



WORLD*Fair*TM 2017

Lysosomal Disease Patient Meeting

We have an exciting meeting planned for this fall. It will have lots of helpful information from three important areas that impact your life:

**Government/Policy Medical
Pharmaceutical Manufacturers**

This year we are proud to have Minnesota Congressman Matt Dean as our keynote speaker. Rep. Dean is a Minnesota leader in rare disease advocacy. He will present information on a Bill to create a Rare Disease Advisory Council for Minnesota. This Bill will work towards bringing U of MN Universities and healthcare systems, Health and Human Services and Patient Advocacy groups together in a collaborative effort to discuss the best strategies to improve understanding of the diagnosis, treatment and impact of rare diseases.

There will be a panel discussion of the many roles pharmaceutical manufacturers provide for patient needs and the various people/positions in the medical community will also participate in a panel discussion. Dr. Jeanine Jarnes will present new information on the management of immune responses to ERT.

As always, there will be opportunities to network with old and new friends, and to discuss your individual needs with company representatives.

Date and Time

Saturday, October 7, 2017

8:00 am – 9:00 am	Registration and Breakfast
8:15 am -- 9:00 am	Raptor Center (3 birds of prey)
9:00 am – 11:45 am	Key Note Speakers
12:00 am--1:30 pm	Breakout sessions

Location

McNamara Alumni Center
U of M Campus
200 Oak St. SE
Minneapolis, MN 55414

FREE reserved parking next to McNamara Center available at:
University Avenue Parking Ramp,
1926 SE 17th Avenue

Register

To register or if you have questions, please call Colleen Blissenbach at 612-617-3565 or email cdoyen1@fairview.org

There is no cost to attend. Breakfast, box lunch, refreshments and snacks included.