider will tell you when you can start having vaginal intercourse again.

If your healthcare provider told you it’s OK to have vaginal intercourse but you feel pain during vaginal intercourse, call them. They can refer you to physical therapy for an evaluation.

Other activities

Before starting any activities such as driving, traveling, or going back to work, talk with your healthcare provider. They’ll help you figure out when you can start doing these again.

Managing your feelings

This surgery will change your body, and it will probably take time for you to get used to these changes. You may feel scared, angry, or worried. You may have questions or fears about how this surgery will impact your sexuality. These feelings are normal, and many people feel the same way.

Everyone gets used to these changes in their own way. For some people, it can take a few months to get used to their changed body image. For others, it can take longer. As time goes on, you’ll get stronger. You’ll be more confident in caring for your incisions and pouch. Your appetite and energy will get better. Eventually, you’ll be able to resume most of your normal activities.

We have many resources to help you. Your healthcare providers will answer your questions. Our social workers, psychologists, psychiatrists, and WOC nurses have helped many people through this change. You can have one-on-one counseling, or you can join one of our online or in-person support groups. We also have clergy available for spiritual support. You may be able to meet with other people who have had this surgery and who can talk with you and answer your questions. Talk with your healthcare provider if you’re interested in this.

To address issues with sexual health and intimacy, you can see someone from our Female Sexual Medicine & Women’s Health Program. For more information or to set up an appointment, call 646-888-5076.
When to call your healthcare provider

Call your healthcare provider if:

- You have a fever of 100.5 °F (38.0 °C) or higher.
- You have chills.
- You're having trouble breathing.
- The skin around your incision is warmer than usual.
- The skin around your incision is getting redder.
- The area around your incision is starting to swell.
- The area around your incision is getting more swollen.
- You have discharge coming from your incisions.
- You have a bad smell coming from your incisions.
- You have pain that doesn’t get better with pain medication.
- You have vaginal bleeding or bad-smelling vaginal discharge.
- You have burning when passing stool or urine.
- You can’t pass stool or urine into your stoma or pouch.
- You have any problems you didn’t expect.
- You have any questions or concerns.

Contact information

Monday through Friday from 9:00 AM to 5:00 PM, call your healthcare provider’s office.

After 5:00 PM, during the weekend, and on holidays, call 212-639-2000 and ask to speak to the person on call for your healthcare provider.
Support Services

This section has a list of support services that may help you get ready for your surgery and recover safely.

As you read through this section, you can use the space below to write down any questions you want to ask your healthcare provider.

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Gynecologic surgery support services

United Ostomy Associations of America, Inc (UOAA)
800-826-0826
www.ostomy.org
Volunteer group who supports people who have or will have an ostomy and provides educational resources.

Women's Cancer Network: Gynecological Cancer Foundation (GCF)
312-578-1439
www.foundationforwomenscancer.org
Nonprofit organization that provides education and supports research on gynecologic cancers.

Wound, Ostomy, Continence Nurses Society
888-224-WOCN (9626)
www.wocn.org
Provides patient resource from nursing members who have special training in the care of people with wounds, ostomies, and continence problems.

MSK support services

For more online information, visit the Cancer Types section of www.mskcc.org

Admitting Office
212-639-7606
Call if you have questions about your hospital admission, including requesting a private room.

Anesthesia
212-639-6840
Call if you have questions about anesthesia.

Blood Donor Room
212-639-7643
Call for information if you're interested in donating blood or platelets.

Bobst International Center
888-675-7722
MSK welcomes patients from around the world. If you're an international patient, call for help arranging your care.
Counseling Center
646-888-0200
Many people find that counseling helps them. We provide counseling for individuals, couples, families, and groups, as well as medications to help if you feel anxious or depressed. To make an appointment, ask your healthcare provider for a referral or call the number above.

Female Sexual Medicine and Women’s Health Program
646-888-5076
Cancer and cancer treatments can have an impact on your sexual health. Our Female Sexual Medicine and Women’s Health Program can help if you’re dealing with cancer-related sexual health challenges such as premature menopause or fertility issues. Call for more information or to make an appointment. We can help you take action and address sexual health issues before, during, or after your treatment.

Food Pantry Program
646-888-8055
The food pantry program provides food to people in need during their cancer treatment. For more information, talk with your healthcare provider or call the number above.

Integrative Medicine Service
646-888-0800
Integrative Medicine Service offers many services to complement (go along with) traditional medical care, including music therapy, mind/body therapies, dance and movement therapy, yoga, and touch therapy.

MSK Library
library.mskcc.org
212-639-7439
You can visit our library website or speak with the library reference staff to find more information about your specific cancer type. You can also visit LibGuides on MSK’s library website at libguides.mskcc.org

Patient and Caregiver Education
www.mskcc.org/pe
Visit the Patient and Caregiver Education website to search our virtual library. There you can find written educational resources, videos, and online programs.

Patient and Caregiver Peer Support Program
212-639-5007
You may find it comforting to speak with someone who has been through a treatment like yours. You can talk with a former MSK patient or caregiver through our Patient and Caregiver Peer Support Program. These conversations are confidential. They may take place in person or over the phone.
Patient Billing  
646-227-3378  
Call if you have questions about preauthorization with your insurance company. This is also called preapproval.

Patient Representative Office  
212-639-7202  
Call if you have questions about the Health Care Proxy form or if you have concerns about your care.

Perioperative Nurse Liaison  
212-639-5935  
Call if you have questions about MSK releasing any information while you’re having surgery.

Private Duty Nursing Office  
212-639-6892  
You may request private nurses or companions. Call for more information.

Resources for Life After Cancer (RLAC) Program  
646-888-8106  
At MSK, care doesn’t end after active treatment. The RLAC Program is for patients and their families who have finished treatment. This program has many services, including seminars, workshops, support groups, counseling on life after treatment, and help with insurance and employment issues.

Social Work  
212-639-7020  
Social workers help patients, family, and friends deal with issues that are common for cancer patients. They provide individual counseling and support groups throughout the course of treatment and can help you communicate with children and other family members. Our social workers can also help refer you to community agencies and programs, as well as financial resources if you’re eligible.

Spiritual Care  
212-639-5982  
Our chaplains (spiritual counselors) are available to listen, help support family members, pray, contact community clergy or faith groups, or simply be a comforting companion and a spiritual presence. Anyone can request spiritual support, regardless of formal religious affiliation.

MSK's interfaith chapel is located near Memorial Hospital's main lobby. It’s open 24 hours a day. If you have an emergency, call 212-639-2000. Ask for the chaplain on call.

Tobacco Treatment Program  
212-610-0507  
If you want to quit smoking, MSK has specialists who can help. Call for information.
Virtual Programs
www.mskcc.org/vp
MSK’s Virtual Programs offer online education and support for patients and caregivers, even when you can’t come to MSK in person. Through live, interactive sessions, you can learn about your diagnosis, what to expect during treatment, and how to prepare for the various stages of your cancer care. Sessions are confidential, free, and led by expert clinical staff. If you’re interested in joining a Virtual Program, visit our website at www.mskcc.org/vp for more information.

External support services

There are many other support services available to help you before, during, and after your cancer treatment. Some offer support groups and information, while others can help with transportation, lodging, and treatment costs.

Visit www.mskcc.org/pe/external_support_services for a list of these support services. You can also talk with an MSK social worker by calling 212-639-7020.
Educational Resources

This section has the educational resources mentioned in this guide. These resources will help you get ready for your surgery and recover safely after surgery.

As you read through these resources, you can use space below to write down any questions you want to ask your healthcare provider.

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Caring for Your Ileostomy or Colostomy

This information will help you care for your ileostomy or colostomy.

An ostomy is an opening created during surgery. A piece of your intestine is brought to the outside of your abdomen (belly) so that bowel movements (stool) and gas can exit your body. The part of your intestine that’s on the outside of your body is called a stoma. Your stoma will look red and moist, similar to the inside of your mouth. Your stoma will be swollen right after surgery, but the size will decrease in 6 to 8 weeks. You won’t feel any pain or pressure within your stoma. You also won’t feel any sensations of heat or cold.

Some ostomies are permanent, while others are temporary. Your surgeon will tell you if your ostomy is permanent or temporary.

Having an ostomy is a big change for most people. We have written this resource with the help of people who have an ostomy. We hope it will help prevent problems and give you tips that can make your daily life easier. Your wound, ostomy, continence (WOC) nurse will go over this information with you.

Types of Ostomies

An ostomy can be made out of the small intestine or colon (large intestine). When a piece of the small intestine is used to create an ostomy, it’s called an ileostomy. When a piece of the colon is used to create an ostomy, it’s called a colostomy.

The consistency of your bowel movements from your ostomy will depend on the part of your intestine that’s used to create the ostomy. Your bowel movements may be liquid, soft, or solid.
The most common types of ostomies are:

**Ileostomy**
- Created from the part of the small intestine called the ileum.
- Bowel movements are loose or watery.
- Stoma is usually located on the right lower side of the abdomen.

**Ascending colostomy**
- Created out of the ascending colon.
- Bowel movements are loose or watery.
- Stoma is usually located on the right side of the abdomen.

**Descending colostomy**
- Created out of the descending colon.
- Bowel movements are usually soft.
- Stoma is usually located on the left side of the abdomen.

**Sigmoid colostomy**
- Created out of the last part of the colon.
- Bowel movements can be soft or firm.
- Stoma is usually located on the left side of the abdomen.

**Loop ostomy**
- Can be made out of any part of the small intestine or colon.
- There are 2 openings in the stoma instead of 1. Many times, only 1 opening can be seen. This type of ostomy is often temporary.

In some surgeries, a second opening is created on the abdomen called a mucous fistula. It secretes left over bowel movements or a mucus-like fluid and leads to
the rectum. If you have a mucous fistula, your WOC nurse will teach you how to care for it while you’re in the hospital. Usually, a small pouch or dressing is placed over the mucous fistula. It will need to be cleaned with water and emptied. Once there is no drainage, the mucus fistula can be covered with a pad and tape.

**Learning to Care for Your Ostomy**

Your WOC nurse and nursing staff will teach you how to care for your stoma and ostomy while you’re in the hospital. When you’re discharged from the hospital, you’ll be given the basic supplies you need to care for your ostomy. During your post-operative appointment, a WOC nurse will re-evaluate your ostomy supplies and will show you how to order them. There’s a list of medical suppliers at the end of this resource.

Your WOC nurse and nursing staff will also arrange for a visiting nurse to come to your home after you’re discharged from the hospital. They will teach you how to care for your stoma and adjust to being home. If you have any questions or problems once you’re home, call your WOC nurse or social worker.

**Pouching systems**

After your surgery, you will wear an odor-proof pouching system over your ostomy. The pouching system is made up of a skin barrier (wafer) and a collection pouch. There are many different sizes and styles of pouching systems. Most pouching systems are either a 1-piece system in which the pouch is attached to the skin barrier, or a 2-piece system in which the pouch can be taken off of the skin barrier. With both types, the pouch attaches to your abdomen by the skin barrier and is fitted over and around your stoma to collect your bowel movements and gas. The skin barrier protects the skin around your stoma from getting irritated by bowel movement leakage.

Your WOC nurse will help you find the best pouching system for your stoma. As the size of your stoma changes, you may need to change your pouching system.

Some people find it helpful to buy a fitted waterproof mattress cover, bed pads, or cloth under-pads to place over the mattress and bed sheets. This protects the bed in case the pouch accidentally leaks during the night.
**Emptying your pouch**

Empty your pouch when it’s almost halfway full. The pouch shouldn’t become overly full. Your WOC nurse will teach you how to empty your pouch.

If you squeeze every bit of air out your pouch, the sides are likely to stick together and make it more difficult to collect bowel movements. There are lubricants to help keep this from happening. This is more of an issue if your bowel movements are thicker. If your bowel movements are thinner, you can use a thickening product in the pouch to help thicken them. Your WOC nurse will help you determine which product you should use.

**Changing your pouching system**

Your WOC nurse will teach you how to change your pouching system. Generally, you should change your pouching system every 3 to 5 days. If you have leakage, change it immediately. This is to protect the skin around your stoma from getting irritated from bowel movement leakage. If you have any leakage or irritation that has lasted for a few days, call your WOC nurse.

**Lifestyle Issues**

**Body image and depression**

After people learn that they have cancer, they often view their bodies in a different way. Surgery, chemotherapy, and radiation therapy can change the way a body looks and works. This may be especially true when someone has a new ostomy. Some people feel disgust. This is normal given the shame in many cultures of talking about bowel movements. Other people feel sadness at the loss or change in their bodies. They view their ostomy with negative thoughts and feelings. However, others view their ostomy more positively and see it as an answer to difficult symptoms, such as bowel incontinence (trouble controlling your bowel movements), or as the path to survival. You may have both positive and negative feelings. As you adjust to living with an ostomy and get used to how it looks and works, you’ll feel better about your body. It may help you cope better if you:

- Look at your stoma to help you get comfortable with it.
- Remind yourself about the positive aspects of your ostomy.
- Talk with other people with ostomies. You may learn good coping techniques and this may help you regain a sense of normalcy.
- Get counseling to help you improve your body image, feelings, and quality of life. At Memorial Sloan Kettering (MSK), counseling is available from social workers and at the Counseling Center. You can reach the Social Work Department at 212-639-7020 and the Counseling Center at 646-888-0200.

**Nutrition**

Your doctor will give you dietary guidelines to follow after your surgery. You’ll also meet with a dietitian while you’re in the hospital to review these guidelines.

Below are some recommended guidelines if you have an ileostomy:

- **Stay hydrated**
  - Drink 8 to 10 (8-ounce) glasses of liquids a day.
  - Limit your intake of caffeine and alcohol. They can dehydrate you.
  - Limit foods high in fiber and high in simple sugars (such as glucose, sucrose, fructose), as they can dehydrate you.

- **Keep in mind that the foods that upset your stomach before your surgery will have the same effect after your surgery.** If you feel bloating and pain after eating a certain food, remove it from your diet. Try eating it again in a few weeks. If you’re still having pain and bloating after eating this food, avoid it.

- **Some people with ostomies choose to limit gas by regulating their diet.** The following can increase gas:
  - Foods such as cabbage and broccoli
  - Carbonated beverages
  - Drinking through a straw
  - Chewing gum

- If you have tried limiting gas and still have a problem, talk with your WOC
nurse. They may suggest using pouches with charcoal filters. The filter can help prevent the buildup of gas and avoid inflation of the pouch. Some people also find it useful to use pouches with filters for airplane travel.

- Don’t prick your pouch to release built-up gas because this will destroy the pouch’s odor proof system.
- Don’t scrub your pouch with water. This will deactivate the filter.

**If you’re constipated (having fewer bowel movements than what’s normal for you):**
- Drink more liquids.
- Eat foods that are high in fiber, such as fruits, vegetables, and bran.
- Walk around.

If you have an ileostomy, don’t take any kind of laxative. It can cause severe dehydration and other problems.

**Medication**

If you have an ileostomy, your body may not be able to completely absorb some medications. This happens because your bowel is shortened and some medications are absorbed slowly. This happens most often with:

- Pills with enteric coating (a special coating that prevents pills from being absorbed until they reach the small intestine)
- Sustained-release or extended release medications (designed to release a medication in the body slowly over an extended period of time)

Ask your doctor if you’re taking these types of medications. If you see a tablet in your pouch that isn’t absorbed, call your doctor.

Don’t crush pills unless your doctor or pharmacist says you can do so. Taking crushed pills can cause problems with digestion.

**Exercise**

Talk with your doctor about what exercise is right for you. For most people,
walking and light stretching are good exercises to do while recovering from surgery. You should avoid lifting or pushing anything heavier than 10 pounds for 2 to 3 months after your surgery. This includes housework such as vacuuming or mowing the lawn. If this will be a problem for you, talk with your doctor.

Don’t do sit-ups, pushups, or any strenuous exercise for 2 to 3 months after your surgery. This will help prevent a hernia (bulging of an organ or tissue) or weakening of a muscle near your stoma. It will also help prevent a stomal hernia (bulging out of your stoma).

**Odor control**

Pouching systems are designed to be odor proof. Unless you’re leaking bowel movements underneath the wafer, there should be no odor while the pouching system is attached. However, if odor is a problem for you, there are deodorants that you can put into the pouch to help. Some examples are drops, charcoal tablets, or lubricating gel. You may need to experiment to find one that works best for you. There are also deodorants you can take orally (by mouth) to decrease your bowel movement’s odor. Check with your doctor or nurse before using any of these.

Many pouches have a built-in charcoal filter. This vents the gas and filters it at the same time to prevent odor. There are also filters that can be bought separately and attached to the pouch, or you can apply a small plastic vent on each pouch. This vent allows gas to be released without opening the pouch.

Some people find it useful to empty the pouch at bedtime. This is done because gas could continue to build overnight and fill the pouch.

**Sexual activity**

Talk with your doctor about when it’s safe to resume sexual activity. Generally, this is after your incision is healed, about 2 months after your surgery. As long as excessive pressure isn’t placed on your stoma, there should be no harm done to it. Do not use your stoma as a sexual orifice.

For many people, returning to a normal sex life will come naturally. This happens as your concerns decrease and your feelings about your body become more positive. Try to stay intimate with your partner as much as possible. Use kind
words, meaningful touch, and hugs.

Check to see if your pouching system fits well and if there’s any odor. Some people prefer to cover their pouching system with a cloth band or wrap, shirt, or lingerie during sexual activity. You can buy ready-made covers from many ostomy supply companies. Some companies also make a pouch with a cotton cover attached to it.

You may also plan your sexual activity for times when you’re less likely to be actively draining or after irrigation. For more information about irrigation, see the section titled “Frequently Asked Questions About Ostomy Care”.

Talk with your doctor or nurse if surgery or radiation therapy caused changes that make it hard to have sex. Examples are erectile dysfunction (difficulty getting or keeping an erection) in men or vaginal dryness, pain, or tightness in women. If these problems don’t get better, you can talk with a sexual health specialist. MSK has a Sexual Health Program to help people address the impact of their disease and treatment on sexuality.

The United Ostomy Associations of America, Inc. (www.ostomy.org or 800-826-0826) has information on sex for both men and women with ostomies.

Other good sources of information are the following resources published by the American Cancer Society. You can get these resources by calling 800-ACS-2345 (800-227-2345) or by going to the links below.


**Showering, bathing, and swimming**

You can shower with your pouching system on or off. However, if you have an ileostomy, you may want to keep the pouch on during the shower due to possible
drainage of bowel movements while showering. You can also use waterproof tape around the edges of the wafer. Depending on your skin and the type of pouch, you can place a piece of plastic over the whole pouching system to keep it from getting wet, but this isn’t always necessary.

When you take a bath or swim, keep your pouching system on. Most people can wear the same swimsuit they wore before their surgery. However, it may not be possible to wear hip-high swimsuits for men or hip-high, two-piece swimsuits for women.

**Work**

You can go back to work as soon as you feel ready and your doctor approves. Unless you tell them, people won’t know that you have an ostomy. It’s highly unlikely that the pouching system will be visible. You will probably be able to wear most of the same clothes you wore before your surgery.

**Travel**

Bring extra pouching systems with you wherever you go, even if it’s only a short trip or a doctor’s appointment. If you’re going to be away from home for several hours, think about where you’re going and mentally prepare yourself. Some people feel anxiety the first few times they travel. If you feel anxious, you may be able to lower your stress by thinking ahead. First, think about where the bathroom is located and what you may need. Second, get the supplies you plan to take with you. Place 2 of every item in a travel bag. This way, if you have an emergency or are away from home longer than you expect, you will be prepared. Remember to bring:

- Pouches
- Wafers
- Clamps (if you have a pouching system with a clamp closure)
- Paper towels

When you’re traveling by air:

- Pack your ostomy supplies in at least 2 places, such as in your carry-on and
• Take extra supplies in case you’re stranded where supplies may not be available.

• You may want to bring a letter from your doctor stating your need for ostomy supplies, as well as a private area be used in case of an extended search by TSA agents.

• If you’re traveling to a foreign country, it’s a good idea to have ostomy information written in that language. One of the 70 member associations of the International Ostomy Association may be able to help with this translation, as well as with locating supplies while visiting their country. For more information, visit www.ostomyinternational.org.

• You may want to carry photocopies of the catalogue pages displaying and explaining your equipment for TSA agents.

• The United Ostomy Association of America web site, (www.uoaa.org) has information and helpful ‘travel tips’ that can be printed and used.

Call Your Doctor or Nurse if You:

• Vomit

• Have a colostomy and haven’t had a bowel movement for 3 days

• Have an ileostomy and haven’t had a bowel movement for 3 to 6 hours

• Develop abdominal pain or symptoms of dehydration (this is more common for those with an ileostomy), including:
  
  ° Increased thirst
  ° Dry mouth
  ° Loss of appetite
  ° Weakness
  ° Decreased urine output
  ° Dark amber colored urine
Muscle, abdominal, or leg cramps
Feeling faint
More bowel movements than usual or changes in consistency
Increased frequency of emptying pouch

**Frequently Asked Questions About Ostomy Care**

**What is irrigation and will I be able to do it?**
Irrigation is a type of enema that eliminates the need for wearing a pouching system over the stoma. This means you wouldn’t have any bowel movements between the irrigations. Irrigation is done every day or every other day. People who have a sigmoid colostomy may be able to irrigate. Most people who have had an abdominal perineal resection have a sigmoid colostomy. **People with an ileostomy should never irrigate because it can cause dehydration.**

You also cannot irrigate:

- If you have a stomal hernia (bulge around the stoma).
- While you’re getting chemotherapy.
- While you’re getting radiation therapy.

Ask your doctor or WOC nurse for more information about irrigation.

**What color should my stoma be?**
Your stoma should be a deep pink or red color at all times. While it's working to move bowel movements out of your body, you may notice a slight change in its color. This should only be for a few seconds to a minute. If your stoma looks dark red, grey, brown, or black, call your doctor immediately.

**Is it normal for my stoma to bleed when I touch it or change my pouching system?**
Your stoma has many blood vessels, so it may bleed easily with touch or irritation.
It’s common to have a small amount of blood on the tissue when cleaning your stoma. The bleeding should stop on its own within a few minutes. If you’re on a blood thinner or aspirin, you may bleed more easily and for a longer period of time. If you bleed a lot or if the bleeding doesn’t stop after 5 to 10 minutes, apply pressure to the area and call your doctor. If you notice blood in your bowel movements or coming from the inside of your stoma, call your doctor immediately.

The area around my stoma seems to bulge. Is this normal?

If you have a change in the curve around your stoma, tell your doctor or WOC nurse. You may have a stomal hernia. This happens because the abdominal wall muscle is weak and causes a bulge to form in the area around the stoma. Your doctor or WOC nurse will examine you to see if you have a stomal hernia.

People with a stomal hernia don’t usually have any symptoms. Surgery isn’t needed unless you have a complication, including a blockage or twisting of the herniated bowel.

If you have any of the following symptoms, call your doctor or nurse:

- Nausea
- Vomiting
- Pain
- Abdominal bloating, and lack of gas and bowel movements in your pouch

If you have a stomal hernia, your WOC nurse can measure you for a custom-made hernia support belt. These belts don’t cure the hernia, but they can prevent it from getting worse by giving it support. A girdle or biking shorts with a hole cut out for your pouch may also work well as an alternative to the custom-fitted hernia belt.

How do I wear a belt over my stoma?

Try to wear the belt above or below your stoma. If the belt is worn directly over your stoma, it can cause:

- Irritation
• Bleeding

• An overgrowth of tissue

While you’re wearing the belt, position your pouch downward or to the side. You can also fold it, depending on your preference.

**What do I do if I have leakage of bowel movements under my wafer?**

If you’re having leakage, tell your WOC nurse. You may need a different pouching system that fits better to avoid this problem.

**Is there anything I should do to prepare for tests, such as scans with contrast dye?**

Always take an extra set of supplies when going to the doctor for tests because you may have to remove your pouching system.

If you live 1 hour or longer away from the hospital, be ready to change or empty your bag during the trip home. The contrast dye may empty into your pouch in 1 or 2 hours, but it can take longer.

**Do I need to wear gloves when I change my pouching system?**

This is a personal choice. There is no risk of infection from changing your pouching system. However, some people prefer to wear gloves because some bowel movements could come in contact with your fingers.

**If I have an ileostomy, is it normal to still pass bowel movements or mucous from my rectum?**

Depending on the type of surgery you had, you may pass left over bowel movements or mucous from your rectum. This should decrease over time, but it may be useful to wear a pad and apply a zinc ointment (such as Desitin®) around your anus to prevent irritation.

**What should I do if I am admitted to the hospital?**

If you’re admitted to the hospital, you may want to bring 1 or 2 extra sets of
supplies with you, just in case they stock a different product or they have run out of the product you use.

**Video Resources**

Please visit [www.mskcc.org/pe/change_ostomy](http://www.mskcc.org/pe/change_ostomy) to watch this video.

This video will teach you how to change your ostomy appliance.

Please visit [www.mskcc.org/pe/empty_pouch](http://www.mskcc.org/pe/empty_pouch) to watch this video.

This video will teach you how to empty your velcro end closure pouch.

Please visit [www.mskcc.org/pe/skin_stoma](http://www.mskcc.org/pe/skin_stoma) to watch this video.

This video will teach you how to care for irritated skin around your stoma.

**Support Groups**

You may find it helpful to join our ostomy group for colorectal patients. This is a live, online support group in which people can share their experiences and provide practical and emotional support for one another. Discussions are guided by a social worker and nurse. The group is open to everyone, including non-MSK patients. The group is held every other month on the 3rd Thursday of the month from 4:00 PM to 5:00 PM. To learn more about the program or to register for the program contact virtualprograms@mskcc.org.

On the United Ostomy Associations of America (UOAA) website, you can search for UOAA-affiliated online and in-person support groups for people with ostomies. To find a group, go to [www.ostomy.org/supportgroups.html](http://www.ostomy.org/supportgroups.html).
Local hospitals may also offer ostomy support groups.

**Ordering Supplies**

Speak with your WOC nurse about ordering supplies. They are familiar with the latest products on the market. You can also call companies to ask for samples of their products.

**Mail order medical supply companies**

Listed below are some mail order medical supply companies. Some accept Medicare and other types of insurance. Call the company directly to ask if they take your insurance. If they don’t, call your insurance company to get a list of suppliers that do. These companies will mail the supplies to you every month. You don’t usually need a prescription to order the supplies. The company will work with your doctor’s office and your insurance company. These companies offer a wide variety of products. Call and ask for their catalogue or view it online.

**Byram Healthcare**
877-902-9726
www.byramhealthcare.com

**CCS Medical**
800-260-8193
www.ccsmed.com

**Edgarpark Medical Supplies**
888-394-5375
www.edgepark.com

**Landauer Metropolitan**
800-631-3031
www.landmetro.com
(Acepts Medicaid in New York State)

**McKesson Medical-Surgical**
800-453-5180
You can also buy ostomy supplies at local pharmacies and medical supply stores. However, the prices may be higher due to a low volume of supplies or because they may not take your insurance.

**Ostomy product manufacturers**

Listed below are companies that make ostomy products such as skin barriers, pouches, and other ostomy accessories (such as pastes, powder, and rings). Many of these companies also have a WOC nurse on staff to answer questions. Feel free to call and ask for samples of products.

**Coloplast Corporation**
800-533-0464
www.coloplast.us

**Convatec Professional Services**
800-422-8811
www.convatec.com

**Cymed**
800-582-0707
www.cymed-ostomy.com

**Hollister, Inc.**
888-740-8999
www.hollister.com

**Marlen Manufacturing & Development Company**
216-292-7060
www.marlenmfg.com

**Nu-Hope Laboratories, Inc.**
800-899-5017
www.nu-hope.com
MSK Support Services

Resources for Life After Cancer (RLAC) Program
646-888-8106
At MSK, care doesn’t end after active treatment. The RLAC Program is for patients and their families who have finished treatment. This program has many services, including seminars, workshops, support groups, counseling on life after treatment, and help with insurance and employment issues.

Social Work Department
212-639-7020
Social workers help patients and their families and friends deal with issues that are common for cancer patients. They provide individual counseling and support groups throughout the course of treatment, and can help you communicate with children and other family members. Our social workers can also refer you to community agencies and programs, as well as financial resources if you’re eligible.

Sexual Health Program
For women: 646-888-5076
For men: 646-422-4359
MSK’s Sexual Health Program helps patients address the sexual impact of their disease and treatment. You can meet with a specialist before, during, or after your treatment.

Resources

Books
100 Questions & Answers about Colorectal Cancer
Authors: David Bub, Susannah L. Rose, and W. Douglas Wong
Boston, MA: Jones and Bartlett Publishers; 2003

The Ostomy Book: Living Comfortably with Colostomies, Ileostomies, and Urostomies
Authors: Barbara Dorr Mullen and Kerry Anne McGinn
Boulder, CO: Bull Publishing; 2008

The Complete Financial, Legal, and Practical Guide for Living With Cancer, HIV, and
**Other Life-Challenging Conditions**
Author: David Landay  

**Positive Options for Living with Your Ostomy: Self-Help and Treatment**
Authors: Craig A. White and Robert W. Beart Jr.  
Alameda, CA: Hunter House; 2002

**The Human Side of Cancer: Living With Hope, Coping With Uncertainty**
Authors: Jimmie Holland and Sheldon Lewis  

**Organizations**

**Cancercare**
800-813-HOPE (800-813-4673)  
www.cancercare.org  
National nonprofit organization that provides free help to people with all cancers. Services include counseling, education, information, and financial assistance.

**Caregiver Action Network**
caregiveraction.org  
Provides support to caregivers of those who are ill, elderly, or disabled. Works to provide caregivers with support, empowerment, education, and advocacy.

**Colon Cancer Alliance (CCA)**
877-422-2030 (toll-free helpline)  
www.ccalliance.org  
National nonprofit patient advocacy organization dedicated to screening, access, awareness, advocacy, and research related to colon cancer.

**Crohn’s and Colitis Foundation of America**
800-932-2423  
ww.ccfaf.org  
Works to find a cure for and prevent Crohn’s disease and ulcerative colitis. Aims to improve the quality of life for people living with these diseases through research, education, and support.
C3Life.com
www.c3life.com
Website dedicated to helping people with ostomies live their lives to the fullest. Online community includes a discussion forum, recipes, blogs, hints, and tips.

Gilda’s Club New York City
195 West Houston St.
New York, NY 10014
212-647-9700
www.gildasclubnyc.org
Provides meeting places for people living with cancer and their family and friends. Gives people a place where they can meet others to build support systems. Offers free support and networking groups, lectures, workshops, and social events.

International Ostomy Association (IOA)
www.ostomyinternational.org
Works to improve the quality of life for people who have ostomies or related surgeries.

National Ovarian Cancer Coalition (NOCC)
888-682-7426
www.ovarian.org
Works to increase awareness of ovarian cancer and to improve the survival rate and quality of life for women with ovarian cancer.

NCI’s Cancer Information Service
800-4 CANCER (800-422-6237)
www.cancer.gov/aboutnci/cis
A federally funded cancer education program that provide accurate, up-to-date, and reliable information on cancer that is easy to understand.

Ovarian Cancer National Alliance (OCNA)
866-399-6262 (toll free)
www.ovariancancer.org
Works to put ovarian cancer on the agenda of policy makers and women’s health leaders.
United Ostomy Associations of America, Inc. (UOAA)
800-826-0826
www.ostomy.org
A volunteer-based health organization that supports people who have had or will have intestinal or urinary diversions. This includes providing education, information, support, and advocacy.

Women’s Cancer Network
312-578-1439
www.wcn.org
Works to inform women around the world about gynecologic cancers. Provides a personalized risk assessment tool to assess risk for gynecologic and breast cancers.

Wound, Ostomy, Continence Nurses Society
888-224-WOCN (888-224-9626)
www.wocn.org
Members are experts in the care of people with wounds, ostomies, and continence disorders.

If you have any questions, contact a member of your healthcare team directly. If you're a patient at MSK and you need to reach a provider after 5:00 PM, during the weekend, or on a holiday, call 212-639-2000.

For more resources, visit www.mskcc.org/pe to search our virtual library.

Caring for Your Ileostomy or Colostomy - Last updated on February 22, 2017
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Common Medications Containing Aspirin, Other Nonsteroidal Anti-inflammatory Drugs (NSAIDs), or Vitamin E

This information will help you identify medications that contain aspirin, other NSAIDs, or vitamin E. It’s important to stop taking these medications before many cancer treatments. They affect your platelets (blood cells that clot to prevent bleeding) and can increase your risk of bleeding during treatment.

Other dietary supplements (such as other vitamins and herbal remedies) can also affect your cancer treatment. For more information, read the resource Herbal Remedies and Cancer Treatment (www.mskcc.org/pe/herbal_remedies).

Instructions Before Your Cancer Treatment

If you take aspirin, other NSAIDs, or vitamin E, tell your healthcare provider. They’ll tell you if you need to stop taking it. You’ll also find instructions in the information about your treatment. Read the “Examples of Medications” section to see if your medications contain aspirin, other NSAIDs, or vitamin E.

Before your surgery

Follow these instructions if you’re having surgery or a surgical procedure. **If your healthcare provider gives you other instructions, follow those instead.**

- If you take aspirin or a medication that contains aspirin, you may need to change your dose or stop taking it 7 days before your surgery. Follow your healthcare provider’s instructions. **Don’t stop taking aspirin unless your**
healthcare provider tells you to.

- If you take vitamin E or a supplement that contains vitamin E, stop taking it 7 days before your surgery or as directed by your healthcare provider.

- If you take an NSAID or a medication that contains an NSAID, stop taking it 48 hours (2 days) before your surgery or as directed by your healthcare provider.

**Before your radiology procedure**

Follow these instructions if you’re having a radiology procedure (including Interventional Radiology, Interventional Mammography, Breast Imaging, and General Radiology). **If your healthcare provider gives you other instructions, follow those instead.**

- If you take aspirin or a medication that contains aspirin, you may need to stop taking it 5 days before your procedure. Follow your healthcare provider’s instructions. **Don’t stop taking aspirin unless your healthcare provider tells you to.**

- If you take an NSAID or a medication that contains an NSAID, you may need to stop taking it 24 hours (1 day) before your procedure. Follow your healthcare provider’s instructions.

**Before and during your chemotherapy**

Chemotherapy can lower your platelet count, which can increase your risk of bleeding. Whether you’re just starting chemotherapy or have been getting it, talk with your healthcare provider before taking aspirin, other NSAIDs, or vitamin E.

**Examples of Medications**

Medications are often called by their brand name. This can make it hard to know their ingredients. The lists below can help you identify medications that contain aspirin, other NSAIDs, or vitamin E.

These lists include the most common products, but there are others. **Make sure your healthcare provider always knows all the prescription and over-**
the-counter (not prescription) medications you’re taking, including patches and creams.

<table>
<thead>
<tr>
<th>Common Medications Containing Aspirin</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aggrenox®</td>
</tr>
<tr>
<td>Cama® Arthritis Pain Reliever</td>
</tr>
<tr>
<td>Heartline®</td>
</tr>
<tr>
<td>Robaxisal® Tablets</td>
</tr>
<tr>
<td>Alka Seltzer®</td>
</tr>
<tr>
<td>COPE®</td>
</tr>
<tr>
<td>Headrin®</td>
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<tr>
<td>Roxiprin®</td>
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<tr>
<td>Anacin®</td>
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<tr>
<td>Dasin®</td>
</tr>
<tr>
<td>Isollyl®</td>
</tr>
<tr>
<td>Saleto®</td>
</tr>
<tr>
<td>Arthritis Pain Formula</td>
</tr>
<tr>
<td>Easprin®</td>
</tr>
<tr>
<td>Lanorinal®</td>
</tr>
<tr>
<td>Salocol®</td>
</tr>
<tr>
<td>Arthritis Foundation Pain Reliever®</td>
</tr>
<tr>
<td>Ecotrin® (most formulations)</td>
</tr>
<tr>
<td>Lortab® ASA Tablets</td>
</tr>
<tr>
<td>Sodol®</td>
</tr>
<tr>
<td>ASA Enseals®</td>
</tr>
<tr>
<td>Empirin® Aspirin (most formulations)</td>
</tr>
<tr>
<td>Magnaprin®</td>
</tr>
<tr>
<td>Soma® Compound Tablets</td>
</tr>
<tr>
<td>ASA Suppositories®</td>
</tr>
<tr>
<td>Epromate®</td>
</tr>
<tr>
<td>Marnal®</td>
</tr>
<tr>
<td>Soma® Compound with Codeine Tablets</td>
</tr>
<tr>
<td>Ascriptin® and Ascriptin A/D®</td>
</tr>
<tr>
<td>Equagesic Tablets</td>
</tr>
<tr>
<td>Micrainin®</td>
</tr>
<tr>
<td>St. Joseph® Adult Chewable Aspirin</td>
</tr>
<tr>
<td>Aspergum®</td>
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<tr>
<td>Equazine®</td>
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<tr>
<td>Momentum®</td>
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<tr>
<td>Supac®</td>
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<tr>
<td>Asprimox®</td>
</tr>
<tr>
<td>Excedrin® Extra-Strength Analgesic Tablets and Caplets</td>
</tr>
<tr>
<td>Norgesic Forte® (most formulations)</td>
</tr>
<tr>
<td>Synalgos®-DC Capsules</td>
</tr>
<tr>
<td>Axotal®</td>
</tr>
<tr>
<td>Excedrin® Migraine</td>
</tr>
<tr>
<td>Norwich® Aspirin</td>
</tr>
<tr>
<td>Tenol-Plus®</td>
</tr>
<tr>
<td>Azdone®</td>
</tr>
<tr>
<td>Fiorgen®</td>
</tr>
<tr>
<td>PAC® Analgesic Tablets</td>
</tr>
<tr>
<td>Trigesic®</td>
</tr>
<tr>
<td>Bayer® (most formulations)</td>
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<tr>
<td>Fiorinal® (most formulations)</td>
</tr>
<tr>
<td>Orphengesic®</td>
</tr>
<tr>
<td>Talwin® Compound</td>
</tr>
<tr>
<td>BC® Powder and Cold formulations</td>
</tr>
<tr>
<td>Fiortal®</td>
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<tr>
<td>Painaid®</td>
</tr>
<tr>
<td>Vanquish® Analgesic Caplets</td>
</tr>
<tr>
<td>Bufferin® (most formulations)</td>
</tr>
<tr>
<td>Gelpirin®</td>
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<tr>
<td>Panasal®</td>
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<tr>
<td>Wesprin® Buffered</td>
</tr>
<tr>
<td>Buffets II®</td>
</tr>
<tr>
<td>Genprin®</td>
</tr>
<tr>
<td>Percodan® Tablets</td>
</tr>
<tr>
<td>Zee-Seltzer®</td>
</tr>
<tr>
<td>Buffex®</td>
</tr>
<tr>
<td>Gensan®</td>
</tr>
<tr>
<td>Persistin®</td>
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<tr>
<td>ZORprin®</td>
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### Common NSAID Medications That Don’t Contain Aspirin

<table>
<thead>
<tr>
<th>Product</th>
<th>Brand Name</th>
<th>Brand Name</th>
<th>Brand Name</th>
<th>Brand Name</th>
</tr>
</thead>
<tbody>
<tr>
<td>Advil®</td>
<td>Duexis®</td>
<td>Mefenamic Acid</td>
<td>PediaCare Fever®</td>
<td></td>
</tr>
<tr>
<td>Advil Migraine®</td>
<td>Etodolac®</td>
<td>Meloxicam</td>
<td>Piroxicam</td>
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<tr>
<td>Aleve®</td>
<td>Feldene®</td>
<td>Menadol®</td>
<td>Ponstel®</td>
<td></td>
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<tr>
<td>Anaprox DS®</td>
<td>Fenoprofen</td>
<td>Midol®</td>
<td>Relafen®</td>
<td></td>
</tr>
<tr>
<td>Ansaid®</td>
<td>Flurbiprofen</td>
<td>Mobic®</td>
<td>Saleto 200®</td>
<td></td>
</tr>
<tr>
<td>Arthrotec®</td>
<td>Genpril®</td>
<td>Motrin®</td>
<td>Sulindac</td>
<td></td>
</tr>
<tr>
<td>Bayer® Select Pain Relief Formula Caplets</td>
<td>Ibuprofen</td>
<td>Nabumetone</td>
<td>Toradol®</td>
<td></td>
</tr>
<tr>
<td>Celebrex®</td>
<td>Indomethacin</td>
<td>Nalfon®</td>
<td>Treximet®</td>
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</tr>
<tr>
<td>Celecoxib</td>
<td>Indocin®</td>
<td>Naproxen</td>
<td>Vicoprofen®</td>
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</tr>
<tr>
<td>Children’s Motrin®</td>
<td>Ketoprofen</td>
<td>Naprosyn®</td>
<td>Vimovo®</td>
<td></td>
</tr>
<tr>
<td>Clinoril®</td>
<td>Ketorolac</td>
<td>Nuprin®</td>
<td>Voltaren®</td>
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</tr>
<tr>
<td>Daypro®</td>
<td>Lodine®</td>
<td>Orudis®</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diclofenac</td>
<td>Meclofenamate</td>
<td>Oxaprozin</td>
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</tbody>
</table>

### Products Containing Vitamin E

<table>
<thead>
<tr>
<th>Product</th>
<th>Brand Name</th>
<th>Brand Name</th>
<th>Brand Name</th>
<th>Brand Name</th>
</tr>
</thead>
<tbody>
<tr>
<td>Amino-Opt-E</td>
<td>Aquavit</td>
<td>E-400 IU</td>
<td>E complex-600</td>
<td></td>
</tr>
<tr>
<td>Aquasol E</td>
<td>D’alpha E</td>
<td>E-1000 IU Softgels</td>
<td>Vita-Plus E</td>
<td></td>
</tr>
</tbody>
</table>

Most multivitamins contain vitamin E. If you take a multivitamin, check the label.

## About Acetaminophen

Acetaminophen (Tylenol®) is generally safe to take during your cancer treatment. It doesn’t affect platelets, so it won’t increase your chance of bleeding. But, talk with your healthcare provider before taking acetaminophen if you’re getting chemotherapy.
Medications Containing Acetaminophen

<table>
<thead>
<tr>
<th>Medication</th>
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</thead>
<tbody>
<tr>
<td>Acephen®</td>
</tr>
<tr>
<td>Esgic®</td>
</tr>
<tr>
<td>Percocet®</td>
</tr>
<tr>
<td>Vanquish®</td>
</tr>
<tr>
<td>Aceta® with Codeine</td>
</tr>
<tr>
<td>Excedrin P.M.®</td>
</tr>
<tr>
<td>Primlev®</td>
</tr>
<tr>
<td>Vicodin®</td>
</tr>
<tr>
<td>Acetaminophen with Codeine</td>
</tr>
<tr>
<td>Fiorcet®</td>
</tr>
<tr>
<td>Repan®</td>
</tr>
<tr>
<td>Wygesic®</td>
</tr>
<tr>
<td>Aspirin-Free Anacin®</td>
</tr>
<tr>
<td>Lorcor®</td>
</tr>
<tr>
<td>Rocicet®</td>
</tr>
<tr>
<td>Xartemis XR®</td>
</tr>
</tbody>
</table>
| Arthritis Pain Formula® Aspirin-
  Free                            |
| Lortab®                        |
| Talacen®                       |
| Xodol®                         |
| Datril®                        |
| Naldegesic®                    |
| Tempra®                        |
| Zydone®                        |
| Di-Gesic®                      |
| Norco®                         |
| Tylemol®                       |
| Endocet®                       |
| Panadol®                       |
| Tylemol® with Codeine No. 3    |

Read the labels on all your medications

Acetaminophen is safe when used as directed. But, there’s a limit to how much you can take in a day. It’s possible to take too much without knowing because it’s in many different prescription and over-the-counter medications. It’s often an ingredient in pain relievers, fever reducers, sleep aids, and cough, cold, and allergy medications.

The full name acetaminophen isn’t always written out. Look for the common abbreviations listed below, especially on prescription pain relievers.

<table>
<thead>
<tr>
<th>Common Abbreviations for Acetaminophen</th>
</tr>
</thead>
<tbody>
<tr>
<td>APAP</td>
</tr>
<tr>
<td>Acetamin</td>
</tr>
</tbody>
</table>

Always read and follow the label on the product you’re taking. Don’t take more than 1 medication that contains acetaminophen at a time without talking with a member of your healthcare team.
If you have any questions, contact a member of your healthcare team directly.
If you're a patient at MSK and you need to reach a provider after 5:00 PM,
during the weekend, or on a holiday, call 212-639-2000.

For more resources, visit www.mskcc.org/pe to search our virtual library.

Common Medications Containing Aspirin, Other Nonsteroidal Anti-inflammatory Drugs (NSAIDs), or Vitamin E - Last updated on September 21, 2020
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Herbal Remedies and Cancer Treatment

This information explains herbal remedies and how they can affect your cancer treatment.

One week before you have surgery or start chemotherapy or radiation therapy, you must stop taking any herbal or botanical (plant-based) home remedies or other dietary supplements. This is because they can:

- Interact with your other medications.
- Increase or lower your blood pressure.
- Thin your blood and increase your risk of bleeding.
- Make radiation therapy less effective.
- Increase the effects of sedation or anesthesia (medications to make you sleepy).

You can continue to use herbs in your food and drinks, such as using spices in cooking and drinking tea. However, you must stop taking herbal supplements before your treatment. Herbal supplements are stronger than the herbs you would use in cooking.

Common Herbal Supplements and Their Effects

These are some commonly used herbs and their side effects on cancer treatments.

Echinacea
- Can cause an allergic reaction, such as a rash or trouble breathing.
- Can lower the effects of medications used to weaken the immune system.

**Garlic**

- Can lower your blood pressure, fat, and cholesterol levels.
- Can increase your risk of bleeding.

**Gingko (also known as *Gingko biloba*)**

- Can increase your risk of bleeding.

**Ginseng**

- Can lower the effects of sedation or anesthesia.
- Can increase your risk of bleeding.
- Can lower your blood glucose (sugar) level.

**Turmeric**

- Can make chemotherapy less effective.

**St. John’s Wort**

- Can interact with medications given during surgery.
- Can make your skin more sensitive to radiation or laser treatment.

**Valerian**

- Can increase the effects of sedation or anesthesia.

**Herbal formulas**

- Herbal formulas contain different herbs. We don’t know their side effects. You must also stop taking these products 1 week before treatment. Do not start taking herbal formulas again until your doctor tells you it’s safe.
This information doesn’t cover all herbal remedies or possible side effects. Talk with your healthcare provider if you have any questions or concerns.

For more information about herbs and botanicals, visit the Memorial Sloan Kettering (MSK) Integrative Medicine Service website at www.aboutherbs.com.

If you have any questions, contact a member of your healthcare team directly. If you're a patient at MSK and you need to reach a provider after 5:00 PM, during the weekend, or on a holiday, call 212-639-2000.

For more resources, visit www.mskcc.org/pe to search our virtual library.
How to Use Your Incentive Spirometer

This information will help you learn how to use your incentive spirometer.

About Your Incentive Spirometer

An incentive spirometer is a device that will expand your lungs by helping you to breathe more deeply and fully. The parts of your incentive spirometer are labeled in Figure 1.

Figure 1. Incentive Spirometer
Use your incentive spirometer after your surgery and do your deep breathing and coughing exercises. This will help keep your lungs active throughout your recovery and prevent complications such as pneumonia.

If you have an active respiratory infection (such as pneumonia, bronchitis, or COVID-19) do not use the device when other people are around.

**How To Use Your Incentive Spirometer**

Here is a video demonstrating how to use your incentive spirometer:

Please visit [www.mskcc.org/pe/incentive_spirometer_video](http://www.mskcc.org/pe/incentive_spirometer_video) to watch this video.

**Setting up your incentive spirometer**

The first time you use your incentive spirometer, you will need to take the flexible tubing with the mouthpiece out of the bag. Stretch out the tubing and connect it to the outlet on the right side of the base (see Figure 1). The mouthpiece will be attached to the other end of the tubing.

**Using your incentive spirometer**

When you’re using your incentive spirometer, make sure to breathe through your mouth. If you breathe through your nose, the incentive spirometer won’t work properly. You can hold your nose if you have trouble.

If you feel dizzy at any time, stop and rest. Try again at a later time.

To use your incentive spirometer, follow the steps below.

1. Sit upright in a chair or in bed. Hold the incentive spirometer at eye level.
   - If you had surgery on your chest or abdomen (belly), hug or hold a pillow to help splint or brace your incision (surgical cut) while you’re using the incentive spirometer. This will help decrease pain at your incision.

2. Put the mouthpiece in your mouth and close your lips tightly around it. Slowly
breathe out (exhale) completely.

3. Breathe in (inhale) slowly through your mouth as deeply as you can. As you take the breath, you will see the piston rise inside the large column. While the piston rises, the indicator on the right should move upwards. It should stay in between the 2 arrows (see Figure 1).

4. Try to get the piston as high as you can, while keeping the indicator between the arrows.
   - If the indicator doesn’t stay between the arrows, you’re breathing either too fast or too slow.

5. When you get it as high as you can, hold your breath for 10 seconds, or as long as possible. While you’re holding your breath, the piston will slowly fall to the base of the spirometer.

6. Once the piston reaches the bottom of the spirometer, breathe out slowly through your mouth. Rest for a few seconds.

7. Repeat 10 times. Try to get the piston to the same level with each breath.

8. After each set of 10 breaths, try to cough, holding a pillow over your incision, as needed. Coughing will help loosen or clear any mucus in your lungs.

9. Put the marker at the level the piston reached on your incentive spirometer. This will be your goal next time.

Repeat these steps every hour that you’re awake.

Cover the mouthpiece of the incentive spirometer when you aren’t using it.

If you have any questions, contact a member of your healthcare team directly. If you’re a patient at MSK and you need to reach a provider after 5:00 PM, during the weekend, or on a holiday, call 212-639-2000.

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