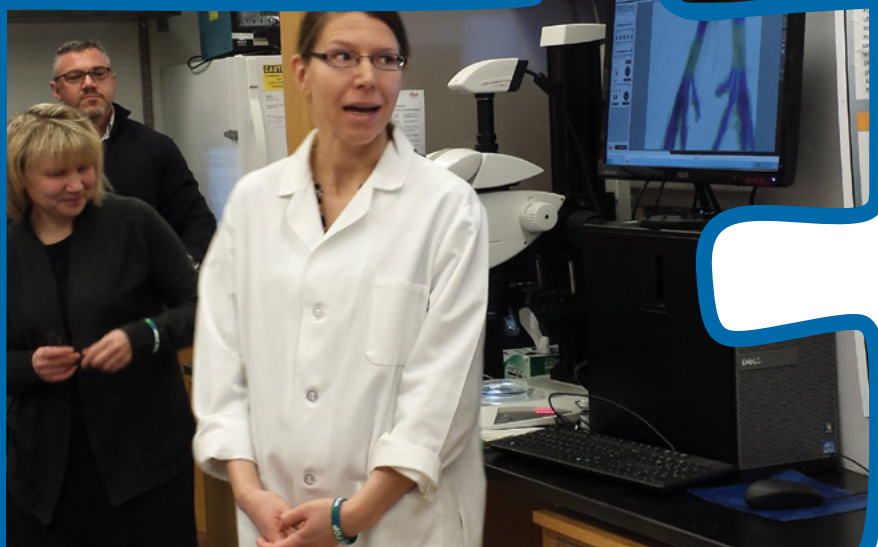


Together,
we can make
a difference!



ifopa

2015 Annual Report

Our vision: A Cure for FOP

Our mission: Fund research to find a cure for fibrodysplasia ossificans progressiva while supporting individuals and their families through education, public awareness and advocacy.

Table of Contents

- 3 A Message from IFOPA Board Chair
- 4 Our Founder, Jeannie Peeper
- 5 The Quest for Treatments and a Cure
The University of Pennsylvania
Competitive Research Grants Program
- 7 FOP Connection Patient Registry
- 10 Family Services
- 12 Advocacy and Awareness
- 13 International Highlights
- 15 Jeannie L. Peeper Award Winners
- 18 2015 Fundraising
- 21 Our Financials
- 23 2015 IFOPA Donor Honor Roll

What is fibrodysplasia ossificans progressiva (FOP)?

One of the rarest, most disabling genetic conditions known to medicine, FOP causes bone to form in muscles, tendons, ligaments and other connective tissues. Bridges of extra bone develop across the joints, progressively restricting movement and creating a second skeleton that encases the body in a prison of bone. There are no other known examples in medicine of one normal organ system turning into another. There is no treatment for FOP.

Symptoms of FOP almost always begin in the first two decades of life, and the majority of patients learn they are affected before the age of ten. Inflamed and painful swellings, usually in the shoulder and back areas, are the first signs of the condition. These swellings eventually clear up, but leave behind an area of mature bone.

FOP patients experience different rates of bone formation, and the exact rate of progression is unpredictable. The bone growth can occur spontaneously, or can be triggered as a result of trauma, an injury, bump or fall. The extra bones grow in the same order that they are first formed in the fetus – from the head, shoulders, arms and progressing to the hips and knees. As the condition worsens, patients eventually have joints locked in fixed positions.

This immobilizing and catastrophic genetic condition strikes approximately one in two million people. It is estimated that 3,500 people worldwide have FOP, although there are only 800 confirmed cases; 285 of which are in the United States.

How would understanding the cause of bone formation in FOP help others?

The information obtained from studying this disease will have far-reaching implications for the treatment of common disorders such as fractures, osteoporosis, hip replacement surgery and other forms of heterotopic ossification that occur in trauma and burn victims.

A Message

from 2015 IFOPA Board Chair
Marilyn Hair



The IFOPA has its eye on the prize of finding treatments for FOP. We are proud of the events and achievements of 2015 that are bringing us ever closer to our goal. New research discoveries along with current and anticipated clinical trials raise our hopes that effective treatments will be available before long that will make a real difference for people with FOP.

IFOPA activities and projects in 2015 included:

- Launching the FOP Connection Registry
- Strategic planning by the Board of Directors that led to the search for an Executive Director
- Adults and children with FOP enrolling in Clementia Pharmaceutical's Natural History Study and Phase II Clinical Trial of palovarotene
- Awarding the first three FOP Competitive Research Grants
- 3,915 individual gifts, 15 family fundraisers and total donations of \$901,500
- Nearly \$3.9 million in net assets

The FOP Connection Registry patient portal was launched in July 2015. One hundred thirty-nine (139) persons with FOP from 28 countries joined the Registry by December 31. The FOP Registry is an invaluable repository of population-wide information that will help FOP researchers, clinicians and pharmaceutical companies better understand FOP. Everyone with FOP can contribute to finding treatments by joining the FOP Registry to tell your story.

As of February 2016, Clementia Pharmaceutical's Phase II Clinical Trial of palovarotene was fully

enrolled. Many clinical trials to test new drugs fail simply because they don't enroll enough participants. Considering that FOP is extremely rare and that patients are spread across the world, this is a true milestone and an accomplishment we can all be proud of.

FOP research continues on an active and promising pathway. There are growing indications, at least in mouse models of FOP, that the disease can be treated in different ways and at different points in the disease pathway.

The IFOPA International President's Council (IPC) comprises 22 leaders of 18 national FOP organizations from across the world. Current efforts of the IPC focus on mobilizing FOP patients worldwide to help develop treatments for FOP. In April, the IPC met in person during the FOP Italia meeting in Rome. National FOP meetings were held in 2015 in Germany, Italy, Russia, Sweden and Spain.

Many thanks to the volunteer Board of Directors who guides the IFOPA. As a nonprofit organization, the IFOPA relies on contributions; we are ever grateful to the many families and friends who conducted fundraising events in 2015.

We are forever guided by the vision of Jeannie Peeper who started a pen-pal group to end the isolation of having FOP. She capably led the organization's growth to achievements even she never imagined.

Thank you for the part you play in the IFOPA. Whether you're a family member, fundraiser, donor, doctor, researcher, staff member, supporter, or have FOP, we need and appreciate you. Your dedication lets us pursue our mission to fund research to find a cure for FOP while supporting individuals and their families through education, public awareness and advocacy.

With hope,

A handwritten signature in cursive script that reads "Marilyn Hair".

Marilyn Hair
Chair, IFOPA Board of Directors 2014-15

Our Founder

Jeannie Peeper



The IFOPA was founded by Jeannie Peeper, a young woman with FOP, who had never met anyone else with the disease. While at the National Institutes of Health (NIH) in 1987, before the days of the Health

Insurance Portability and Accountability Act (HIPAA), Jeannie was given the names of 17 other people who had FOP. Jeannie wrote to each of them, and 10 wrote back, forming a pen pal group which became the International FOP Association in 1988.

A native of Ypsilanti, Michigan, Jeannie's life began to change at an early age with the onset of FOP. Instead of allowing the condition to overwhelm her, she began a journey that allowed her to earn a college degree, and create an organization that has ended the isolation formerly associated with the disease by connecting and supporting persons with FOP and their families.

In 1989, Jeannie collaborated with Drs. Michael Zasloff and Fred Kaplan at the University of

Pennsylvania to initiate the FOP Collaborative Research Project. Since our founding in 1988, and thanks to the generosity of fundraisers and donors, the IFOPA has funded over \$9.1 million dollars in FOP research to find treatments and a cure!

The list of IFOPA achievements under Jeannie's leadership are multiple, and include hosting international symposiums and gatherings for families, clinicians and researchers; establishing a website; participating in many media projects; hosting strategic planning meetings; visiting with donors; and providing valuable counsel to FOP families. Jeannie served as President of the IFOPA from 1988 until she retired in 2013 with the title Founder and President Emeritus.

Isolation was typical before the IFOPA was formed, and Jeannie's goal was to bring people with FOP together. Today, the IFOPA is the umbrella organization for people with FOP worldwide and the place to come for education and support. The IFOPA gives support to 1,000 individuals representing over 50 countries. Over 450 of these individuals have FOP; over 280 of them are from outside the United States.

Jeannie's vision and commitment were at work back in 1988 and are still alive today as she provides support and encouragement to those living with FOP and their families. Her life's work are an inspiration to clinicians and researchers working to cure FOP worldwide.

2015 Board of Directors



Back row left to right:

Moirä Liljesthröm, Chris Bedford-Gay, Rory Otto, Gary McGuire, Amy Gordon, Gail Weakland

Front row left to right:

Paul Brinkman, Karen Munro, Marilyn Hair, Nancy Sando

Not pictured:

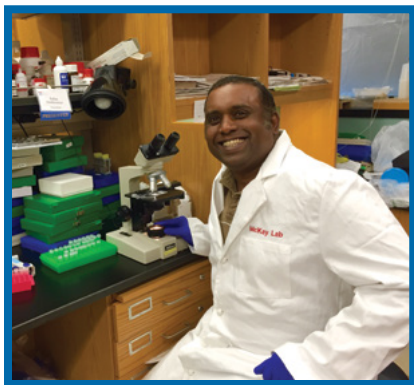
Brian Harwell

The Quest

for Treatments and a Cure

Funding basic research to grow knowledge of FOP and translational research to develop treatments and, one day, a cure for the disease are at the core of the IFOPA's mission.

The University of Pennsylvania's Center for Research in FOP & Related Disorders



Dr. Salin Chakkalakal at The Center for Research in FOP & Related Disorders

The Center for Research in FOP & Related Disorders was established in 1997 by Drs. Fred Kaplan and Eileen Shore, with the generous support of The Cali Family, at one of the most prestigious medical and research institutions in the

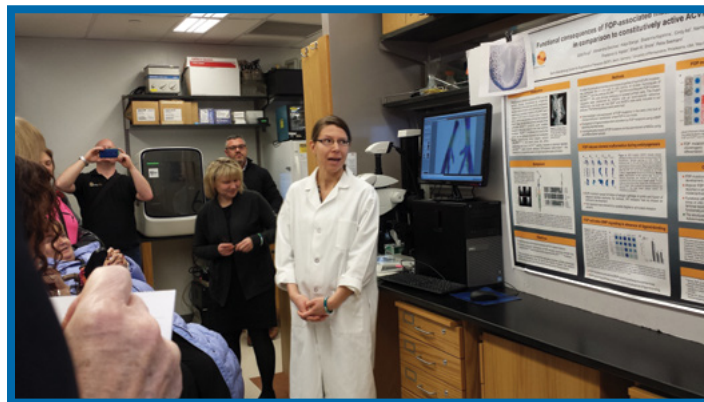
country – The University of Pennsylvania School of Medicine in Philadelphia.



IFOPA Board visits the FOP Lab at the University of Pennsylvania

In April 2006, the research team pinpointed a single gene mutation – one letter out of six billion in the human genome – that causes the runaway bone growth of FOP. This was a watershed event in the 300-plus year history of FOP. The discovery immediately opened

the floodgates of research as the primary cause and ultimate target of the disease were instantly revealed.



Julia Haupt explains FOP research being done at the University of Pennsylvania to IFOPA Board members.

In 2015, the IFOPA funded nearly \$500,000 for nine research projects at The University of Pennsylvania. Fundraisers held throughout the year by families of FOP individuals are the primary source of this funding

Competitive Research Grants (CRG) Program



Sona Brinkman

In 2015, the IFOPA, through a fundraising campaign led by the family of Sona Brinkman, launched the Competitive Research Grants Program; grants focused on discovery and advancement of new therapeutic approaches to FOP

with a disposition toward approaches likely to have near-term clinical or translational relevance.

Applications were reviewed by a voluntary board with relevant and appropriate expertise related to FOP to evaluate proposals, and that was free of conflict of interest.

CRG Scientific Advisory Board

- Vicki Rosen, PhD, Chair
Department of Developmental Biology
Harvard School of Dental Medicine
- Karen Lyons, PhD
Department of Molecular, Cell and Developmental Biology
UCLA/ Orthopaedic Hospital Department of Orthopaedic Surgery
- Ernestina Schipani, MD, PhD
Department of Orthopaedic Surgery
University of Michigan
- Michael Whyte, MD
Division of Bone & Mineral Diseases
Washington University School of Medicine
- Michael Zasloff, MD, PhD
Department of Surgery
Georgetown University Medical Center

In 2015, nearly \$125,000 in grants were awarded to three institutions.

1. Assessment of small agents in FOP primary fibroblast cultures to explore new therapeutic targets

Principal Investigators: Gerard Pals, PhD and Marelise Eekhoff, MD, PhD

Institution: VU University Medical Center, The Netherlands

Award Amount: \$55,000



Gerard Pals



Marelise Eekhoff

2. The role of exercise in the progression of FOP

Principal Investigators: Paul B. Yu, MD, PhD and Yue Shen, MBBS

Institution: Brigham and Women's Hospital and Harvard Medical School, United States

Award Amount: \$28,000



Paul B. Yu



Yue Shen

3. Validation of novel diagnostic and targeted prophylaxis for FOP related heterotopic ossification

Principal Investigators: Yuji Mishina, PhD and Benjamin Levi, MD

Institution: University of Michigan, United States

Award Amount: \$41,800



Yuji Mishina



Benjamin Levi

FOP Connection Patient Registry

The FOP Connection Registry is an observational research study spearheaded by the IFOPA that collects and reports specific medical information on individuals living with FOP. The Registry will ultimately have two portals – a patient portal and a medical portal. As the name suggests, the patient portal includes data submitted by a person living with FOP, or a family member or other caregiver with their input. When available, the medical portal will include data submitted by the clinician caring for an individual with FOP. It completes the story of an individual's disease progression to have both types of data.

The FOP Registry meets the needs of all stakeholders involved in the drug development process: patients, doctors, and academic and pharmaceutical researchers. A unified, global and coordinated approach through a single registry provides the most comprehensive data on FOP, ultimately enabling better and faster development of therapies. Because of the generosity of the FOP community, the IFOPA was able to take the lead and develop a single worldwide registry for all FOP individuals. In 2015, the IFOPA invested more

than \$230,000 to complete the design and build of the patient registry.

The FOP Registry:

- Organizes the entire FOP community for potential clinical trials and studies
- Empowers FOP patients and the community as a whole
- Collects data to better understand FOP and its impact on the body and physical function, as well as the emotional impact of the disease
- Advances the understanding of FOP treatment outcomes when they become available

The FOP Registry was launched on July 29, 2015 in English; other translations will be available at a later date. After the launch in July, the FOP Connection Registry was represented at the November 2015 FOP Stichting Nederland patient meeting in Amsterdam, The Netherlands.

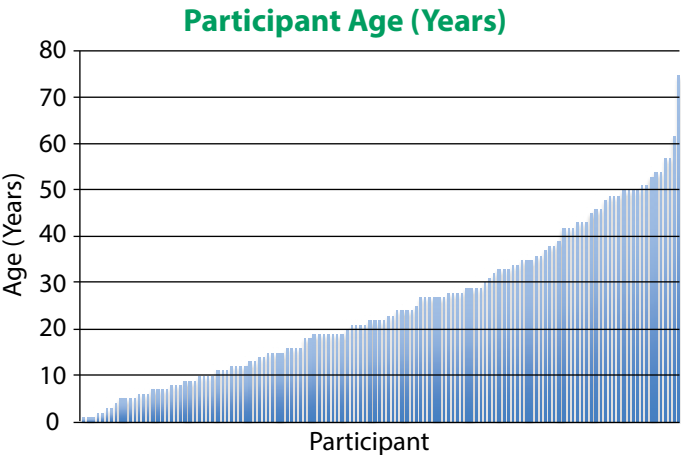
The Registry is governed, in part, by a Medical Advisory Board (MAB). The MAB is a group of clinicians with experience in FOP who, among other tasks, will design the Registry's medical portal. They also convened their first meeting at The Netherlands meeting.



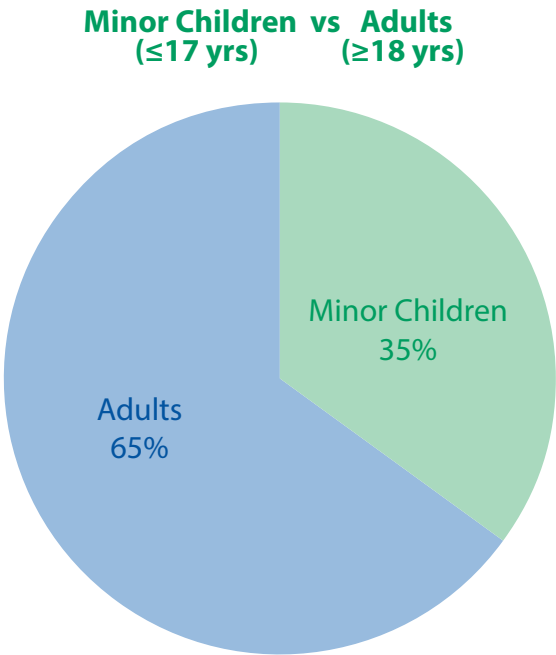
"Knowing I have personally contributed information that will help researchers and drug developers better understand FOP and eventually find multiple treatments . . . that's what it's all about, I can make a difference." Nancy Sando, First FOP Registry Enrollee

Registry Data Highlights as of December 31, 2015

- 139 registered participants from 26 countries on six continents; reflecting a truly global FOP community
- Average age of participants is 24.9 years, with a range of 1 year to 75 years (see chart)

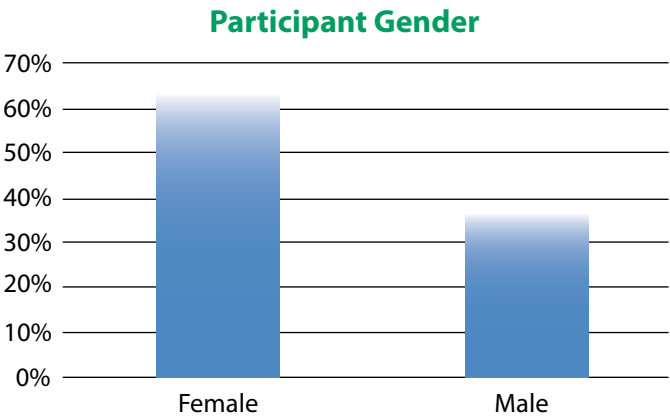


- Among the 139 participants, 35% are minor children; 65% are adult participants (see chart)



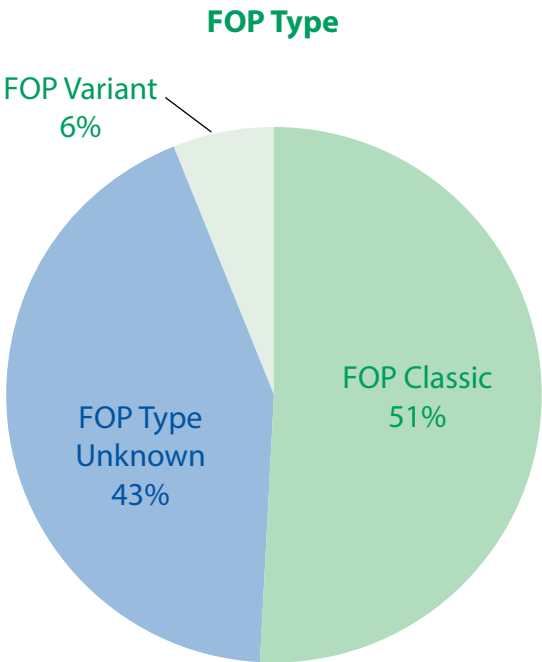
- Among the minor children, 46% are females and 54% are males; almost an even split between the genders

- 63% of participants are female; 37% are male (see chart). There is no known relationship between FOP disease and gender, so the data suggests that males overall are under-represented in the Registry.



- Of the female participants, 25% are minor children (age 17 years or younger)
- Of the male participants, 51% are minor children; a disproportionally higher percentage of minor male children compared to minor female children, suggesting that adult males are under-represented in the Registry

- 86 participants provided information about their FOP type with 51% (44/86) having reported FOP Classic, 43% (37/86) having reported FOP Type Unknown, and 6% (5/86) having reported FOP Variant



FOP Connection Registry Medical Advisory Board

Academic Medical Advisors

Genevieve Baujat, MD
Hôpital Necker-Enfants Malades
France

Matt Brown
University of Queensland
Australia

Carmen de Cunto, MD
Hospital Italiano de Buenos Aires
Argentina

Patricia Delai, MD
São Paulo School of Medicine
Brazil

Marelise Eekhoff, MD, PhD
VU Medical Center, Amsterdam
The Netherlands

Nobuhiko Haga, MD
University of Tokyo
Japan

Edward Hsiao, MD, PhD
UCSF School of Medicine
United States

Frederick Kaplan, MD
University of Pennsylvania
United States

Richard Keen, MB, BS, BSc, PhD, MRCP
Royal National Orthopaedic Hospital
United Kingdom

Rolf Morhart, MD
Klinikum Garmisch-Partenkirchen
Germany

Robert Pignolo, MD, PhD
University of Pennsylvania
United States

Maja di Rocco, MD
Gaslini Institute
Italy

Christiaan Scott, MBChB, FCPaed
Red Cross Children's Hospital
South Africa

Keqin Zhang, MD, PhD
Shanghai Tongji University
China

Pharmaceutical Industry Advisors

Eric Bachman, MD, PhD
Alexion Pharmaceuticals, Inc.

Donna Grogan, MD
Clementia Pharmaceuticals, Inc.

Xiaobing Qian, MD, PhD
Regeneron Pharmaceuticals, Inc.

Family Services

Through the generosity of donors, the IFOPA is able to provide support for individuals living with FOP and their families through a number of programs and services at no cost.


Available resources and services include:

Guidebooks

- *What is FOP? A Guidebook for Families* provides a comprehensive look at FOP answering many questions commonly asked by families and medical professionals
- *What is FOP? Questions and Answers for the Children* explains FOP in easy-to-follow language for children ages 8 to 12 and is recommended for younger children with adult supervision

Medical Binder

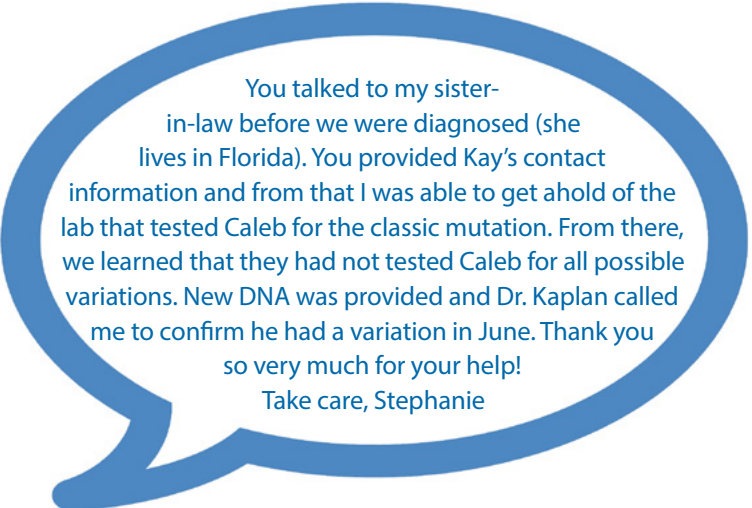
- Stores important personalized information all in one place
- Valuable for communication with doctors
- Easy reference to emergency contact numbers, past flare-up and medical history, and current and past prescriptions



Just a note to say thank you very much for the medical information on your website that enabled me to safely anaesthetize a middle-aged patient with advanced FOP at short notice. I found the information very helpful in my anesthetic management and advising critical care on his post-operative medical management.
Yours sincerely, Dr. Ruth S. Newton,
Consultant Anesthetist, U.K.

Emergency Tools

- Individuals living with FOP can request items to help others understand FOP in a medical emergency situation
1. Emergency card – Wallet-sized and laminated, contains key treatment information
 2. USB flash drive – Contains pre-loaded personal medical information forms
 3. Portable pill holder – Recommended waterproof container for an emergency dose of Prednisone (pill holder is not childproof)



You talked to my sister-in-law before we were diagnosed (she lives in Florida). You provided Kay's contact information and from that I was able to get ahold of the lab that tested Caleb for the classic mutation. From there, we learned that they had not tested Caleb for all possible variations. New DNA was provided and Dr. Kaplan called me to confirm he had a variation in June. Thank you so very much for your help!
Take care, Stephanie

Online Support Groups

Support groups allow individuals living with FOP and/or their parents to connect with one another in a safe environment. These groups are used to ask questions and get advice, share concerns, provide practical suggestions for everyday life and situations that may arise, and simply be connected to others with FOP.

These groups are generally administered by FOP individuals or, in the case of Parents with FOPers, by a parent of a child with FOP. To join one of these groups, send an email to together@ifopa.org and we will forward the request to the page or email group administrator.

Facebook Groups

1. IFOPA'ers – Group for individuals living with FOP and/or their parents
2. Parents of FOPers – Group for parents raising a child with FOP
3. FOP Ladies – Group for women with FOP where no topic is off limits

Email Groups

1. FOP Online – Group for individuals living with FOP and/or their parents.
2. Pray for FOP Healing – Diverse group that comes together in the Judeo-Christian tradition of prayer, encouragement and friendship sharing inspirational quotes and prayer requests and celebrating members' milestones; any denomination is welcome
3. Ladies of FOP – Group of women age 13 and up who discuss issues pertinent to ladies with FOP, including puberty, relationships and other issues

Mentoring Program

- Provides a family-to-family connection
- Volunteer mentors may be a patient, parent, sibling or caregiver
- Mentors listen and share their experiences
- Mentors provide open communication for families
- Mentors are familiar with IFOPA programs and services

Quality of L.I.F.E. Award (Living Independently with Full Equality)

- Monetary award designed to assist individuals with FOP to live more independently
- Award can be used for items that improve a patient's health, welfare and/or independence
- Award is a lifetime maximum grant of \$1,500

- Each application is reviewed by the L.I.F.E. Award Committee

e-Newsletter, Website and Social Media

- FOP Connection is our monthly e-Newsletter with information about living with FOP and includes research news and updates and ways you can connect, advocate and support the IFOPA. It is available to anyone who cares about our vision to #cureFOP.
- FOP Community Updates is our monthly e-Newsletter for individuals living with FOP and includes birthdays, condolences, celebrations and other community event announcements and reminders
- Thousands of people visit our website and social media pages to learn more about FOP, IFOPA events and to support our work to cure FOP each year.



ifopa.org



twitter.com/ifopa



facebook.com/IFOPA

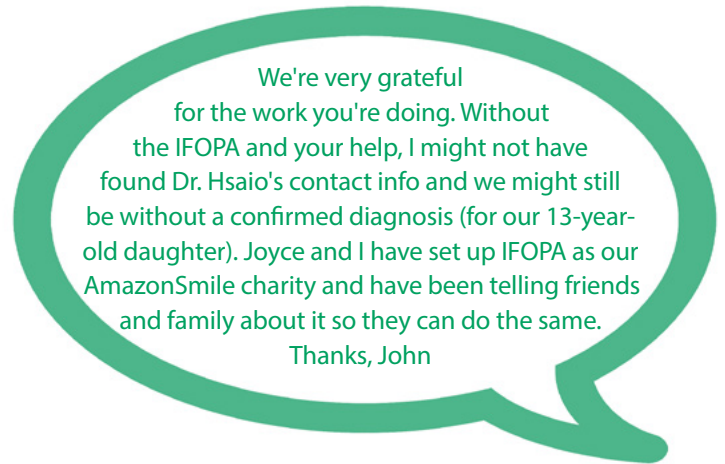
Advocacy and Awareness

Rare Disease Day 2015

- The IFOPA and Dr. Kaplan were pleased to work with writer Christin Melton who wrote “Fibrodysplasia Ossificans Progressiva: Before You Biopsy, Look at the Toes” for Rare Disease Report to raise awareness of FOP among healthcare professionals.



- The “Role of the FOP Patient Community” webcast hosted by Cementia Pharmaceuticals paid tribute to the FOP community in recognition of Rare Disease Day. The IFOPA, FOP individuals, leading researchers and physicians shared how the patient community enhanced their knowledge of the intricacies of FOP and furthered efforts to find treatments and a cure.



International FOP Awareness Day 2015

In celebration of the announcement of the FOP gene discovery nine years ago on April 23, the IFOPA launched the FOP Competitive Research Grant Program to provide funding to scientists studying FOP for the discovery and advancement of new therapeutic approaches. See article on page 6 to learn more.



International Highlights

The IFOPA embraces the worldwide FOP community in many ways – one of which is through the International President's Council (IPC). The IPC is an IFOPA committee made up of 22 leaders of 18 FOP national organizations around the world. In May 2015, FOP Australia was established and joined the IPC.

Current efforts of the IPC focus on mobilizing FOP patients worldwide to help develop treatments for FOP. The IPC meets in person each year at the FOP Italia annual meeting. The support of generous donors allowed the IFOPA to fund travel expenses for nine of the 14 FOP national organization leaders who attended this important meeting.

2015 highlights of the FOP national organizations include:

- Annual German FOP Meeting
- Annual Family Meeting in Spain
- 2nd Annual FOP Russia Meeting
- FOP Sverige Summer Family Meeting



Moscow excursion during FOP Russia Meeting:
(left to right) Anastasia Vorobiova (interpreter), Igor Zahvatov (Heads Baltic countries for FOP Russia), Nadezda Zahvatova (Igor's mom), Dmytriy Medvedev (volunteer), Olesya Radushko, Vladislav Grachev, Rada Borzova (Heads Siberian branch for FOP Russia), Nadezhda Borzova (Rada's mom), Vladimir Borzov (Rada's dad)

FOP Italia 9th Annual Meeting Held in Rome, Italy

Nearly 100 people were in attendance, March 20 to 21, including physicians and researchers from Italy, United Kingdom, The Netherlands, Germany, Japan, and the United States, as well as representatives from two pharmaceutical companies – Clementia and Regeneron. In addition, representatives from the IFOPA and International President's Council participated in the global event that focused on research and patient issues.

Presentations on various research approaches and tools to find treatments for FOP were given and Betsy Bogard, IFOPA Global Research Development Director, updated attendees about the FOP Registry that would be launched in summer 2015.

At the conclusion of the scientific meeting, a FOP clinic for Italian patients was held and the International President's Council convened its annual in-person meeting. The fourteen IPC leaders who attended from Argentina, Australia, Canada, France, Italy, Poland, Russia, The Netherlands, South Africa, Sweden, United Kingdom and the United States gave updates about the work they are doing in their countries. Leaders discussed how the IFOPA and IPC should coordinate working with pharmaceutical companies, the importance of translating the FOP Registry into many languages, and collecting biological samples in one or more bio banks.



IFOPA International President's Council meeting in Rome, Back left to right: Enrico Cristoforetti, Marilyn Hair, Carrie Connell, Marie Hallbert Falhlberg, Irene Snijder, Massimo Alfieri, Julie Collins and Mihail Belyaev. Front left to right: Moira Liljesthröm, Chris Bedford-Gay, Katarzyna Ziaja, Marie-Emmeline Lagoutte and Chris Scott

#FOP23April – FOP Meeting in Eskilstuna, Sweden

Marie Hallbert Fahlberg, leader of the FOP national organization in Sweden, held a special meeting on International FOP Awareness Day, April 23, to inform patients, families, doctors and the media about the palovarotene clinical trial for FOP which came to Europe in 2015.

A short feature on Swedish TV about FOP, the clinical trial, FOP Awareness Day and interviews with Marie and her son Hugo Fahlberg who has FOP, aired several times during the day and ended with a longer report in the evening news.



Hugo Fahlberg with his grandfather Per Hallbert in Eskilstuna, Sweden



Annual German FOP Symposium



Annual Family Meeting in Spain

2015

Jeannie L. Peeper Award Winners

We are grateful to each award recipient for their continued loyalty to help the FOP community. We appreciate your passion and dedication to assist the IFOPA in its mission to fund research to find a cure for FOP while supporting individuals and their families through education, public awareness and advocacy.

Lifetime Leadership Award

Jennifer Snow



Jennifer (Jen) Snow is one of the leading pioneers of the IFOPA. She has worked tirelessly for the past 21 years to help the FOP community and has actively supported the

IFOPA's mission. Along with her family, they have held many amazing grassroots fundraising events, including the Find a Cure BBQ. This 17-year event holds the record for the longest running fundraising event in IFOPA history, raising more than a million dollars for the IFOPA. Since her daughter Stephanie's FOP diagnosis in 1994, Jen has shared the FOP story publicly in both local and national media and at events. She has served on the IFOPA Board of Directors in many capacities over the years, including Vice Chair in 2011. Our Founder and President Emeritus Jeannie Peeper notes, "Jen, you have played a key role in so many IFOPA projects, wearing many different hats; it's hard for me to name them all." She served on and led many IFOPA committees, including those that supported past special events such as the 3rd & 4th International Symposiums, and more recently

helped design the patient panel at the first-ever FOP Drug Development Forum. Jen has played a key role in research approaches from the beginning and helped form the current IFOPA research and drug development vision as past Co-chair of the Research Committee. Jen always advocates for FOP patients everywhere. Her passion for the FOP community and her dedication to the patient voice being heard is reflected in all that she has accomplished. Thank you Jen for your dedication, love and generosity.

Outstanding Community Involvement Award

Individual – Michael Man



Michael (Mike) Man has been a pioneer in the IFOPA's fundraising efforts. His commitment to the IFOPA's mission began when his son Spencer was diagnosed with FOP, and Mike's generosity continued long after Spencer's death in 1991 at the age of 14. Prior

to Spencer's death and continuing well after, Mike's endeavors in supporting the IFOPA are historic. The IFOPA has been able to expand its research and member services because Mike was consistently committed for so many years. The IFOPA benefited from the Spencer B. Man Memorial Foundation which Mike created in honor of his son in 1994, and the IFOPA received funding from the Foundation until its closure in 2012. Mike personally donated up to a \$4,000 match for each Spencer Man Memorial Drawing event that the IFOPA held. Mike's fundraising efforts also included his memorable donation of a classic car that the IFOPA auctioned off and IFOPA member, Sharon Fitts, and her husband, Henry, won. Mike also provided consulting to the IFOPA on fundraising as the association was growing and he served as a Board Member from April 2009 through Dec. 2010. Mike's dedication and extraordinary passion are remarkable.

Group – Lincoln's Legacy: Doug & Marilyn Wheelock Mark & Robin Gambaiana



Marilyn and Doug Wheelock with
grandsons Hudson and Lincoln

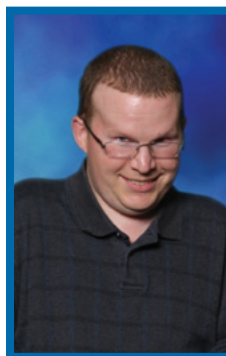


Robin and Mark Gambaiana

The Lincoln's Legacy events started in 2009, in honor of their grandson Lincoln Wheelock, shortly after his diagnosis at age three. Both sets of grandparents and their families immediately became involved in the IFOPA and held a dinner plus silent auction, inviting friends and family to learn about FOP and raise funds for FOP research. They have continued to host biennial dinners and auctions, with each event becoming larger. During the years they don't host the dinner and auction event, Lincoln's Legacy does a very large letter writing campaign to raise FOP awareness and funds to support FOP research. In addition to these larger-scale events, the families have created a variety of other smaller-scale activities including "Leap for Lincoln," a Trike-a-thon, holiday shop, Sioux City Bandit (indoor football team) game volunteers, and Living History Farms "Mud Runs." The "Mud Run" in November 2014, had a total of 29 runners participating in honor of Lincoln and collectively generated \$32,500 for FOP

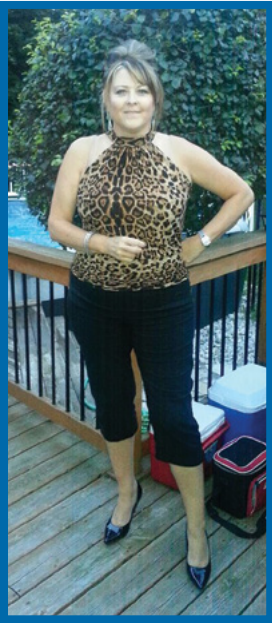
research. Each family has had birthday parties or Boss's Day events where donations were made to the IFOPA in lieu of presents. To date, Lincoln's Legacy events have raised more than \$500,000 and we are thankful for their continued support to find a cure for FOP.

Inspiring Leadership Award Patrick Doerr



Patrick (Pat) Doerr is an IFOPA community member who has FOP. Since he became a member in 1996, Pat has demonstrated exceptional commitment to improving the quality of life of individuals in the FOP community and his personal achievements in the face of adversity are outstanding. He has served on the IFOPA Board of Directors as Secretary and Vice Chair. Presently, Pat is actively serving on two IFOPA committees, the IFOPA Website Redesign Committee (co-chair) and the IT Committee (co-chair). He has also given a hand to the Mentoring Committee from time to time. Professionally, Pat works as an accountant and his co-workers honor him annually by raising money for the IFOPA. He has encouraged others to donate to the IFOPA through online beauty pageants, local businesses, are regular donors, and his family orchestrated a bicycle event – Unswerving, The Trans-Wisconsin Bicycle Ride to honor him. Pat is faithful in representing the FOP community and the IFOPA. He has worked with Gary Whyte to further FOP and rare disease efforts in the legislative arena, and has attended annual fundraising events for many years including the Comedy Show and Bingo For A Cure! Pat's support includes attending FOP symposiums, speaking at the Scientific Workshop: Strategies for the Treatment of FOP, participating in the IFOPA's Teen and Adult Meeting, and 25th Anniversary Celebration. Leading by example, living independently with determination, exhibiting a positive attitude and a zest for experiencing life to the fullest, Pat is an inspiring role model to our FOP community.

Outstanding International Leadership Award Carrie Connell



Carrie Connell lives in London, Ontario, Canada. After her daughter Brooke was diagnosed with FOP in 2007 at age six, Carrie joined the IFOPA, and was disappointed to learn that there was no national FOP group in Canada. She decided, therefore, to form one herself, and in 2009 she became the founder of the Canadian FOP Network (CFOPN). Carrie has served as President since the group's inception, and has been the driving force behind the organization's activities. Under

her leadership, the CFOPN became a not-for-profit corporation and obtained charity status, the latter of which enables Canadians who donate to support FOP research charitable tax receipts, allowing deductions for income tax purposes. Carrie has spearheaded and organized CFOPN family conferences in 2009 and 2013, both of which were deemed a great success by attendees. These events included opportunities for FOP patients to be assessed by FOP medical specialists Drs. Kaplan and Pignolo and by FOP dental experts Drs. Nussbaum and Friedman. Carrie has provided logistical assistance through the CFOPN to families wishing to run charitable fundraising events, including helping organize online donation websites. While leading CFOPN, Carrie has also participated in annual conferences run by the Canadian Organization for Rare Diseases (CORD), as well as recently attending the IFOPA's 2015 International President's Council meeting in Rome, hosted by FOP Italia. Her role as the IFOPA's International President's Council Canadian representative is vital to helping the IFOPA successfully implement our education, awareness and advocacy programs to support our Canadian FOP members and their families.

2015

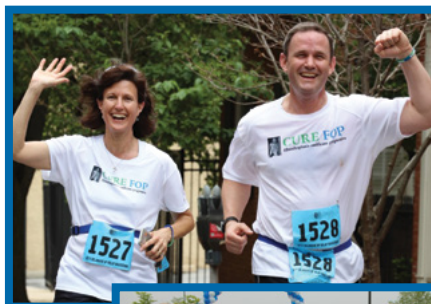
Fundraising

Each year individuals embrace their passion and creativity by hosting unique fundraising events and doing various fundraising activities to support the IFOPA's mission. We could not be more grateful to the family fundraisers and their networks of friends and colleagues who supported them so generously. Thank you!

Events that raised \$5,000 or more in 2015

All for One, One for All – Delaware Marathon

In honor of Justin Henke
Wilmington Riverfront, Delaware
\$9,716



Left: Justin's parents Wendy and Kevin Henke



Middle: Pre-race cheer by UPenn FOP Lab staff



All for One, One for All group with Justin Henke in the front row center

Awake for a Cure

In honor of Ashley Martucci
Delmar, New York
\$38,094



Dusk to dawn fun for middle school students playing carnival games at Awake for a Cure

Bingo for a Cure!

In honor of Joshua Scoble
Allentown Fairgrounds, Pennsylvania
\$92,000 grant from Joshua's Future of Promises, 501c3



Dr. Fred Kaplan speaks to the large crowd at Bingo for a Cure!



Bingo for a Cure! event brings together FOP friends and families

Back row left to right: Jeannette Bordeau, Lisa Bordeau, Christine Lichtenberg, Lindsay Ruiz, Ashley Kurpiel, Patrick Doerr, Kathy Ford

Front row left to right: Karina Chaikhoutdinov (seated), Joey Hollywood, Dr. Fred Kaplan, A.J. Gonzales, Bobby Johnson, Joshua Scoble

Charity Ride for FOP

In honor of Ashley Martucci
Staten Island, New York
\$5,259

Lincoln's Legacy Dinner, Drawing & Silent Auction

In honor of Lincoln Wheelock
Sioux City, Iowa
\$56,427



Lincoln's Legacy Dinner Event

Back row left to right: Marilyn & Doug Wheelock, Dr. Fred Kaplan, Trish Wheelock, Robin & Mark Gambiana
Front row left to right: Lee, Hudson and Lincoln Wheelock

Lincoln's Legacy Mud Run

In honor of Lincoln Wheelock
Urbandale, Iowa
\$16,450



6th Annual Lincoln's Legacy Mud Run was a snow run!

Midnight Sun Color Run

In honor of Dilyn Martin
Kotzebue, Alaska
\$5,000



Color toss at Midnight Sun Color Run in Kotzebue, Alaska



Dilyn's family with fundraisers

Back row left to right: Ann Sieh of OTZ Telephone, Jade's sister, Derek Martin (Dilyn's Dad)
Front row left to right: Maija Lukin, Lucy McConnell, Jade Hill (Dilyn's Mom) and Dilyn Martin

ZipperQ4 BBQ, Auction & Jeep Drawing

In honor of Zip Gordon

Claremore Fairground, Oklahoma

\$120,344



Amy and Zip Gordon give BBQ awards at ZipperQ4

Other Family Fundraising Activities that raised \$5,000 or more

In honor of Sona Brinkman

Letter campaign donations to support the
Competitive Research Grant Program

\$153,347

In honor of Ashley Martucci

Letter campaign donations

\$13,700

In honor of Natalie McGuire

Letter campaign donations

\$391,665

In honor of Sienna Otto

Sienna's Flower Garden donations

\$36,792

In honor of Maria Wray

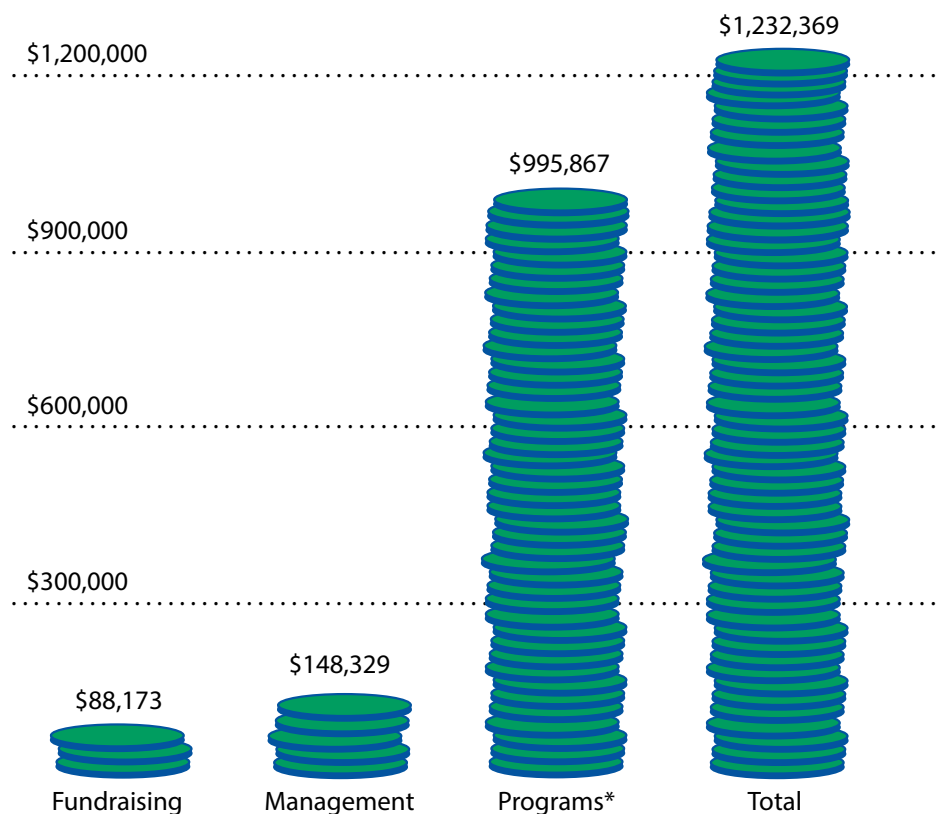
Various fundraising activities

\$11,857

Our Financials

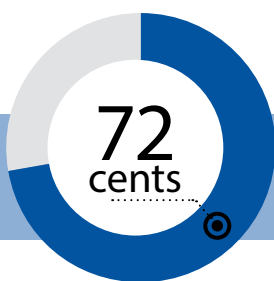
Where Your Money Went

Fiscal Year 2015



*Programs

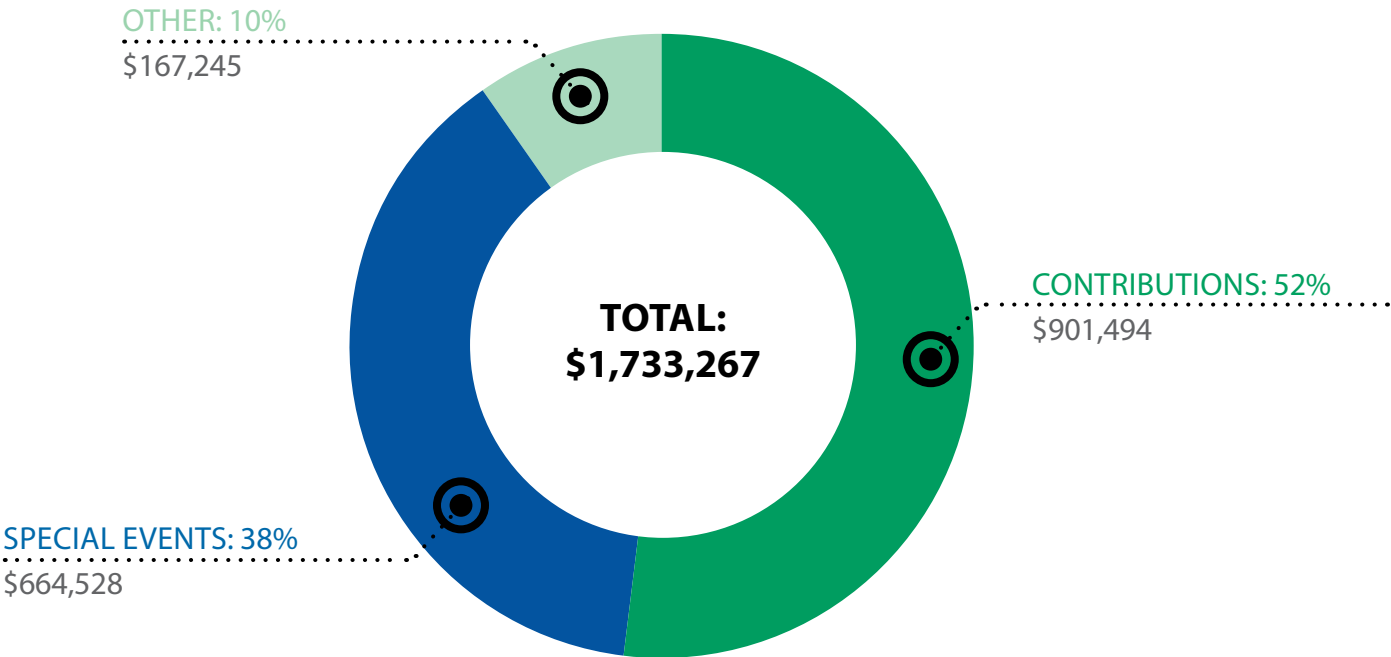
Medical Research: \$900,938 Education & Support: \$56,008 Public Awareness: \$38,921



of every dollar goes to medical research.

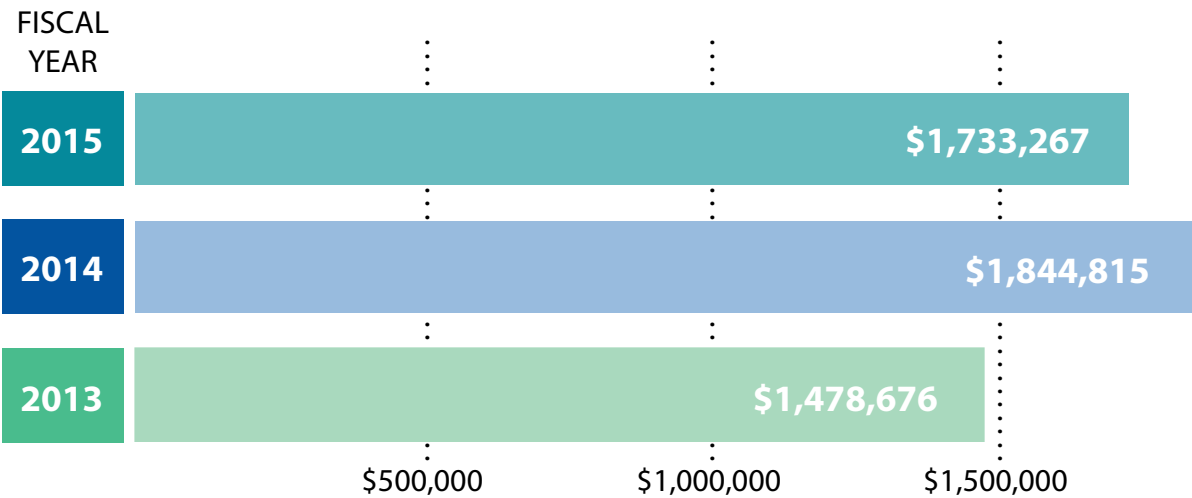
	Fiscal Year 2014	Fiscal Year 2015
Medical Research	\$757,962	\$900,938
Education & Support	\$41,116	\$56,008
Public Awareness	\$38,037	\$38,921
Fundraising	\$60,975	\$88,173
Management	\$94,259	\$148,329

Where Our Money Came From



Annual Revenues and Financial Position

Annual Revenues



Financial Position: December 31, 2015

ASSETS		LIABILITIES		NET ASSETS
\$4,200,660	–	\$312,827	=	\$3,887,833

2015 IFOPA Donor Honor Roll

Thank you to the many wonderful families, individuals, organizations, corporations and foundations that supported the IFOPA with generous contributions between January 1 through December 31, 2015.

Diamond

\$25,000 and above

Anonymous
Clementia Pharmaceuticals, Inc
Jesse David Hendley Foundation
Joshua's Future of Promise, Inc.
Donna Martucci-Segal & Gary Segal
Vanguard Charitable

Jefferson County School District
Patrick Lynch
Allison & Nino Marakovic
Angela & Peter Martucci
Halsey & Natalie Otto
Societe Generale
United Way of Delaware
Doug & Marilyn Wheelock
Wheelock & Bursick Dentistry
Cecil & Gilda Wray

Brady Walker
Ed & Gail Weakland
William Blair & Company Foundation
Melinda & Taylor Wray

Platinum

\$10,000 - \$24,999

Bridgewater Associates, LP
Rob & Priscilla Brinkman
Fidelity Charitable Gift Fund
FOP e.V.
FOP France
Goldman Sachs Gives
Goldman, Sachs & Co.
Claudia & Kerry Hueston
Resat & Tugce Korkmaz
Atul Madan
Mildred & Joseph Martucci
Phoebe Snow Foundation
Powell Broadcasting
Prestige Brands
George & Pamela Rohr
Svenska FOP Foreningen

Silver

\$2,500 - \$4,999

Monica & Kriston Beck
Jared & Stephanie Bennett
BNSF Railway Company
Canyon Partners
Frank & Joyce Casavo
Cedar Hollow Inn
Central States Crane & Hoist LLC
Cisco Foundation
Goldman Sachs Asset Management
Scott Gordon
Kenneth & Denise Henke
Jack Krakower
Lawrence Maguire
Michelle Maguire
Delmar & Mabel McWilliams
Robert & Sandra Melton
Craig Merrigan
Larry & Peggy Nilson
Pelco Structural, LLC
Corby Rust
Robby & Erin Sawyer
Sioux City Bandits
Skelmir, LLC
Tyson Stanely
The Morgan Fund
Triam Fund Management, L.P.

Bronze

\$1,000 - \$2,499

Gina Addeo-Sullivan & Michael Sullivan
Ally Financial
August Packaging
Avery Outdoor
Ayco Charitable Foundation
Carol Ayotte
Bennett's Route 66 Pharmacy
Elizabeth & Robert Blanchette
Anita Bonanni
Mark Brungardt
John & Amanda Cali
Roger & Joy Caudron
Gregory Cerchione
CFG Community Bank
Christine & Clarence Cibella
Douglas & Sharon Coleman
Concord Village Lions Club Services
Foundation Inc
Cut Rate Auto Purchasing Corp.
Srikanth Dasari & Anitha Kosanam
Frank & Joy Degennaro
Doctors Express Mgmt of Central Texas
LLC
DPGroup GC Inc.
Aris Economides
Mary Francioso
Paul & Dixie Gambaiana
GE Foundation
General Mills
Jacob & Amy Gordon
Jack Gordon Jr. & Elise Gordon
Minor & Jane Gordon

Gold

\$5,000 - \$9,999

AQR Capital Management, LLC
Deans Knight Capital Management, Ltd.
Laura Willhoite
Rusty & Amy Edmiston
Mark & Robin Gambaiana
Herman Goldman Foundation

Greater Horizons
 Gryphon International Investment Corporation
 Carol Ann & Allan Hagen
 Marilyn Hair & Richard Steele
 Nancy & Joseph Hair
 Harford Chapter No. 83 OES
 Sharon & Lance Holm
 HSBC Philanthropic Programs
 Institutional Capital LLC
 Eric Jauquet
 Kelly Johnson
 Pierre & Colleen Jraiche
 Jon & Kara Keenze
 James & Fran Kennedy
 Kathryn Kennedy
 Kiwo, Inc.
 Linda & Reed Kneebone
 Peter & Constance Lacaille
 Lakewood High School
 Loomis Sayles & Co.
 William Luong
 Richard Martucci
 Sally & Bill Meadows
 Memorial Machine
 Boyd & Lenore Meyer
 Midtown Animal Hospital P.C.
 NISA Investment Advisors, LLC
 Northrop Grumman
 David & Julie Olson
 Pasha Lake Cabins
 PayPal Giving Fund (Missionfish)
 Perella Weinberg Partners
 Prudential Financial
 Doyle & Bridget Queally
 Samuel Ramirez
 Gilles Randot
 Shailaja Reddy & Mike Oswald
 Jean Sand
 Schoen Family Charitable Trust
 Schwab Charitable Fund
 Securities Finance Trust Company
 Jerry Shapiro
 Jordan & Melista Sligh
 William & Ruth Ann Smith
 St. Nicholas Roman Catholic Church
 Paula Stafford
 Sheldon Stone

Chris Stratton
 Christy & Travis Terral
 The CarMax Foundation
 The Home Collection, Inc
 The Stocker Foundation
 Robert & Terri Thomas
 Thomson Reuters
 Towers Watson Matching Gift Program
 Tulsa Community Foundation
 U.S. Charitable Gift Trust
 United Way of Midland County
 Triveni & Sadhana Upadhyay
 W.K. Kellogg Foundation
 Trisha & Lee Wheelock
 Will Rogers Stampede Rodeo
 Helen & Chris Wray
 Mohamed Hafez Zaher

Cooper

\$500 - \$999

Albany Podiatry
 Alex & Ani, LLC
 Amazon Smile Foundation
 America's Auto Auction Tulsa
 BancFirst
 Edith Beck & Kathleen Bartelt
 Marvin L Bellin
 Vinni Bhandari
 Big Fish Ontario
 BlackRock Matching Gifts Program
 William & Jeannie Blank
 Jerrold & Thelma Blumberg
 Perry & Karen Bourne
 Brass Eye Center
 Briar Cliff University
 Sharon Burke
 Randall L Burnight D.D.S.
 John & Bobbie Cary
 Warren & Gloria Chapman
 Paul & Kathy Coffin
 Constellation Energy Group Employee Fund
 George & Carolyn Cox
 Cross-Bo Construction
 Charlie Crumpton
 Debbie & Mark Davis

Decker, Sbuttoni, Boghosian, Dicerbo & Lawless
 Delmar Pediatrics PLLC
 Double Rafter D Enterprises
 Mary Drechsler
 David DuBard
 Jason & Celesta Easton
 Fred Erbes
 Mike & Emily Ferrell
 Robert Flanigan
 Andrea Fleser
 FOP Stichting Nederland
 John & Helen Franklin
 Fright-Rags, Inc.
 Robin & Ivo Garcia
 Gateway Mortgage Group, LLC
 Heather Gaynor
 Sandra Gordon
 Patrick Gordon
 Garret & Deborah Green
 Dave Guggenmos & Bill Book
 Dason Gwartney
 Beth & Matthew Hardison
 Harry Bruno Salon
 Elena High & Dr. David High
 Rapture Hill
 Donna Holloran
 Hughes Optician, Inc
 Douglas Hulsey
 iGive
 Indian Springs Country Club
 Industrial Process Design
 James Elam Fishing, LLC
 JD Beaven's Martial Arts
 Jimmy Makes Pizza, LLC
 Cindy & Roger Johnson
 Jim & Lynn Johnson
 Kiwanis Club of Santa Maria
 Kneifl Electric, Inc
 Laura & Kevin Kuechenmeister
 Ladco Inc.
 Jeremy Ladd
 Vicki & Pat Langin
 Lawrence & Sandra Post Family Foundation
 Brittany Lee
 Linda Lepak
 Moira Liljestrom & Federico Robert

James & Tisha Lindgren
 Love Air Conditioning LLC
 Joanne & Anthony Maracic
 Robert Martin
 Joseph & Diane Martucci
 Mazzotta, Sherwood & Vagianelis, P.C.
 Christopher McCarrick
 Britta McSparen
 McWilliams Farm
 Larry & Patty Melton
 Jennifer Melton
 Christopher Metkus
 Linda Miller & Timothy Duperron
 Mondrian Investment Partners, Inc
 Loren & Connie Mulder
 Mullin Services, Inc
 Karen Munro & Peter Friz
 Odessa L Ofstad
 John Palmer
 Peoples Bank
 Perception Travel LTD
 Performance Physical Therapy, LLC
 Tim & Anna Petersen
 Patricia Pinkham & Larry Albee
 Tracy Pomerson
 Premier Consulting Partners
 Project Visual
 Herbert & Rita Raiffe
 Carlos Rangel
 Red, White & Bounce
 Lynn Refer
 RegentBank
 Renaissance Charitable Foundation, Inc
 Donald & Doris Ribble
 Steven Rose & Sandy McAgee
 Phisanu Sermchaiwong
 Justin Sheperd
 Sioux City Country Club, Golf Shop
 Sioux City Explorers
 Robert Sparling
 Spectrum Paint
 Jason Spurgeon
 St Peter's Health Partners
 St. Andrew United Methodist Church
 Wendy & Mark Stevenson
 Jonathan Stichbury
 Talbots, Inc
 The Kresge Foundation

The Ross Group Construction
 Corporation
 Sven Thorson
 Trattoria Fresco
 Delayna & Adam Trease
 Tyson
 Union Bible Study Association, Inc.
 Karuna Venter
 Jacques & Marcelle Vergellati
 David & Gloria Wadd
 Stephen & Miranda Wade
 Wagner, Kuntz & Grabouski, PC
 Amanda Walker
 Betty Wheelock
 Linda & Richard Wille
 David Willis
 Julie Wingerter & Seth Lieberman
 Kris & Art Winters
 Woodbury County Conservation Board
 Dick & Judy Yamane
 Yellowhouse Machinery Co.
 Zazove Associates, LLC
 Zebco

Associates

\$100 - \$499

Harvey Abrams
 Absolute Massage & Bodywork Spa
 Acadian Asset Management LLC
 Michael & Andrea Adamczyk
 Curt & Joann Adams
 Kathy & Bill Adkins
 Catherine Adlerman
 Sabrina Aguilar
 Timothy Akanmu
 Elisa Albert
 Eric Alby
 Salma Al-kindy
 James & Letitia Allen
 Camille Allen
 Moosa Al-Moosa
 Julia Althoff
 amazon.com
 Mary & Mark Amodeo
 Frederica & Marvin Amstey
 William Anderson

Brent & Mary Anderson
 Shirley Andre
 Tom & Kathy Andrews
 David & Kathleen Angle
 Aon Foundation
 Leslie Araskog
 Archie's Wayside Restaurant
 Claudine Armand
 Allison Armour-Garb
 Margaret Armstrong
 Beverly & Donald Arnold
 Andrea Arrigo
 Arthur Murray Dance
 Corrine Astrup
 Auto Definitions of Claremore, Inc
 Joy Axelrad
 Catherine Baase
 Mark Bachman
 Cheryl & Boyd Bahney
 Teresa Bailey
 Kelly Bailey
 George & Patricia Baker
 Matt Balanda
 Jeff Baldus
 Gladys Ball
 Bonnie Ballard
 Jennifer Baniak-Hollands
 Dana Bannan
 Larry Banzet
 Bernard & Christine Barbell
 Ellenor Barish
 John Barker
 Daniel Barnes
 Liz Baroody-Solomon
 Michelle Barriere
 Richard & Sera Barry
 Rita Bartholomew
 Constance Bartholomew
 Tony & Lisa Barton
 Peter Bassler
 John & Lynna Beaven
 Darlene Beccaria
 Chris & Helen Bedford-Gay
 Kenneth Beekley & Patricia Olson
 Avraham Belizon
 Bella Zen Spa
 Paul Benzing
 Crystal Berger & Daniel Leo

Beatrice Bertrand
 Gordon & Cheryl Besser
 Aaron Beutler
 Billy Sims BBQ
 Karel & Robert Birnie
 Lavonne Blanco
 Blessed Hands Massage Therapy
 Bob & Beth Blowers
 Bob Roe's Point After
 Michelle & Roger Bogard
 James & Jeanne Anne Bonanni
 Chris & Lori Bonanni
 Joseph Bond
 David & Karen Bond
 Melissa Boni
 Vassiliki Bonness
 Annette Bonzelaar
 Shari Book
 Patricia Borusiewicz
 Floanne Bowen & Linda Titus
 Dorothy & David Bowles
 Alan Brandt
 Thomas & Marianne Brasel
 Eileen & Rich Braun
 Bravelets LLC
 Charles & Janice Bray
 Alysa Brennan
 Jonathan & Rachel Bress
 William & Carole Brice
 Don Brister & Diane Bennett
 Ellen Brodock
 Ann Elizabeth Brooks & Donald Mann
 Brother Nature Charter
 William Brown
 Debra Brunner
 Bucks Creek Foundation
 Bobby Buonomo
 John & Barbara Burgdorger
 Brian Bursick, D.D.S. & Kristy Bursick
 Bush Dry Cleaners
 Delio Bustos
 Nicole & Vance Butler
 Louis & Steven Butterworth
 Dana Byron & Allison De May-Byron
 C. E. Moody Construction
 Cable One Advertising
 Kevin Caggiano
 Augusta Caggiano

James & Lynn Cain
 Misty Cain
 Chris Calabrese & Stella Plutino
 Susan Caldwell & Timothy Johnston
 Jeremy & Mikayla Callison
 Susan & Al Camarillo
 Chad Campbell
 Kathryn Campion
 Karen & Charles Campise
 Brian & Debra Cannon
 Cardinal Farms
 Cardinal Health Foundation, Inc
 Robert Carlson & Aileen Lee
 James & Cheryl Carney
 Bill & Karen Caron
 John Carpinello
 Steve and Bobbi Carriger
 Gordon Carriger
 Dr. & Mrs. John L Carroll, Jr.
 Molly Carson
 Marie M. Carter
 Brandon & Shannon Cathey
 John & Margaret Cavanagh
 Paul Cavazos
 CCK Strategies
 Marat & Irina Chaikhoutdinov
 Lonnie & Pat Chambers
 Brad & Melody Chambers
 Chevron Humankind
 Christie Fishing
 Mario & Anna Chuliver
 Dr. Abe Chutorian & Helen Chutorian
 Ricard Ciccone, M.D. & Natalie Ciccone
 Beth Clagg
 Claremore Tag Agency
 Dorothy & Larry Clausen
 Becki Clauss
 Clayton State University
 Patricia L Cleary
 Jackie Clement
 Randy & Jamie Clyma
 Kim Cochran
 Jerome & Sherry Cohen
 Cole Chiropractic Clinic
 College of Dentistry, University of Iowa
 Markham & Becky Collins
 Allison Collis
 Steven & Barbara Condon

David & Diane Connaughton
 Tom Connell
 Judith & Bruce Connery
 George & Irma Contrades
 Gianni & Adelina Cotognini
 Counterpart International
 Cozy Cottage
 Crisafulli Bros
 Sharon Criss
 Lacy & Derrick Crissup
 Eileen Crowell
 Edward & Jennifer Crum
 Rex & Sherry Crumpton
 Crystal Clean Car Wash
 Jeffrey Cucunato
 Larry Cunningham
 Custom Sewing Solutions
 D P Sales
 Cathy & Mike Dailey
 Dakota Dunes Country Club
 Henry & Susan Daley
 John & Patsy Daniel
 Lori & Kenneth Danzer
 Jerrold & Lisa Danzer
 Darwood Associates, Inc
 Amy Davenport
 Scott & Robin Davis
 Craig & Barbara Davis
 Thomas & Kathy Dawson
 Anthony De Casaris
 Adam & Brita de Malignon
 Tom De See
 Robin De Vito
 Sherri Dean
 Roxann & David Debeau
 Tulip & James DeJulio
 Rick Del Mastro
 Elaine Delmonte
 Deloitte
 Norma & Robert Dempsey
 Nancy Dennings
 James & Lisa DePalma
 Gale & Michael DePaolis
 Marian Derkse-Disco
 L George Detweiler
 Timothy Devlin
 Maria Di Giambattista-Shafer
 Robin Diamonte

Phillip & Kimie DiBello
 Danny & Ruth Dick
 Heather & Kevin Dick
 Stephanie DiGiovanni
 Sean Dinn
 Distinguished Programs Insurance
 Brokerage, LLC
 Tara DiTore
 Melissa Dodge
 Lucille Doege
 Richard & April Dollarhide
 Andrea Dominowski
 Richard & Caroline Donaway
 Tammy Donnelly
 Terri Dooley
 Milan & Shraddha Doshi
 John & Jolynn Dotson
 Neva Doyle
 Jillane Doyle
 Dr. Timothy Suorsa Optometry
 Ainslie Drake
 Paula J Drechsler
 William & Sharon Drilling
 Drilling Morningside Pharmacy
 Donald Dugger
 Dunkin' Donuts
 Mark & Laura Durkovic
 Karrie Dvorchak
 East Woodstock Cong. Church
 Breece Eberly
 Sharon Eckart
 Dale & Marla Edwards
 Edwin Evers, Inc.
 Gary & Valere Egnasko
 Edgar & Barbara Einhorn
 David & Judi Einhorn
 Elements of Beauty
 Embrasse-Moi
 Elizabeth Emery
 Benard Emery
 Pam Emery
 Jennifer Emery
 Adrienne Emery
 William & Judy Engstrom
 Erdody Studios
 Andrea & Larry Erskine
 Robert & Sarah Erwin
 Ran Ever-Hadani

Tania Fago
 Raouf & Mona Farid
 Brent Farmer
 Everett & Phuong-Nga Farr
 Fathree & Fathree P.C.
 Fred Feldman
 Cynthia Ferrari & Bryan Thomas
 Jacqueline Fielding
 Fiera Capital Corporation
 Isabel Filippone
 Beth Fillmore
 Gregory & Kim Finkbeiner
 Brady Finton
 Rick Fishbein
 Nichole Fisher
 Ryan Fitzpatrick
 William Flanagan & Mary Lou Truitt
 Sherri & Steven Flaugh
 Timothy & Julie Fleetwood
 Michael Fleisher
 Kathryn Ford
 Laura Ford
 Forefront Medical Services, PLLC
 Steven & Julie Forney
 Brittany Fortna & Jeff Cooper
 Brent Foster
 Kenneth & Donna Franklin
 Alan & Pat Fredregill
 Jeff Freedman
 Stephani Freeman
 Mark Friedel
 Mark & Karen Friedman
 Christine Friga
 Scott Frink & Susan Beidler
 Froman Welding & Fabrication Inc
 John William Frye, Jr & Catherine Frye
 Dr. Edwin Galkin & Roberta Galkin
 Jeanne Galla
 Kathleen Gallagher
 Daniel Gallagher
 Primo Gallanosa
 June & Matt Gambaiana
 Tom Gambaiana
 Mindy Gandin
 Jason Gans
 Robin Gantz
 Andrew Garb
 Fred & Joan Gardner

Mr. Jimmy Garner
 Jon Garrison
 Margaret & James Gavin
 Anna Gavrilov
 Jeffrey & Shay Gebauer
 John & Elizabeth Gianis
 Steven & Jessica Giardini
 Beth Giarrusso
 Ted Gibson
 Robby Gibson
 Linda Gicewicz
 Brita Gieryic
 Robert Giese, Jr. & Laura Giese
 Elizabeth Gillin
 Brooke Gillman
 James & Anne Gilmore
 Connie & Thomas Girard
 Adam Glasser
 John Goebel & Ann Rehan
 Ahmet Gokcen
 Dan Goldberg
 Jonathan & Deborah Goldman
 Elizabeth Goldstein
 Martha Gomez
 Kathryn Gooch
 Christine Gordinier
 Felice Gordis
 Richard Gordon
 Donna Gordon
 Casey Gowler
 Donna Grabow
 Grant Gingerich State Farm
 Reagan Gray
 Christopher Green
 Green Valley Nursery
 Greenberg's
 Lisa & Gregory Greves
 Craig Gries
 Amy Griffin
 Leon & Lois Grothe
 Luanna & Roger Gruber
 Lynda Gudmundson
 Guidepoint Global
 Dustin Gunnens
 Rishi & June Gupta
 Gymnastics City
 Sally Hair & Brian Reid
 Hair Relations

Kay Hall
 Ken & Melissa Hall
 Rose Hall
 Rosie & James Halpin
 Greg & Ivy Hamerly
 Lyle & Mary Hamstra
 Denise Hansen
 Aarti Harper
 Danielle Harris
 James Harshaw
 Daniel & Sally Hartley
 Melinda & Thomas Hartzell
 Sally Hawke
 Kevin & Debbie Hazlett
 Lirong & Sunny He
 Trudy & Amanda Hedgpath
 Marcia Helentjaris
 Tracy Hembree
 Jackie Henderson
 Lacey Hendrix
 Eric Henry
 Michael Heraty
 Debra Herber
 Anthony & Bonita Hermes
 Terry & Linda Hersom
 Phyllis Hess
 Stanley & Nina Heyn
 James Heys
 High Plains Welding
 Don Hill
 Carl & Judy Hischka
 Hitches, Trailers & More
 Camilla Hjerl
 Leona Hodge
 Konstantinos & Ivana Holiastos
 Ginger Holley
 Eric Holmquist
 Josephine E. Holzer
 James & Barbara Homer
 Charles Hong
 Robert & Pamela Hopper
 James Hough
 Martin & Kathryn Howard
 Edward Hsiao
 Mark Huamani
 Leslie A Hughes
 Ingrid Hughes
 Rebecca Hughes

Robert Hunkeler
 Jeff & Tammy Hunter
 JoAnn Huth
 James Hyler
 Denise & Dana Ianniello
 IBM Employee Services Center
 Jess Ihrke
 Integrated Associates, LLC
 Integrated Practice Management, Inc.
 Iowa-Nebraska State Bank
 Heather Isaacs
 Daniel Ishac
 J & J Pharmacies, Inc.
 Reid & Sandra Jacobsma
 Eric & Angela James
 James R Vozar Insurance Agency, Inc
 Sabine Jansen
 Kimberly Janson
 Austin & Carissa Janssen
 Stave & Martha Jany
 Mike & Judith Jayne
 Nancy & Richard Jayne
 Carol & Dean Jeffery
 Jeffrey A. Bright, D.M.D., P.A.
 Jeff's Sno & Blow
 Bryan Jendretzke
 Anne Jennings
 Jerry's Pizza
 Jesco Product, Inc
 John Deere Foundation
 John Williams Company
 Susan Johns
 Carl Johnsen
 Neil Johnson
 Trent & Sherry Johnson
 Stacia Johnshon
 Roxanne Jolley
 Natalie & Jeff Jones
 Jacob & Chrystal Jones
 Zac & Kori Jones
 Michelle & Matt Jones
 Gregory Jones
 Jones Pest Control
 James & Marilyn Jordan
 Jenny & Craig Jorgensen
 Scott Jorgensen
 Annalena & Martin Josefsson
 Morgca Joule

Reginald Joule
 JP Morgan Chritable Giving Fund
 Virginia Judge
 Mary Jusun
 Daniel Kaler, D.D.S., P.C.
 Dr. Harvey Kaplan & Gay Kaplan
 Robert Kapphahn
 Valerie & Chad Kapphahn
 George Karkanias
 Corey Kautz
 Paul Kavanagh
 Ronald Kazyak
 KCAU TV
 Mike Kedzior
 Diane & David Kehm
 Linda & David Kellen
 Lisa & Charles Keller
 Sally Anne & James Kellogg
 Phil Kendall
 Murray & Kimberly Kennedy
 Thomas Kent
 Kiki Kenworthy
 Michael Kerran
 Pam Kilmain
 Misong Kim
 David W. Kimball
 Fiona & Ryan Kimber
 Scott & Andrea Kimmelman
 Alasdair King
 King's Pointe Water Park
 John Kirton
 Dr. Joseph Kitterman & Kathleen Kitterman
 Elbert Klein
 Brandon Klinge
 Adam Klucevsk
 Alisha & Matthew Knapp
 George & Gloma Kneebone
 Knova's Carpets, Inc.
 Daniel & Kelly Kobosko
 Gina & Tim Koch
 Kate Koetting
 Kenneth Kolkebeck
 Paul & Kristen Kolsrud
 Janice Korver & Dr. David Korver
 Steve Kuhn
 Christian Kurth
 Paula & Allan Kutner

Shivin Kwatra	Dr. Aarti Madan & Kyle Brinkman	Michigan Agricultural Commodities, Inc
Trieneke Kylstra	James Madison	Randy & Lindley Miedema
Carrie & David Kysar	Frank & Carol Magliery	Kevin & Brenda Miedema
Khushi Ladwa	Krista Maki	Miles Inn
Lakeport Pottery	Stephanie & Tory Malorzo	Lyndsay Miller & Tim Terveer
Vincent & Barbara Lamark	Julie Maltzman	Rhonda & Blake Miller
Jared & Julie Lamb	Maria Wray Events	David Miller & Heather Williams-Miller
Heather Lang	Marina Inn Conference Center	Larry Miller
Raymond & Marie Lanza	Deborah Marko	Cynthia Miller
Rene & Cindie Lapierre	Kenneth & Christel Marks	Rebecca Miller-Riley
Denis & Catherine Laplante	Martin Marks & Lori Klein	Lisa & Michael Mingino
David & Carolyn LaRue	Meegan Maroney-Pelafigue	Mike & Kelly Moffitt
Charlene & Allan Law	Maxine Martin	Lori Monico
Lawn Pros	Tara Martin	Morgan Stanley
Jesse Leach	Don Martindale	Morningside College
Susan Leader	Evelyn Martinelli	Multicare Physicians Group
Kim & Jim LeBaron	Lisa Martinelli	Chris Munch
Jane & Jeremy Ledbetter	Carlos Martinez	Kelley Murray
Richard & Loree LeFleur	Begona Martinez de lecea	Joseph Murray
Ryan & Michelle Lenerz	Jeffrey & Laura Martino	Gregory & Beth Murtha
Leonard & Charlotte Lenzen	Michele & Jim Marum	NANA Regional Corporation, Inc
Jean & Rob Leonard	Darya & Kris Mastronardi	Haley Nash
Mark & Sheryl Leonard	Kathy & Nathan Maxwell	National Christian Foundation
Sarah Lepak	James & Mary Maxymillian	Chloe Naylor
Letters By Laurie	Edward Mazurkiewicz	Renee Nearpass
Lyslie Lewellen	William & Kathryn McCabe	Lars & Meghan Nedwick
Patricia M Lewis	Joellen & Sean McCarthy	Stephen & Terrie Neely
Elizabeth Leyden	Lucy McConnell	Paul & Jonnie Neely
Lightening Car Wash	Mike & Irene McCoy	Neely Financial Services Inc
Heather Link & Kevan Sternberg	Perry & Peggy McGehee	Nancy Nelson
Sandrya Link	Donald & Lee Ann McGill	Sheila Nelson
Bettie Linker	Jerry McGlothlin	Gary Nelson
Thomas & Stacy Livingston	Scott & Colette McGuire	Network Systems+, Inc.
Sanantha & Greg Lombardo	Sharon McKenna	John Neuman
Bryan & Debra Long	Stephen McKinney	New Life Christian Gift Shop
Kristen Elizabeth Long	Mike & Gretchen McMinn	Kenneth & Joyce Newcomb
Lonnie's Window Tint, Mufflers & Accessories	Ariel McNichol	Martin Newman
Jessica Lopez	McSharry & Associates Realty LLC	Mike Nice
Louis F & Mary Tagliatela Foundation	Stephanie & Herbert McSpadden	Beth Nicholas
Barbara & Earl Lovelace	Margaret & Curtis McWilliams	Joseph E Nicolas, MD & Catherine Nicolas
Sue & Linn Lovin	David & Gail McWilliams	Todd & Cathy Nilson
John Lucia & Marianela Olivares	Medved Running and Walking Outfitters	Steven & Holyce Nissen
John Lucia	Robert Melton	Joe Norman
Erik & Susan Lundquist	Ralph & Carmella Meno	Northwest Arctic Borough
Jared & Lori Lusk	Niels Mesotten	Beth Novick
Kathleen & Mike Lutito	David High	Burton & Diane Nussbaum
James Macfarlane	Bob & Maureen Metz	NW Electric Inc
	Michael & Christine Metzging	

Timothy & Denise Oakley
 Patricia & Lawrence O'Hara
 Kimberly Olive
 Megan Olsen
 Olaf Olsen
 Randy & Paula Olson
 Orange Development
 Adam Oravec
 Nicole O'Shaughnessy
 Gary & Lynn Oswald
 Eric & Rory Otto
 Charles & Wendy Otto
 OTZ Telecommunications, LLC
 OTZ Telephone Cooperative, Inc
 Treena & John Owens
 Francine & Donald Padgett
 Troy & Kelly Paino
 Diane Paletz
 Carol Palmer
 Katarzyna & Pawel Panasiewicz
 Marvin Pansegrau
 Barbara Pantuso
 Susan Parker
 Timothy Parker
 Pathway Services, Inc
 Andrea & Nicholas Patti
 Dylan Paty
 Whitney Pebsworth
 Diana Herrera
 Rob & Bobbie Penney
 Arlene & Allan Peper
 Leslie Perrin
 Perry Creek Animal Hospital
 William & Cheryl Peters
 Jeanne & James Peters
 Richard Petersen
 Virginia Peterson
 James Peterson
 Marlene Petrides
 Linda Pfingsten
 Janna Pfluger-Boucher
 Nancy & Butch Phillips
 Jonathan Pierce
 Dr. Robert J. Pignolo
 PIMCO Foundation
 Sue Ping
 Pioneer Pickers
 Robert Pirtle

Barbara Pittman
 Christopher Place
 Art Pluim
 Don Pollak
 Fernando Pombeiro
 Javier Masuet Pombo
 Roxanne G. Poole
 Stephanie Powell
 Andrew Powell
 Matthew & Grace Powers
 Brandon & Julie Pregler
 Shirley & Harry Prellwitz
 Patrick & Jane Premo
 Michael Prosser
 Elaine Quinn
 R & C Robinson LLC
 R G & A M Douglas Foundation IMA
 Eric & Michelle Rahn
 Roxanne Raiola
 Sanjev Rajaram
 Steven & Heather Ralph
 Jennifer Ramsden
 Shelly Ransom
 Carolyn Rants
 William & Carol Raper
 G. L. & D. D. Ratliff
 Colleen Raventos
 John & Alene Reeve
 Randy Reichert
 Lisa Reid
 Maida Remmer
 Thomas & Jenny Rench
 Jeff & Nancy Rendinaro
 John Rendinaro
 Gerard & Dorothy Rendinaro
 Resurgent Capital Services
 Rhett Minson Agency, Inc
 Robert & Dorothy Rice
 Charley Richey
 Rick Hanson Plumbing, Inc
 Frank & Barbara Rickabaugh
 Riddles Jewelry
 Kathy & Rodger Riggle
 Toni Riley
 Robert Burnett Memorial
 Bryan Robinson
 Sean Robinson
 Carrie Rocke

Carol Roeder
 Walt Rogers
 Ellen Roller & Paul Selver
 Pedro Romero
 Marvin & Luella Rorick
 Rose Hill Junior High School
 Beau & Kristin Roseberry
 Ryan Rosensweig
 Sandra Roth
 Grace Russell
 Rustic Cuff
 Jim Ruud
 Michelle Ryan
 Peter Sacchitello & Marie
 Bruzzese-Sacchitello
 Stuart & Jodi Sacks
 Sacred Life Ayurveda
 Jeannine Salvesson
 Gregory Sanchez
 Amy Sanders
 Molly & Peter Sanders
 Nancy Sando
 Anthony Sardella
 Irene Sasaki
 Masahiro Sato
 Kelly Sauders
 Fred Schachinger & Sarah Lochner
 Marilyn Schaeffner
 Marilyn Schafbuch-Tanner
 Roberta Schiller
 Bernhard Schmid
 Thomas & Kathleen Schmidt
 Thomas & Rachel Schmiedel
 Michael & Lindsey Schmitz
 Eric & Maureen Schnabolk
 Louis Schornstein
 Lee & Robin Schroeder
 Bob Schulte
 Pamela & Rob Schultz
 Scott Schweighauser
 Robert & Karen Scott
 David & Georganne Scott
 Mary & Kelly Scott
 Scott L Temple Property
 Scott Miller Salon
 Maria Scrivens
 Marsha & Lewis Seavey
 Craig & Crystal See

Carol & Jonathan Segal	Ingrid Sorensen	Mitzi Thomas
Sandra F. Selby	Stuart Soroka	Rita Thompson
Sally Selquist	Susie Sours	Dorothy Thompson
Tammy Seres	South Mountain Middle School	Craig Thompson
Charlie Shaffer	Joe Spanier	Daniel & Susan Thornton
Robert & Dawn Shaffer	Susan & Reid Sperber	Donna Thummel
Denis Shanahan	Lyle & Donna Spieler	Tiger Rock Martial Arts
Siddhartha & Anand Sharma	Jack & Mary Lee Spinks	Carol Tilley
Barbara & Jim Sheahan	Sara Springmeyer	Carol Tims
Keith & Connie Shellhammer	Joyce Spurling	Tires, Tires, Tires
Carol & William Shortridge	Jeffrey Stammen	Celeste Tirro
Alexandra Shrake	Jennifer Stanbrough	Scottie & Lauri Titsworth
Margaret Sichau	Taralyn & Phyllis Stearnes	David Tjen
Ann & Eric Sieh	Joe Steccato	Shirley Todd
Karla Sigler	Sandra Stecker	Laura & Barry Tofteland
Alan & Barbara Sikes	Les & Gail Steele	Kristin Tollin
Matthew & Drew Sikes	Martha Steidley & Marcie Gaudette	Beverly Tomlin
Silicon Valley Community Foundation	Caryn Sternstein	Tool's Restaurant
Art & Eileen Silva	Andrea & Scott Stevenson	Donald & Jennifer Torey
Silver Maple Landscaping & Nursery, LLC	Evelyn Stockinger	Thomas Trail
Lynsey Simmons	Paul & Julie Stone	Tri State Certified, LLC
John & Madison Sinuc	Stonebrook Day Spa	David & Andrea Troiani
Sioux Air, Inc	Stoney Creek Inn	Audrey Troy
Sioux City Endodontics, PLLC	John & Teri Stoops	Donald & Donna Truby
Sioux City Musketeers Hockey team	Joshua & Jaimie Stough	Truman Heartland Community Foundation
Sioux City Police Officer Assn.	David & Helen Stratton	June & Terry Tucker
Sioux City Symphony	Richelle Streater	Cheri & Andrew Turner
Siouxland Golf Charities, Inc.	John Stuart	Daniel Turner
Rod & Janie Sisovsky	Studio 427	Michael & Laraine Turner
Rosemary Sixbey	James Sullivan	John & Kim Ubben
Richard & Karen Sjolund	Sunnybrook Community Church	United Way of Massachusetts Bay & Merrimack Valley
Daniel & Shawn Skelton	Sunrise Pediatrics	United Way of Metropolitan Chicago
Michael & Morgan Skogland	Superior Realty & Auctions, Inc	United Way of the Greater Lehigh Valley, Inc.
Skyco Painting	Manpratap Suri	Jackie Upton
Jamie Slack	Eric Swanson & Lisa Brooks	Wojciech Urban
Jim Slattey	Amy Swanson	Jill Urner
David & Kathy Sligh	Barry Switzer	John & Michele Vagianelis
Mary & Pearce Sloan	Sherry & Werner Szilas	Joseph & Karen Valentine
Melissa Small	T & W Tire	Greg & Lynn Van Ballegooyen
Lance Smith & Jami Hill-Smith	Talbots - Pittsford	Grada Van Ballegooyen
Deon & Mindy Smith	Satomi Taniuchi	Barbara & Dave Van Hare
Steve & Linda Smith	Gretchen & Aadi Taute	John & Denise Van Otegham
Nancy Snith	Robert Taylor	Robert & Joan Vanderhorst
Wade & Beth Smythe	Heidi & Scott Templeton	Melissa & Damon Vangelis
Donna & Thomas Snider	The Barcus Family Grantor Trust	Mary & Kostantinos
Larry & Sara Snider	The Benevity Community Impact Fund	
Jenny Somers	The District on Main	
	Ajit Thomas	

Paul & Antoinetta Venables
Teresa & Ben Venditti
Pablo Vengoechea & Tim Boyland
Kathy & Dom Vicchio
Denise S. Vietti
Shirley & J E Vincent
Francesca Vitali
VIVE Salon
Loren & Ann Vogel
Jason & Kristin Vogel
Dick Wagner
David & Patricia Walker
Judith Walker
Sallyanne Wallace
Terri Meta Walsh
John Walston
Vicki & Gary Wanderscheid
Esta & William Wannamaker
Therese Ward
Mark Warford
Stephen & JulieAnn Warner
Susan & Andrew Warrington
Ronald & Brenda Washburn
Ernest & Linda Washington
Paula & Lester Waston
Thomas Weakland
Kelly Weaver
Linda & Stephen Webb
Andrew & Krista Webb
Marcia Weller Weinhold
James & Deborah Weishan
Ilene Weiss
Judy Weiss
Jan & Boyce Welch
Robert & Kathie Welch
John Weller
Mark & Barbara Wellisch
Well's Dairy Inc
Wells Enterprises, Inc.
Brian & Kristin Weltman
Blanche Weltman
Kurt Wenrich & Becky Kaplan
Matt Wexler
Emilie Balfour Weyl
James & Beverly Wharton
Adam & Gretchen Wheelock
James White
Justin White

Naomi White
Kathryn Wiegand
Allyson Wiener
Lindsie Wilde
Susan & Loras Wilgenbusch
Curt & Angela Willett
Karen Willhoite
Jeffrey & Tiffanie Williams
John & Penny Williams
Carol & Tom Williams
Patrick & Barbara Williams
David Williams
Joanne & John Wills
Wilson Sports
Wine Styles
Winna Vegas
Kris Winters
Kathy Witherspoon
Woodbury County Emergency Services
Cassie Woods
Charles K Wray
Felicia & Taylor Wray
Dr. James Wright & Helen Wright
Tina & Todd York
Dr. Paul B. Yu
Brenda & Aric Yurko
Karen Yutsus
Suzanne & Berton Zbar
Henry & Kerry Zenich
Christopher & Jennifer Zonnas