



The LGDA cordially invites you to help support its mission while celebrating the Winter Olympics



by staying home!

Our Non-Event

We're offering the gold medal of fundraising events: the one that doesn't even happen.

Let's Celebrate!

The Winter Olympics are here!

We're inviting all our supporters to watch the best athletes in the world compete.

From the comfort of their very own homes.

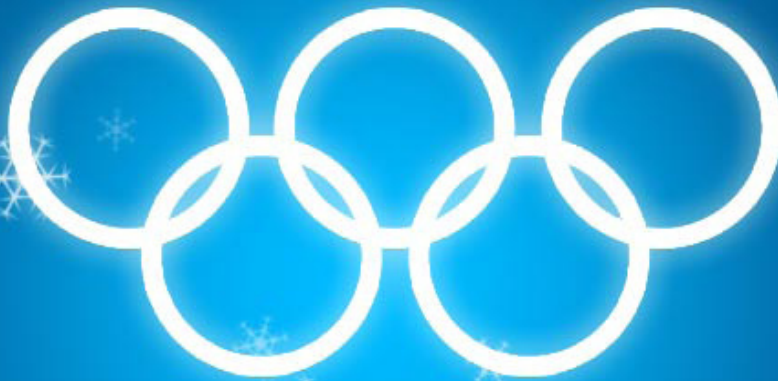
Or do something entirely different. It doesn't really matter.

Do Not

- Save a date for our event. *There isn't one.*
- Dust off your sharp digs for said event.
- Spend money on auctions and table games.

Do

- Save money on the fundraising dinner we're not holding.
- Watch the Olympics. Or do anything else!
- Return our non-event donation form and share with your friends.



W I N T E R O L Y M P I C S



Why we're doing this

It's Efficient

Our non-event has almost no overhead costs. We're not renting a hall, paying a caterer, buying advertising or twisting people's arms to get volunteer help.

Besides, it's almost impossible for an international organization like ours to gather supporters together. Oceans are in the way.

It's Effective

100% of your donation **directly benefits the LGDA** to help fulfill its mission of providing help and hope to patients (we call them Warriors) and their families.

It's Easy

Stay home and enjoy the Olympics or partake in an activity of your choosing.

All you have to do is mail in a donation or donate online at www.lgdalliance.org/donate-now.



*This, ladies and gentlemen,
is what fundraising should look like.*



Where your donation goes

Our Mission

The Lymphangiomatosis and Gorham's Disease Alliance:

- Provides information and a support network for its patients and families;
- Supports science and clinical research to help develop treatment, diagnosis and management;
- Advocates for public policies that benefit those Warriors who have these diseases.

History

The LGDA was founded in 2007 by Warrior Jana Sheets in Florida. Lymphangiomatosis took her from us in 2010. For over 20 years, Jana never met anyone who had either disease. Through the LGDA, she met dozens of others. It is estimated that less than 500 people across the world have been diagnosed with either disease.

Progress

We've started a patient registry to develop a critical mass of research to help find treatment and a cure, and we're organizing our second international patient conference set for July. Small local fundraisers have been held across the world as diagnosis and public awareness expands.

LGDA Non-Event Donation Form

Name _____

Address _____

City _____

Email _____

Phone _____

CC# _____

Exp. Date/CVC# _____

For checks: Please make payable to LGD Alliance.
Donate online at www.lgdalliance.org/donate-now.

The LGDA is a 501(c)(3) nonprofit organization. All donations are tax deductible.

Please return by March 1 to:

LGD Alliance
19919 Villa Lante Place
Boca Raton, FL 33434



Lymphangiomatosis & Gorham's Disease Alliance
To give hope and find help

www.lgdalliance.org