IN THIS ISSUE...

- Web Page Launched: www.stuttersfa.org.
- YEARS OF SERVICE
- Genetic Study Update
- NCCD Award to Malcolm Fraser

WINTER 1997-98

The Stuttering Family Research Project—An Update

by Dennis T. Drayna, Ph.D.

Rockville, MD - The Stuttering Family Research Project at the National Institutes of Health is completing its first year with some exciting developments to report. First, within the NIH the project has received increased support from the National Institute on Deafness and Other Communication Disorders. Stuttering research is now an important priority at this Institute, and continued funding of the project is assured for the coming year.



Dennis T. Drayna, Ph.D.

The response to our appeal for volunteers has been excellent. We have heard from over 1,000 people. From these responses, we have been able to identify over 400 qualified families

which represents one of the largest groups of families with stuttering ever identified. More than 150 of these families have signed up all their qualified family members, and these family members are now enrolled and participating.

In addition, the project has now been able to identify enough qualified families to move into the next phase of the study which involves speech evaluation for each of the volunteer participants. This process is being carried out with videotaped speech samples from each volunteer, and is being done through the mail. These speech samples will allow us to begin building up the important clinical information needed from each participant.

Right now, one of our most important goals is to get *all* the members of qualified families to join our study. Participation from all the qualified members of our existing families would bring us long way toward meeting our No One Stutters in Cyberspace

Those who stutter find comfort and help on the Internet.

That's because the Internet lets the more than three million people who stutter communicate using their computer keyboards instead of their voices.

"It's changed life and made it really positive for stutterers," said Barry Guitar,

Ph.D., professor of communication sciences at the University of Vermont. People who stutter frequently feel self-conscious about their speech impairment, and the Internet provides a new way for them to express themselves.

"Those who stutter often end up feeling that what

they have to say isn't worthwhile," Guitar explained. But on the Internet, stutterers can access electronic bulletin boards and join in discussions in "chat rooms" without worrying about their stutter. "It gives them an opportunity for growth," added Guitar.

Still, experts warn that those who stutter should not use the Internet to avoid speaking.

"The more you avoid, the more powerful stuttering becomes," said Peter Ramig, Ph.D., professor of speech pathology at the University of Colorado in Boulder.

However, Ramig said the Internet contains abundant information stutterers can use to help them better manage their speech difficulties.

"I think the advantages of the Internet outweigh the disadvantages," Ramig said. "The amount

said. "The amount of information already out there is impressive."

One important source of information is the Stuttering Foundation of America's Internet site on the World Wide Web (www.stuttersfa.org). The site contains updated information on the Foundation



Stuttering Foundation of America's Web Page.

tion's books, brochures and videotapes available to the public and professionals, a nationwide resource list, and online resources.

The Stuttering Foundation, based in Memphis, Tenn., is the first nonprofit charitable organization in the world dedicated solely to the problem of stuttering. It has distributed over 3 million publications to professionals and the public since its founding 50 years ago.

Continued on page 3

World Congress on Fluency Disorders a Resounding Success

by Eugene B. Cooper, Ph.D.

Three hundred and fifty individuals from twenty-eight countries attended the International Fluency Association's Second World Congress August 18-22, 1997 in San Francisco, California. The International Fluency Association (IFA) was founded in 1991 to facilitate world-wide interchanges between researchers, clinicians, and consumers regarding the latest developments in

Continued on page 5



Dr. Phil Schneider at SFA booth with Karen Eikeland from Norway.

Continued on page 3

Stuttering Foundation Reaches Out Through New Venue

New York - Two thirteen year-olds recently helped SFA continue to educate children and adults about the nature and treatment of stuttering problems! On August 7th, Katie McHugh and Kevin Gebraski, accompanied by speech-language pathologist June Campbell, flew from Chicago to New York City to be interviewed by Children's Express.



Katie McHugh, June Campbell, and Kevin Gebraski at Rockefeller Center.

Created in 1975, Children's Express (CE) is a national nonprofit youth development and leadership organization. Children's Express uses journalism to give children a significant voice in the world. It is a news service which issues news releases. These releases are researched, reported, and edited by CE reporters ages 8 through 13 and teen editors for audiences of all ages.

Children's Express weekly column is syndicated to newspapers around the country. The following is their interview:

Children's Express "By Children...for everybody"

Kids who stutter speak out

CE News Team: Steve Craft, 18 and Asher Lindo, 12 Edited by: Julia Kim, 15 and Stella Lee, 16

More than 3 million people in the United States have some sort of speech problem; stuttering is one that many Americans, both kids and adults, have to deal with. Kids with stuttering problems are not abnormal and can do all the things that other kids can do. Sometimes inconsiderate people hurt stuttering kids because they don't know how it feels.

Children's Express conducted a roundtable discussion with two kids who stutter. We found their experiences very moving. We realized that although stuttering is a problem, people who suffer from it are not much different from anyone else. We sympathized with their hurtful experiences, and learned that something must be done about the situation.

Steve Craft, 18, CE: I myself have a stuttering problem, and I find that I stutter most when I'm doing public speaking or that kind of stuff. When does stuttering most affect you?

Katie McHugh, 13: Mostly when I'm nervous or in a tense situation, or when I think I'm going to stutter. It's this whole psychology thing in your head.

Kevin Gebraski, 13: When I'm nervous or reading out loud. One day I can read maybe a whole paragraph to the class, the next sentence it takes one minute to do it.

CE: Kids get teased quite often. How do you handle it?

Katie: I don't get a lot of teasing. I went to a very small school. There's only 10 kids in my class, so we all know each other. They know I'm sort of sensitive about that—even though about half of them are complete jerks.

Just ignore it. I don't really respond to it, because it's going to be a really tough situation unless you prepare yourself with a comeback.

Kevin: Most people who know you and like you won't do that because it's not worth it.

Katie: Certain kids tease in order to make themselves feel better. It's pretty



The interview at Children's Express.

much all the same reasons why kids tease other kids. Sometimes they might understand a little better if they would see themselves in our shoes, and see that it does hurt to be teased.

Kevin: They don't know what it's like. They'll never know.

Katie: I have been working on it for about three years now and it's a whole lot better. There are occasions where it gets worse, but you just have to practice to make it easier.

Kevin: Although my mom goes

SIDEBAR

June Campbell is a speech language pathologist who specializes in working with children and adults with stuttering problems. She is part of the *Stuttering Foundation of America*, an organization that educates both parents and other adults on how they can best interact with children with stuttering problems to help prevent the problem from ever developing.

Children's Express recently talked with Campbell about her job and what the Stuttering Foundation is doing to help out stuttering kids all across America. with me to my therapy classes and my therapist is very supportive, I think that self-motivation is most important.

Katie: My mom goes with me to all of my classes in therapy, but I think the person that really encourages me is myself. I always want to get better.

'Stuttering feels like when you swing a bat in water, really slow.'

Kevin: Stuttering feels like when you swing a bat in water, really slow, that's kind of how it is. Therapists are great. They know how children feel and they help correct it, but they don't know what we feel like unless we tell them. I was so bad, I couldn't even talk to you. It's gotten better and better over the years. I read (aloud) much better.

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Katie: It's not a problem that would affect your mind. There are perfectly normal people that sutter.

Kevin: Like Marilyn Monroe was a stutterer.

Katie: And Carly Simon and James Earl Jones.

Everyone gets nervous and stutters sometimes. We need to get this message across to people, that stuttering is not abnormal and it's not something to make fun of. Not making the person feel inferior is the most important thing that can be done for them to regain their self-esteem. It's something that cannot so much be cured but rather treated with the proper care and time.

CE: What causes stuttering?

Campbell: There have been years of research done, and there are different kinds of stuttering and different reasons for it. There may be hereditary factors as well as psychological factors. There are also emotional overtones to stuttering. When someone fears they're going to stutter, they tend to stutter even more.

CE: What would you hope to accomplish

in speech therapy?

Campbell: We would never say we're curing stuttering but that we go through a Continued on page 5

No One Stutters in Cyberspace

Continued from page 1

But just as there is good information on the Internet, those who stutter should also be wary of being misled by false claims. The problem has long been a target of charlatans with quack cures because anything which distracts the person who stutters may reduce stuttering temporarily.

"Although adults who stutter can be dramatically helped by speech therapy, there are no instant miracle cures for stuttering nor is therapy an overnight process," said Jane Fraser, president of the Stuttering Foundation.

"We hope our Web site will answer frequently asked questions about stuttering and provide in-depth information to those who need help, including parents and teachers of children who stutter," Fraser said. If you don't use the Web, call SFA toll-free at 1-800-992-9392 for information.

Stuttering Family Research

Continued from page 1

goals. Participation in our research is completely voluntary, so we are relying on the "volunteer spirit" within our families to help us reach our goal. To help make sure we get there, we are still seeking additional families, especially stuttering individuals with brothers and sisters who also stutter.

Over the past year, advances in genetics research have continued at an astonishing rate. In the past year alone, genes have been identified which contribute to disorders such as asthma, Parkinson's Disease, and diabetes. Other important discoveries have been made in studies of genetic factors in speech and language disorders, and studies to be published soon indicate that speech and language disorders can yield to the same type of approaches that have been used successfully to understand these other medical conditions.

The tools available for genetic research have advanced equally fast. Scientists continue to discover better and faster ways to look at the genetic differences that exist between all people, and to correlate these differences with inherited disorders in families. Although our project is challenging, we are gratified to see these advances become available to us in our research, and we are more hopeful than

Malcolm Fraser Honored Posthumously at **National Communication Awards Ceremony**

Actor James Earl Jones Co-Hosted 16th Annual Ceremony

Washington, DC - Malcolm Fraser, founder of the Stuttering Foundation of America, was honored posthumously

at the 16th annual National Council on Communication Disorders (NCCD) Awards Ceremony at the Lansburgh Theater in Washington, D.C. on September 9, 1997.

For his commitment to help people who stutter, Fraser was honored with the Charles Van Riper Award. Established by the American Speech-Language-Hearing James Earl Jones and SFA President Fraser. Association (ASHA)

two years ago, the Van Riper Award was named in honor of the late Charles Van Riper, pioneer in the field of speech pathology.

Fraser founded the Stuttering Foundation in 1947. Today SFA staffs a stuttering hotline, publishes 23 books and eleven brochures on stuttering, and distributes over 500,000 publications



and videotapes a year to those who stutter, their families, speech-language pathologists, and other professionals.

> Jane Fraser, daughter of Malcolm Fraser and current president of the Stuttering Foundation of America, accepted the award from Van Riper's son John.

> NCCD, the umbrella organization for ASHA and 32 other national organizations serving people with communication disabilities, recognizes those who devote their time and energy to increase public awareness of

communicative disorders as well as those who have met challenges presented by speech, language, and hearing disabilities and who have inspired others by their achievements.

Special guests included Senator Trent Lott, Senator John Glenn and Mrs. Annie Glenn, and co-host actor James Earl Jones.



Fraser accepting award for her late father, Malcolm Fraser.



Malcolm Fraser shown receiving NCCD Award from Dr. Kathleen Griffin in 1984.



Fraser with ASHA President Charlena Seymour and Executive Director Frederick Spahr.

ever that our research will be able to give us an understanding of some of the causes of stuttering.

We want to particularly thank all the people who have contacted us and offered to help in our research. We are well on our way to meeting our goal of 500 families enrolled. If you think you and your family may be qualified and you haven't contacted us yet, please

take a minute to contact us now. Every family counts!

Editor's note: If you believe that you and your family qualify, contact Dr. Dennis Drayna, National Institute on Deafness and Other Communication Disorders, National Institutes of Health, 5 Research Court, Room 2B44, Rockville, MD 20850. Tel: (301) 402-4930; Fax: (301) 480-8019.

New England Workshop Lauded and Oversubscribed!

Second Annual New England Workshop Held June 26-30, 1997

by Suellen Wedmore, M.Ed, CCC, SLP

Boston, MA - "Excellent! This was the best professional workshop I've ever attended!" said Vicki McCready, M.Ed., SLP, a participant of the second annual Stuttering Foundation of America New England Workshop.

Here twenty-two speech-language pathologists from the eastern United States and the U.S. Virgin Islands, selected from over 450 applicants, spent five days in Boston, Massachusetts, honing their skills in diagnosing and treating early childhood stuttering.

The participants represented a wide range of therapy settings and experience, but all showed a strong interest in stuttering. "It was so valuable. And I learned a lot from the participants themselves", said a recent graduate in the field who works in a public school.

The workshop began with an overview of the theoretical framework of stuttering presented by Dr. Susan Dietrich, Ph.D., of the University of North Carolina at Greensboro, and Sheryl Gottwald, Ph.D., of the University of New Hampshire. Next, the presenters shared practical ideas for diagnosis and therapy.

Dr. Gottwald presented The Capacities and Demands Model to help under-

practice in Boston, presented information on stuttering and problems which co-exist with it.

Barry Guitar, Ph.D., from the University of Vermont, encouraged students to develop their own metaphors to help to better understand stuttering and thus to gain control over it.

"All of the presenters were knowledgeable and interesting as well as being empathetic human beings," said participant Susan Brooks, M.A., of Harvard, Massachusetts. A highlight of the workshop was meeting Jane Fraser, president of Stuttering Foundation of America who presented a history of the Foundation, founded by her late father Malcolm Fraser in 1947. "I was moved that she traveled so far to be with us," added Brooks, "and by the great effort she made to get to know each of us as individuals.'

One afternoon participants alternated roles as therapists and parent consul-

stand and explain disfluency in young children, and Dr. Dietrich presented a direct treatment model, Tension Control Therapy, that provides visual and proprioceptive feedback that students can use to control their speech. Maureen Tardelli, M.Ed., in private

tants for preschool and school-age children. "I really appreciated the time devoted to intervention and to real therapy situations with hands-on practice. I would recommend this workshop to anyone who is going to work with stut-terers," said Vicki McCready. And Mary Beth Coltharp of Richmond, Maine, echoed this enthusiasm, saying, "It was an incredible experience. I'd go again in a heartbeat!'



Sheree Reese, Perry Flynn, Susan Brooks.



Rachel Baumgarten, Susan Lewis, Barry Guitar.





Attendees at reception on the last day.

Back row standing: Susan Brooks, Frances Chang, Carolyn Coss, Patricia Ottaviano, Perry Flynn, Alice Berger, Sheree Reese, Suellen Wedmore, Jennifer Walton, Helen Anis, Rachel Baumgarten, Susan Lewis, Marie Fournier, Ann Moosman. Second row: Ruzhena Sikharova, Dianne Kudrez, Laura Specyal, Cheryl Cadmus, Barry Guitar, Vicky Williams, June McMillen-Leach, Mary Beth Coltharp, Vicki McCready. Third row: Sheryl Gottwald, Susan Dietrich, Maureen Tardelli.

The Stuttering Foundation of America is a tax-exempt organization under section 501(c)(3) of the Internal Revenue Code and is classified as a private operating foundation as defined in section 4942(j)(3). Charitable contributions and bequests to the Foundation are taxdeductible, subject to limitations under the Code.

Sertoma Announces Sixth Annual Communicative Disorders Scholarship Program

(Kansas City, MO - December 1, 1997) Sertoma International announced today that it will continue its financial support of master's students of audiology and speech-language pathology. Currently in its sixth year, the Sertoma Communicative Disorders Scholarship Program has provided more than \$400,000 in educational assistance, making Sertoma the largest source of funding for master's students studying communicative disorders.

"Our annual \$75,000 scholarship program accomplishes two very important things: It helps ensure that qualified professionals will be available in the future to assist people who have communicative disorders. Also, it fulfills and perpetuates the *SER*vice *TO MAnkind mission*," said Jon C. Haapajoki, president of the 86-year-old, nonprofit volunteer organization.

Sertoma will award thirty \$2,500 scholarships for the 1998-99 school year to citizens of the United States, Canada, and Mexico. Qualified applicants must have at least a 3.2 minimum grade point average on a 4.0 scale. The application deadline is 5:00 PM Central Time on Friday, March 27, 1988.

Students interested in the awards, funded by the Sertoma Foundation, should send self-addressed, stamped envelope to Communicative Disorders Scholarships, Sertoma International, 1912 East Meyer Blvd., Kansas City, MO 64132.

World Congress on Fluency Disorders

Continued from page 1

the understanding, assessment, and treatment of stuttering.

During the Congress' five days, one hundred and fifty presentations ranging from day-long seminars to brief reports





Participants from Northwestern Workshop for Specialists attending the IFA Congress are rounded up by Newsletter Editor Nina Reardon for group photo.

were made not only by scientists and clinicians who treat stuttering but by those who stutter who are involved in support and self-help groups. One such



Speech-language pathologists from Argentina, Beatriz de Touzet, Karina Couselo-Rios, Mariela Ginhson meet with SFA's Fraser.

presentation was a fifty year history of the Stuttering Foundation of America by SFA President Jane Fraser. The Foundation also hosted a reception, giving attendees the opportunity to exchange ideas in a social setting.

At the opening ceremony, Congress President Eugene B. Cooper, Distinguished Professor at Nova Southeastern University in Fort Lauderdale, Florida, applauded what he perceives to be a dramatic shift in the thinking of researchers and clinicians in recent years.

Cooper noted that both researchers and clinicians are viewing stuttering as a



Authors Marty Jezer and Fred Murray confer at SFA reception.

Anders Lundberg of Sweden and Marie-Pierre Poulat of France meet at SFA reception.



syndrome. As a result, they are no longer focusing exclusively on the act of being disfluent, but are attempting to identify the affective, behavioral, and cognitive components. With stuttering no longer being viewed as a unitary behavioral disorder, clinicians are addressing their stuttering clients' feelings and attitudes as well as the problem's behavioral aspects.

The Third World Congress on Fluency Disorders will be held in Copenhagen, Denmark, in the year 2000.

SIDEBAR

Continued from page 2

process designed to diminish the problem as much as possible. We would let the child know what our goal is because if they understand what it's all about, it's going to mean more to them and they'll be more cooperative. Our goal is to decrease the stuttering as much as possible, and also to help the person be less embarrassed about it.

CE: Is it harder for adults or children to stop stuttering?

Campbell: It's harder for adults if they have never had speech therapy before. And yet with children it's different because the adult has to help the child understand what's happening. We firmly believe in educating parents on how to talk with their child early on. We encourage them to speak slowly yet naturally, and not to rush the child in even non-

verbal ways. Early prevention and early intervention is recommended. Most pre-school children while learning to talk go through a stage that sounds like stuttering and it may be totally normal. We don't want the adults in their lives to be pushing them too hard about what to say and how to say it. Instead they should model a relaxed manner of speaking.

CE: Do kids who stutter have harder problems in school?

Campbell: Not necessarily as far as their intellect or their potential in class, but they may have a harder time in school when it comes to feeling uncomfortable with verbal interactions. Stuttering doesn't make kids abnormal or inferior and this should be a realized fact. We should increase education and promotion of stuttering with things like this article, newspapers or TV, all factors of media. Hopefully once the public knows of the issues, problems such as harassment can be eliminated, and kids

can feel that they belong.

There's no guarantee that the teasing will stop though because some kids want to be mean and that's their prerogative, while others just don't care. Kids put other kids down to hide insecurities about themselves, and that makes them feel better. Also, if everybody's making fun of someone else and they want to be part of the crowd, they may join in, succumbing to peer pressure. But kids who stutter can join speech classes that will help and the best thing is not letting it get to you, because if there are friends and people who care for you and support you, then you'll be okay.

The Stuttering Foundation of America can be reached at 1-800-992-9392.

Children's Express is a news service reported by children ages 8 to 18. For more information call us at (212) 741-4700 or visit our website at http://www.ce.org

NEWSBRIEFS

- This year's SFA conference on the school-age child who stutters will be co-sponsored with the California Speech-Language-Hearing Association and held in Newport Beach, California, on June 5th and 6th, 1998. For more information, contact SFA at 1-800-992-9392. Conference coordinator will be Dr. Judy Montgomery.
- The annual "Stuttering Therapy: Workshop for Specialists" co-sponsored by the Stuttering Foundation and Northwestern University will be held at Northwestern University in Evanston, Illinois, July 13-24, 1998. For more information about the workshop, contact the Stuttering

Foundation at 1-800-992-9392, or Dr. Hugo Gregory, Stuttering Programs, Northwestern University, 2299 Campus Drive North, Evanston, IL 60208. This unique workshop brings together speech-language pathologists from all over the world.

- The Stuttering Foundation of America New England Workshop, "Diagnosis and Treatment of Children Who Stutter: Practical Strategies," directed by Dr. Sheryl Gottwald, will be held at Boston University, Boston, Massachusetts, July 28th through August 2nd, 1998. For application and information about traineeships, call 1-800-992-9392.
- The SFA Southeastern Workshop, "Diagnosis and Treatment of Children Who Stutter: Practical Strategies," directed by Dr. Susan Dietrich, will be held at University of North Carolina-Greensboro, in Greensboro, North Carolina, June 10th through June 14th, 1998. For more information, call SFA at 1-800-992-9392.
- For those interested in joining Toastmasters International as a way to improve fluency, communication or public speaking skills, their address is: Toastmasters International, Inc., Attention: Membership Department, P.O. Box 9052, Mission Viejo, CA 92690, Telephone: (714) 858-8255; Fax: (714) 858-1207.
- For those looking to obtain a copy of Self-Therapy for the Stutterer in Japanese, write to Dr. Shokichi Nakajima, 2-21-1 Ogawa Machida-shi, Tokyo 194, Japan, telephone/fax: 0427 (96) 5092.
- For those looking to obtain a copy of Self-Therapy for the Stutterer in French, write to the Association des Begues du Canada, 7801 Rue Ste.

Stuttering Foundation Reaches Pediatricians at Annual AAP Convention

Because many parents turn first to their pediatrician for help when they hear their child begin to stutter, making sure pediatricians have up-to-date information on stuttering is of utmost importance to SFA.

Speech-language pathologists June Campbell, M.A., and Diane Hill, M.A., were on hand at the Stuttering Foundation booth in New Orleans for this year's convention of the American Academy of Pediatricians.



Diane Hill discusses publications with two pediatricians.

They fielded questions about stuttering and shared Foundation books, brochures, and videotapes with the more than 9,000 pediatricians and allied medical professionals in attendance.



Speech-language pathologists June Campbell and Diane Hill confer with pediatricians at Stuttering Foundation booth in New Orleans, November 1-3, 1997.

The Foundation also reached out to those in the nursing profession in 1997 with informational booths at the conventions of the Association of School Nurses and the National Association of Pediatric Nurses.

Claire, Montreal, Quebec, Canada, H1L 1V8. Please enclose \$15.00 Canadian to cover printing, postage, and handling costs.

■ Dr. Anne Foundas is looking for adults who stutter in the greater New Orleans area interested in participating in a project to study neuro-anatomical models of stuttering. If you are interested in participating or have any questions or comments, contact Dr. Anne Foundas at the following address: Department of Neurology, Tulane University School of Medicine, 1430 Tulane Avenue, New Orleans, LA 70112-2699. Phone: 504-595-5605 (Voice Mail and evenings, ask for Cassie Browning), 504-588-5734 (Days, ask for Laura Cardin).

New Books on Stuttering Available from Bookstores:

- Clinical Decision Making in the Diagnosis and Treatment of Fluency Disorders by Walter H. Manning, Ph.D., published by Delmar Publishers, Albany, New York.
- Nature and Treatment of Stuttering: New Directions by Richard F. Curlee, Ph.D. and Gerald M. Siegel, Ph.D., published by Allyn and Bacon,

Needham Heights, MA.

- Special Children, Challenged Parents, by Robert A. Naseef, Ph.D., published by Carol Publishing Group, Secaucus, New Jersey.
- Straight Talk on Stuttering, Information, Encouragement, and Counsel for Stutterers, Caregivers, and Speech-Language Clinicians by Lloyd M. Hulit, Ph.D. The publisher is Charles C. Thomas, Springfield, Illinois.
- Stuttering: A Life Bound Up in Words by Marty Jezer. Published by Basic Books, 1997. This book may also be purchased through the NSP, 1-800-364-1677.
- Stuttering and Science by William H. Perkins, Ph.D. It is published by Singular Publishing Group, Inc., in San Diego, California.
- Stuttering: A Search for a Cause and a Cure by Oliver Bloodstein, Ph.D.. It is published by Allyn & Bacon, Needham, MA.
- Does My Child Have a Speech Problem? by Katherine L. Martin. Chicago Review Press, Inc., Chicago, IL.

New Videotape Offered Free to Public Libraries

The new 38-minute videotape, Therapy in Action: The School-Age Child Who Stutters, is getting rave reviews from speech-language path-

ologists, parents, teachers and physicians. The tape is an excellent resource and is certain to further the understanding of stuttering and what can be done to help the schoolage child.

It was produced by Edward G. Conture, Ph.D., of Vanderbilt University, Barry E. Guitar, Ph.D., University of Vermont, and Jane Fraser of SFA in collaboration with June H. Campbell, M.A., Northwestern University, Hugo H. Gregory, Ph.D., Northwestern University, and Patricia M. Zebrowski, Ph.D., University of Iowa.

For the past four years, the Stuttering Foundation has sent its videotapes free to thousands of public libraries. "There are few more distressing problems for teenagers than stuttering. We are delighted to be able to make this invaluable resource, Do You Stutter: Straight Talk for Teens, available to them," said Mary Louise Abrams, many people of Panola County. The SFA will continue to offer free

tapes to public libraries, and the newest videotape for the school-age child is no exception. If your library would like a copy, have them call the Foundation at 1-800-992-9392, or write to the address

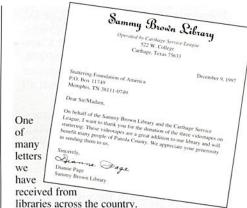
on the back of the Newsletter.

Once the library has shelved the tape, SFA will send a press release to local newspapers to apprise people of its availability.



director of the Paramus Public Library. Sue Corcoran of the Ontario March of Dimes was also enthusiastic. "I believe that this video is a valuable addition to our present collection."

Dianne Page of the Sammy Brown Library adds, "I want to thank you for the donation of the three videotapes on stuttering. These videotapes are a great addition to our library and will benefit



National Stuttering Awareness Week May 11-17, 1998















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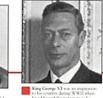
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stop you.











you are definitely in good company!

Don't put off ordering your NSAW brochures for National Stuttering Awareness Week 1998. The brochures feature thirteen famous people who stutter and are an addition to any clinic or classroom. For information on how to order, call 1-800-992-9392.

Happy New Year from the Stuttering Foundation of America



Back row: Pat Hamm, Roberta Brugge, Anne Edwards, Tracy Grever. Front row: Jane Fraser, Leslie Harrell. Not Pictured: Don Edwards, Joe Fulcher, John Bell.



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