Thanks to all of you who have cared, loved, and used your minds so brilliantly to help us.

Without you, we wouldn't stand a chance.

- Belinda Miller

WE ARE HONORED THAT YOU CHOSE TO SUPPORT US.

FSH muscular dystrophy, a debilitating disease affecting nearly a million men, women, and children worldwide, has long lacked any treatment, but because of you, our families can look forward to having a therapy by the year 2025, if not sooner.

BECAUSE OF YOU, thousands of families can get support from local chapter leaders and become activists in their communities.

YOU HAVE CATALYZED BREAKTHROUGH RESEARCH

on stem cells, CRISPR, new treatment strategies, and more than \$100 million of private investment in FSHD therapies.

THANKS TO YOU, we are launching the FSHD Therapeutic Accelerator Project to remove barriers to FDA approval of promising treatments.

Together, we have journeyed incredible distances. With your continued support, we will reach our goal!

- MARK STONE

President & CEO



FSH Society

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2018 **ANNUAL REPORT CARD**

Education

PARTICIPANTS

1,900

1 FSHD CONNECT CONFERENCE **3** FAMILY DAY CONFERENCES **4** WEBINARS

YOUTUBE VIEWS

27,000



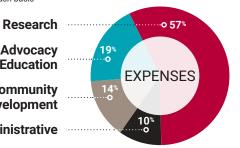
Financials

Cash basis

Patient Advocacy and Education

> Community **Development**

Administrative



Results that matter

Our vision is a world free of the suffering caused by FSH muscular dystrophy (FSHD). In 2018, we saw remarkable progress toward treatments and greatly expanded our ability to serve our members. Here is what we have accomplished, thanks to your dedicated support.

www.fshsociety.org

Accelerating Research

RESEARCH INVESTMENT

REVENUE RAISED

\$2.8

INTERNATIONAL RESEARCH CONGRESS ATTENDEES

CLINICAL TRIAL RESEARCH NETWORK SITES



2018 NIH FUNDING OF FSHD RESEARCH

THERAPIES IN DEVELOPMENT

BIOTECH & PHARMA PARTNERS

Engagement



NEW 2,300

VOLUNTEERS

813

SUPPORTERS 3.281

CHAPTERS LAUNCHED

MEMBERS

WALK & ROLL FUNDRAISERS

> **NUMBER OF** WALKERS

FUNDS RAISED

\$239,000



