



RARE DISEASE DIVERSITY COALITION

VIRTUAL CONVERSATIONS IN SUPPORT OF RARE DISEASE DAY

PANEL ONE

MONDAY
FEBRUARY 28, 2022
12 PM EST

HOSTED BY



Nischelle Turner
Co-Host
Entertainment Tonight

Knowledge is Power:

A CONVERSATION WITH THE TATE BROTHERS AROUND RARE DISEASE

Far too many Black, Indigenous, and People-of-Color patients have a family history of disease or ailments of which they are unaware or do not communicate to their health care provider. This is especially problematic in the case of rare diseases that are harder to diagnose and more likely to go untreated. Join us for an insightful conversation with the Tate brothers about the importance of knowing one's family medical history and passing that information on to future generations. We will also discuss the Tate brothers' connection to advocacy around sickle cell and other rare diseases.

SPEAKERS



Lahmard Tate
Actor & Producer
Tate Brothers Foundation



Larenz Tate
Actor & Producer
Tate Brothers Foundation



Larron Tate
Actor, Producer & Writer
Tate Brothers Foundation

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PANEL TWO

MONDAY
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HOSTED BY



Richard Lui
Anchor & Journalist
NBC News/MSNBC

How to Advocate for Your Health as a BIPOC Patient

Too often the symptoms of a rare disease may be dismissed by a physician due to lack of knowledge and/or medical bias. This is especially true in Black, Indigenous and People-of-Color communities. Join us for a conversation with key experts that will provide practical tips on how to best advocate for oneself as a BIPOC patient and how to get access to early detection and treatment options for rare diseases.

SPEAKERS



Olga Lucia Torres, JD, MS
Professional Patient Advocate
& Lecturer
Columbia University



Yves Duroseau, MD, MPH
Chair, Department of
Emergency Medicine
Lenox Hill Hospital



Sachet Walker
Caregiver for a
Son with FSGS

LIVE ON YouTube

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