

Questions?

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A RESEARCH COLLABORATION BETWEEN THE
MEDICAL COLLEGE OF WISCONSIN AND NMDP

CMS Innovation Center's Cell and Gene Therapy (CGT) Access Model for Sickle Cell Disease (SCD)

This study collects information about how cell and gene therapies **impact your life** and is for patients receiving **Medicaid**.

You and your parent or guardian would **answer questions on a survey** about how you're feeling, costs of your treatment, and information about you.

The surveys take **15-25 minutes to complete** and **can be filled out online or on paper**. Every question is optional, and you can say no at any time. You will be compensated \$25 for the first survey before treatment and \$10 for every survey you complete after treatment for up to 5 years if funding is available.

Next Steps

1. Agree to be contacted by CIBMTR for research on a form your treatment center will share with you.
2. The center where you get treatment will give us your contact information.
3. You or your parent will get a call from the study team or an email from Surveys@CIBMTR.org with more information and links to join this **free and optional study**.

Sharing knowledge, sharing hope.

By sharing some of your health information, you may help people with sickle cell and other diseases. Information from volunteers has helped us find out **which treatments help people live longer**, when to use different therapies **and more!**



We protect your privacy.

CIBMTR's database is secure. CIBMTR follows US and international privacy laws. We cannot share information about your identity.

To learn more, visit CIBMTR.org or scan the QR code.