



**Dedicated to finding a cure and better treatments for Juvenile Myositis and improving the lives of families affected by JM.**

**Cure JM serves families and patients of ages affected by Juvenile Dermatomyositis (JDM) and Juvenile Polymyositis (JPM)**

Cure JM is the leading funder of Juvenile Myositis disease care and research. Here's how we're changing the lives of children and young adults with JM:

- Establishing the Cure JM Clinical Care Network, which will expand excellent JM care to children around the country
- Pursuing promising new JM treatments
- Funding clinical trials
- Funding Centers of Excellence
- Investing over \$19 million to drive JM research forward
- Providing vital resources and connections to families and patients



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*"We are deeply grateful for the tremendous role Cure JM has played in raising awareness of JM and advocating for children and families. Through Cure JM's outstanding commitment to the well-being of these young patients and its generous donations to research, Cure JM has touched the lives of many individuals."*

— Lauren M. Pachman, M.D. Professor of Pediatrics,  
Northwestern University's Feinberg School of  
Medicine Division of Pediatric Rheumatology, Ann &  
Robert H. Lurie Children's Hospital of Chicago Head,  
Cure JM Program of Excellence in Juvenile Myositis  
Research, Stanley Manne Children's Research  
Institute



### How can I Get Connected?

- Join us at [www.curejm.org/quickjoin](http://www.curejm.org/quickjoin)
- Text or call (202) 596-6267
- RSVP for an upcoming event [www.curejm.org/events](http://www.curejm.org/events)
- Email [info@curejm.org](mailto:info@curejm.org)

### Other resources:

View educational recordings at [www.curejm.org/videos](http://www.curejm.org/videos)  
Join your local chapter at [www.curejm.org/quickjoin](http://www.curejm.org/quickjoin)

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