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[Ewing Sarcoma - Childhood - Coping with Side Effects](#) [1]

This section has been reviewed and approved by the [Cancer.Net Editorial Board](#) [2], 05/2015

ON THIS PAGE: You will find out more about steps to take to help your child cope with physical, social, and emotional side effects. This page includes several links outside of this guide to other sections of this website. To see other pages, use the menu on the side of your screen.

[Fear of treatment side effects](#) [3] is common after a diagnosis of cancer, but it may help to know that preventing and controlling side effects is a major focus of your child's health care team. This is called palliative care, and it is an important part of the overall treatment plan, regardless of the stage of disease.

There are possible side effects for every cancer treatment, but patients don't experience the same side effects when given the same treatments for many reasons. That can make it hard to predict exactly how you will feel during treatment.

Common side effects from each treatment option for Ewing sarcoma are described in detail within the [Treatment Options](#) [4] section. Learn more about [cancer and different treatments, along with ways to prevent or control them](#) [5]. Side effects depend on a variety of factors, including the cancer's stage, the length and dosage of treatment(s), and your child's overall health.

Talking with the health care team about side effects

Before treatment begins, talk with your child's doctor about possible side effects of each type of treatment your child will be receiving. Ask which side effects are most likely to happen, when they are likely to occur, and what can be done to prevent or relieve them.

And, ask about the level of caregiving your child may need during treatment and recovery, as family members and friends play an important role in the care of children and teens with Ewing sarcoma. Learn more about [caregiving](#) [6].

In addition to physical side effects, there may be emotional and social effects as well. Patients and their families are encouraged to share their feelings with a member of the health care team who can help with [coping strategies](#) [7], including concerns about [managing the cost of cancer care](#) [8]. These strategies can be different based on the age of your [child](#) [9] or [teen](#) [10].

During and after treatment, be sure to tell the health care team about the side effects your child experiences, even if you feel they are not serious. Sometimes, side effects can last beyond the treatment period, called a long-term side effect. A side effect that occurs months or years after treatment is called a [late effect](#) [11]. Treatment of both types of effects is an important part of survivorship care. Learn more by reading the [Follow-Up Care](#) [12] section of this guide or talking with your child's doctor.

The next section in this guide is [Follow-up Care](#) [12], and it explains the importance of check-ups after cancer treatment is finished. Or, use the menu on the side of your screen to choose another section to continue reading this guide.

Links

[1] <http://www.cancer.net/cancer-types/ewing-sarcoma-childhood/coping-side-effects>

[2] <http://www.cancer.net/about-us>

[3] <http://www.cancer.net/node/24492>

[4] <http://www.cancer.net/node/18804>

[5] <http://www.cancer.net/node/25238>

[6] <http://www.cancer.net/node/25009>

[7] <http://www.cancer.net/node/21>

[8] <http://www.cancer.net/node/24865>

[9] <http://www.cancer.net/node/25114>

[10] <http://www.cancer.net/node/25116>

[11] <http://www.cancer.net/survivorship/long-term-side-effects-cancer-treatment>

[12] <http://www.cancer.net/node/18807>