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[Myelodysplastic Syndromes - MDS - Coping with Side Effects](#) [1]

This section has been reviewed and approved by the [Cancer.Net Editorial Board](#) [2], 11/2015

ON THIS PAGE: You will find out more about steps to take to help cope with physical, social, and emotional side effects of MDS. 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 MDS, but it may help to know that preventing and controlling side effects is a major focus of your 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 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 MDS are described in detail within the [Treatment Options](#) [4] section. Learn more about [the most common side effects of MDS and different treatments, along with ways to prevent and control them](#) [5]. Side effects depend on a variety of factors, including the subtype of MDS, the length and dosage of treatment(s), and your overall health.

Talking with your health care team about side effects

Before treatment begins, talk with your doctor about possible side effects of each type of treatment you 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 you may need during treatment and recovery, as family members and friends often play an important role in the care of a person with MDS. 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 their health care team, who can help with [coping strategies](#) [7], including concerns about [managing the cost of your medical care](#) [8].

During and after treatment, be sure to tell the health care team about the side effects you experience, 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](#) [9]. Treatment of both types of effects is an important part of survivorship care. Learn more by reading the [Follow-up Care](#) [10] section of this guide or talking with your doctor.

The [next section in this guide is Follow-up Care](#) [10] and it explains the importance of check-ups after 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/myelodysplastic-syndromes-mds/coping-side-effects>

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

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

[4] <http://www.cancer.net/node/19387>

[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/25396>

[10] <http://www.cancer.net/node/19390>