

5P- News

Family Support Group for Children with Cri du Chat Syndrome

Mission Statement: To encourage and facilitate communication among families with a member who has 5p- Syndrome and to spread awareness and education about the syndrome to these families and their service providers.

5p- Society, PO Box 268, Lakewood, CA 90714-0268 (562) 804-4506 toll free (888) 970-0777 fax: (562) 920-5240
email director@fivepminus.org

President's Message *by Greg Abbruzzese*

Dear Five P Minus Society

I would like to extend a special thanks to the following Board Members who were able to attend the meetings in Dallas this past February: Paul Bakken, Gloria Griffin, Micah Hoerning, Kent Nicholls, Eileen Sherman, Georgia Richey, and Melynda Commiato (our parent representative from Dallas). Please keep in mind that these Board Members travel on their own time and at their own expense and selflessly give of their time to the Society. Thank you all!

I have received some very positive feedback from Society members who were very interested in some interactive Q&A sessions on the internet. I would encourage more members to contact me directly so that we can help improve our abilities to serve you better. We cannot improve our services unless we hear feedback. Please contact me, or any Board member directly - gabbruzzese@converse.com.

As the 2007 Minneapolis conference draws closer, we are once again looking for donations directed toward helping the conference. Paul Bakken and his family have done an excellent job securing the site, speakers, and volunteers to help make everyone's visit to Minneapolis that much more special. In fact, Paul's mother

and a friend are making over 300 embroidered bags for our families. They (we) are seeking additional assistance for room and or event sponsorship. Basically, if you would like to sponsor a breakfast, maybe a room (playroom, adult/children lounge room, etc.), it will go a long way and help defray the growing costs of running a conference. Please send any monetary assistance direct to the 5 P-Society and put a comment in the notes section that you would like these funds directed to this event. Furthermore, if you are living in the Minneapolis area, please feel free to contact Paul, or any Board member, if you would like to help with any of the logistical issues that are still outstanding.

Some of the key speakers at this year's conference will be Dennis Campbell, AmySue Reilly, Joan Henley, and Dr. Simmons (dentist). In addition Gloria Griffin and Jim and Fran Bodle will be running the grandparents workshop.

Looking ahead to 2008 IN Virginia Beach, which is being run by Christy Blount and Anne Zetlemoyer, things are still going very well. Fundraising is going smoothly with two more 10K events, another summer rental raffle (see info on page 7), and several other fundraising initiatives. We are

very grateful for all of Christy and Anne's continuous efforts!

As I had mentioned earlier, we have decided on the Dallas area to host the 2009 conference. With the assistance of Melynda Commiato we hope to have an equally successful event in 2009 as well. Families in and around the Dallas area who wish to assist with the conference, please feel free to contact Melynda at (903)396-2288, melynda@commiatohitek.com or any of the Board members.

I look forward to seeing you all in Minneapolis!

Greg Abbruzzese

Resources

- Message board: Post a message or respond to another parent's questions at http://groups.yahoo.com/group/Cri-Du-Chat_Syndrome/ Need to join group to participate. New members joining this active group daily.
- Clothing: <http://www.special-clothes.com/>
- Supported Living Services www.supportedlife.org
- Sign Language Vocabulary <http://commtechlab.msu.edu/sites/aslweb/brower.htm>
- Signing Videos and DVDs: <http://twolittlehands.com>
- Speech Therapy ideas: <http://www.speechtx.com/index.htm>
- Speech/Language and Feeding tools: <http://talktoolstm.com>
- Specialty toys – Abilitations: <http://www.schoolspecialtyonline.net>
- Identification bracelets: <http://www.stickj.com>
<http://www.mypreciouskid.com>
<http://www.911destiny.com>
<http://www.laurenshope.com>

Inside this issue:

President's Message	1
Parent's Pages	2,3
Conference Information	4,5
Contributions and Fundraising News	6,7
2007 Conference Registration	8,9
T-shirt Order Form	10

CONFERENCE
REGISTRATION
AND T-SHIRT
INFORMATION
AVAILABLE IN THIS
ISSUE OF 5P-NEWS

WHAT ARE YOU DOING MOM? NOTHING! *By Linda Mischley*

“What are you doing Mom?”

“Nothing” I replied to my oldest child.

“Then why is Bobby going to Respite this weekend if you are doing nothing?”

RESPITE. It sure is! Respite is a time to relax, sit back, and do nothing if you wish.

Respite for me started when my child with Cri-du-chat syndrome turned 14-years-old. In Oakland County Michigan a severely mentally impaired children, age 4-26 , can go to a Respite house for a weekend or a week every month.

Prior to the age of 14, I never used this Respite service because of the guilt my family put on me. In their mind, I am the Mother and Bobby is MY JOB. What else would I want to do, seems to be what they were all thinking. My family only relied on each other as babysitters for my child with Cri-du-chat syndrome. Our extended family was all out of state or 6 hours away from our home. We didn't have any family to go to for weekend respite help. So, we just muddled along.

What changed was I got mad. One weekend I had planned to fly out of town to a soccer tournament with my teenage daughter. The babysitter/staff, that was going to stay for the weekend with Bobby, backed out. That was the last straw.

I had 3 children – not one child. I wanted to be there for all my children – not just the one child. They all needed my attention. That week, Bobby was signed up for the Respite Pro-

gram.

Every month from then on I “booked Bob” and I always found “nothing” to do.

The monthly respite got Bobby used to being without me and used to the fact that I came back to get him. He had fun with other kids at Respite. The house took the kids to concerts, shows, circus, Sesame Street Live, bowling, parks, and a variety of outings in the community. He was active while at the Respite home. He didn't have time to miss me.

We never know what we are teaching our kids, and ourselves, for the next “season” of our life. I am sure that because I used the Respite house, I didn't get burned out caring for my son's daily needs. I looked forward to my freedom once a month to be just me. Bobby is still home with me at the age of 26 years old. But soon, he will move to his own house.

I am now a widow. My two older children are married and onto their own busy lives. That Respite house, that my family didn't want me to use, saved my sanity during the time of my husband's death. They made room for Bobby when I called during that difficult time. Respite ended when Bobby turned 26 years old.

Find out from your Community Mental Health or ARC if you have some sort of respite program or a respite camp program for severely mentally impaired children. I encourage you to try it and just do “nothing”.

MY DAY IN CLAYTON *by 3-year-old Gabrielle Strozier told by her mom Alaine*

Last Friday my mom, big brother Isaiah and I went to Clayton to meet some other children with Cri-du-Chat (5p minus). I was really shy when we got to Mrs. Amy's house, but after about an hour and a snack, I felt more comfortable. I played with my new friend Mason and got to gently touch PJ, my other new friend who is still just a tiny baby.

Mason is six years old, and he can growl really loud like a bear. After I heard him growl a few times, I started to do the same thing; everyone laughed. Mason was diagnosed with CDCS while he was in the hospital. Mrs. Amy, his mom said that the nurse was really good about taking care of him in the NIC-U. Mason had a hard time with acid reflux when he was a baby, but he is okay now. He is in speech therapy, OT and PT just like me. Mason also goes to school.

PJ is only ten weeks old and he is really, really cute. Mrs. Jenna found out that PJ had CDCS while he was still living in her belly. When he cried he sounded just like me when I was a baby; just like a cat. Cri-du-chat means “cry of the cat.” PJ's mom Mrs.



Jenna, let me touch him gently and I really enjoyed that. PJ is on oxygen all day to help with his breathing, and he also has a feeding tube because he has not learned to suck from a bottle just yet. I got to watch his mom feed him; Isaiah, my brother felt sad for PJ.

When I was born the doctors did not know anything was different about me. My mom knew though! I was eight months when the doctors told my mom and dad that I had CDCS. I did not have any problems when I was a baby; just a tiny bit of reflux every now and again. Most of my problems came when I got older. My muscles were weak so I started going to PT, OT and speech therapy. I am doing well just like Mason and PJ. We are all conquerors!!!

My mom was so happy to meet other families with children that have CDC. The ladies were really nice to my mom, and invited her to the CDC Convention in Virginia next year. Mom is not sure if she will go.

I had a great day in Clayton meeting my new friends.

WHAT WORKS FOR MY 5P CHILD *By Linda Mischley*

What a December day! It was snowing like crazy and I was busy with our two school age children off on a snow day. My water broke. This is not good. My due date was not for another 6 weeks. Bobby decided to be born December 3, 1980 and from that moment on, life would not be what we imagined.

Bobby was premature, Cri-du-Chat syndrome, with an inability to suck or swallow. We tube fed him for two years. During this time is when I started learning about nutrition. I learned that our bodies are made up of minerals. Without the proper daily minerals our bodies will become diseased. So I was on a mission to find the right minerals and vitamins that would help to heal Bobby's very poor health. In 1980, doctors did not recognize REFLUX as they do in 2007. So what happened was Bobby would eat, lay down, and the gasses would come up his esophagus, into his throat, and to his tiny ear canals and cause constant daily green puss ear infections. The only relief he got was being on daily antibiotics which would mask the problem until he got off the antibiotics and the cycle would start all over again. He had daily fevers. He was a very sick child. Yearly the ear infections would lead to pneumonia or dehydration and he would end up in the hospital for a week. In 1999, by the age of 17 years old, Bobby was diagnosed with pre-cancer of the Esophagus, Barrett's Esophagus. All the text books stated no known cure.

What is my point you may be asking. All during this time I was at the health food stores, reading nutritional books, buying all types of pill vitamins, minerals, proteins. Whatever I heard from friends, relatives, coworkers, radio, or television I would try. I needed to ease his daily pain which caused very poor eating. January 2000, my brother called with a new product for me to try. Naturally I said yes to ship it to me. The doctor's drugs for Barrett's Esophagus did nothing to relieve any of his pain symptoms or lessen his ear infections. He still had daily fevers, sleep apnea, choking spells, and missed a lot of school.

When the Eniva products arrived, I started using them immediately. How Eniva is different is that it is a cell-ready liquid. When the liquid enters the body, it is assimilated by the body to act as "food" for the body. Food with all the proper minerals and vitamins the body needs to heal itself. I put the Eniva products into Bobby's first morning cereal every day. He does not know he is taking anything. But his body does. From 2000 until the present, I can see daily healing.

Bobby's last Esophagus scope, October 2006, reported no sign of Barrett's Esophagus. His esophagus cells are back to normal cells. Bobby no longer goes to the hospital every year. Bobby no longer has sleep apnea. He is now sleeping through the night and so is MOM, me. His legs have straightened, his pain lines across his eyebrows are gone, and he no longer has fevers every day. He no longer is on daily antibiotics. He is no longer biting his arm out of anger because he is not mad any longer. He is calm and pays attention at school.

If you are like me, you will want to try Eniva's Jr. VIBE or Regular VIBE for 3 months. At the end of the three (3) months, you will see a difference in your child. At the end of three months, contact your Medicaid case worker and request to be reimbursed for your child's nutritional supplements. The Medicaid budget wording states they cover "nutritional supplements". My state Medicaid (Michigan) has been paying for the Eniva products (\$100 a month) for four years. Last month, the case worker decided to cut out Bobby's nutritional supplements budget because she thought any other cheap product would do. I appealed with the Judge in a state hearing and won. I proved to the Judge that VIBE is a one a day liquid vitamin/mineral that is unlike any other product I have tried for my son. It is listed in the doctors PDR reference books in the doctor's office. VIBE is 100% absorbable as our bodies think we are eating food.

We all try to get our Cri-du-Chat children to eat nutritionally. It is not easy for any of us.

VIBE just made it a little easier. Investigate the Eniva website at www.eniva.com.

Click on "ENIVAQUALITY" (on the left hand side of the webpage). You will learn why the products work in our body. It sure gave me HOPE for a healthier future for my Cri-du-Chat child and I think it will give you hope and health benefits too. If you are interested in trying VIBE email me at lmischley1980@wowway.com. Linda Mischley Eniva member #2993.

WELCOME TO MINNEAPOLIS 2007 *by Paul and Alison Bakken*

Hello from snowy Minnesota! My family and the other host families are looking forward to the 2007 conference, and hope that we see many familiar friends returning as well as many new families attending their first conference. The last few host committees have put on wonderful conferences, and we are striving to meet that high standard!

The information about the hotel and area attractions are on the Society website (www.fivepminus.org). I encourage everyone to check it out and plan ahead to make the trip more enjoyable. There are plenty of fun things to do in the area, such as the zoos, water parks, science museum and, of course, the Mall of America.

As for the conference itself, we will be following the format from past conferences pretty closely, with just a few tweaks based on feedback from last year's conference.

Thursday evening will kick off with a welcome reception and sibling mixer. Board Member Kent Nicholls has graciously agreed to offer his introduction session for new families that evening, too.

Friday morning, we will have breakout sessions until lunchtime. We will have a Friday lunch event with a state fair theme. We plan to have a burgers and hot dogs type of menu, and hope to have some carnival type games for kids of all ages. Childcare rooms will be available during the morning breakout sessions. A sibling workshop will also be available in the morning.

Friday after the lunch event will be family free time – people can visit the Mall of America or other local attractions on their own. The hospitality suites will be open in the evening.

Saturday will have breakout sessions in the morning and parent panels in the afternoon. Due to the popularity of the parent panels, the time allotted for the panels has been extended. Hopefully we will have enough time to get to all of those questions you have!

Childcare rooms will be available during both the morning breakout session and the afternoon parent panels. We will also offer a chaperoned, full day sibling outing to the Mall of America, where the kids will take advantage of the Park at the Mall, which is an indoor amusement park. The cost per child is \$25. This fee includes unlimited rides, but does not include lunch, so please send some extra spending money with your kids if they plan to attend this event.

Saturday evening we will have the traditional dinner/dance/raffle. We will have a photographer available for families who wish to purchase portraits. Hospitality suites will be open after dinner.

Sunday morning, we will have the usual wrap up session – giving families a chance to get those last minute questions answered, and to say goodbye to all of their new friends!

Please take advantage of the pre-registration form contained in this newsletter. Not only does pre-registration help us better plan for an effective conference, it will save you some money! Please also indicate if family members will be participating in the sibling workshop or outing, using the childcare rooms, or purchasing T-shirts.

Once again, we look forward to seeing you in Minnesota! The host committee is working hard to make sure that your stay is a pleasant and productive one, and we will also make certain that all of the snow is gone by the time you get here in late July!

Regards,

Paul & Alison Bakken

T-SHIRT CONTEST

HEY ALL YOU T-SHIRT DESIGNERS! WE'RE LOOKING FOR A DESIGN FOR THE 2007 MINNEAPOLIS CONFERENCE. WHAT DO YOU THINK OF WHEN YOU THINK ABOUT MINNEAPOLIS? SUBMIT YOUR ORIGINAL DESIGNS BY MAY 1ST. ALL AGES WELCOME. SUBMIT TO 5P- SOCIETY, PO BOX 268, LAKEWOOD, CA 90714.

2007 CONFERENCE INFORMATION

DATES: July 26-29, 2007

LOCATION: Sheraton Bloomington Hotel, Minneapolis south, 7800 Normandale Blvd, Minneapolis, MN 55439. (952) 835-7800.

RATES: The Society has been able to obtain an excellent room rate of \$105.00 per night (+ applicable taxes). When you call to make your reservations tell them you are with the 5p- Society. Reservations must be made by July 1st to get this great rate. Room reservations can be made either by calling the hotel directly (952) 835-7800, the toll free reservation line at (800) 325-3535 or online by clicking the online reservations button for preferred online booking on the 5p- Society website at www.fivepminus.org/annconf07.htm. **At the writing of this newsletter all Cabana rooms have been booked.** Remaining guest rooms are available. If you have any problems with your room reservations please contact the 5p- Society office at 888.970.0777 and we will help you in making your reservations.

REGISTRATION: See attached registration page (8 and 9) for rates.

T-SHIRTS: See order form for T-shirts on page 10 of this newsletter.

SCHOLARSHIPS AVAILABLE: We are pleased to announce that we have received a generous donation for the conference and are able to offer five (5) scholarships. These scholarships are for hotel accommodations for three nights at the Sheraton Bloomington Hotel and for the basic registration fee (\$150.00) for up to two adults and two children. Applicants must be members of the Society. Applications are now available and are being accepted. You may obtain an application by contacting the 5p- Society at director@fivepminus.org, or by phone (888) 970-0777 (toll free), or (562) 804-4506. We'd like to thank the Arango family for this opportunity.

SPEAKERS: The following speakers have committed to the 2007 Annual Conference:

Dr. Dennis Campbell
Dr. AmySue Reilly
Dr. Joan Henley
Dr. Kirt Simmons
Dr. Geoff Towers
Protected Tomorrows

Still waiting on commitments from:

Dr. Monica Bessler
Mr. John Halloran, Prentke Romich

SCHEDULE (times are approximate and subject to change)

Thursday, July 26, 2007

Registration 5—11 PM
Welcome Reception — 5—7 PM
New Family Orientation — 7—9 PM
Sibling Mixer — 7—8 PM
Hospitality Suite Open — 7—11 PM

Friday, July 27, 2007

Child Care Registration 7:30—9:00 AM
Registration 8—9 AM
Opening Session 9—9:30 AM
Breakout Sessions 9:30 AM —12:30 PM
Friday Fun Event—including lunch —1—4 PM
Grandparents Session—4—6 PM
Dinner on your own
Board of Director's meeting 8—10 PM
Hospitality Suite Open—7—11 PM

Saturday, July 28, 2007

Child Care Registration 7:30—9:00 AM
Registration 8—9 AM
Opening Session 9—9:30 AM
Morning Breakouts 9:30 AM—12:30 PM
Sibling outing 9:00 AM—5:00 PM
Lunch on your own
Afternoon Parent Panels—2:00—5:00 PM
Dinner, Dance, Photos, Raffles, General meeting — 6:30—11:00 PM
Hospitality Suite Open— 8:30—11:00 PM

Sunday, July 29, 2007

No Childcare on Sundays
Closing Session—10—11:30 AM

SEE YOU IN NORFOLK, VIRGINIA IN 2008

THANK YOU TO ALL OUR 2007 CONFERENCE DONORS

Many thanks to the 2007 Conference Committee for their many hours in planning and preparing the upcoming conference. In particular the numerous donations that they have received from their families, friends and businesses. Hopefully we haven't missed anyone!!

Phil & Judy Rogers
 Susan Johnson
 Mary Steininger
 Jeff & Lynette Jensen
 All About Children Pediatrics, P.A.
 Andrea Belz
 Lynn & Alice Lober
 Dave & Molly VanBronkhorst
 Don & Linda McDowell
 Greg & Kay Madson
 James & Kathryn Rosenberger
 Nancy Ciaranello
 Tim & Sarah Wagner
 Laura Wagner
 Pat & Ron Jensen
 Tom & Laura Mayer
 Judy Kocis
 Jennifer & Terry Young
 Carl & Cheri Jensen
 Rebecca Weber
 Steven & Lois Law
 Flatau Family
 Bernard & Betty Fae Nusinow
 Andy & Andrea Schanzer
 Beverly Allen
 James and Jan Ness
 Lynne Miller
 Vivian Rogers
 Meg Tilley
 Karen Crew
 Ann O'Reilly
 Lois Gosley
 Doak Mahlik
 Charles and Kimberly Fiss
 Mark and Kim Hanson
 Bethel Lutheran Church
 Claire Boettcher
 Paul & Alison Bakken
 First Weber Group Foundation, Inc.
 Wendell Smith
 Alliance Bernstein (employee match)
 Virginia Lord
 Kimberly Clark (employee match)
 Stephen & Katherine Jensen
 Oliver's Fund
 Thomson West (employee match)
 Katherine Zuccaro
 Burnsville Commercial Properties, Inc.
 Linda Berreau

Matthew & Amy Burton
 Patricia Zweber
 Thomas McCormick
 Pam Christiansen
 Robert and Sharon Spinharney
 Florence Laino
 Michael & Michele Karich
 Marian Anderson
 Corrine Bergstrom
 Mary Ann Valesano
 Debbie Soular
 Cynthia Thomas
 Louis Dechambeau
 Cynthia Nordstrom
 Susan Lohrke
 Reiss Living Trust, Elaine Reiss TTE
 George Velcheff
 Angeline Andrew
 Jennifer Fischer
 Elsie Grivette
 Julie Jorgenson
 Susan Lorch
 Tinamarie D'amato
 Jackie Soular
 Danielle Amundson
 RE Biskowski
 Kevin & Kristine Peck
 David & Kathryn Rennie
 David & Rose Eckert
 Philip Seaver
 Elizabeth Jepson
 Gislason & Hunter, LLP
 Michael Dove
 Beth Forbord
 Patricia Hol
 Susan Richard
 Bonita Reiss
 Ardeth Wensel
 Barry Grivette
 Vyna Grivette
 Mark Andrews
 Amy Laine
 Richard Velcheff
 Tammy Warren
 Donald Soular
 Susan Chelstrom
 Suzanne Dechambeau
 Hoyt Lakes Mini Storage

Over \$20,000.00 has been raised to help offset the conference expenses. Do you have an employer, friends and families that would like to assist? Still looking for donations for Welcome Reception and possible continental breakfast.

2nd ANNUAL NAGS HEAD HOUSE RAFFLE AND 10K RUN

Once again our friends at Atlantic Realty have donated a one week vacation house rental as a fundraiser for the 2008 Annual Conference in Norfolk, Virginia.

Ticket price \$10.00 a piece, limited to 1,000 tickets printed

To view house online: Go to www.atlanticrealty-nc.com; find your Outer Banks vacation rental; choose #3 advance search; \$368 "The Painted Lady."

Raffle to be held on June 27th

Please email Christy Blount at bill3125@dlba.hrcocxmail.com to purchase raffle tickets for a free week rental at this lovely five bedroom house with pool and Jacuzzi.

KaleRunning is pleased to announce [5p- Society](#) as the charity for the 2007 Chesapeake Bay 10k & Children's 1 Mile.

A running event to be held April 28, 2007. 8:00 AM at the Ocean View Beach Park that will benefit the 2008 Annual Conference in Norfolk, Virginia. For more information go to www.chesapeakebay10k.kalerunning.com. Make an online pledge donation for more info contact team@kalerunning.com.

Her Name is Montel

A story of acceptance from a father's perspective. Casey Evans tells us his story of accepting his daughter with Cri du Chat Syndrome. Casey tells us of his struggle to accept Montel and found it difficult to believe that he was the only person in the world with such apprehension. He thought it was important to tell the story of how he changed. He wanted to let others know that if they were struggling with acceptance of their son or daughter they were not the only ones to have done so, and that it is perfectly okay, they will get through it. But perhaps more importantly than how he changed, he wanted to document who changed him. It was the power of the pure and loving spirit of his own daughter, Montel. Casey hopes that new parents someday may be able to gain from his experience and realize that while it may not feel like it at the time, their own son or daughter with Cri du Chat truly is a very special human being, and quite possibly one of the greatest blessings they will ever receive in their lives.

You may obtain the 84 page book at Barnes and Noble (www.barnesandnoble.com) and Amazon (www.amazon.com). A portion of the proceeds from the sale of each book benefit the 5p-

Check out your local WalMart for a donation to the 5p- Society.

Many thanks to Georgia Richey for letting us know that individual WalMart stores will donate to various charities \$500.00. We've already received three \$500.00 donations from different WalMarts. If you'd like to assist with fundraising this would be an easy and great way to help. Contact Georgia at Georgia@richey.cc for more information.

Lilly's Lightbug

Another father, John Morrow, has written a story for children with special needs. It teaches the kids with any limitation that it is ok to learn at your own pace. This wonderful story emphasizes the importance of patience, persistence, and especially love. This story was inspired by John's daughter with Cri du Chat Syndrome, Megan.

Please email John at trcpublishing@yahoo.com and let him know if you are interested in purchasing this book and how many you would like. This way John will know how many books to print. The money will not be collected until 3 weeks before printing the books. Cost per book \$18.95 (inc shipping).

ANNUAL QUILT RAFFLE

We will once again have two quilts to raffle off at the Annual Conference; the first is our Cri du Chat Children Quilt that portrays drawings from our children with Cri du Chat Syndrome. Thank you to Carolina Kretschmar for her continued support, time and energy in producing this classic one-of-a-kind quilt.

SECOND QUILT

Our second quilt is designed and produced by Grandmother Frances Bodle. A surprise design awaits us. Raffle tickets are \$1.00 each for both quilts.



2007 ANNUAL CONFERENCE AND GET TOGETHER MINNEAPOLIS, MINNESOTA

Registration form

Yes, I plan to attend the Annual Conference and Get Together

Members—Registration fee \$150.00 USD (two adults and two children) _____

Non-Members—Registration fee \$200.00 USD (two adults and two children) _____

Each Additional Adult—\$50.00 USD each (banquet cost) _____

Each Additional Child—\$25.00USD each (banquet cost) _____

No, I am unable to attend, please accept my donation to help offset the cost of the conference _____

Total Enclosed _____

Name _____ Phone Number _____

Address _____ Email Address _____

City, State, Zip _____

Names of Adults who will be attending the conference	Relationship to 5p- Syndrome Child/Adult
_____	_____
_____	_____
_____	_____
_____	_____

Names of Children attending conference	Relationship to 5p- Syndrome Child/Adult	Age	Childcare	Sibling Event
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____
_____	_____	_____	_____	_____

Please let the hotel know if you need special items such as cribs, roll-a-ways or refrigerators. There will be an additional charge for the roll-a-ways and refrigerators.

Items requested directly from Hotel: Refrigerators _____ Cribs _____ Roll-a-way _____

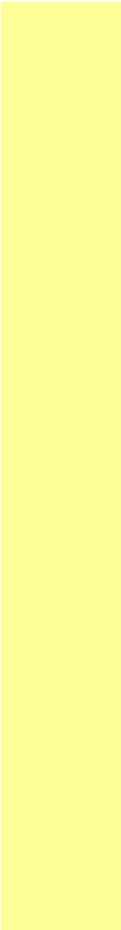
Will you be needing a high chair for the banquet? Y/N

Providing us with this information will not guarantee that you will receive a high chair, however for planning purposes we will alert the hotel of the amount of highchairs needed for the event.



5p- Society
PO Box 268
Lakewood, CA 90714-0268

Return Service Requested



T-SHIRT ORDERS

Pre-order your 2007 conference t-shirts. Order before June 15 to guarantee your shirt size. We will be having a t-shirt design contest. What do you think of when you think of Minneapolis, Minnesota? Contest is for all ages. Please send your t-shirt design to the 5p- Society by May 1st.

Volunteers of the 5p- Society, including Board Members, Conference Committee members, Welcome Committee and child care volunteers will have a different color t-shirt. Please indicate your size and color below. T-shirts are \$10.00 each. If you want to order them and have them shipped to you please add \$5.00 shipping fee (for up to 5 shirts). Please call the office if you want to have more than 5 shirts shipped.

	White	Volunteer
YS	_____	_____
YM	_____	_____
YL	_____	_____
AS	_____	_____
AM	_____	_____
AL	_____	_____
AXL	_____	_____
AXXL	_____	_____

No. of Shirts _____ x \$10.00 = _____
Shipping fee (if any): _____ Total Amount enclosed _____

Name: _____

Address: _____

Phone Number: _____

Paid by ___ check ___ credit card Visa/Mastercard (please circle)

Card # _____ Exp. Date _____