Thank you to everyone who made the 2007 Challenge Grant a success! You not only met the challenge, it was exceeded – AGAIN! Over $25,000.00 was received towards the challenge. Combined with the anonymous benefactors’ donation of $10,000.00, we have over $35,000.00 received as a direct result of this opportunity!

The Wilson’s Disease Association will issue its third Request for Proposal (RFP) for basic science research! The WDA will fund $30,000.00 in basic science research in 2008! Building upon the success of our previous RFP’s, the WDA will accept grant proposals for innovative basic, clinical or translational research relevant to the cause or treatment of Wilson disease. The WDA grant program is designed to provide seed funding to investigators for the testing of initial hypotheses and collecting of preliminary data to help secure long-term funding by the National Institute of Health and other major granting institutions.

Twisted – The rest of the story…

Many of you know our WDA Secretary, Carol Terry and her long history of involvement with the Association. On this very special celebration of the 25th Anniversary of the WDA we are pleased to share the rest of her story with you.

Please make your conference travel plans accordingly so that you will not miss this special event on Friday evening, April 25. Dr. Joel Herskowitz, will present his original dramatic reading, “Twisted: The Carol Terry Story” in conjunction with the Welcome Reception. We are pleased to include this performance in the 2008 Annual Conference agenda for no additional charge.

If you plan to attend “Twisted” on Friday evening, you will want to be sure to register for the Conference Banquet being held on Saturday evening. Dr. Herskowitz, who is the Keynote Speaker, will share more of Carol’s story and attendees will discover what inspired him to create this portrayal of Wilson disease. Please indicate if you would like to attend on the Conference registration form. There is an additional fee to attend this event.

Please register to attend these events on the 2008 Conference Registration Form included in this issue of The Copper Connection.
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The Wilson’s Disease Association is a nonprofit 501 (c)(3) organization.

The Copper Connection is a quarterly newsletter published by the Association that informs members of findings in the area of Wilson’s Disease. There is no copyright. Newsletters and other publications can disseminate any information in The Copper Connection. Please cite attribution to the Association and the author.

The Copper Connection  
1802 Brookside Drive  
Wooster, OH 44691  
888-264-1450  
kimberly.symonds@wilsonsdisease.org  
www.wilsonsdisease.org

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**Our Mission Statement**  
The Wilson’s Disease Association funds research and facilities and promotes the identification, education, treatment and support of patients and other individuals affected by Wilson’s Disease.
HOTEL RESERVATIONS:

HYATT REGENCY O’HARE
800-233-1234 (Reservations)
(*Request: Wilson Disease Assn Group Rate*)
847-696-1234 – Hotel direct number.

GUEST ROOM RATES:

$139 single occupancy  
$149 double occupancy  
$154.00 triple occupancy  
$159.00 quad occupancy

All above rates are exclusive of 13% tax (taxes subject to change). A credit card is required to guarantee your room. Rooms are on a first come, first serve basis.

CONFERENCE RATES:

$100.00 per person*  
Please make checks payable to the Wilson Disease Association.  
Registration and checks are due by Friday, April 3, 2008. After this date, registration is $125.00 per person.

*Registration includes the Friday night reception, breakfast, lunch and a break on Saturday.

25 YEAR ANNIVERSARY BANQUET:

$75.00 per person  
Join us in celebrating the WDA’s 25th anniversary on Saturday night, April 26!

(This is an optional and not inclusive of the $100 conference fee.)

AIRPORT:

Chicago O’Hare – there is a free airport shuttle to the hotel

FUN CONTEST!

For years several WDA members have claimed they are the oldest living WD patient. We decided to find out the answer and settle this debate once and for all. The winner will receive “The Oldest Living WD Patient” award at the 2008 Annual Conference Banquet. If you wish to enter, please complete the entry form below and mail to the WDA office, or include the information requested in an e-mail to: Kimberly.symonds@wilsons disease.org, with the word “Age” in the subject line. Deadline for entry is April 15, 2008. Good luck!

Name: ____________________________________  Date of Birth: _________________

Address: ________________________________________________________________

Phone: ________________  Email Address: ____________________________________
WDA 2008 ANNUAL CONFERENCE AGENDA
“25 YEAR WDA FAMILY REUNION”
Hyatt Regency O’Hare, Chicago Illinois

Friday, April 25
8:30am – 4:30pm  WDA Board of Directors Meeting
4:30pm – 7:00pm  Welcome Reception
Mingle with other attendees while enjoying appetizers and beverages then join us for a very special performance of “Twisted”, a dramatic reading by Joel Herskowitz M.D.

Saturday, April 26
8:00 – 8:30am  Registration and Continental Breakfast
8:30am – 8:45am  Kimberly F. Symonds - Executive Director, Wilson’s Disease Association
                    Mary L. Graper – President, Wilson’s Disease Association
8:45am – 9:05am  “Studies of Microbes and Mice to Understand the Roles of Copper in Health and Disease”
                    Dennis J. Thiele PhD. Vice Chair Department of Pharmacology and Cancer Biology
                    Duke University Medical Center, Durham, NC, WDA Medical Advisory Committee
9:05am – 9:25am  “Characterization of Several Disease-Causing Mutations in Wilson Protein”
                    David L. Huffman PhD. Associate Professor, Department of Chemistry, Western Michigan University, Kalamazoo, MI. 2006 WDA Research Grant Recipient
9:25am – 9:45am  “Newborn Screening for Wilson Disease”
                    Kimberly Symonds WDA Executive Director; Secretary, National Organization of Rare Disorders (NORD) Board of Directors; Board member, Save Babies Through Screening Foundation
9:45am – 10:05am Q & A of Previous Speaker Panel – Drs. Thiele, Huffman, and Ms. Symonds
10:05am – 10:15am Break
10:15am – 10:45am  “A Practice Guideline on Wilson Disease”
                    Michael Schilsky M.D. Medical Director - Liver Transplantation, Yale-New Haven Organ Transplant Center, New Haven, CT. WDA Medical Advisory Committee Chair
                    Eve A. Roberts M.D. FRCPC. Professor of Medicine, Pediatrics, and Pharmacology, University of Toronto, Canada. Hepatologist, The Hospital for Sick Children, Toronto. (On Sabbatical). WDA Medical Advisory Committee.
10:45am – 11:05am  “Wilson’s Disease Management: Special Situations Including Nutrition & Pregnancy”
                    Fred Askari M.D., PhD. Associate Professor of Internal Medicine; Director of Wilson’s Disease Clinic, University of Michigan. WDA Medical Advisory Committee
11:05am - 11:20am Q & A of Previous Speaker Panel – Drs. Askari, Roberts & Schilsky
11:20am – 11:40am  Wilson’s Disease Association Business Meeting
11:45am – 1:15pm  Lunch
1:30 – 1:50pm  “Pediatric and Adolescent Acute Liver Failure and the Need for Transplant”
                    Simon Horslen M.D. Professor of Pediatrics, University of Washington; Medical Director - Liver & Intestine Transplantation, Children’s Hospital & Regional Medical Center, Seattle, WA.
1:50 – 2:10pm  “Decreased Bone Density in Wilson Disease Patients”
                    Karl Heinz Weiss M.D. Wilson’s Disease Clinic, Dept. of Internal Medicine, University Hospital of Heidelberg, Heidelberg, Germany
2:15pm – 2:30pm  “Rehabilitation of Movement Disorders in Wilson Disease”
                    Tanya Simuni, M.D. Associate Professor of Neurology; Director, Parkinson’s Disease and Movement Disorders Center, Northwestern University, Chicago IL.
2:30pm – 2:45pm  Q & A of Previous Speaker Panel – Drs. Horslen, Simuni & Weiss
2:45pm – 3:00pm  Break
3:00pm – 5:00pm  Breakout Sessions – Facilitated by the Speakers
6:30pm – 9:30pm  25th Anniversary Conference Banquet
                    Keynote Address - “Twisted” The Carol Terry Story
                    Joel Herskowitz M.D. Division of Pediatric Neurology, Boston Medical Center, Boston, MA.
                    Presentations and Awards – Medical Student Writing Competition and Volunteers

Sunday, April 27
8:00am – Noon  WDA Fundraiser Walk for Wilson Disease, “WOW – Walk Out Wilson’s”
                    Lincoln Park, Zoo Entrance - 2400 N. Stockton Drive, Chicago
                    Transportation available from the hotel
                    Registration: 9 – 10:00 am  Kick-Off: 10:00am
WILSON’S DISEASE ASSOCIATION
2008 ANNUAL MEETING REGISTRATION

Due on or before April 3, 2008

Name:___________________________ Email address:_____________________________________

Address:___________________________________________________________________________

City:___________________ State:______________ Zip:___________ Country: _________________

Telephone #:______________________ WDA member:   Yes  No         Membership Level:_________

May we publish your address, telephone and email on the meeting attendee list:  Yes   No

Name of others attending with you: Relationship:   If under 21 – age:
1._____________________________________ ____________________ ________________
2._____________________________________ ____________________ ________________
3._________________________________________________________ ________________

Payment Information:

If registered by April 3, 2008 - $100.00 per person
If registered after April 3, 2008 - $125.00 per person

_____ No. of Conference registrations @ $100.00 (or $125.00) ____________________
        Membership Deduction ____________________
_____ No. Attending Friday Night Reception ____________________
_____ No. of Banquet registrations @ $75.00 ____________________
        Total amount: ____________________

_____ Please check here and circle your meal request, if necessary:     Kosher    Vegetarian
_____ Number requesting WDA provided transportation to the Walk-A-Thon on Sunday

Check #______Amount: $__________ (Check payable to the Wilson’s Disease Assn)

_____ Visa #________________________________ Exp. Date:_____________ CID: ______________

_____ Mastercard #__________________________ Exp. Date:_____________ CID: ______________

Name on credit card:___________________________________________________________

I authorize the WDA to charge my credit card in the amount of $_______________________________

Signature__________________________________________________________________________

Mail registration form and payment to: Wilson’s Disease Association
                      Attn: Kimberly Symonds
                      1802 Brookside Drive, Wooster, Ohio 44691
                      Fax No.: 330-264-0974

DUE BY APRIL 3, 2008
For Your Information

**WDA Wish List:**
- Donated Office Supplies – copy paper
- Donated Printing Services
- Donated Air Miles
- Family Run Fundraisers
- Newsletter Support

**Newsletter Deadline:**
The Copper Connection welcomes, and would appreciate, any articles that members would like to submit for publication consideration. If you would like to submit an article, the deadline for the June newsletter is May 16, 2008. Please e-mail your article to the WDA office at: wda@sssnet.com

The Wilson’s Disease Association is a charitable organization which relies on donations to do its work. Please help us! Tax-deductible donations may be sent to:

Wilson’s Disease Association
1802 Brookside Drive
Wooster, OH 44691

**Join our Online Community**
The Wilson’s Disease Association partnered with Inspire to bring our members a place for open dialogue among all Wilson disease members – no matter what the relationship is to Wilson disease. Join us by participating in a discussion, start your own discussion, create a blog, or vote in a survey. There are many ways to participate. Go to the WDA website, www.wilsonsdisease.org and click the button on the front page. Hope to see you there!

**Shopping!**
Shop iGive.com and find everything you need from paper goods and decorations to clothing and accessories to music and entertainment; the perfect gift for everyone on your list. Plus WDA benefits from your shopping! There are hundreds of stores on the iGive site including: Disney Store, Eddie Bauer, Spiegel, JC Penney, Walmart, Barnes and Nobel, Gap, Home Depot and Starbucks! Each store gives a percentage back to WDA if you register and shop through the iGive.com site.

**Stock Donations**
Please consider the Wilson’s Disease Association for a 2008 stock donation. For more information, please contact the national office at 888-264-1450.

**Foundation Assistance Needed**
WDA members help us in so many ways. Here is one more thing you can do to help. Put us in touch with Foundations who might be willing to help fund WDA’s mission. Some of you may have access to a Foundation through your company, family members, or friends. While it is true that many Foundations have very specific missions, they are often amenable to proposals that come to them from a Director or Trustee.

If you know of a Foundation that may be willing to assist us, please contact the WDA office (888-264-1450 or wda@sssnet.com). We can work with you on the best approach and proposal that would be most suitable. Who knows, your efforts in this regard could pay enormous dividends!

**Membership Mailing**
The 2008 Membership Renewal applications have been mailed. Please make sure you return your application, even if you can’t afford the minimum donation. We need to know you still want to receive our information. Thank you!

**Thank You To Our Wonderful Volunteers**
Dvora Konstant • Dr. Michael Schilsky
Lisa Sniderman-King • Carol Terry
William W. Graper

**National Disease Research Interchange (NDRI)**
Please take the time to register with NDRI. Your blood and tissue donation might be the one that helps find a cure, or a better treatment. To donate blood or tissue, please contact a Rare Disease Coordinator at raredisease@ndriresource.org, or by phone at 800-222-6374.
Dear Madam:
How are you madam, and your family member’s also. I hope everybody fine. Here I am and my son also fine. Really I would like to say thanks because you had helped my son get medication. We never forget your help. The medication reached to my son in India safely. Thanks a lot. Everyday I pray for your family to get good.

Yours sincerely,
Abdul Muthali

The Conference concluded with a unanimous decision to form an organization for Wilson’s disease and electing me as the President/Chairman to take this forward and translate the resolve into reality. A team of medical professionals too were chosen to be an active part of the organization which was to include patients and their families.

My alliance with the very special people with Wilson’s dates back to almost a decade when I first discovered that my daughter Ambuja had Wilson’s. I have been trying to cope up with all the challenges posed and to be associated with as many Wilson’s families as possible but this is the opportunity which God has entrusted upon me to serve, share and help more and more people who are living with Wilson’s.

The Wilson’s disease International has been extremely helpful to all stakeholders and me in particular. I have been in regular touch with the Ms Mary Graper, a most worthy President of WDA who has been ever so helpful in arranging to send the frightfully expensive Syprine and making it available for Ambuja. She has been an inspiration for me and I have tried to live up to the high standards she has set to reaching out to people and make a difference in their lives.

I am extremely grateful to WDA International and look up to replicate the organizational structure of a most successful parent body. May I request you to send me a copy of the constitution/memorandum rules and bye-laws for me to customize it to Indian needs and register our organization in India?

We, in India, look forward to your support and stronger bonding between WDA International and W.I.S.H. (Wilsonsdisease India Support and Help.) I am joining you for the 25th Anniversary Celebration of the Wilson’s disease Association in Chicago and have already made my bookings. I would like to participate in the walk and would also like to raise funds and am open to bringing some Indian handicrafts (in silver for the Silver Jubilee?) if you think they can be leveraged for any which way WDA thinks it can be used. I am so excited to meet all the wonderful people with whom I have been in communication on the List.

Warm Regards

Dr. Ashish Sabharwal
President Designate
W.I.S.H.

---

Dear Ms. Symonds

Leadership, they say, is taking initiative. Your email to the organizers of the first ever National Conference on Wilson’s disease in India has possibly changed everything for thousands of WD patients in India and perhaps many more in the region.

Thanks to your email, I received an invitation to attend the National Conference on Wilson's Disease from Dr. Eapen, the organizer and Professor of Gastroenterology at Christian Medical College, Vellore, a small township in South India, 2000 kms from my abode in New Delhi.

I had been toying with the idea to launch the Indian arm of the Wilson’s disease Association on the lines of WDA International for a very long time and this invitation compelled me to book a passage to Vellore. I put in a request to be allowed to speak at this scientific meeting which was accepted as I submitted my intention of sharing my resolve to begin an association/foundation/society to help patients and families of Wilson’s disease in India. Though I am a medical professional, I cannot profess any claims towards treating Wilson’s disease and strongly believe that Wilson’s patients can derive benefit if professionals who are dedicated to managing the disease are actively involved in the organization. The organizers were gracious enough not only to allow me, a parent, to speak at their predominantly medical deliberations but also allowed another parent, Mr Limaye, whom I had urged to come over to Vellore. You had introduced Mr Limaye to me over the web and we had exchanged a few mails before we met for the first time in Vellore.

The day long conference was a great success with quality interaction amongst the participants. We sat through the entire deliberations and bonded extremely well as a group intent on bringing out the best for our common cause.
Walk-Out-Wilson’s in Chicago!!!!

The WDA is pleased to support Nancy Marsala in organizing the First Walk-Out-Wilson’s in Chicago, IL. April 27, 2008 the Walk-a-thon will take place in the Lincoln Park Zoo area. The WDA will provide transportation from the Hyatt (conference hotel) to the Zoo area.

To register for the walk please go to www.firstgiving.com/wda or use the link provided on our website. Fundraising is an important part of Walk-Out-Wilson’s in Chicago. It will help support the important programs already established by the WDA. You may choose to collect pledges from family, friends, or co-workers or create your own personal fundraising Web page after you’ve registered. It’s an easy and effective way to ask family and friends for their support.

As a special incentive for all Chicago high school students – Nancy has arranged for you to receive service learning hours. To learn more about this program, please contact Nancy directly.

It isn’t too late! Nancy is still looking for volunteers to help make this event a success. Please contact her at nkmeow@sbcglobal.net or 773-777-1020. To set-up a team, or make a donation, please go to www.firstgiving.com/wda.

New Director: FDA Office of Orphan Products Development

Timothy Coté, M.D., M.P.H. was named FDA’s new Director of the Office of Orphan Products Development (OOPD). He replaces Dr. Marlene Haffner who retired in 2006.

Dr. Coté previously served as a director of CDC’s health programs in Rwanda. He managed a staff of U.S. government and contractual employees, as well as locally employed staff with a budget of $119M directing programs in HIV/AIDS, malaria, and avian influenza. WDA’s Executive Director, Kimberly Symonds, met with Dr. Coté last fall at the National Organization for Rare Disorder’s fall conference.
Thanks for Your Support!

**Challenge Grant Donations**
John and Cynthia Adams • Calvin Akin  
Raymond Alessandrini • Melissa Asgaonkar  
Frederick & Juanita Aukeman • Brenda Sue Baker  
Anvar & Zoukra Bash • John & Loni Becker  
Daniel & Haleigh Banayan  
Alta Black (In honor of Cody Gorel)  
Mary Lou Blasius • Eugene Brown  
Mei Lan Chen • David & Diana Chipkin  
Bryan & Paula Clay • David & Naida Cohn  
Douglas Coleman • Maryann Coppolino  
Gordon Crim • Edna Cross • Barbara Eckland  
Jeanne Fink • Barbara Frayer • Ellen Freedman  
Douglas Goostree • Martine Greene  
Stephen Heiniger • Berna & Joseph Heyman  
David & Marilynn Hickam • Stanley & Ruth Hill  
Dr. Fred Hirt • Philip & Cecelia Horink  
Jeremy Hubball • Reed Hughes  
Christine Hunter • Bonnie Inwood  
Lori Janow (In memory of Carole Janow)  
Marianne Joyner • Joshua & Stefanie Kaplan  
Henry & Marcia Kaplan (In honor of Michael Schilsky, MD)  
Emmet Keeffe • Chris King • Shari Kurzrok Schnall  
Sharon Lagas • Mary Ann Lehnerer  
Suzanne Lemoine • Barry Levinson  
Robert & Beverly Linkins • Rose & Robert Malaczewski  
Ronald Marchese • Nancy Kesler Marsala  
Edward Martin (In honor of Mr. and Mrs. Fetzer Mills)  
Duane & Beverly Mattheis • Pauline McNulty  
Pat Meadows • Fetzer & Penny Mills  
James & Helen Moseley (In memory of Erika Lynn Moseley)  
Vui & Th Nguyen • Kelly Oboyle  
Elizabeth Osborne (In memory of Kelsey Fink)  
Mary L. Oster Granite • Betty Perinka • Rose Plautz  
William Popp • Jess Porres  
Doug & Mary Powell (In memory of James Badamo)  
Len & Diann Ptak • Michael Rabin  
Edward & Miriam Rabin • John Randall  
Dolores Roessl • Steven Rubenstein  
Delia & David Ruiz • Joyce Russell • Charles Rust  
Theresa Rutherford • Linda Schwebke  
Mr. and Mrs. Vincent Scire • Aarti & Arjun Sehgal  
Mildred Shannon • Peter & Gracia Sheptak  
Ruthann Siebert • William Siegert  
John & Gail Sisti • Bonnie Smelser  
Natalie Smith • Eileen Smith • Timothy Smith  
Michael Sobkowski • Susan Tarquinio  
Carol Terry • Frank & Bernadine Wall  
Walter & Tori Wall (In memory of Rick Wall)  
Lynn Wehmueller • Laura Weinberger  
Robert Whittier • John Witort  
Paricheher Yomtoob • Paul Zandhuis  
The Katz Foundation • Infinite Possibilities Foundation  
The Evictor • Geico Philanthropic Foundation  
David Novick • Kristin Ulibarri

**General Contributions:**  
Katherine Basch • Duane Mattheis • Sharad Wagle  
Pei Chao • Chien-Hui Chung • Michael Cohen  
Maryann Coppolino  
Brasilia Dimuro (In honor of Justin Garlepp)  
David Gamba • James Hines  
Joyce Hoggard (In honor of Debra Hoggard)  
Duane & Beverly Mattheis • Mark Mirkin  
Washtenaw United Way • Lewis Katz  
Aton Pharma • Trinity Partners, LLC

**Honorarium:**  
Nathan Chinn (In honor of Justin Garlepp)  
James & Vicki Harker (In honor of Carrol “Butch” Cross  
Dr. Selma Knobel (In honor of Nancy Schwebke)  
Ken Knobel (In honor of Justin Garlepp)  
Gail Nunn (In honor of Geoffery Nunn)  
Concept Components (In honor of Kayla Paturzo)  
The Foundation (In honor of Justin Garlepp)

**Membership:**  
Bernice Gilchrist (In honor of Travis & Molly)  
Joyce Hoggard (In honor of Debra Hoggard)

**M.A.P**  
Anonymus • American Commercial Mortgage  
Duc Cao • Vladimir Grigor

**Marketplace**  
Ruth Brandt • Debra Collier • Kay Robinson  
Bernadette Dqoraczyk

**Silver Membership:**  
Shai Dobrusin • Sharon Lagas

**Copper Membership:**  
Justin Garlepp

**Memorial:**  
Mike Gardner (In memory of Mark Waring)  
Charles Aubuchon (In memory of David Yomtoob)  
Dorie Badamo (In memory of James Badamo)
WILSON'S DISEASE ASSOCIATION

MEMBERSHIP APPLICATION

☐ New Member  ☐ Renewing Member  ☐ Mr.  ☐ Mrs.  ☐ Miss  ☐ Ms.  ☐ Dr.

Name__________________________

Address__________________________

City__________________________ State________ Zip________ Country__________________________

Home Phone____________ Work Phone________ Email__________________________

MEMBERSHIP DUES:

☐ Basic Membership – $35
  • Receive quarterly WDA Newsletter via e-mail – (Please provide e-mail address above)

☐ Basic Plus Membership – $60
  • Receive quarterly WDA Newsletter via U.S. mail or e-mail – (Please provide e-mail address above)

☐ Silver Membership – $120
  • Receive quarterly WDA Newsletter via U.S. mail or e-mail – (Please provide e-mail address above)
  • 10% discount on Annual Meeting registration for up to two registrants

☐ Gold Membership – $250
  • Receive quarterly WDA Newsletter via U.S. mail or e-mail - (Please provide e-mail address above)
  • 15% discount on Annual Meeting registration for up to four registrants
  • Special recognition as a Gold Member in WDA Newsletter

☐ Copper Membership – $1000
  • Receive quarterly WDA Newsletter via U.S. mail or e-mail – (Please provide e-mail address above)
  • 20% discount on Annual Meeting registration for up to four registrants
  • Special recognition as a Copper Member in WDA Newsletter
  • 10% discount for Annual Meeting program ad
  • No annual dues

I WISH TO MAKE A DONATION TO WILSON'S DISEASE ASSOCIATION:

Name__________________________

Address__________________________

City__________________________ State________ Zip________ Country__________________________

I am making a tax-deductible donation of $__________

☐ In honor of__________________________  ☐ In Memory of__________________________

Send acknowledgement to: Name__________________________ Address__________________________

City__________________________ State________ Zip________ Country__________________________

PAYMENT INFORMATION

Membership Fee  $________  ☐ Visa ☐ Mastercard  ☐ Check or Money Order attached

Donation  $________  Card #__________________________

Total  $________  Expiration Date:_____________  CID#__________________________ (3 digits on back of card)

Signature__________________________
Pill Box Timer
Remember to take your medication on time with WDA's new Pill Box Timer. These great timers have, in addition to the normal display, an alarm and stopwatch to ensure that you don't miss a single dose. Holds 15 aspirin-size tablets; has attached lanyard for use around neck, but small enough to fit in a pocket or purse. 3.5" L X 2" W X 5/8" Deep. Two colors available: purple and green. Donation requested is $15.00 plus shipping and handling ($2.50).

Number of Pill Boxes __________

WDA Awareness Bracelet
You asked for them...we got them! WDA Awareness bracelets! They are copper in color and are designed to bring awareness to Wilson’s disease. The bracelets are packaged in groups of 5. The suggested donation per bracelet is $2.50, that comes to $12.50 per package. Copper color, inscription to read: STOP COPPER! SUPPORT WDA wilsonsdisease.org

Package of Bracelets___________

2005 Annual Resource Guide
Exceptional Parent magazine publishes a comprehensive Annual Resource Guide for the special needs community. This is an indispensable reference tool for physicians, allied health care professionals, teachers, families and caregivers. EP has extended a special offer to the WDA! The resource guide retails for $19.95, WDA will distribute this guide for a donation of $10.00 plus shipping and handling ($2.50). WDA has a limited number available.

Number of Guides __________

W.O.W. Walk-A-Thon T-Shirt
We have a number of the 2007 Walk-A-Thon shirts left for those of you who would love to have one. The shirts are bright yellow with black lettering. A limited number of Mediums and Larges are all that remain. Suggested donation is $20.00 per T-Shirt, shipping and handling included.

Number of T-Shirts
M _____ L _____
Total: $__________

WDA Bumper Sticker
Join in bringing awareness about Wilson’s disease! These stickers are 3" x 10" and have Copper color graphics with black text. Thank you to members Melissa and Patrick for designing them. And, a special thank you to Patrick for donating them to the WDA in honor of his son Jeff! Don’t like sticking these things to your bumper? How about your front door, boat, work cubicle or “just about anywhere things will stick!” Suggested donation is $5.00 per sticker, shipping and handling included.

Number of Stickers __________

Penny Card
WDA's Penny Card Campaign continues to be a great success. To join in the fun, order your cards today. It is a suggested donation of $15.00 per card...what a great awareness tool for your community!

Number of Penny Cards ________

WDA's Penny Card Campaign continues to be a great success. To join in the fun, order your cards today. It is a suggested donation of $15.00 per card...what a great awareness tool for your community! To Order:
Contact the WDA office by phone or e-mail to order.
WILSON'S DISEASE ASSOCIATION
Telephone: 888-264-1450
E-Mail: wda@sssnet.com

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Number of Stickers __________

To Order:
Contact the WDA office by phone or e-mail to order.
WILSON'S DISEASE ASSOCIATION
Telephone: 888-264-1450
E-Mail: wda@sssnet.com
Wilson’s Disease Association
Volunteer Profile

Name: ________________________ Connection to Wilson’s Disease: ________________________
(Please include professional designations: e.g. M.D., Ph.D.)
Spouse or Significant Other’s Name: ________________________
Home Address: ________________________ Home Telephone Number: ________________________
Fax: ________________________
E-Mail Address: ________________________ Business Title: ________________________
Business Address: ________________________ Business Telephone Number: ________________________
Fax: ________________________
Occupation and Job Responsibilities: ________________________
Company has a matching gift program (circle one): Yes  No
Board Memberships and Professional Organizations: ________________________
Social Affiliations/Clubs and Organizations: ________________________
Personal Interests/Hobbies: ________________________
Areas of Experience or Expertise:
  ______ Auditing  ______ Writing
  ______ Legal - Nonprofit Experience  ______ Media
  ______ Government Affairs  ______ Graphic Arts
  ______ Marketing  ______ Board of Directors
  ______ Public Speaking  ______ Computer Technology
  ______ Fundraising:  ______ Web Site/Internet
    ______ Special Events  ______ Newsletter
    ______ Foundations  ______ Local Support Group Organizing/Leading
    ______ Corporations  ______ Office Work
  ______ Other (specify) ________________________