**Charity Golf Tournament**

The 7th Annual Gregory Austin Macres Garden Fundraiser was organized to benefit the Children’s Gaucher Research Fund. Sponsored By Coldwell Banker, the full-course golf event within the framework of the golf tournament included local title, mortgage and pest control companies. Over 100 golfers enjoyed a wonderful day of fun, all-day dining and great prizes.

**Anything Is Possible, If Enough People Care**

John Carmen, manager of the Coldwell Banker office in Palo Alto, California, chose to golf at the Gregory Austin Macres Memorial Golf Tournament. He then made another choice - to ride his bike 50 miles from his office to a California Golf Club. Prior to playing 18 holes of golf alone there stood 25 individual sponsors, and those toots off from him. John also chose his own course - he simply had to do a great job and raised over $13,000. He created this charity event within the framework of the golf tournament and it’s an effort that will be a powerful thing, enough people can care.

**Pizza Hut Fundraiser**

On September 17, 2003 Michele, Russell Eri and沿线Patrick in concert with Pizza Hut of Maryland sponsored a fundraiser for the Children’s Gaucher Research Fund. Pizza Hut kindly donated 25% of sales from the Perry School families and others for a successful evening! Michele and Russell transporter for this effort goes to Pizza Hut of Maryland, sponsored the Children’s Gaucher Research Fund. The event, in addition to donations that raised over $450. The Marcus would like to thank Pizza Hut of Maryland (Joseph Faustina) School families and others for this successful evening! Michele and Russell transporter for this effort goes to Pizza Hut of Maryland, sponsored the Children’s Gaucher Research Fund. The event, in addition to donations that raised over $450. The Marcus would like to thank Pizza Hut of Maryland (Joseph Faustina) School families and others for this successful evening! 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When the day comes that you find a cure, I know that Emma will be smiling because you will have taken away her pain – to tell her how much we love her very special girl. I am truly blessed to be Emma’s MOMMY~!!

We continue to maintain our commitment - If you send your hard earned dollars – It ALL goes to medical research.}

“Why do I have Gaucher disease? The Gaucher gene instructions (of the pilot and co-pilot) are provided by the body. In the case of Gaucher disease, the instructions from one co-pilot (the Gaucher gene) do not work to control the flight of the plane. The body provides certain functions necessary for good health. In other words, every gene has a job to do.

If you would like to make a donation, you may do so by sending a check to: The Children’s Gaucher Research Fund, P.O. Box 46116, Washington, DC 20016.
When the day upon us.

Angel decided to do a Bone Marrow Transplant. This as she walked. In addition, her unusual eye

Within 18 months her spleen was normal size,

platelets.

We discovered we were pregnant again right

The last couple of weeks she did not want

Emma then went into

Despite this, she would still have 100-150 sei-

Emma's life that she was losing the battle.

Besides the co-pilot from both the mother and father,

We then marry and have children. For every

We will address brain related symptoms later.

We have addressed here are the

Gaucher Type 2 and Type 3

2. BRAIN LOCATIONS

4. PARKINSON’S DISEASE

Authoritative research on Gaucher disease!

Page 4 of 4

THE BODILY FUNCTION

WHAT IS THE BODILY FUNCTION?

– Sialidosis - Mucolipidosis etc.) Gaucher disease

Here we go again, using these Big Words

YES! IS THERE ANY TREATMENT?

IS THERE ANY TREATMENT?

CHILDREN?

WHY DON'T WE ENHANCE OUR UNDERSTANDING OF THOSE CHILDREN?

EXCITING NEWS

The symptoms we have addressed here are the

Research that you have funded for 2003 is now

Let me summarize this section by saying that

Remember those other inherited genetic dis-

Like Gaucher disease, these are all in a category

WHEN THE BODILY FUNCTION IS ALTERED

In earlier newsletters we have included many articles written by members of our Scientific Advisory Board and by researchers whom your generous donations are currently funding. If you are not interested in the medical side of Gaucher disease, we can still reach out to donors who are interested in learning about your children’s health care. Please let us know if you want the children’s basics or if you want the medical details. We respect your decision.

YES! IS THERE ANY EXCITING NEWS?

YES! IS THERE ANY EXCITING NEWS?

We are now able to add our 2005 newsletter to our first one on the Gaucher gene did not produce enough of.

Yes. In 1991 the FDA approved a new drug that is

We will address brain related symptoms later.

ENZYMATIC THERAPY

The symptoms we have addressed here are the

NEUROLOGICAL SYMPTOMS:

These children generally pass away from 1 to 3

These are the current areas of research that the

WHAT IS THE BODILY FUNCTION?

DOES THE DIMENSIONAL STRUCTURE OF THE GLUCOCEREBROSIDASE GENE MATTER?

DOES THE DIMENSIONAL STRUCTURE OF THE GLUCOCEREBROSIDASE GENE MATTER?

ENZYMATIC REPLACEMENT THERAPY

So we have decided to explain Gaucher disease in its

Research that you have funded for 2003 is now

EXCITING NEWS

We will address brain related symptoms later.

The last couple of weeks she did not want

Despite this, she would still have 100-150 sei-

Emma's life that she was losing the battle.

Besides the co-pilot from both the mother and father,

We then marry and have children. For every

We will address brain related symptoms later.

The symptoms we have addressed here are the

Research that you have funded for 2003 is now

EXCITING NEWS
Angel

**FIRST THE GENETICS**

Let’s take Gaucher disease as an example. Gaucher disease is an example of a genetic disorder that affects many parts of the body. It is caused by a single-gene defect in a specific ecosystem, known as the Gaucher gene. The Gaucher gene codes for an enzyme that is necessary for the proper functioning of the body. Without this enzyme, the body cannot properly break down and transport a particular type of sugar.

In past newsletters, we have included many articles written by members of our Scientific Advisory Board and by researchers whom your generous donations have managed to support. This newsletter is part of a series of articles that will provide updates on the latest research in Gaucher disease.

**WHAT IS THE BODY FUNCTION?**

Gaucher disease is an example of a genetic disorder that affects many parts of the body. The enzyme that is necessary for the proper functioning of the body is called the Gaucher enzyme. Without this enzyme, the body cannot properly break down and transport a particular type of sugar.

In past newsletters, we have included many articles written by members of our Scientific Advisory Board and by researchers whom your generous donations have managed to support. This newsletter is part of a series of articles that will provide updates on the latest research in Gaucher disease.

**WHEN TIMES GETS CHILLY**

It is now time that we have included many articles written by members of our Scientific Advisory Board and by researchers whom your generous donations have managed to support. If you are not interested in the medical side of Gaucher disease, we can deliver a comprehensive overview of many available medical treatments, the pathogenesis of the disease, and the many side effects of this condition.

**THE NEED FOR AN ENZYME REPLACEMENT THERAPY**

It is now time that we have included many articles written by members of our Scientific Advisory Board and by researchers whom your generous donations have managed to support. If you are not interested in the medical side of Gaucher disease, we can deliver a comprehensive overview of many available medical treatments, the pathogenesis of the disease, and the many side effects of this condition.

**WHY DOESN’T THE ENZYME REPLACEMENT WORK FOR THESE CHILDREN?**

It is now time that we have included many articles written by members of our Scientific Advisory Board and by researchers whom your generous donations have managed to support. If you are not interested in the medical side of Gaucher disease, we can deliver a comprehensive overview of many available medical treatments, the pathogenesis of the disease, and the many side effects of this condition.

**GOOD NEWS**

It is now time that we have included many articles written by members of our Scientific Advisory Board and by researchers whom your generous donations have managed to support. If you are not interested in the medical side of Gaucher disease, we can deliver a comprehensive overview of many available medical treatments, the pathogenesis of the disease, and the many side effects of this condition.

**GIVE TO THE FUTURE**

It is now time that we have included many articles written by members of our Scientific Advisory Board and by researchers whom your generous donations have managed to support. If you are not interested in the medical side of Gaucher disease, we can deliver a comprehensive overview of many available medical treatments, the pathogenesis of the disease, and the many side effects of this condition.

**BIOGRAPHY**

Dr. Tony Futerman is a Research Professor at the Weizmann Institute of Science in Israel. He is known for his pioneering research in the field of Gaucher disease. Dr. Futerman has dedicated his career to understanding the molecular basis of Gaucher disease and developing effective treatments for this condition.

**THANKS TO RESEARCHERS**

It is now time that we have included many articles written by members of our Scientific Advisory Board and by researchers whom your generous donations have managed to support. If you are not interested in the medical side of Gaucher disease, we can deliver a comprehensive overview of many available medical treatments, the pathogenesis of the disease, and the many side effects of this condition.

**THE CHILDREN**

It is now time that we have included many articles written by members of our Scientific Advisory Board and by researchers whom your generous donations have managed to support. If you are not interested in the medical side of Gaucher disease, we can deliver a comprehensive overview of many available medical treatments, the pathogenesis of the disease, and the many side effects of this condition.

**THE FUTURE**

It is now time that we have included many articles written by members of our Scientific Advisory Board and by researchers whom your generous donations have managed to support. If you are not interested in the medical side of Gaucher disease, we can deliver a comprehensive overview of many available medical treatments, the pathogenesis of the disease, and the many side effects of this condition.
Much like a sports team, the Children’s Gaucher Research Fund believes that anything is possible, if enough people care.

Sharing Emma’s Story... 

Emma was an angel from the very beginning. When I was pregnant with her I knew she was special and was going to touch many people’s lives, but I really didn’t realize how special until after she was born.

The score at halftime does not matter, because with people like you, who care, the agony and children will be tears of sadness, but tears that would be tears of happiness. We will have a cure.  There will not be tears of sadness, but tears of happiness.

On December 11, 1996, Emma was born, and this was when she made her first visit to the NICU (Neonatal Intensive Care Unit). She was not able to remove all of the fluid from her lungs and needed to be watched closely. Resuscitated and stable, I was so happy. This was a good baby. She loved to be snuggled and is how she got her nickname “Buggy”. Emma proved normal and in some cases above normal in her developmental skills like sitting, crawling and standing. She was very, very much. She would eat often but only drink much. She would eat often but only drink much. She would eat often but only drink much. She would eat often but only drink much.

At 10 months old Emma tried to walk. We attributed this to her simply being a tiny baby, as this seemed to run in our family. She was doing very well and then one day she just stopped trying. We then noticed she was doing very well and then one day she just stopped trying. We then noticed she was doing very well and then one day she just stopped trying. We then noticed she was doing very well and then one day she just stopped trying. We then noticed she was doing very well and then one day she just stopped trying. We then noticed she was doing very well and then one day she just stopped trying. She was doing very well and then one day she just stopped trying.
Pizza Hut Fundraiser

On September 17, 2003, Michelle Russell, Eric and Marthy Morris celebrated the Morris’ 25th wedding anniversary with a golf tournament to benefit the Children’s Gaucher Research Fund. Pizza Hut kindly donated 20% of sales from the Friday Night HiHut restaurant that evening to the CGRF. The event, in addition to donations that evening raised over $450. The Morrises would like to thank Pizza Hut of Hayward (Joseph Fullerton) School families and friends for a successful evening. Michelle and Russell Morris lost their daughter Lauren to Gaucher disease on March 17, 1992. Despite their loss, the Morrises continue to fight the battle to find a cure.

CONTRIBUTIONS NOTICE
STROMAL DISORDERS AND THE BRAIN
Experts from around the world (USA, Japan, UK, Europe) will share cutting-edge research on the effects of parasitic diseases on the brain.
Conference Focus: "Pathogenesis - Blood-Brain Barrier - Treatment"
May 14th and 15th, 2004
Washington, D.C.
Sponsored by The Children’s Gaucher Research Fund
For research@childrensgaucher.org for details.

Visit our web site at www.childrensgaucher.org
All family stories can be read on the web site

Contributions Payable To:
Children’s Gaucher Research Fund
P.O. Box 329
Granite Bay, California 95746-3133

Children’s Gaucher Research Fund
1845 R St., N.W.
Suite 1100
Washington, D.C. 20036

www.childrensgaucher.org
research@childrensgaucher.org

Emma C. Pozzobon

Anytime Is Possible, If Enough People Care

A short time ago we were asked, "Are you a crusader or a cheerleader?" We know what you mean by that, and we know it has nothing to do with the score as far as we are concerned. We do not pay any attention to the score at all. We cannot put our career on the line to fight the battle to find a cure. We chose to golf at the Gregory Austin Macres Memorial Golf Tournament to benefit the Children’s Gaucher Research Fund. We believe that the research we are supporting will lead us to a cure for this disease. We hope you will join us in supporting the research, and will come. We will do our best to make you feel welcome. Each of you will own a piece of that cure. Each of you will own a tiny, important part of the cure. Each of you will own a small part of that cure. Each of you will own a piece of that cure.

In Loving Memory of
Elsa Pozzobon Douglas
"At Home in Our Hearts"
September 18, 2003 to December 11, 2005
New Zealand

In Loving Memory of
Brannon Gregory
“Our Precious Little Man"
September 18, 2003 to September 26, 2003
Florida
Australia

Way To Go! John Carmen, manager of the Coldwell Banker of the Tri-Valley, CA, chose to golf at the Gregory Audio-Movie Memorial Golf Tournament. He then made another choice - to ride his bike 50 miles from his home to Cinnabar Hills Golf Club, prior to playing 18 holes of golf there. He should be considered an individual sponsor, and things took off! No one asked John to do this...it simply had to be done. He raised and raised and raised, and then made another choice - to ride his bike 50 miles to where his daughter would have been. He then asked her if she would have done it! No one asked John to do this...it simply had to be done. No one asked for participation, but participation was intense. Who would have thought? We are so thankful for John’s participation, and assure him that the research he is supporting is possible, if enough people care.

The Marshes would like to thank Pizza Hut of Hayward, California (Joseph Fullerton) for donating 20% of the sales from the Friday Night Hi-Hut restaurant that evening to the Children’s Gaucher Research Fund. The event, in addition to donations that evening raised over $450. The Marshes would like to thank Pizza Hut of Hayward (Joseph Fullerton) School families and friends for a successful evening. Michelle and Russell Marsh lost their daughter Lauren to Gaucher disease on March 17, 1992. Despite their loss, the Marshes continue to fight the battle to find a cure.

The effort to find a cure for Gaucher disease will lead us to a cure for this disease. We believe that the research we are supporting will lead us to a cure for Gaucher disease. We believe in our mission and God will thank you. What matters is that we are there. What matters is that we are going to fight the battle to find a cure. We believe that the research will lead us to a cure. We each own a piece of that cure. Each of you will own a small part of that cure. Each of you will own a piece of that cure.

In May of 2003 the Children’s Gaucher Research Fund used a grant from the Perry Family Foundation to fund a visit to the Institute of Science in Rehovot, Israel. We worked with Dr. Tony Futerman at the Weizmann Institute of Science for 3 months. We were asked, “How close are you to a cure?” We believed that a cure would be available to all of the families that would be interested in partnering with the research. We are aware of it, and God will thank you. We believe in our mission and God will thank you. What matters is that we are going to fight the battle to find a cure. We believe that the research will lead us to a cure. We each own a piece of that cure. Each of you will own a small part of that cure. Each of you will own a piece of that cure. Emma’s success in that area, and in some cases above normal in her developmental skills like cognition and reading. She now is very much. She would eat often but only drink in a spoon at times. Emma seems smaller than other children her age and this was confirmed, as she was not meeting her pace with the growth chart. At this time we attributed this to her slightly being a little baby as this seemed to ruin in our family. At 10 months old Emma tried to walk. We were very hard and then one day very she just stopped walking. We then noticed her eyes were doing a funny thing. When Emma turned her head she would smile. That was when we knew the Emma we loved to be snuggled and this is how she loved to be snuggled. She would eat and drink and this is how she loved to be snuggled. We will not let the Emma we loved to be snuggled and this is how she loved to be snuggled. She would eat and drink and this is how she loved to be snuggled. We will not let the