



FSHD is

**the most common form of muscular dystrophy,  
but there is **no** treatment.**

One in 14,000 people are estimated to have FSHD,  
and **the actual number may be twice that.**

Help **“Unlock the Code”** to treat FSHD

**Yes, I want to help!**

October 1, 2011, Harwich, Massachusetts

**Major Sponsor: \$1,000**

Your company name on a banner at the event

Your company name mentioned in correspondence and advertising for the event

Your company logo on the Walk 'n' Roll T-shirt

Your company name listed on signs along the walk route

Your company name listed as a sponsor at the event and on the event's Facebook page

**Sponsor: \$500**

Your company logo on the Walk 'n' Roll T-shirt

Your company name listed on signs along the walk route

Your company name listed as a sponsor at the event and on the event's Facebook page

**Partner: \$250**

Your company name listed on the Walk 'n' Roll T-shirt

Your company name listed as a sponsor at the event and on the event's Facebook page

**Friend: \$100**

Your company name listed as a sponsor on the event's Facebook page

**Just want to make a tax-deductible donation of \$ \_\_\_\_\_**

**Payment may be made by check or credit card to the FSH Society:**

AMEX

VISA

MasterCard

Discover

Card number \_\_\_\_\_

Expiration date \_\_\_\_\_ Signature \_\_\_\_\_

**Please return this form with payment by July 31 to:**

FSH Society, 64 Grove St., Watertown, MA 02472

**THANK YOU!** For more information on this disease, please go to [www.fshsociety.org](http://www.fshsociety.org)