

October 20, 2005

To Whom It May Concern,

On behalf of SickKids Foundation, we are extremely pleased that Lucy and Enio Liegghio will be holding the "5th Annual Life for Luke Dinner Gala" on March 4, 2006, with proceeds supporting SickKids Foundation and Ara Parseghian Medical Research Foundation.

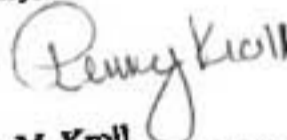
We are most appreciative of the Life for Luke Fundraising Committee's efforts as the funds raised are directed towards the research of Niemann-Pick Type C, a genetic disease affecting children that currently has no cure. The future promise of "SickKids" rests upon the availability of more funds to provide modern equipment and support services for new advances in diagnosis, treatment, care and hopefully some day - a cure. The proceeds from this event will help us work towards fulfilling this promise.

If you are able to lend your support to this event, please accept our deepest gratitude. Please note this letter is valid for one year from the date issued. Should you have any questions, please call me at (416) 813-7321.

Please direct all donations to the event organizer:

Life for Luke Foundation
c/o Lucy & Enio Liegghio
20 Airdrie Drive
Woodbridge, ON L4L 1C6

Sincerely,



Penny M. Kroll
Development Coordinator

555 University Ave
Toronto, Ontario
Canada M5G1X8

Phone 416 813 6166



Thank you for your generosity.

It has enabled these foundations to make advancements in the battle against Niemann-Pick Type C and other genetic diseases.



National Niemann-Pick Disease Foundation, Inc.
415 Madison Avenue; Post Office Box 49
Fort Atkinson, WI 53538-0049
920-563-0930; Fax #: 920-563-0931
Web Site: www.npdf.org
E-mail: npdf@idcnet.com

October 26th, 2005

To Whom It May Concern:

This letter is to inform you that, Lucy and Enio Liegghio, of Vaughan, Ontario, are active members of the National Niemann-Pick Disease Foundation, Inc. (NPDF). The Liegghio family has held events supporting the work of the NPDF in the past and plan on sponsoring another event on Saturday March 4th, 2006. The upcoming "Life for Luke Dinner Dance and Silent Auction Gala" will be their 5th annual event. Through their ongoing efforts and enduring spirit, the Liegghio family has lifted the term of "family fund raiser" to a new level. Their annual "Life for Luke" dinner gala has raised thousands of dollars for the urgently needed research required to delve into the mystery of Niemann-Pick Disease. Tragically, a treatment and cure, wasn't in time to save their precious little boy Luke but that hasn't diminished the efforts of the Liegghio's. They continue to raise funds for NPC research so that no other child has to address the tragic consequences of this disease.

The NPDF is an international, voluntary, non-profit organization made up of parents, medical and educational professionals, friends, relatives and others who are committed to finding a cure for Niemann-Pick disease. NPC Type C is a rare, genetic, degenerative, neurological terminal disease. Currently, there is no treatment or cure. Individuals with this illness rarely survive past their late teens. The Foundation was established in February of 1992 by parents of children suffering from Niemann-Pick Disease. Concerned with cutbacks in funds for medical research and knowing that without research there was little hope for their children, these parents united to form an organization, which would eventually become the National Niemann-Pick Disease Foundation, Inc.

The heart of the National Niemann-Pick Disease Foundation (NPDF) is the strength of our family members working together toward the common goals of family support and raising funds for research. The foundation continues to grow and flourish due to continued involvement and support of our families. The steady growth of the foundation has been spirited on through many different avenues in the past few years. Awareness campaigns and fundraising events, initially born out of a family's commitment to their children, have spawned numerous NP sponsored activities.


Our primary goals are to:

- Promote medical research into the cause of NPD in order to eventually find a cure
- Provide medical and educational information to assist in the correct diagnosis and referral of children with NPD
- Provide support to families of NPD patients
- Facilitate genetic counseling for parents who are known carriers of NPD
- Encourage the sharing of research information among scientists
- Support legislation that positively affects patients and families with NPD

Despite the tragic loss of Luke, Lucy and Enio Liegghio continue to be extremely involved in the commitment to try to raise funds to support research to find a cure for this devastating disease. They have worked with the foundation to raise public awareness through their volunteer efforts throughout their community. If you have any further questions please feel free to contact me.

Thank you for your support and attention to this matter.

Sincerely,



Nadine M. Hill
Director - NPDF

October 28, 2005

Mr. and Mrs. Enio Liegghio
20 Airdrie Drive
Woodbridge, Ontario
Canada L4L 1C6



Dear Enio and Lucy:

On behalf of the Ara Parseghian Medical Research Foundation—our family and thousands of others like yours who have been deeply affected by the ravages of Niemann-Pick Type C (NP-C) disease—I want to thank you sincerely for planning to host your fifth annual fundraising event on March 4, 2006.

Your past four events including a dinner gala, dance and silent auction have been very successful thanks to the wonderful generosity of your family, friends and community...as well as a lot of hard work on your part! We are thankful to be one of the recipients of the proceeds from your event in our quest to fund medical research to find a treatment for all children with NP-C. Through your events you have contributed thousands of dollars to the Parseghian Foundation to help us fund medical research.

When three of our four children were diagnosed with this fatal, genetic disease in 1994 we knew medical research was our only hope in saving their lives and the lives of other children. With the help of my father-in-law, Ara Parseghian, the legendary Notre Dame football coach, we launched the Foundation in 1994.

It has been a sad year for your family, having lost Luke to complications of NP-C disease, and for our family with Marcia's passing in August. We have both lost our precious children.

By mustering the courage and energy to hold another fundraising event to benefit NP-C research in Luke's memory shows the commitment you have to all of the NP-C families whose children can still run with ease, laugh without falling to the ground, and who still have hopes and dreams for their futures. We share the same commitment and pray no other families will have to experience what we have. Thank you for continuing the fight...we simply must.

We hope many individuals and businesses in your community will continue their wonderful generosity by becoming an event sponsor, donating items for your auction, giving gifts-in-kind needed for your event, and by attending the gala itself. Best wishes for another successful event in 2006. Thank you.

Sincerely,



Cindy K. Parseghian
Volunteer President