

ASAP CONNECTIONS

Providing Research, Education and Support since 1988

2014 ASAP Conference Site

In 1675 a Quaker missionary from England, encouraged by New Jersey proprietors John Lord Berkeley and Sir George Carteret, arrived to establish a settlement near the Delaware River, which was inhabited by the Lenni-Lenape Indians. The Keith survey of 1685 established the western boundary of Middlesex and Somerset counties and later, the Township of Princeton. Today Keith's Line is recognized as Province Line Road. With the laying of the cornerstone for Nassau Hall in 1754, Princeton began its development as a location for quality education. Nassau Hall was named for William III, Prince of Orange-Nassau. This simple stone edifice was one of the largest public buildings in the colonies and became a model for many other structures in New Jersey and Pennsylvania.

On July 23, this historic center will become the meeting place of the largest gathering of Chiari and syringomyelia patients and specialists in the world. Activities will take place at the Westin Princeton at Forrestal Village. Come early or stay longer to take advantage of the shopping and wineries while enjoying the fabulous amenities of The Westin such as the indoor pool, restaurants, Westin Heavenly Bed®, and more.



Set in a beautiful suburban setting, The Westin Princeton at Forrestal Village is situated in the picturesque Princeton Forrestal Village Shopping Center. This shopping center features the Koi Spa and Can Do Fitness Center as well as restaurants and select retail stores. Our convenient location on the Route 1 Corridor of Princeton puts us halfway between Philadelphia and New York.

ASAP has reserved a block of rooms at the Westin at a special guest room rate of \$135. Use one of the following to make hotel reservations:

- Call 888-627-7036 and say you are attending the ASAP Chiari & Syringomyelia Conference
- Or make reservations at <https://www.starwoodmeeting.com/Book/asapconference2014>

Attendees are responsible for their room, tax and incidentals. Events start Wednesday evening, July 23 with onsite registration and reception, and end Saturday afternoon.

Meet the 2014 Conference Host



Erol Veznedaroglu, MD, FACS, FAANS, FAHA, also known as Dr Vez, is director of the Capital Institute for Neurosciences. He is also chairman of the Department of Neurosurgery at Capital Health. Responsible for the Capital Institute for Neurosciences and the rapid development of Capital Health as a regional referral center for complex brain and spine conditions, Dr. Veznedaroglu is an innovator in developing new ways to deliver neuroscience care to patients.

His philosophy in building the Institute has been to bring together some of the most experienced neuroscience professionals from across the country, combine them with the most advanced technology and procedures, and provide the highest level of neuroscience care available in a personal, comfortable and accessible environment traditionally not found at large academic medical centers.

Dr Veznedaroglu completed his fellowship training in cerebrovascular/neurointerventional neurosurgery and neuro-critical care at Thomas Jefferson University Hospital for Neuroscience. His expertise includes the treatment of aneurysms, arteriovenous malformations, stroke, chiari malformations and other potentially fatal conditions of the brain and cerebrovascular conditions.

He is one of a select number of neurosurgeons in the country trained in both traditional neurosurgical approaches as well as endovascular approaches, enabling his patients to have the full range of treatment options available to them under the direction of the same physician.

Dr Veznedaroglu has been recognized by both peers and patients as a Top Doc, appearing frequently in SJ Magazine's Top Doc issue and as part of South Jersey Magazine's Top Docs as voted by readers feature. He was also recognized as a finalist in the category of Best Physician by NJ Biz in its annual Healthcare Heroes awards program.

Read more about Dr Vez's accomplishments at www.ASAP.org under Annual Conference.

Kids For A Cure Club

by Patricia Maxwell

Laurie Swihart and Kaylee Tutrow have volunteered to work with ASAP children. Inserted with this newsletter is a new edition of Kids' Connections.

For the second year in a row, Annie Chapman is organizing the Kids For A Cure Club Convention to be held in Princeton, New Jersey, July 23-26. She has been in touch with a number of different organizations gathering information and lining up speakers. Children 5-16 can participate while their parents attend the general sessions of the ASAP Chiari & Syringomyelia Conference.

Below is a picture of some of the kids during the 2013 meeting. As you can see from the smiles, they have a great time. Come experience the fun for yourself this July in Princeton.



Visit us on the web

www.ASAP.org

Please Note: Articles in this newsletter are not intended as a substitute for medical advice and do not necessarily represent the viewpoints of the editor, Medical Advisory Board or Board of Directors. Please contact your doctor before engaging in any new therapy or medication.

New Support Groups Formed

Kerry Chu has served as Chair of the Support Groups Program for several years. For information on support groups, contact her at Kerry_Chu@ASAP.org or 516-330-7152.

Delaware Valley ASAP Chiari/Syringo Support Group

Landing, NJ

Leader: Sharon Sweeten

email: Delaware_Valley_Support@ASAP.org

phone: 856-672-0602

ASAP's North Central MA Chiari/Syringo Group

North Boston area

Leader: Kevin Mixon

email: MA_support@ASAP.org

phone: not available

ASAP's Tulsa OK Chiari/Syringo Support Group

Tulsa, OK

Leader: Lauri Edwards & Charyl Turner

email: usmcmom69@gmail.com & charylturner@gmail.com

phone: 928-710-9144 & 918-688-2305

South Louisiana Support Group

Lafayette, LA

Leader: Rhea Dugas

email: south_LA_support@ASAP.org

phone: 337-852-5945

Huntington WV Chiari/Syringo Group

Huntington, WV

Leader: Angela Holley

email: WV_support@ASAP.org

phone: 304-638-3432

DFW Chiari/Syringo Group

Dallas /Ft. Worth, TX

Leader: Simone Key

email: dfwsupport@ASAP.org

phone: 214-640-0651

ASAP U.P. of Michigan Chiari & Syringo Group

Gwinn, MI

Leader: Stephanie Carlson-Ballone

email: stephecb@rocketmail.com

phone: 906-235-4740

ASAP's Dakota Chiari/Syringo Support Group

Pierre, SD

Leader: Halley Lee

email: shlee@pie.midco.net

phone: 605-295-0168

A Word From Our President



I can't believe we are getting ready for another conference. Where does the time go? ASAP's Chiari & Syringomyelia Conference is July 23-26 at The Westin in Princeton, New Jersey. At the meeting we plan to announce next year's location and host. We will also be getting updates on the four new research projects we funded in addition to the ongoing outcome studies and genetics research projects.

- Developing a Preoperative Chiari Severity Index to Stratify Prospective Clinical Studies for Chiari Type I Malformation by David Limbrick, MD, PhD
- Comparative Analysis of Posterior Fossa Volumes in Symptomatic versus Asymptomatic Chiari Malformation by Alan Siu, MD
- Multicenter in vitro Assessment of 4D PC MRI for Quantification of Cerebrospinal Fluid Motion in Chiari Malformation and Syringomyelia by Bryn A Martin, PhD
- Is CSF Flow the "Holy Grail" to Determining Clinicopathological Significance of Chiari Malformation? by Mark Quigley, PhD
- The Genetics of Chiari Type I Malformation (CMI) with or without Syringomyelia by Allison Ashley-Koch, PhD
- Outcomes in Patients Undergoing Surgical Intervention for Chiari Type I Malformation with Syringomyelia by Bermans Iskandar, MD

On a different note, I must say here in North Carolina I am happy to see winter end. The birds are singing, trees and flowers are blooming and the days are longer. I know for many of us the seasonal changes are not easy. Spring comes with rain and, if you are like me, you can tell the weather a day ahead. My children joke about me being the weather woman.

It's been a very busy spring. Recognizing that we had to do more for syringomyelia awareness I decided to have a walk dedicated solely to SM. The walk is titled Syringo My What? Steps Across the States. We are also continuing Together We Can Walk and Roll for a Cure.

Social Media and ASAP

Tweet, share and pin. Like us on FaceBook, join our Pinterest page and follow us on Twitter. Go to www.ASAP.org, you will see the social media icons on the upper right side of the page. Click the icon and go directly to the ASAP page.

At this time we have 13 spring walk locations. Many thanks to Beth Brixius, Heather Leitner, Tabitha Peters, Michael & Bridget Borys, Lauri Edwards, Charyl Turner, Lee Ann Clutter, Margaret Leisenheimer, Cathy Tufts, Laurie Swihart, Lori Tutrow, Anna Tannreuther, Rain Deskin, Gina Huffman and Cat Becker. Please contact me at 585-747-9651 if you are interested in being a walk coordinator. I am now booking for the fall. We work closely with our volunteers so you can have a successful walk. I also believe it gives the coordinator a source of comfort knowing they can always talk to someone. I feel like I made many new friends and I thank each and every one of you for a job well done. You provide both awareness and funds for research and education.

We completed our 25th year celebration. I would like to thank everyone for their continued support I am thankful to our volunteer Board of Directors, Medical Advisory Board and Research Committee, our community of volunteers and our staff. We may be a small group but we all work well together and get the jobs done. I would however like to build our committees so if you have a few hours a month, please contact me.

I hope to see many of you at the conference and look forward to meeting you. For those who earned the conference through fundraising, congratulations!

As I close I would like to request that everyone keep one of our board members and her family in your prayers. Renee Tobias was forced to resign due to a leukemia diagnosis that required immediate chemotherapy treatment. We will miss Renee and hope in time she will be back with us again.

Gentle Hugs always,

Patrice
Patrice_Schaublin@ASAP.org



ASAP's Chiari & Syringomyelia Conference
July 23-26, 2014

Fundraising and Awareness

ASAP volunteers have been busy this spring bringing awareness to many parts of the country by hosting events in their area. Special thanks to:

Beth Brixius - Syringo-My-What? Steps Across the States - April 5 in Great Falls, Virginia

Heather Leitner - 1st Annual Walk & Roll for a Cure 5k - April 12 in Hurricane, Utah

Tabitha Peters - Syringo 5k Walk & Roll and Kids Lollipop Walk & Roll - April 19 in Peru, Illinois

Yogi Borys - 1st Annual Yogi's Walk & Roll - April 26 in Geneva, New York

Lauri Edwards & Charyl Turner - Syringo-My-What? Steps Across the States - April 26 in Tulsa, Oklahoma

LeeAnn Clutter - Syringo-My-What? Steps Across the States - April 26 in Kingston, Oklahoma

Margaret Leisenheimer - Syringo-My-What? Steps Across the States - April 27 in Middle Village, New York

Laurie Swihart and Cathy Tufts - 3rd Annual ASAP's Sugar Land Texas Walk and Roll for a Cure - May 3 in Sugar Land, Texas

Rain Deskin - Rain's Day 1k Walk & Roll for Syringo & Chiari - May 17 in Christiansburg, Virginia

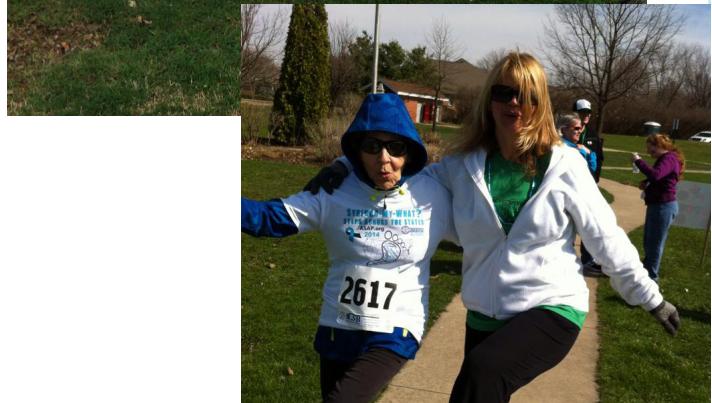
Tutrow family - 5th Annual ASAP Walk & Roll for a Cure - May 24 in Greenfield, Indiana

Gina Huffman - 1st Annual Walk & Roll Lewis Center Ohio - May 31 in Lewis Center, Ohio

Cat Becker - ASAP's Media, PA Together We Can Walk & Roll for a Cure - May 31 in Media, Pennsylvania

Anna Tannreuther - 3rd Annual Clyde's Chiari Awareness Walk - June 7 in Middletown, Ohio

Sunshine Matthews - Sunshine's Charity Run - June 27-29 in Ferguson, North Carolina



Paying It Forward

Hello everyone,

My name is Laurie Swihart. I am the proud mom of four kids with Chiari and syringomyelia. I am also happy to be the chair of this year's Auction and Raffle Committee for our 2014 ASAP Conference in Princeton, New Jersey this summer.

I am excited to have this opportunity to try and give back to ASAP by helping at this year's annual conference. For me ASAP was the first organization I found after my daughter was diagnosed with Chiari and syringomyelia 8 years ago. I was scared and at a loss because I had never heard of either of these conditions. Through ASAP.org I was able to connect with other parents who were going through the same or similar things with their children on the message boards.

I found comfort knowing I was not alone and I had others to connect with on bad or good days who would understand. ASAP.org is how we found the best doctor for not only my daughter but for all four of our children. For my family ASAP has been priceless in helping me not only educate myself through the years but it also helped me find a connection through its support groups in my area.

Two years ago my friend who is a fellow Chiari mom and I wanted to find a way to give back to ASAP. We decided to host a Walk and Roll. Since then we have had the privilege

of hosting three successful Walk and Rolls to raise funds for research and awareness in our community.

The work ASAP does is so very important. As many of you know, the research and education about these conditions are key. I am writing this letter in hopes that you might be willing to donate an item for either the conference auction or raffle. I helped with the auction last year and I loved seeing all the wonderful items that people donated such as; handmade items, jewelry, game night baskets, gift cards, weekend hotel stays, trips, purses, Chiari and syringomyelia related items, and hand crafted art work, just to name a few. Please consider helping us with a donation this year. I know that your gift will make a difference in the lives of those who have Chiari and syringomyelia.

To contribute, we ask that you take a picture of your donation, including the value and a brief description, then email to me. This way if you are joining us for the conference you can bring it with you, or as we get closer to our date, it can be shipped directly to the hotel to avoid multiple shipping costs. If you do not want to email, just give me a call with a description and value of the item. With this information we will be able to better plan our raffle and auction. Thank you for your support of ASAP. Please feel free to contact me at laurie@sugarlandpc.com or call 832-754-5581 regarding donations or questions.

Medical Advisory Board Member Takes New Position

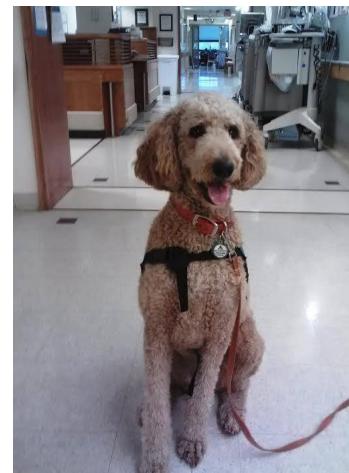
John A Jane, Jr, MD joined the ASAP Medical Advisory Board in 2009. His recent promotion as Director of the Residency Program at the University of Virginia School of Medicine and Medical Director of both its inpatient and outpatient services has led him to step down as a member of the ASAP Medical Board.

Dr. Jane wrote, "It has been my pleasure serving as a member of the Medical Advisory Board for ASAP. This is a wonderful patient advocacy organization and I have learned a great deal from my interactions at the ASAP meetings. There is a family atmosphere that I appreciate greatly and have enjoyed getting to know everyone."

We have been honored by Dr Jane's involvement with our organization as a pediatric neurosurgeon. We extend our congratulations and best wishes for the future.

Pet Therapy

Meet Ben E - the poodle who will be offering pet therapy for the ASAP Kids For A Cure program this year in Princeton, New Jersey. Register now for ASAP's Chiari & Syringomyelia Conference July 23-26, 2014! Use the registration form inserted with the newsletter or go to www.ASAP.org. Under Great Expectations' drop down menu click on Annual Conference. On the annual conference page you will find a side menu listing registration. Click on registration, fill out the form, submit, choose options and use PayPal to complete.



Remember ASAP...

When It's Time to Remember Loved Ones

Our appreciation to everyone who made a recent donation to ASAP on behalf of their friends / loved ones.

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Becky Wood

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Jeremy & Jeanna Welch

Glenda Schroeder

James Shearer

Jerry & Ethel Bradford

Kelly & Leslie Yauck

Lyndon & Joann Imke

Mary Ann Christopher & family

Ray & Elaine Dickerson



ASAP History

After hearing a television news story in 1988, Barbara White discovered the Bobby Jones Open, an annual golf tournament held to raise money for SM research. That summer, Barb and Don attended the Open in Michigan. Four people diagnosed with syringomyelia were present and it was the first time most had met another person with the disorder.

Robert A Jones, the BJO founder later served on the ASAP Board of Directors for many years. The organization supported ASAP in multiple ways through the years and continues to fund the 800 line, college scholarship program and assistive equipment grant. It is a partnership that has lasted for 26 years.

Every year since 1979, Bob Jones from around the world have gathered to play golf in honor of the "original" Bob. Over 350 Bobs from 38 states and 4 countries have participated in the tournaments. The Bobby Jones Open is committed to providing continuing support for syringomyelia research, rehabilitation or related subjects in a relentless search for a cure for this crippling disease.

The 36th annual Bobby Jones Open will be held at The Golf Club Star Ranch in Hutto, Texas (just north of Austin) on June 8-11. For more information visit www.bobbyjonesopen.com.

Contact Information American Syringomyelia & Chiari Alliance Project

Mailing Address:	PO Box 1586 Longview TX 75606-1586
Physical Address:	300 North Green Street, Suite 412 Longview, Texas 75601
Phone:	903-236-7079
Fax:	903-757-7456
Toll-free:	800-ASAP-282 (800-272-7282)
Staff:	Patricia Maxwell & Jamie Mayhan
Email:	info@ASAP.org Patricia_Maxwell@ASAP.org Jamie_Mayhan@ASAP.org

AZ (Arizona) Syringo & Chiari Support Group

Contributed by Kathleen Hall

The AZ Syringo & Chiari Support Group, a statewide organization founded in the late 1980s, has grown to over 200 members, with our membership rolls adding new people weekly. Some attend regularly and some have visited and want to remain on our email list to receive information as an option.

In 1988, the first ASAP conference in Colorado was attended by several people from Arizona who, prior to their meeting at the conference, had not met others with their condition. Returning home to Arizona with their knowledge and plans to meet and discuss their neurological conditions, they gathered for the first few years at the home of a member until larger accommodations were needed.

Through the diligence, persistence and hard work of our two founders, Judy Dragiewicz and Shelley Norris, our support group was reorganized in 2000, and through Shelley's efforts, we obtained a larger meeting room at the St. Joseph's Outpatient Rehab Building. Shelley had experienced several surgeries elsewhere and after getting to know the rehab staff where she was sent for her recovery, thought the facility and location in Central Phoenix ideal.

In 2006, Judy and Shelley both decided to "step back" and take on advisory roles as needed. We designed a blog www.azsyringochiari.wordpress.com (the current one). A committee worked on a trifold brochure, which has our local group featured with our own designed logo, a very cool ribbon, and zipper! Our AZ Syringo & Chiari Support Group's September Awareness Proclamation is a permanent document issued in our AZ State Senate, 2008. We must now work on obtaining the same for a Syringo Walk in 2015.

In January of 2014 we were granted a meeting room in the main building of St. Joseph Hospital and Medical Center. The rooms may alternate, if we have to adjust our new schedule due to a holiday weekend, there will be an ample time schedule arranged. The rooms are larger and easy to access and a bonus is the ability to have Wi-Fi and large screen viewing for all attending. Parking is available in Parking Garage 6, also Garage 3, plus there is a reduced fee for parking on Saturdays. Family and friends are encouraged to attend with you and learn who and what we are about!

We also have featured speakers covering a variety of topics who attend our meetings. Please join us and share your questions, concerns and thoughts. We look forward to meeting you!

In addition to growth of membership over our group's time span, we have seen many changes. Diagnostic procedures, types of disorders and conditions added to our list of suspected causes of chronic pain. We are only now seeing what lies out there and trying to educate ourselves and

others, including physicians who refuse to acknowledge our illness! We felt unique and alone with these disorders in the 70s, 80s and 90s but now find that they are not rare anymore as we meet more people each year. Very young children, some infants, to the older "young at heart" make up our ranks, and we welcome you all!



From that first annual ASAP conference in Colorado in 1988, our group has grown, as has ASAP's prestige and reputation because of the quality of membership of its lay people, Board members and Medical Advisory Board who back the organization all the way.

We as support group members and leaders are here at the local level to convey ASAP's outreach in any activity at any opportunity that presents itself. As for attendance at our meetings- for us, it is reaching our members and prospective members at every possible level: our blog, by phone, email and FaceBook as you will never know what media touches a person or motivates their interest into finding positive support. Attendance for us is never about the amount or number, it is simply being present for anyone who wanders in looking for a friend, support or information, or all three. Chronic conditions take a horrendous toll on all of us, the diagnosed and the caretaker. Learning to adapt and cope is an ongoing process, some of us have dealt with for many years but can always pick up new tricks from those fresh from decompression or diagnosis.

If you ever visit Phoenix during one of our scheduled meetings, please join us. You can contact us at arizonasyringochiari@gmail.com. Our meetings take place in January, April, July, October, and December- the third Saturday of the month unless that weekend is a holiday. (We'll meet the Saturday before the holiday weekend).

Time: 11:00 am to 1:00 pm
Location: St. Joseph's Hospital
Dining Room A or we will notify
350 W. Thomas Rd.
Phoenix, AZ 85013

Email or check our blog for our next meeting date. We also have "Meet and Greets" during our off meeting months. Meeting date reminders posted on ASAP email blasts, and via our FaceBook page invites!

[Editor's Note: **The July meeting has been cancelled. Next meeting date is June 21.**]

American Syringomyelia & Chiari Alliance Project Inc.
P.O. Box 1586 Longview, Texas 75606-1586

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ASAP's Mission: to improve the lives of persons affected by syringomyelia, Chiari malformation and related disorders while we find the cure.

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ASAP Connections

The American Syringomyelia & Chiari Alliance Project (ASAP) is a tax-exempt 501(c)(3) organization. Our goals include providing a clearinghouse for information on syringomyelia (SM), Chiari malformation (CM), and related conditions.

We offer a supportive network of programs and services and fund research to find better therapies and cures. ASAP is supported by tax deductible donations.

ASAP Connections is published quarterly for ASAP members. Your contributions of articles, letters, and photos are encouraged. The editor reserves the right to edit any article in order to accommodate space.

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Longview TX 75606

ASAP Connections Editor: Patricia Maxwell

KIDS' CONNECTION



Welcome to the New Kids' Connection

Hi, my name is Laurie Swihart. I am very excited to let you know that Kaylee Tutrow and I will be taking over the Kids' Connection newsletter. We are both looking forward to hearing from you all and we hope you will enjoy these newsletters. If you have any questions or ideas, please email us. Here are some interesting facts about me. I am a mom of 4 awesome kids. I have two girls, Caroline, 14, Isabelle, 12, and two boys, Charlie, 12, and Nicholas, 10. They all have Chiari, syringomyelia and EDS. I am married to a really neat guy we call Mr. Charles. We live in Texas which means we love football and we are huge Texans fans! We have three pets, our puppy Sam, Leo our cat, and a fish named Swimmy. I have been involved with ASAP for several years. I had the privilege of meeting Kaylee and her family two years ago at my very first ASAP conference. The picture to the left is of the two of us at last year's conference. I think one of the best things about ASAP is its help in making connections between people that can turn into a lifetime friendships. Here are a few words from Kaylee about herself.

Hi everybody! My name is Kaylee, I am 17 years old and I live in Indiana (Go Colts!). I live with my mom, dad, and my younger brother Keegan who is 15. Here are some things about me. My favorite color is purple, I have a beagle named Fred, and I love riding horses. I'm so excited to hear from all of you and get to know you! I was diagnosed with Chiari malformation in fifth grade. Both my brother and I have the condition. Since I was diagnosed, I have had surgery and been diagnosed with other conditions like Ehlers-Danlos syndrome. I have been involved in ASAP for a few years now, and especially enjoy going to conference and hanging out with the other kids.

~ASAP 2014 Conference Corner~



Every year, ASAP hosts a conference where doctors, families, and friends get together. This year, the conference will take place July 23-26, at the Westin Princeton at Forrestal Village. My favorite part of the whole event is spending time in the Kids For A Cure Club room. While the doctors talk to the parents, we all spend some time together in another area. In the past, we have done arts and crafts, danced, been visited by therapy dogs, and had a private meeting with a brain specialist who answered our questions. There is something for everybody to do! If you want to come to conference this year and join us in the kid's room, have your parent register at ASAP.org.

Pen Pals

We are trying to start a pen pals program so that all of us kids can talk to one another and hopefully even make some friends! If you want to join the program, please email us. You could start by telling us your name and a little about yourself. What is your favorite color? What condition do you have? What do you like to do for fun? How old are you? Once you send your email, we will put all of the kids together on a list and we can all email or write back and forth. We're really looking forward to hearing from you! If you just want to chat or have suggestions for the newsletter, please email.

We hope you all are having a great school year and are excited to talk with you!

Just for Fun

H	A	F	Z	R	A	P	C	S	T	R	E	N	G	T	H	X	Q
O	W	V	C	U	R	E	K	F	Y	N	E	T	Z	M	N	S	O
P	A	X	Q	D	W	E	F	J	K	V	S	Y	P	A	L	Q	P
E	W	Y	X	O	P	B	E	L	I	E	V	E	Q	R	H	N	U
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C	E	D	N	C	T	Q	O	J	F	U	Y	O	N	P	C	S	Z
A	S	Z	W	Q	F	Q	J	C	O	U	R	A	G	E	Q	E	C
V	S	E	C	T	S	Z	U	X	Q	V	P	D	F	P	W	M	G
Q	Z	W	O	P	A	I	L	E	Y	M	O	G	N	I	R	Y	S

CURE

HOPE

COURAGE

CHIARI

LOVE

STRENGTH

FIGHT

BELIEVE

AWARENESS

POWER

HAPPY

PURPLE

SYRINGOMYELIA

Coming up next in Kids' Connection...

In the next newsletter, we will be interviewing a child life specialist and asking her your questions. You can ask her about your disorder, your friends or family, or suggestions for fun things you can do. We will compile a list of as many as we can. The questions and answers will appear in the newsletter!

Send your questions to

Kaylee at
kidsclub2@ASAP.org

Laurie at
kidsclub1@ASAp.org



ASAP's Chiari & Syringomyelia Conference

July 23-26, 2014

Attendee Registration Form

Pre-conference registration closes July 10

Contact Information

Name: _____

Address: _____

Phone: _____ Email: _____

Additional Attendees

Please add contact information if different from above, including ages of children.

Name: _____

Name: _____

List additional attendees and contact information if needed on back.

Fees	Number	Sub-total
Adult registration by July 10 (includes banquet):	_____ x \$160.00	_____
Children's registration: ages 6 - 15 (includes banquet):	_____ x \$ 85.00	_____
Additional closing banquet tickets:	_____ x \$ 65.00	_____
Family pack: 2 adults, 2 children (ages 6-15) (Includes banquet)	_____ x \$450.00	_____

Number of vegetarian entrees banquet: _____

Special diet request: _____

Note: By submitting this form and/or attending the conference, you agree to allow ASAP to use photographs and/or videos taken of you in ASAP's promotional materials. You understand that ASAP will not publish or record any personally identifiable information without your express written consent.

Payment Information

I want to make a donation. Scholarship Fund \$_____ Conference Sponsor \$_____

Payment Type: Check Credit Card Total Amount Enclosed (see sub-total above): \$_____

Credit Card Type: Visa MasterCard Discover American Express

Cardholder Name (please print): _____

Credit Card Number: _____ Expiration Date: _____

Cardholder Signature: _____

Mail to:
ASAP Conference Registration
PO Box 1586
Longview TX 75606