

Collaboration. Innovation. Transformation.

The *Coalition for Clinical Trial Equity (CCTE)* is seeking members who share our mission to *collaborate* on developing *innovative* strategies that *transform* the landscape of clinical trial participation for Black patients.

# What is the Coalition for Clinical Trial Equity?

CCTE is a collaborative initiative launched by the Foundation for Sarcoidosis Research (FSR) to address disparities in clinical trial participation, particularly among Black Americans. Our mission is to foster a more inclusive research landscape by dismantling barriers and increasing representation of Black patients in clinical trials. Through collective efforts, the coalition seeks to pave the way for more equitable access to clinical trials and ultimately improve health outcomes for Black patients.

We have a strong team of organizational partners, esteemed academic institutions, patients and world-renowned clinicians, researchers and bioethicists leading the coalition's efforts.

### JOIN FOR CLINICAL TRIAL EQUITY

#### Together we've:

- Partnered with corporations to provide paid time off for clinical trials participation.
- Secured a groundbreaking clarification from the U.S. Department of Labor ensuring FMLA benefits provide protected leave for employees participating in clinical trials.

#### **Get Involved!**

#### **CCTE has three pillars of focus:**

- FMLA & Corporate Infrastructure to confront jobrelated barriers to participation.
- Patient Education & Engagement to close the gap in knowledge and transparency through community-building.
- Patient Benefit & Compensation to expand benefits and compensation for patients.

**Join the Coalition** for Clinical Trial Equity to collaborate on driving systemic change with targeted, pragmatic strategies focused on advocacy, education, and support!

Visit **www.ignorenomore.org** to *learn more* about the coalition and how you can join.

Sponsored by Mallinckrodt Pharmaceuticals, Boehringer-Ingelheim, and made possible in part by a grant from 23andMe.



#### Meet our Founding Steering Committee Members:



#### **Bioethicists**

Arthur Caplan, PhD Jennifer Miller, PhD Sonya Charles, PhD

#### Patients & Care Partner\*

Kathryn Washington, MPH Jessica Propps\* Mary Oldham Purvis Hunt, MA

# **Membership Opportunities**

# **Benefits**

- Logo posted on coalition's website.
- Opportunity to join one of three committee workgroups to lead efforts in clinical trials diversity.
- Acknowledgement on public-facing materials (presentations, website, publications, etc.).
- Receive coalition badge to add to your website highlighting your commitment to equity.
- Recognized as partner on groundbreaking accomplishments like the FMLA benefit for participants.

# **Membership:**

### **Non-Profit**

Non-profit Membership is <u>complementary.</u>

## **Industry/ For-Profit**

Annual membership fees are tiered by company budget:

- \$1,000 (Less than \$1M)
- \$2,500 (\$1 \$5M)
- \$5,000 (\$5 \$10M)
- \$7,500 (\$10 \$25M)
- \$10,000 (Over \$25M)

Ask about how you can support coalition initiatives like our paid time off initiative. This initiative partners with employers to provide paid time off for employees participating in clinical trials. Contact: ignorenomore@stopsarcoidosis.org

Visit www.ignorenomore.org to learn more.

