## Who We Are

### Meet our Board of Directors

**Dr. Leslie Cooper, MD – Co-Founder** – Chair, Enterprise Dept. of Cardiovascular Medicine for all of Mayo Clinic

**Dr. Jack Price, MD** – Professor of Pediatrics at Baylor College of Medicine, Cardiologist at Texas Children's Hospital

**Dr. Bettina Heidecker, MD** – Head of Heart Failure and Cardiomyopathies Charite Hospital, Berlin, Germany

**Candace Moose, Co-Founder** Giant Cell Myocarditis Survivor and Transplant Recipient

**Joseph Rumore** – Viral Myocarditis Survivor and former Managing Director of a national insurance company

**Louis Romano,** Owner of Home Well Care

**Francine Andrea**, Vice-President for Enrollment Management, Student Affairs and Chief Compliance Officer for Felician University

**Joel Aranson,** Founder and Chairman of National Sporting Good Corporation and father of a Giant Cell Myocarditis victim

**Giustina Schiano,** Mother of a Viral Myocarditis victim and Family Advocate for the Myocarditis Foundation

**Michael A. Linn,** Sales Leader for the Instruments Division of the Stryker Corporation

**Stephanie Kennan,** Senior Vice President of Federal Affairs at McGuire Woods Consulting

**Gary Kubera**, Former Chemical Industry Executive and CEO



How You Can Help

The Myocarditis Foundation is seeking support to further its awareness efforts and to expand its research program. Please contact us if you

wish to help the Myocarditis Foundation save more lives from sudden cardiac death brought upon by the disease myocarditis.



#### The Myocarditis Foundation

3518 Echo Mountain Drive Kingwood, Texas 77345

(281) 713-2962 www.myocarditisfoundation.org



# The Myocarditis Foundation



### **?**

Working to end sudden death from myocarditis through research, awareness, and advocacy



# What is the Myocarditis Foundation?

The Myocarditis Foundation is a 501(c)3 nonprofit organization dedicated to providing information about myocarditis to medical professionals, patients, and families with the goal of saving more lives.

Myocarditis is often misdiagnosed and is a poorly understood disease, which can rapidly progress to heart failure and death.

The Myocarditis Foundation works to fund research to find better ways to diagnose, treat and prevent myocarditis from having devastating effects on patients and their families around the world.

Since their establishment in 2005, the Myocarditis Foundation has awarded 21 research fellowship grants to young doctors in world-renowned university research facilities, with the shared goal of improving the care and medical outcome for patients affected by this devastating disease.

In addition to funding research, the Myocarditis Foundation takes great strides to increase the awareness of the disease myocarditis and its link to sudden cardiac related death in otherwise young, healthy people.

## Who does the Myocarditis Foundation serve?

The Myocarditis Foundation provides information and support to patients diagnosed with myocarditis, families who have lost a loved one to myocarditis, and medical professionals who are seeking information for their patients with myocarditis.

#### How does the Myocarditis Foundation engage the medical community and the general public?

- Sponsors conferences, workshops, and lectures world-wide directed at patients, families, medical professionals, or professional heart disease groups
- Created a private and secure online research network to bring together and encourage the collaboration of researchers and physicians interested in and currently working on myocarditis research
- Maintains the only free and official myocarditis-specific website with features such as FAQ's, research findings, blog, newsletters, and real-life stories

# How does the Myocarditis Foundation spread awareness?

- Holds and supports fundraisers and awareness events around the world
- Sends myocarditis information to the medical community and the general public
- Uses newsletters, blogs, and social media to provide the most current information