

# Transplant and Globoid Cell Leukodystrophy (GLD)

Learning more about your child's (or your) treatment options can help you make informed medical decisions.

## This fact sheet **tells you:**

- What GLD is
- What the transplant steps are
- When to see a transplant doctor
- Questions to ask your doctor

## What is **GLD**?

GLD, also called Krabbe Disease, is an inherited disease. It affects how the body breaks down food into energy. In GLD, the body is missing an important protein to break down certain fatty substances, or fats. When these fats aren't broken down, they build up and damage the protective covering around nerve cells, called myelin.

## Symptoms of GLD include:

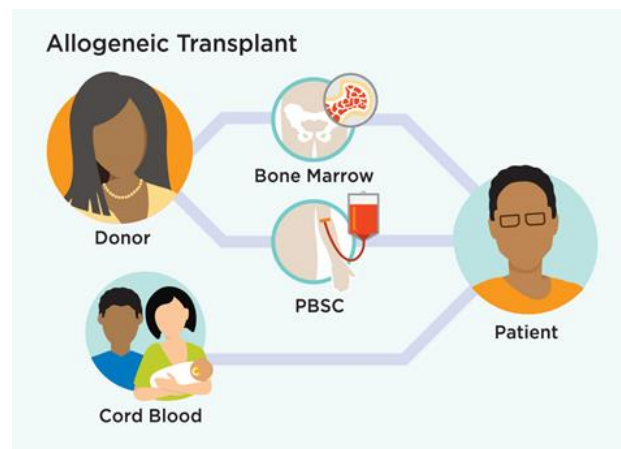
- Difficulty with memory, learning, speaking, and understanding
- Poor muscle control
- Muscle weakness
- Stiff joints

Children with GLD often have symptoms when they're only a few months old. Teens and adults can have GLD but it's less common.

## What is a **blood or marrow transplant (BMT)**?

BMT, also called bone marrow transplant, replaces unhealthy blood-forming cells with healthy ones. The new cells have the right proteins so the body can break down fat. Transplant prevents more damage to the myelin, but it can't fix any damage that has already happened.

The type of BMT that treats GLD is called an **allogeneic transplant**. It uses healthy, blood-forming cells donated by someone else to replace the unhealthy ones. The healthy cells can come from a family member, an unrelated donor, or umbilical cord blood.



## Allogeneic transplant steps

1. First, your doctor searches for your child's donor. This can take weeks to months.
2. Then, your child will get chemotherapy, and possibly radiation, to kill the unhealthy cells. This can take up to 2 weeks.
3. On transplant day, the donated cells are given to your child through an intravenous (IV) infusion. The new cells travel to the inside of their bones.
4. After a few weeks, the donated cells begin to make healthy blood cells.
5. Then, your child will need many months to recover. Your child will spend some of this time in or near the hospital.

Doctors, advanced practice providers, nurses, dietitians, and social workers will closely care for your child during the entire process to prevent and treat any side effects or complications.

## When to **see a transplant doctor**

You should see a transplant doctor **as soon as possible after a diagnosis of ALD**. BMT can stop the disease from progressing but it can't fix damage that has already taken place.

## Your first appointment with a transplant doctor

The transplant doctor will:

- Review your child's medical history
- Talk with you about treatment options and their risks and benefits
- Recommend the best time for your child to get a transplant and prepare for treatment
- Start searching for a donor, if BMT is recommended
- Schedule appointments with other doctors, like a neurologist

## Questions to ask your doctor

- What are the chances transplant will stop the GLD from causing more damage?

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- What are the possible side effects of transplant? How can they be reduced?

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- How might my child's quality of life change over time, with or without transplant?

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## AT EVERY STEP, WE'RE **HERE TO HELP**

Be The Match has a team dedicated to providing information and support to you before, during, and after transplant. You can contact our Patient Support Center to ask questions you may have about transplant, request professional or peer support, or receive free patient education materials.

CALL: **1 (888) 999-6743** | EMAIL: **patientinfo@nmdp.org** | WEB: **BeTheMatch.org/one-on-one**



Every individual's medical situation, transplant experience, and recovery is unique. You should always consult with your own transplant team or family doctor regarding your situation. This information is not intended to replace, and should not replace, a doctor's medical judgment or advice.