A Message From the President

Hi everyone,

My name is Cameron Penn. I became board president for the OI Foundation in July 2019 but this is the first opportunity I’ve had to speak to all of you. It’s an honor to follow two stewards who effortlessly steered the foundation for several years. I’d like to thank Ken Gudek and Mark Birdwhistell for their selflessness, their remarkable consistency and their extraordinary dedication. But I would be selling their leadership just a bit short if I didn’t recognize their families and especially their wives, Teresa Gudek and Martha Birdwhistell. While Ken and Mark were the faces of the board, peel back the curtain and you quickly realize their indelible impact wouldn’t have been possible without incredible support.

As we begin to celebrate a half-century of the OIF’s commitment to all the deserving members of our community, we’re not only reflecting on our history, we’re envisioning what’s next for all of us. Given the current global uncertainty, it has become increasingly apparent that no year in the storied history of the OIF may have a greater, broader impact than 2020.

Right now, we’re generously supporting the brightest minds and biggest hearts in research, we’re expanding fundraising platforms and we’re constantly improving our outreach, through social media and by adding more regional conferences in new cities all over the country. Grants have grown. Membership has grown. The number of researchers funded has grown, Corporate sponsorships for the national conference have grown. And the overall investment in our community has grown. In short, we’re doing everything we can to invest even more in you.

But nothing we do exists in a vacuum. In this critical year, we’re asking each of you to take an active part in the direction and future of the OI community, starting with our pivotal PCOR Project. Patient Centered Outcomes Research is designed with every member of the OI community in mind and it’s directed by Dr. Laura Tosi, our CEO Tracy Hart and our steering committee leads, Michelle Duprey and Mark Birdwhistell. The goals of the OIF PCOR Project are simple: 1.) We want to create an

The mission of the OI Foundation is to improve the quality of life for those living with osteogenesis imperfecta through research, education, awareness and mutual support.
environment where you can better understand treatment options; and maybe most importantly, 2.) We want to empower each and every one of you with a bigger, more influential role in your own care.

To make this dream a reality, we not only want your voice, we need to hear from you. Share your story with us, share your thoughts and opinions. The quickest and easiest way is to join the OIF Registry. Go to oif.org/PCOR. It only takes a few minutes. You can also follow us on Facebook and Instagram @OIFoundation.

In return, we’ll make this promise to you: we’ll do the exact same thing. The information you share with us is invaluable to our researchers and medical professionals. And when they make discoveries, when new trends appear—you will be the first to know. We know research can sometimes seem like a black hole: you share your vital, personal information and no one ever explains why, how or if it helps. We’re changing all that, and we’re changing it quickly.

So, as the 50th Anniversary of the OIF gets in full swing, I want to extend my heartfelt thanks to each of you for your continued support of each other. You never cease to amaze us. We on the board also owe a sizable debt of gratitude to our entire OIF staff, our researchers and medical professionals, fundraisers big and small, and everyone else who supports our mission in countless ways.

As we begin to navigate this rest of this unpredictable year and rapidly-changing world, I want each of you to know that we will continue our unbroken commitment to each of you.

Warmest regards,

Cameron Penn
OIF Board President
#MeMyselfAndOI
Awareness and Education

In September 2019, the OI Foundation launched its newly redesigned website which allows users to more easily access and share information. As the only voluntary national health organization dedicated to supporting families living with OI, it is our goal to provide information in an easily accessible format for OI community members, medical professionals, and caregivers. The OIF website receives an average of 27,000 unique visitors each month.

With the help of OI community members across the nation, National OI Awareness Week 2019 was officially proclaimed in 16 states across the United States and the OI Foundation’s #SHAREforAWARENESS posts reached over 200,000 people on Facebook. Throughout the week, more than 60 OI community members created Facebook fundraisers to raise more than $9,000 in support of the OI Foundation. The next National OI Awareness Week will take place May 2-9, 2020. National OI Awareness Week is scheduled annually around Wishbone Day (May 6th) – the international day for OI awareness.
OIF Annual Scientific Meeting

On April 10-12, 2019, the OI Foundation gathered over 77 researchers, clinicians, and medical professionals at the 19th Annual OIF Science Meeting in Chicago, IL. Co-chaired by Dr. Joan Marini and Dr. Cathleen Raggio, this important meeting hosted speakers and attendees from the National Institutes of Health (NIH), the Brittle Bone Disorders Consortium (BBDC) sites, and many other centers and labs from across the continent. Speakers presented data on topics ranging from the Effect of Rapamycin on Bone Mass and Strength in Amish Mice to an Update on treatment of adults with OI. The OI Foundation thanks Dr. Marini and Dr. Raggio for their outstanding work as meeting co-chairs, and the speakers and participants for making this a productive meeting. The OI Foundation especially thanks the Buchbinder Family Foundation who makes the Annual OIF Science Meeting possible every year.

Rare Bone Disease Meetings

The OIF successfully co-sponsored a Rare Bone Disease Working Group meeting at the American Society of Bone and Mineral Research (ASBMR) annual meeting in September 2019. The meeting gathered more than 200 scientists and medical professionals. The meeting was chaired by Dr. Laura Tosi, a member of the OIF’s Medical Advisory Council, and included topics such as Management Pearls to Enhance the Care of Patients with Rare Bone Diseases, New Disease Models, and Clinical Trial Updates.

MEDICAL ADVISORY COUNCIL (continued)

Cathleen Raggio, MD
• Hospital for Special Surgery, New York, NY, Orthopedic Surgeon
• Co-Director of the Kathryn O. and Alan C. Greenberg Center for Skeletal Dysplasias
• Principal Investigator, Brittle Bone Disorders Consortium site in New York, NY

Frank Rauch, MD
• Canadian Shriners Hospitals for Children, and McGill University Faculty of Medicine, Montreal, Quebec, Canada
• Director, Biomedical Laboratory, Shriners Hospitals for Children, Montreal, Quebec
• Principal Investigator, Brittle Bone Disorders Consortium site in Montreal Qc, Canada

David Rowe, MD
• University of Connecticut Health Center, Farmington, CT
• Director for Regenerative Medicine and Skeletal Development, University of Connecticut

Sandy Sandhaus, MD, PhD
• University of Colorado School of Medicine, and National Jewish Health, Denver, CO
• Clinical Director at the Alpha-1 Foundation, Denver, CO
• Medical Director and Executive Vice President of AlphaNet
• Medical Director of AlphaNet Canada
• Dr. Sandhaus applies his experience with internal medicine, pulmonary disease, critical care medicine and the biopharmaceutical industry to the study of lung disease

Jay Shapiro, MD
• Dr. Shapiro is Consultant to the Endocrine and Diabetes Dept. at the Walter Reed National Military Medical Center, Bethesda, MD
• Endocrinologist experienced in the treatment of genetic and metabolic bone disorders in children and adults
Regional Conferences

Since its launch in 2015, the OIF’s Regional Conference program, Uniting Unbreakable Spirits, has reached more than 1,100 members of the OI community, half of whom had never attended an OI event before. In 2019, the OIF hosted a record four regional conferences in Chicago, Illinois; Atlanta, Georgia; Sacramento, California; and Wilmington, Delaware.

NIH Brittle Bone Disorders Consortium (BBDC)

The OIF is pleased to announce that the BBDC has been funded for an additional five years! For the past five years the BBDC has been funded as part of the National Institutes of Health’s Rare Diseases Clinical Research Network. The goal of the initiative is to better understand all genetic forms of OI, expand treatment options and train the next generation of physicians and scientists to study OI. As the lead patient advocacy organization partner, the OIF will expand on the outreach to medical professionals and constituents through the various online learning portals housed on the OIF’s website as well as continue to provide opportunities for scientists and OI researchers to connect and collaborate.
Eugene Washington PCORI Engagement Award

In the Spring of 2019, the OIF was approved for a two-year funding award through the Eugene Washington PCORI Engagement Awards program, an initiative of the Patient-Centered Outcomes Research Institute (PCORI). PCORI is an independent, nonprofit organization authorized by Congress in 2010 to fund comparative effectiveness research that will provide patients, their caregivers, and clinicians with the evidence needed to make better-informed health and healthcare decisions. The goal of patient-centered outcomes research (PCOR) for OI is to provide doctors and care providers with information that is relevant to the needs of the OI community. This allows clinicians to provide better care and empowers the OI community to advocate for themselves. Until recently, OI research has been focused on fractures, but the characteristics of OI go beyond bone and includes ear, lungs, eye, and heart problems. PCOR filled this gap in, addressing the needs of the OI community that have previously gone unheard.

Rare Bone Disease TeleECHO Clinic Series

Starting in the summer of 2019, the OIF, in collaboration with the Rare Bone Disease Alliance and OIF Medical Advisory Council member Dr. Laura Tosi of Children’s National Hospital, helped launch the inaugural Rare Bone Disease TeleECHO Clinic Series. ECHO (Extension for Community Healthcare Outcomes) is a digital medical education program that increases a physician’s access to expert knowledge to help them improve their quality of care.

The Rare Bone Disease TeleECHO Clinic Series helps to spread knowledge of rare bone conditions like OI to physicians and medical practitioners through monthly video tele-mentoring sessions. By increasing access to expert medical knowledge for physicians in this cutting edge and cost-effective model, the OIF can help increase the standard of care for people with rare bone conditions.
2019
Members
Joan Ablon, Ph.D.
Cynthia Adanene
Waheeda Afridi
Michael Ahern
Diana Alaniz
Gunner Albitre
Tillie Allen
Paul Andrews
Cindy Assmann
Wanda Bachman
Cathy Bailey
Bruce & Dee Baraw
Jeffrey & Cynthia Barnett
David & Marita Barth
Kathleen Battle
Michael Bellas
Armand & Lisa Benassi
Jerry & Patricia Bennett
Daphna Berman
Norman & Ethel Berman
James & Donna Bernie
Robert Bernstein
Joan Berry
Steven Biagini
Carolyn Bickham
Krystal Bickham
Helga Binder, M.D.
Steven & Cynthia Blind
Bonnie Bolden
Carolyn Boone
Linda Boryczka
Cathy Bowen
LeRoy & Marlene Boyer
John & Nancy Braitmayer
Karen Braitmayer & David Erskine
John & Denise Brill
Edward & Ruth Brosh

Impact Grants
In partnership with the Children’s Brittle Bone Foundation, the Jeanie Coleman Impact Grant program was designed and established to provide funding for items that will significantly improve the quality of life for a person who has OI and who has limited financial resources. In 2019, the OIF was able to fund more than $100,000 of items and services for 24 families. The 2019 grants covered a range of items including hearing aids, a wheelchair, outdoor ramp for access to a home and travel expenses for surgery.

OIF National Information Center
The Foundation provides medically verified information related to OI. Topics range from medical issues such as genetics, diagnosis, and treatments to daily living strategies such as school and employment. Educational materials are available in print and electronically through the OIF website. This past year, OIF staff responded to more than 11,000 direct inquiries for information. OIF also sponsors a network of support groups across the United States. Support group activities provide opportunities for mutual support and increased community awareness. Currently, there are 38 active groups in 32 states.

The OIF continues to improve its web-based information center, www.oif.org/informationcenter, which covers information about OI and issues that affect children and adults, as well as provides online resources for providers who see few OI patients. The site is also host to the OIF podcasts (24 to date) on a variety of topics including; spine issues and basilar invagination in OI; dental; surgical interventions; pregnancy and OI; hearing loss and OI; mental health and the OI patient; treatment options for adults and children; and an overview of the diagnosis and treatment of OI.
Social Networking

The OI Foundation manages three official social networking sites: the OIF Facebook Page – followed by 12,390 Facebook users; the OI Foundation Twitter page (@OIFoundation) - followed by 2,528 Twitter users, and the OIF Instagram page (@oifoundation) – followed by 656 Instagram users. The OIF Forum page, a Facebook group with 10,247 members, is for OI community members to ask questions, share information and experiences, and connect with other OI community members.

International Collaborations

The OIF had the opportunity to be a part of an international meeting on quality of life in November of 2019. The Quality of Life 4 OI Conference held in Amsterdam and coordinated by Care4BrittleBones, was a conference designed to examine ways to measure quality of life in all areas of OI research and care and to improve outcomes for people with OI. The conference was supported by the OIF and included speakers and session leaders from our Medical Advisory Council. OIF CEO Tracy Hart led a session with her colleagues Patricia Osbourne, from the Brittle Bone Society in the UK, and Ingunn Westerheim, President of the OIFE on patient engagement and empowerment across borders. Also this year, the OIF continued promoting the multi-language OIFE Passport, which serves as a handy communication support for individuals affected by OI who are traveling internationally or living abroad. This pamphlet contains the statement that the bearer has OI, a brief explanation of what that means and how a person with OI should be treated in emergency situations – in 22 languages! And lastly, the OIF is helping to plan the OI International Meeting in Sheffield, UK in September 2020.
OIF Patient-led Listening Session at the U.S. Food and Drug Administration (FDA)

In September 2019, the OIF brought together a group of ten OI parents, adults and medical professionals for a patient-led listening session at the FDA. The objectives of the session were to discuss with the FDA the impactful symptoms of OI, the experience of living with OI, as well as the impacts that current treatments have on daily life. The discussion also covered the importance of including endpoints other than fracture in future research. Staff members from several FDA divisions attended the meeting, including; the Office of the Commissioner, Center for Drug Evaluation and Research and the Center for Biologics Evaluation and Research.

Thank you Sharon Mutnick, Dr. Kara Ayers, Erin and Fisher Ortiz, Jessica Finkel Minor, Michael Stewart, Carolyn Tipton and Dr. Laura Tosi for participating on the OIF’s panel during a Listening Session about osteogenesis imperfecta at the U.S. Food and Drug Administration.
New Events

During 2019, the OI Foundation held seven Fine Wines Strong Bones galas, three golf tournaments and the new Medics on the March event. These volunteer led events are critical to funding the programs and resources of the Foundation. Money raised from these fundraising events will help the OI Foundation provide scholarships to the OIF National Conference, create new resources, and fund new research. In 2020, we are looking forward to the return of the OIF’s National Unbreakable Spirit® Walk-n-Wheel to kick-off the National Conference weekend! During fall 2020, the OIF will host its second annual Medics on the March event, where 15 of the OIF’s medical experts will walk 50 miles to raise awareness for OI. This year’s walk will begin in downtown Washington DC and follow the historic C & O Canal Towpath into Maryland. You can follow their progress on www.oif.org/campaign/medicsonthemarch and the OIF’s Facebook page!

The OIF’s first Medics on the March gathered 15 doctors and researchers who walked 50 miles to raise awareness about osteogenesis imperfecta.

Camping out on night one of Medics on the March!

2019 MEMBERS (continued)

Patrick & Bernadette Gleason
Meredith Goldberg
Naomi Goldstein
William Gombeski
John & Paula Goodrich III
John & Barbara Goodrich Jr.
Corinne Gottman
Paul & Dina Granger
Virginia Greene
Stephanie Greenwood
Mae Gribble
Peggy Grigg
Carla Gruber
Juanita Gruenloh
Charla Gualano
Stephen Gudek Sr.
Stephen Guin
Jean Haffenreffer
Donald & Barbara Hagy
Anat Haimovici
James Hall
Jerry Hall
Steven Hardesty
Jim & Kathy Harmer
Brian & Kristen Harpish
Marilee Harrald-Pilz
Gerald Harris, PhD, PE
John & Jennifer Hartnett
Donald Hoffman
Michelle Hofhine
Robert Hogan
Mark and Theresa Holden
Douglas Holleman
Ingrid A. Holm, M.D.
Jason & Emily Holub
Helen Holubnyczyj
Jeanette Hosseini
Ruth Hovden
Wallace Howarter
Fundraising Events  
July 1, 2018 – June 30, 2019

As a small organization for a rare disorder, the OI Foundation relies on volunteer events held around the country to help raise awareness and funds to support the OI Foundation’s mission to improve the lives of people living with osteogenesis imperfecta. Thank you to all the hosts and attendees!

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<thead>
<tr>
<th>$100,000 and greater</th>
<th>$2,500 to $4,999</th>
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<tr>
<td>Facebook Fundraisers</td>
<td>Alle Shea Project Events</td>
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<tr>
<td>Fine Wines Strong Bones DC</td>
<td>OI Carnival</td>
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<td>OI Golf Classic</td>
<td>Atkinson Yard Sale</td>
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<td>Strong Bones Gala Houston</td>
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<tr>
<td>Fine Wines Reception Naples</td>
<td>Dogfish Head Alehouse Fundraiser</td>
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<td>Strong Bones Gala Boston</td>
<td>Jacaranda Golf Outing</td>
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<td>Medics on the March</td>
<td>Ludwig Fundraiser</td>
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<td>National Unbreakable Spirit®</td>
<td>Paint for OI</td>
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<td>Walk-n-Wheel Baltimore</td>
<td>Sacks Family Events</td>
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<td>Riley’s Gathering Place Golf</td>
<td>Sweat for Sammie</td>
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<td>Beef &amp; Brew for Better Bones</td>
<td>Eccleshall Fundraiser</td>
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<td>Blue Jeans for Better Bones</td>
<td>Elkton United Church</td>
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<td>Bone China Tea</td>
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<td>Strong Bones Gala Tampa</td>
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<th>$5,000 to $9,999</th>
<th>$100 to $499</th>
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<tr>
<td>Alameda FOE Dinner Dance</td>
<td>DQ Operation Brain Freeze</td>
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<tr>
<td>Batting For Addison</td>
<td>Florida Panthers OI Nights</td>
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<tr>
<td>Fine Wines Reception NYC</td>
<td>Potorski Dinner</td>
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<td>Last Break Dance</td>
<td>Tri for OI</td>
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<td>Miracle Michael Golf</td>
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REVENUES FY2019

CONTRIBUTIONS AND GRANTS $695,369
EVENT REVENUE $679,343
INVESTMENT INCOME $177,182
DIRECT MAIL REVENUE $358,085
CONFERENCE REVENUE $111,670
PRODUCT SALES, LESS COST OF GOODS SOLD $2,673
IN-KIND CONTRIBUTIONS $47,346
TOTAL SUPPORT AND REVENUE $2,089,668

DISTRIBUTION FY2019

PROGRAM SERVICES (INCLUDING RESEARCH)
PERCENTAGE 70%
AMOUNT $1,471,938

FUNDRAISING
PERCENTAGE 10%
AMOUNT $219,207

ADMINISTRATION & GENERAL
PERCENTAGE 20%
AMOUNT $418,605

TOTAL EXPENSES
PERCENTAGE 100%
AMOUNT $2,109,750

TOTAL NET ASSETS $1,496,703
2019 Members (continued)

Elizabeth Howe
Susan Howland
Larry & Pat Hufford
Paul Hurst
Faith Hutin
Harvey & Judy Irvin
Araya Isaacson
Kevin & Deborah Isensee
Dwight & Evangeline Jaggar
The Jane M. And Bruce P. Robert
Charitable Foundation
Jean Jasinski PhD
Eamon & Mary Jo Jennings
Dennis & Jean Johnson
Jennifer Jolivette
David & Mary Kahn
Andrew & Keri Kendall
Tara Keppel
Carol Kessler
Edward & Dev Kiedaisch
P.J. & Jennifer Kimball
Thomas & Sandra King
Tina Kingery
Carl & Mary Knapp
Marie Konopacky
Joseph Korfhage
Lea-Rachel Kosnik, Ph.D.
Gloria Kowbel
Christine Koziol
Wm. Kroener
Paul Kuester
Nancy L'Estrange
Daniel LaBarge
Mary Lane & Philip Cheney
Larry & Deborah Lapkin
Deborah Lecours
Brian & Kristin Lenahan
Elsie Leon
Peter & Cheryl Lewis
Lawrence Lindvig

2019 United Way/CFC/
Employee Workplace Giving/
Online Charity Giving

Abbvie Employee
Engagement Fund
Adobe
Albertsons Safeway
Allstate-The Giving Campaign
Amazon Smile
America’s Charities
American Express Company
Employee Giving
American Express Foundation
Mr. Andrew Anaya
Andersen Corporation
Mr. Christopher Assmann
AT&T United Way/Employee
Giving Campaign
Automatic Data Processing, Inc
Mr. Brian Baldwin
Bank of America Charitable
Foundation, Inc.
Bank of America United Way
Campaign
Benevity’s Giving Tuesday
$100,000 Charity Giveaway
Bergen County’s United Way
Best Buy Employee
Giving Program
Ms. Lori Boatright
Boeing Employee Good
Neighbor Fund
Ms. Debra Brace
Bright Funds Foundation
Lane & Carol Brundieck
The CarMax Foundation
Mr. Edward Carrasco
Chevron Workplace Giving
Community Health Charities
Ms. Suzanne Davison
Mr. Derek Dees
Dun & Bradstreet
Mr. Michael Eggertson
Exelon Corporation
FHLBank Topeka
FMware Foundation
Mr. Scott Georgetti
Give With Liberty
Michael & Carol Goebel
Goldman Sachs Gives
Ms. Stephanie Goode
Frederick & Jenan Hicks
Intel Foundation
Ms. Joanne Jackson
Mr. Robert Johnson
Johnson & Johnson Family of
Companies
JPMorgan Chase Foundation
Mr. Joel Keith
Kimberly-Clark Foundation
Ms. Carol Kirchhoff
KLA Foundation
Liberty Mutual Group, Inc.
Douglas & Michelle Major
Fred & Jean Major
Ms. Erin R. Mantz
Mr. Chase Marino
Marta Employees Charity Club
Mr. Joseph Martin
Ms. Melissa McConnell
Medical Research Agnc. of
America (CFC)
MidAmerican Energy Foundation
Saurabh Mittal
MRAA-CFC
National Instruments
Workplace Giving
Neighbor to Nation
Network for Good
Mr. Joe Newman
Nextera Energy
Mr. Troy Nordberg
John & Lindsay Painter
Palo Alto Networks
Ms. Karen Pellicci
PepsiCo Foundation
Pfizer Foundation Matching
Gifts Program
Mr. Robert Pizza
Pledgeling Foundation
Price Waterhouse Co
UNITED WAY, COMBINED FEDERAL CAMPAIGN, WORKPLACE & ONLINE CHARITY GIVING (continued)

Ms. Janice Puckett
Randstad Giving
Mr. Charles Rhoades
Mr. Christopher Riley
Roche Diagnostics
Mr. Patrick E. Ronan
Sanofi US Charity Gift Program
Mr. Kevin Schlott
Ms. Christina L. Schrier
Ms. Kayleen Schultz
State Employees’ Community Campaign
Jerome & Sandy Strubhart
Thrivent Funds
TIAA Charitable
Ms. Diane M. Tierney
United Airlines
United Way
United Way Mesa
United Way of AL
United Way of Central Maryland
United Way of Central Texas
United Way of Dane County
United Way of Delaware
United Way of Metropolitan Chicago
United Way of San Diego County
United way of SD
United Way of Tucson & Southern AZ
United Way of West Central Connecticut
United Way of Yellowstone
United Way Truist
UnitedHealth Group Employee Giving
US Bank Foundation
VMware Foundation
Walmart
Waters Corporation
Wells Fargo Community Support Campaign
Mr. Jeffrey Werner
Mr. David Whelan
Mr. Nicholas Whitmore
Ms. Tessa Widman
Robin Wolfecone
Ms. Regan Young
Kenneth & Carol Hollar Zwick
Anthony Robert & Christie Zyskowski

2019 MEMBERS (continued)

Michael & Caren Loguercio
Carlos Long
John & Mandi Lowery
Brian J. Mach
Frederick Maloney
Betsy & Bob Manela
Wallace & Deborah Mangum
Edward & Alice Margolis
Jean Martin
Cristina Martinez
Charlene Maus
Sally McElroy
Thomas McGarry
Betsy Ann McGee
James McGowan
Rita McGrinder
Lisa McIlwain
Charles McKay, M.D.
Caroline McLain
Dick & Margie McLaughlin
Peter & Kathleen McMenamin
Bill McNutt
Mary Merz
Martha Ann Messinger
Gene Messner
Nancy Miller
Christina Mills
Abdul Mohammed
Paula & Tony Moniz
Shirley Moody
Mike Clark & Sarah Morian
Kathryn Morrison
Pat Morse
Richard & Kathie Morse
Gene & Beth Mulroy
Mark Papich & Marcy Murphy
Jonathan & Sharon Mutnick
Helen Nahrstedt
Joseph & Charlene Nanista
### Honorariums 2019

**January 1, 2019 – December 31, 2019**

Lucas Robson Alfonso  
Corrine Merzeraud

Nicholas Atkinson  
Kathleen Allen  
Kim Atkinson  
Carol Casten  
Frank & Robin Falkow  
Erik & Stephanie Molden

Mr. & Mrs. Bain, Matthew & Grace  
Sylvia Leaf

Cynthia Barger  
Kerri Reilly

Ruth Barringer  
W.C. and Evelyn Barringer  
Jane Douglass

Gwena Claire Bateman  
Ron & Karen Bateman

Claude Battaglia  
Betty Battaglia

Tony Benish  
Daniel McHugh

Luis Bermeo  
The CarMax Foundation

Ms. Bobbie, John Strelow & Marylou  
Sylvia Leaf

Karen Braimayer  
The Karina Foundation

Mr. & Mrs. Evan Brook, Bailey, Dylan & Pets  
Sylvia Leaf

Mr. & Mrs. William Brown & Family  
Sylvia Leaf

Alec Cabacungan  
Chris Chulos  
Jay & Barbara Wittenbaum

Jodi Carp  
Sylvia Leaf

Mr. & Mrs. Eric Casem, Children & Family  
Sylvia Leaf

Mr. & Mrs. Andrew Chapkis  
Sylvia Leaf

Miss McKenzie Coan  
Mike & Steph Maley

Audrey Cook  
Charles Cook

Dr. Joanne Cyganowski  
Alexander Hogan

Frans Deblois & Michelle Ronan  
Mt. Airy Matress

Dr. Michael D’Egidio & Family  
Sylvia Leaf

Alanah Demko  
Wanda Pfeiffer  
Benjamin Stafford  
Donald & Kathleen Woikoski

Julian DiGeorge (2nd Birthday)  
Courtney Gaines  
Asher Griffin  
Holly Moher  
Bernice Russo  
Sandra Stimpson-Russo  
Jessica Weiner

Tim Dombro  
Joseph Musker  
Jeffrey Gerber & Shannon Rudisill

Dorian & Family  
Sylvia Leaf

Sara & Earl Dyke  
Trey Lipsitz

Jane & Jim Early  
Lawrence Leventhal  
Michael & Sara McFadden

Jacob Everett  
Doris Everett  
Robert & Debra Harwell

Jay & Denise Gagne  
Karen Gagne

Kelly Gagnon  
KLA Foundation

Miss Audrey Garfield  
David & Jacqueline Sices

Deborah Glickman  
Mitchell & Maxine Selig

Miss Skylar Goldberg  
Joel Feldman

Conner Goldhammer  
Kent & Lesa Tegeler

Stephanie Greenwood  
Richard Wharton

Anna Gualano  
Sammie Leach  
Jan & William Stowe

Jacob Grys  
Elizabeth Grys

Flora Hand  
Jo Ann Berkenbush

Chris Harkman  
Derek Dees

Harper  
Thomas & Johanna McCarthy  
Nick & Leslie Ream

---

**2019 MEMBERS (continued)**

- Lynn & Erah Newman
- Diane Nier Spurgeon
- Raymond & Maureen Noeth
- Scott and Christie Nohre
- Joseph & Sharon Nucci
- Claudia & Sean O’Brien
- Elizabeth O’Conor
- John & Sheryl O’Meara
- Earl & Betty Olmscheid
- Andres Osornio Ocaranza
- Ann & Borden Painter
- Mark Palen
- James Palestrini
- Frank & Maureen Palestrini Jr.
- Marybeth Parke Robbins
- Stephen Patterson
- Jeffrey & Angela Paul
- Eugene Paykel
- Cameron & Lauren Penn
- Nancy Pesce
- Nick & Karlee Petruic
- Raymond Phillips
- Audrey Phillips
- Judy Picciallo
- Dick & Karen Pinney
- Henry & Rose Piper
- Joseph & Marjorie Polerecky
- Don & Jacquelyn Powell
- Melinda & Daniel Preston
- Dennis & Kathi Primus
- Catherine Ann Pusateri
- Melanie Rak M.D.
- Irene Ratto
- Roger Gray Renegar
- Gail Ann Renetty
- Edward & Roxanne Rios
- Conrad & Mary Rivers
- Kathryn Roberson
- Guy & Meisie Roberts
HONORARİUMS 2019 (continued)

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Tanner Adam Hirniak
Stefan Hirniak
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Sharon Lee Envall
Walter Holman
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Sharon Ferguson
Emma Johnston
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Patricia McPherson
Jackson Koal Jones
Jeff & Kelly Povroznik Jones
Kathy Jones
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Karsen
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Imran Katchi
Susan Mathai & Chris Sigakis
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Sylvia Leaf
Corey Leaf
Sylvia Leaf
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Sylvia Leaf
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Sylvia Leaf
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   Consolidated School-Quarters
   For Causes
   Marcone Maia
Amber Robson
   Amanda Woloshyn
Dr. Melissa Rose
   Barrie Fligelman
   Sherrie Krangel
Rossi Family
   Bob & Leslie Turner
P.A. Russo
   Robert Levendakes
Miss Jaden Sacks
   Hunter & Stefanie Dec
   Horace Mann School-Fifth Grade Class
Allison Schwartz
   Christy Barry
Lawrence Schwartz (Birthday)
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Leah Sensenbrenner
   Lisa O’Connell
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   Dwight & Evangeline Jaggard
Noah Shapiro
   Geoffrey Stern
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   Marie Moarek
Mrs. C.Z. Sherman, cousins, caretakers & pets
   Sylvia Leaf
Matthew Smith
   Candace Godin
Arianna Snow
   Michael and Nancy Raymond
Dani Spann (Birthday)
   Kevin & Deborah Isensee
   Janet Kilkelly
Kevin Solorio
   Lance Kirchner
Addison Stephens
   Batting For Addison
Shannon Strauch
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Jenny Stup & Joey Reed
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Molly Sullivan
   Kacey Ragsdale
Alexi & Avery Suri
   Mary Peterson-Suri
Barbara Taylor
   The Society of Irish Women
Marty Wagner
   JC & Neil Milner
Skylar Walsh (Birthday)
   Julia Lopez
   Alanna Rowan
   T.J. Schuck
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   Christopher Batterby
Holly Weilbrenner
   Martha Johnston
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   Sylvia Leaf
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   Sylvia Leaf
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January 1, 2019 – December 31, 2019

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July 1, 2018 to June 30, 2019

The OI Foundation gratefully acknowledges the generous individuals, foundations and corporations whose support helps us respond to thousands of inquiries each year, produce new information resources, fund research, organize National and Regional Conferences and facilitate the operation of support groups across the country. This list represents donations made from July 1, 2018 to June 30, 2019. We regret any error or omission and ask that corrections be brought to our attention.

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<thead>
<tr>
<th>Name</th>
<th>Organization/Role</th>
</tr>
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<tbody>
<tr>
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<td>Fibrous Dysplasia Foundation</td>
</tr>
<tr>
<td>Melissa &amp; Doug Fordyce</td>
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<tr>
<td>Jeanne Franzone</td>
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<td>Don H. Gardner</td>
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<td>Mary Gardner</td>
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<td>James Geisman</td>
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<td>Douglas Gibson</td>
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<td>Michael &amp; Carol Goebel</td>
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<td>Stephanie Goode</td>
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<td>Goodnow Real Estate Service</td>
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<td>Christine Gorman</td>
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<td>Granite State Glass</td>
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