

Questions?

Surveys@CIBMTR.org

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CIBMTR®

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 data about how cellular and gene therapies **impact quality-of-life** and is eligible for patients receiving **Medicaid**.

You would **answer surveys** which include multiple choice questions about how you're feeling, financial impacts of your treatment, and information about your background.

The surveys take **15-25 minutes to complete** and **can be filled out online or on paper**. Every question is optional, and you can opt out 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. Your treatment center will provide your contact information to us.
3. Expect an email from Surveys@CIBMTR.org or get a call from the study team 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 determine **which treatments help people live longer**, when to use transplant or other cell therapies, how closely donors need to match patients, **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.

