

You are your child's best advocate – so learn all you can about thalassemia!

Doctors and nurses want your child to get the *best possible care*. But a child may not always know what information they need to tell their health care team.

It's often up to you to fill in the missing information, to ask questions, and to make sure that there is effective communication about any issues that can affect your child's health. And to do that, you need to *know as much as you can* about thalassemia, its treatment, and the possible complications that can occur.



Speaking up for your child helps to ensure that they get the help they need to live a stronger and healthier life.

Download the free “A Guide to Living with Thalassemia” at
<http://tinyurl.com/CAFGuide>

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Cooley's Anemia
FOUNDATION
Leading the Fight Against Thalassemia

www.thalassemia.org