

## Questions?

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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](mailto: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](https://CIBMTR.org) or scan the QR code.**