



February 25, 2017
www.VegasCaresAboutRare.org

Sponsorship Information and Application

Our Mission:

*Enhance the quality of life for children diagnosed with rare, life-limiting,
and complex medical conditions in Southern Nevada*



info@littlemisshannah.org
www.littlemisshannah.org

10624 South Eastern Avenue, Suite #A847
Henderson, Nevada 89052
Phone: 702-608-2488
Fax: 702-541-9947



Show your support for Childhood Rare Disease Awareness with a 5k and 1 mile fun run!

Thank you for your consideration in sponsoring our upcoming event, our *Vegas Cares About Rare 5K*. This community event is scheduled to take place on February 25, 2017, in Las Vegas, in recognition of World Rare Disease Day.

Since our first community fundraising event in 2012, we have raised more than \$60,000 to support our mission to help enhance the quality of life for young children diagnosed with life-limiting rare diseases, undiagnosed complex medical needs, and children who have been placed in hospice or palliative care in Southern Nevada. Our goal for this event is to raise \$20,000 to continue to help our special families.



Our volunteer-staffed organization offers financial assistance for families by way of our Medical and Therapy Equipment Grant program. We also organize support groups and family activities as well as hold various toy and hospitality drives to support local Las Vegas area pediatric hospital units.

But we need YOUR help. We can only continue to provide these support services and activities by obtaining the necessary funding. We offer various levels of participation and sponsorship to fit your resources.

Please help us provide support and comfort for our local families who live daily with the financial, emotional, physical, and social struggle that caring for a medically complex child can bring. We can't change their inevitability, but we can help them focus on enhancing the quality time they have with their ill child. *We can't do it without the support of our community.*

Attached you will find information regarding our organization and this event as well as sponsorship information. Please contact me at robert@littlemisshannah.org or 702-818-0140 to discuss the available options.

We look forward to your support and hope you join us on this very special day!



**Little Miss Hannah
FOUNDATION**

Robert Ostrea
Co-Founder and President
Little Miss Hannah Foundation

info@littlemisshannah.org
www.littlemisshannah.org

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February 25, 2017
8:15 am Race Time
Kellogg Zaher Park

**Show your support for
Childhood Rare Disease Awareness
with a 5k and 1 mile fun run!**

**Create or Join
a Team
Save \$5!**

Join us for an inspirational
race in recognition of
World Rare Disease Day!

5K-Timed Individual Pricing
includes custom medal,
T-shirt, and bib

\$30 – 8/1 thru 9/30

\$35 – 10/1 thru 11/30

\$40 – 12/1 thru 1/31

\$45 – 2/1 thru 2/23

Team discount pricing available

Register today at
VegasCaresAboutRare.org

Did you know...

- There are 3500+ rare diseases that affect children under 18 years old
- 30% of children diagnosed with a rare disease do not live to see their 5th birthday



All proceeds benefit the Little Miss Hannah Foundation, a 501(c)(3) organization committed to enhancing the quality of life for children diagnosed with rare, life-limiting, and complex medical conditions in Southern Nevada. EIN: 45-3993921

For sponsorship information, please contact Robert Ostrea
robert@littlemisshannah.org / 702-608-2488



SPONSORSHIP OPPORTUNITIES

February 25, 2017

Show your support for
Childhood Rare Disease Awareness
with a 5k and 1 mile fun run!

	\$2500 Rare Champion	\$1500 Rare Rockstar	\$1000 Rare Advocate	\$750 Finish Line	\$500 Water Station
Logo on Back of Shirt	✓	✓	✓		
Main Event Banner Placement	✓	✓	✓		
Social Media / Website Inclusion	✓	✓	✓		
Packet Pickup Inserts	✓	✓	✓		
Prominent Place of Logo on Back of Race Shirt	✓	✓			
Logo on Printed Materials	✓				
Free Race Entries	10	6	4	2	2
Logo and Name on Banner at Station				✓ One available	✓ Three available

Your tax-deductible donation via sponsorship will help us support the families of children diagnosed with rare, life-limiting, and complex medical conditions in Southern Nevada.

Please contact Robert Ostrea at robert@littlemisshannah.org for more information.

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Vegas Cares About Rare 5K / 1M February 25, 2017 – Las Vegas Sponsorship Form

Please return via email to robert@littlemisshannah.org or fax to 702-541-9957

Sponsorships and Gift Bag insertion items should be mailed to:
Little Miss Hannah Foundation, 10624 S. Eastern Avenue, Suite A847, Henderson, NV 89052

Sponsor Information

Corporate Name: _____

Sponsor Contact: _____ Title: _____

Address: _____

Phone: _____ Email: _____

Website: _____

Participation Commitment

Event Sponsorship

(must be received by February 1, 2017 to be included in promotional materials)

- ☐ Rare Champion - \$2,500
- ☐ Rare Rockstar - \$1,500
- ☐ Rare Advocate - \$1000
- ☐ Finish Line - \$750
- ☐ Water Station - \$500

Payment Options

- ☐ A check payable to Little Miss Hannah Foundation is attached
- ☐ A credit card authorization to Little Miss Hannah Foundation is attached
- ☐ I will be sending a check payable to Little Miss Hannah Foundation

Signature: _____ Date ____ / ____ / ____

One Time Credit Card Payment Authorization Form

Sign and complete this form to authorize the **Little Miss Hannah Foundation** to make a one time debit to your credit card listed below.

By signing this form you give us permission to debit your account for the amount indicated on or after the indicated date. This is permission for a single transaction only, and does not provide authorization for any additional unrelated debits or credits to your account.

Please complete the information below:

I _____ (full name) authorize **Little Miss Hannah Foundation** to charge my credit card account indicated below for \$ _____ (amount) on or after _____ (date). This payment is for _____.

Billing Address _____

City, State, Zip _____ Phone _____

Email: _____

Account Type: ☐ Visa ☐ MasterCard ☐ AMEX ☐ Discover

Cardholder Name: _____

Account Number: _____ CVID: _____

Expiration Date: ____/____/____

SIGNATURE _____

DATE _____

I authorize the above named business to charge the credit card indicated in this authorization form according to the terms outlined above. This payment authorization is for the goods/services described above, for the amount indicated above only, and is valid for one time use only. I certify that I am an authorized user of this credit card and that I will not dispute the payment with my credit card company; so long as the transaction corresponds to the terms indicated in this form

Offering Hope, One Child At A Time



We work with families of children with life-limiting rare diseases, undiagnosed complex medical needs, and kids with debilitating neurologic, metabolic, and genetic conditions.

Contact us for info on how to become a LMHF family!

LITTLE MISS HANNAH'S MISSION:

To enhance the quality of life for children diagnosed with rare, life-limiting, and complex medical conditions in Southern Nevada



Equipment Program



Be the Sunshine Campaign

Childhood Rare Disease Facts

- There are more than 3,500 rare diseases that affect children
- 30% of children diagnosed with a rare genetic condition will not live to see their 5th birthday
- 55% of families will incur direct medical expenses not covered by insurance



Get Involved!

Maximizing a child's happiness and enabling families to enjoy time with their child is impossible without your support.

"Be the Sunshine in a Child's Life" and join the Little Miss Hannah community today!

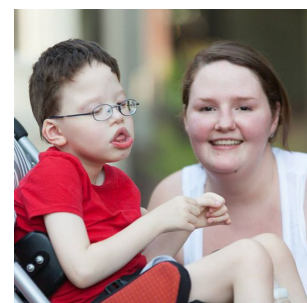
- * Volunteer
- * Host a Donation Drive
- * Become a corporate sponsor



PICU Care Bags Program



Vegas Cares About Rare



Family Activities



info@littlemisshannah.org
www.littlemisshannah.org

Find us online!



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Little Miss Hannah Foundation is a 501(c)3 tax-exempt nonprofit corporation - EIN#45-3993921

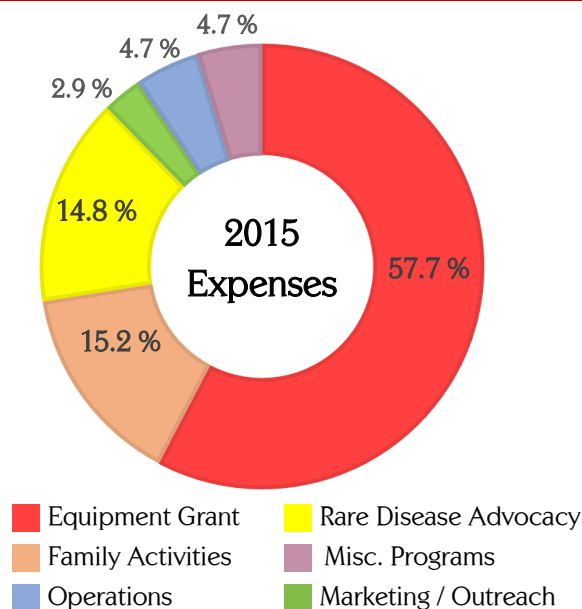
2015 Child Impact Report



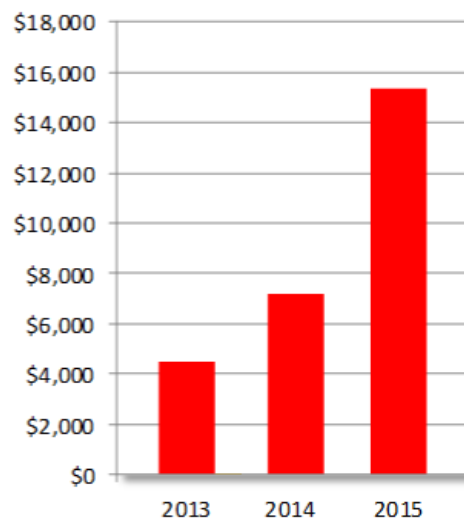
Thank you Southern Nevada for helping us enhance the quality of life for local children diagnosed with rare, life-limiting, and complex medical conditions.

As we look back at 2015, we can see what an amazing and productive year we have had! Since 2012, we have:

- Raised over \$50,000 to help local children and families
- Purchased 50+ new pieces of equipment including:
 - 5 special needs car seats
 - 4 special needs tricycles
 - 5 special needs bathing chairs
 - 3 communication software programs
 - 6 special needs feeding chairs
 - 10 special needs chairs
- Worked with 112 families with children diagnosed with 40+ rare diseases
- Welcomed over 1250 people at Town Square for our annual Rainbows in the Wind Festival Fundraiser in July 2015
- Delivered 125 care bags to local area pediatric intensive care hospital units in 2015
- Participated in Rare Disease Day in Washington D.C., Nevada Disabilities Conference, School World Rare Disease Day activities, and RARE Patient Advocacy Summit for childhood rare disease awareness and advocacy.



Expenditures for Medical/Therapy Equipment Grant Program



We are a family-run, 100% volunteer supported organization!



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