



2022 ANNUAL REPORT



July 1, 2021 - June 30, 2022

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Our Mission



The Alpha-1 Foundation is committed to finding a cure for Alpha-1 Antitrypsin Deficiency (Alpha-1) and to improving the lives of people affected by Alpha-1 worldwide. Our initiatives in four main areas- Research, Education/Outreach, Awareness/Fundraising, and Public Policy/Advocacy are paving the way for important discoveries, new therapies, and increased awareness and understanding of this rare genetic condition.

Our accomplishments and accreditations reflect our unparalleled commitment to medical research at its finest: we have invested over \$91 million to support Alpha-1 research and programs at 123 institutions in North America, Europe, South America, the Middle East, and Australia. These innovative, impactful projects have built a solid infrastructure for promoting ongoing research and the development of new therapies. We have also fostered collaborations with investigators throughout the United States and Europe, working closely with the pharmaceutical industry, the National Institutes of Health (NIH), the U.S. Food and Drug Administration (FDA), and Alpha-1 patients and families.

There are more advancements on the horizon than ever before and our commitment to fund vital research needed to make breakthroughs in Alpha-1 is greater than ever. The Alpha-1 Research Registry is growing with enthusiastic Alphas who are ready to advance the Foundation's research goals.

During this fiscal year, we welcomed a new President and Chief Executive Officer, Scott Santarella. Scott joined the Foundation in September 2021, with over 25 years of experience as a senior executive. Scott's vision for the Alpha-1 Foundation included the continuation of achievement through forward momentum, the optimism of forward-thinking through innovation, and the advancement of impact through forward progress.

"Now is the time to accelerate our efforts across the spectrum and maximize our resources to expedite progress toward our ultimate goal –a cure," said Santarella. "It is here that patients play the most vital role. The key to solving the Alpha-1 puzzle will be found in the patient."

We utilized innovative and creative ways to connect the community through this part of the pandemic. Hybrid event options transformed the way we served the community in all of our programs. A COVID event grid was created to provide a framework on how to resume in-person events while keeping the safety of Alphas a top priority. We were able to host the annual Alpha-1 National Conference in a hybrid fashion in 3 cities across the United States, assuming a unique blend of online presentations using our robust virtual environment with interactive features. This was just the first step in our path to a "new normal."

Alpha-1 Outreach & Education

Our website, www.alpha1.org, is a clearinghouse for information on Alpha-1 providing educational resources, highlighting news and upcoming events, and creating opportunities for virtual connection within the global Alpha-1 community. We maintain content targeted toward several different audiences, including newly diagnosed Alphas, family members and caregivers, healthcare providers, researchers, and industry partners. The website received over 484,000 visits from over 234,000 unique visitors in FY 2022. The Alpha-1 Foundation's website is most often the first contact that newly diagnosed Alphas and their family members have with the Foundation and in many cases, their first source of information about Alpha-1 Antitrypsin Deficiency.

We connect with the Alpha-1 community across multiple social media platforms. In addition to promoting new Foundation initiatives, community education events, and fundraising campaigns, our presence on Facebook, Instagram, and Twitter has directly increased traffic to our website, where users can access the information they need.

The monthly Community Currents electronic newsletter keeps the Alpha-1 community and volunteer leadership updated on all Alpha-1-related news, upcoming events, research, programs, fundraising activities, and the Alpha-1 Research Registry.

The Alpha-1-to-One Magazine is a valued resource produced three times a year for the Alpha-1 community offering practical advice, personal experiences, and pertinent news for people touched by Alpha-1. In FY 2022, we distributed 30,000 printed copies of each issue, including bundles to Clinical Resource Centers (CRCs) and healthcare providers; each issue is also posted to a digital publishing platform on our website, where it is accessed by nearly 2,000 more visitors a year. In the Fall issue of the magazine "All In For Alpha-1" spotlighted the 2021 Virtual Alpha-1 National Conference. In the Winter issue "Riding for a Reason" focused on the 25th anniversary of Escape to the Cape Bike Trek. The Spring issue's theme "Forward Momentum" welcomed Scott Santarella as the new President and CEO of the Alpha-1 Foundation.



Alpha-1 Advocates

Our advocates are a voice to improve outcomes for Alpha-1 patients. Together, we are fighting to improve the fact that Alpha-1 is under-recognized, under-diagnosed, and unknown to most people by ensuring our community's priorities are factored into public policy and legislation. Our advocacy program targets policymakers, regulatory agencies, the public, and the media with campaigns to promote research and raise awareness of the challenges faced by Alpha-1 patients and care providers. Our FY 2022 policy priorities included access to care, medical research, therapeutic development, screening and detection, federal and state funding, blood product safety, and education and awareness.

We worked with Congress to find a solution to the patient access problem facing Medicare beneficiaries who are not able to obtain a home infusion. In March 2022, the John W. Walsh Home Infusion Act introduced a permanent benefit for Alphas to receive augmentation therapy in the home setting through Medicare Part B. This process is ongoing, and we continue to work diligently with advocates and legislators to meet the needs of the Alpha-1 community.

We work strategically with federal agencies, memberships, and coalitions—focusing on the issues that matter most to Alphas—to inform legislators of the burdens of the patient community and to advance our public policy agenda for the benefit of the Alpha-1 community.

Additionally, working closely with the U.S. Food and Drug Administration (FDA), Critical Path Institute (C-Path), and National Institutes of Health (NIH).

- November 10, 2021, Fall Virtual Lobby Day
- Fall Advocacy Training Series
- 2022 Rare Disease Day



Alpha-1 Education Days



FY 2022 Alpha-1 Education Days:

October 2, 2021

November 6, 2021

February 12, 2022

March 26, 2022

April 30, 2022

VIRTUAL ALPHA-1 EDUCATION DAY

Saturday, February 12th
10:30 am - 2:30 pm ET

[REGISTER NOW](#)

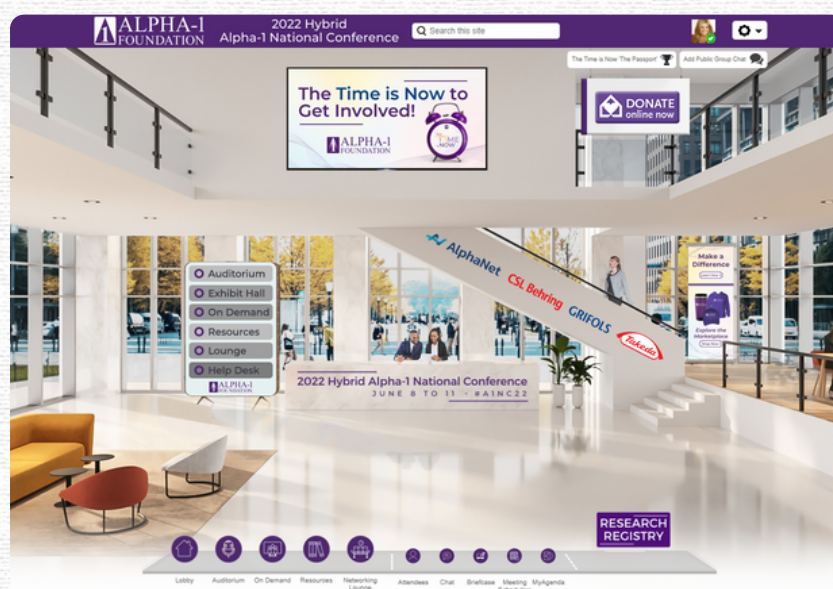
The Alpha-1 National Education Series is comprised of regional Alpha-1 Education Days for Alphas, their families, and caregivers. These one-day educational programs provide access to current medical information, timely topics affecting Alphas, resources from leading Alpha-1 experts, and opportunities to connect with the Alpha-1 community.

In 2022, we continued our use of a virtual event platform, enabling those who were unable to attend an Alpha-1 Education Day in person to participate and connect with the Alpha-1 community. Extending our education series into the virtual space has also created community-building and engagement opportunities through more direct access to social media, polls, and surveys.

More than 2,300 people registered for the FY 2022 Alpha-1 Education Days.

Alpha-1 National Conference

As the largest gathering of the Alpha-1 community, the National Conference provides important opportunities for networking, education, advocacy, and industry publicity. Over the course of four days, attendees had access to scientific presentations, meet-and-greet and Q&A sessions with Alpha-1 experts, and other events hosted by sponsors.



After two years of completely virtual programming, we were proud to announce that the first-ever Hybrid Alpha-1 National Conference was held on June 2022. The format of the Hybrid Alpha-1 National Conference was meticulously crafted in a way that sought to combine the best aspects of the traditional in-person format while also continuing to provide the accessibility and flexibility offered by a virtual format. Three sites were selected to represent major regions in the United States: New York City, Miami, and Oregon. Each site is also home to one of the Foundation's Clinical Resource Centers (CRCs), specialized locations that give patients access to doctors who are familiar with Alpha-1. The four-day agenda was packed with engagement and interaction, including three Platinum Sponsor live events, three phenomenal keynote speakers, eighteen open forum question and answer breakout sessions, three live panel sessions, and on-demand education sessions throughout the conference. Presentations from the conference were available to view on-demand on the Foundation's Vimeo channel.

Stats: 1,195 registered to attend

667 attended from 47 states and 10 countries

510 patients and family members

53 medical professionals

23% of first-time attendees



Meetings and Conferences

Scientific meetings and conferences organized by the Alpha-1 Foundation generate awareness of Alpha-1, identify new areas of research interest, and establish the Foundation's credibility within the rare disease community as a resource for serious research, education, and advocacy.



The 23rd Gordon L. Snider Critical Issues Workshop entitled "Liver Biomarkers and the Path to Alpha-1 Liver" was held on November 19, 2021. The overall goal of the workshop was to bring together researchers, clinicians, and industry representatives to focus on identifying biomarkers to support the path to drug approval in liver disease. Co-chaired by Virginia Clark, MD, MS from the University of Florida and Pavel Strnad, MD from University Hospital/Aachen.

"Outstanding speakers and the co-chairs brought into sharp focus both challenges and pathways to drug approval in a new era of drugs to treat Alpha-1 Antitrypsin Deficiency related liver disease," stated Dr. Mark Brantly, Alpha-1 Foundation Scientific Director.

We reach thousands of physicians, academic researchers, and industry partners each year through our exhibitions at major national and international medical conferences, where we share insights into new technologies, research findings, and priority research areas.

The CRC Forum took place on June 30, 2022, and was an opportunity to engage Alpha-1 physician specialists and leaders in the field of Alpha-1 research and the patient community to learn about new Foundation programs, research initiatives, and updates from industry sponsors.

Alpha-1 Awareness

Annual campaigns help facilitate community support and improved approaches to care.

#Alpha1Awareness- November is Alpha-1 Awareness Month: 30 days of awareness-focused initiatives, through which the Alpha-1 community shares stories and resources while raising funds to support the search for a cure. A dedicated webpage, online training, digital resources, media outreach, community events, and a social media calendar are available throughout November specifically for awareness. The 2021 Alpha-1 Virtual Walks allowed anyone who wanted to make a difference in the Alpha-1 community to contribute in their own style, time, and pace. Over 700 participants tracked miles on STRAVA and raised over \$72,000 for Alpha-1 in just one month.



#RareDiseaseDay - During the month of February, we collaborated with the National Organization for Rare Disorders (NORD) to spread awareness of Alpha-1 on Rare Disease Day. The goal was to get patients interested and involved in rare disease research.

Family Testing

Alpha-1 Coded Testing (ACT) Study has enrolled over 47,000 individuals since collecting data on the impact of at-home confidential genetic testing. The ACT study is a confidential testing program aimed at those who are at risk of Alpha-1, it provides an opportunity for individuals and their family members to be tested for Alpha-1 in the privacy of their homes, without the concern for health and life insurance issues associated with a diagnosed pre-existing condition.



The study screened 1,897 people in FY 2022 and 52% of those tested were identified as carriers of Alpha-1 and 5% were diagnosed as ZZ Alphas.

Community Fundraisers

Our Building Friends for a Cure (BFC) community fundraisers are designed to increase awareness and promote research. Support from the Alpha-1 community is a critical component of our goal to raise awareness of the disease and treatment options. Through the BFC program, we steward relationships with donors to boost awareness of Alpha-1 and get the community involved in working towards a cure.



Building Friends for a Cure consisted of a series of community fundraising events this year in support of Alpha-1 research, including:

- Riding for a Reason
- Escape to the Cape
- Alpha-1 Virtual Walks
- Celtic Connection
- Walks, Run, and Marathons
- Alpha Hours
- Climbs and Hikes
- Video Game Streaming Marathons
- Golf Tournaments

Research Grant Awards

The Alpha-1 Foundation directed more than \$2 million in peer-reviewed research grants in FY 2022. These grants support innovative research spanning the spectrum of scientific discovery. The Foundation prides itself on being the #1 funders in Alpha-1 research having invested over \$90 million at 128 institutions in 14 countries worldwide. Research funded by the Alpha-1 Foundation has improved our understanding of the mechanisms underlying the clinical manifestations of Alpha-1 and identified novel drug targets likely to lead to new therapeutic solutions.

The Foundation is proud to announce its in-cycle grant awardees for 2022, with 11 grant recipients receiving over \$2 million for their projects. Of this year's grant recipients' fields of study, 45.5 % were lung focused, 36.4 % were liver-focused and 18.2 % were focused in both areas. The research was focused 72.7% on basic research and 27.3 on translational research, keeping a well-rounded portfolio.



Brian Hobb, MD
Brigham and
Women's Hospital

*"Inflammatory Protein Biomarkers of
Reduced Lung Function in AATD"*



**Joseph
Kaserman, MD**
Boston University

*"Mechanisms of Hepatic
Heterogeneity and ATF6 Mediated
Metabolic Dysfunction in AATD"*



**Jarrett
Morrow, PhD**
Brigham and
Women's Hospital

*"Blood and Lung Microbiome in
Alpha-1 Antitrypsin Deficiency"*

*"Mitochondrial Dysfunction in Alpha-1
Antitrypsin Deficiency Associated
Liver Disease"*



**Pasquale
Piccolo, PhD**
Telethon Institute of
Genetics and Medicine

Research Grant Awards



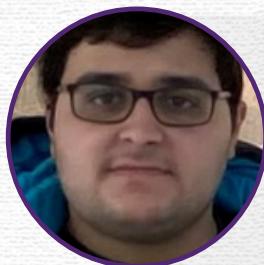
Francesco
Annunziata, PhD
Telethon Institute
of Genetics and
Medicine (TIGEM)
Naples, Italy

*"Identification and Modulation of
Local and Systemic Environmental
Factors in AATD"*

*"Severity and progression of airway
dropout in AAT knockout transgenic
ferret model of COPD"*



Shah
Hussain, PhD
University of
Alabama at
Birmingham



George
Sophocleous, PhD
University College
London

*"Structural and biophysical dissection
of the alpha-1 antitrypsin
polymerization pathway"*

*"Modulating the Co-Translational
Misfolding and Polymerization of
Antitrypsin"*



Lisa
Carita, PhD, BSc
University College
London



Tomas
Carroll, PhD
Royal College of
Surgeons in Ireland

*"A Family Affair- Clarifying the Risk of
Lung Disease in ZZ AATD"*

*"Longitudinal Evaluation of Lung
Disease Progression in PiMZ Patients"*



Monica
Goldklang, MD
Columbia
University Irving
Medical Center



Valerie Gouon-
Evans, PhD
Boston Medical
Center

*"Engineered and Edited Patient-
Derived iPSC for AATD-Associated
Liver Disease Cell Therapy"*

Statement of Financial Position

Current Assets	2022	2021
Cash and cash equivalents	\$12,377,362	\$8,614,083
Investment securities	\$16,836,191	\$18,369,431
Accounts receivable	\$675,117	\$982,721
Other current assets	\$167,500	\$21,728
Total Current Assets	\$30,056,170	\$27,987,963
Investment securities- non-current	\$5,039,394	\$6,432,525
Investment in TAP	\$363,069	\$136,115
Property and equipment, net	\$12,983	\$21,749
Other assets	\$25,000	\$25,134
Total assets	\$35,496,616	\$34,603,486
Current Liabilities	2022	2021
Accounts payable and accrued expenses	\$1,908,720	\$1,951,672
Refundable advances	\$505,000	\$130,000
Deferred revenue	\$210,000	\$286,000
Total current liabilities	\$2,623,720	\$2,367,672
Net Assets	2022	2021
Without donor restrictions	\$28,669,884	\$29,114,292
With donor restrictions	\$3,303,012	\$3,121,522
Total net assets	\$32,872,896	\$32,235,814
Total liabilities and net assets	\$35,496,616	\$34,603,466

Statement of Activities & Changes in Net Assets

Changes in net assets without donor restrictions	2022	2021
Contributions, bequests and grant revenues	\$6,211,034	\$5,385,961
Donated securities	-	\$10,230
Fundraising revenues	\$49,150	\$52,684
Net realized and unrealized investment (losses)/gains	(\$3,267,543)	\$4,778,951
Interest and dividends	\$465,634	\$460,729
In-kind rental income	\$210,000	\$200,000
Net assets released from restriction	\$2,447,606	\$1,816,820
Other Income	\$374,788	\$248,788
Total support/revenue without donor restrictions	\$6,490,669	\$12,954,163
Expenses	2022	2021
Research and Clinical Studies	\$3,450,654	\$3,713,401
Patient Education	\$1,197,628	\$1,075,527
Patient Advocacy	\$429,678	\$288,876
Total program services	\$5,077,960	\$5,077,804
Administrative and general	\$555,657	\$492,150
Fund raising	\$628,414	\$550,275
Total supporting services	\$1,184,071	\$1,042,425
Total expenses	\$6,262,031	\$6,120,229
Change attributed to income (loss) from TAP	\$226,954	\$1,427
Increase in net assets without donor restrictions	\$455,592	\$6,835,361
Changes in net assets without donor restrictions	2022	2021
Contributions, bequests and grant revenues	\$2,629,096	\$2,343,702
Net assets released from restriction	(\$2,447,606)	(\$1,816,820)
Increase in net assets with donor restrictions	\$181,490	\$526,882
Increase in net assets	\$637,082	\$7,362,243
Net assets, beginning of year	\$32,235,814	\$24,873,571
Net assets, end of year	\$32,872,896	\$32,235,814

Programs and Services

The Alpha-1 Foundation is committed to finding a cure for Alpha-1 Antitrypsin Deficiency (Alpha-1) and improving the lives of people affected by Alpha-1 worldwide. The Alpha-1 Foundation has invested more than \$91 million to support Alpha-1 research and programs at more than 123 institutions in North America, Europe, the Middle East, and Australia.

The Alpha-1 Research Registry is a confidential database of people with Alpha-1 (Alphas) and Alpha-1 carriers. It is a resource for investigators seeking Alphas to participate in clinical trials, surveys, and other scientific and medical activities. For more information, email alpha1registry@alpha1.org or call toll-free 1-877-228-7321 ext. 252. <https://www.alpha1.org/alphas-friends-family/resources/participate-in-research/>.

Alpha-1 Coded Testing provides free, confidential testing administered through a research study that evaluates the perceived risks and benefits of genetic testing. For more information, email alpha1lab@alphaone.ufl.edu or call toll-free 1-855-476-1227 <https://www.alpha1.org/healthcare-providers/testing-and-treatment/alpha-1-coded-testing-study/>.

The Support Network is comprised of over 80 support groups nationwide that provide guidance and education to Alphas and family members, create awareness in local communities, and advocate for national and state issues that affect Alphas. Four Virtual Support Groups provide a forum for topics such as Alpha-1 Kids, Pre & Post Transplant issues, Caregivers, and Timely Topics. To find a support group near you, visit alf.org/support-groups.

The Patient Information Line 1-800-245-6809 is available free of charge to anyone affected by Alpha-1 and provides support and answers to topics such as Alpha-1 testing, emotional impact, and physician and support group referrals.

The Genetic Counseling Program 1-855-476-1227, based at the University of Florida, is a free and confidential service that provides direct contact with a genetic counseling service to provide information on the risks and benefits of testing for Alpha-1, interpreting test results, understanding the genetics of Alpha-1, and other issues associated with having a genetic illness. <https://www.alpha1.org/alphas-friends-family/support/genetic-counseling/>

The Peer Guide Program 1-877-346-3212 connects newly diagnosed Alphas with other Alphas who have a similar set of circumstances to provide emotional support, discuss resources available, and assist in making choices about health.

Programs and Services

The Oxygen Fund provides oxygen equipment for Alphas in financial need to travel to physicians, hospitals, Alpha-1 educational events, and other activities. For more information contact Joy Collins at jcollins@alpha1.org or call 1-877-228-7321 ext. 251. <https://www.alpha1.org/alphas-friends-family/resources/oxygen-resources/>

Patient Education Programs include an annual National Education Conference which brings together over 600 Alphas, caregivers, industry representatives, clinicians, allied healthcare workers, and scientists to discuss a wide range of Alpha-1-related topics. For more information, visit alf.org/alphalconference. Additionally, a series of Alpha-1 Education Days are held each year in various cities throughout the U.S. to bring patient education to a regional audience. For more information, visit alf.org/educationdays. Recordings of these events are available via E-Education at alf.org/education-videos.

Building Friends for a Cure is a program designed to nurture ongoing friendships in the Alpha-1 community. The goal is to increase awareness and raise funds for research and related programs which will ultimately lead to a cure for Alpha-1. By becoming involved in fundraising efforts, such as virtual events, letter-writing campaigns, and other social and sporting events, you are making a difference. Visit alf.org/building-friends, contact amcbride@alpha1.org, or call 1-877-228-7321 ext. 233 for more information.

The Alpha-1 Kids Program is geared to the special needs and concerns of parents of children with Alpha-1. A committee of volunteer parents helps direct the program, which consists of a hotline, age-appropriate informational books, a virtual support group, and a parent peer guide program. For more information, visit alf.org/alpha1-kids or contact Cathey Henderson at chenderson@alpha1.org.

Educational resources about Alpha-1 and related topics are available for Alphas, their families, caregivers, and healthcare providers. Informational brochures such as 'What is Alpha-1?', 'Am I an Alpha Carrier?', 'The Liver and Alpha-1', and 'It's All In The Family: Family Testing' are available to download and order through our website, alf.org/a1-publications, or by calling toll-free 1-877-228-7321 ext. 251. The Alpha-1 to One Magazine is published three times a year and includes timely articles about Alpha-1 research and Alpha life, as well as answers from Alpha-1 experts to questions from the Alpha-1 Community. Join the mailing list here at alpha1.org/alphas-friends-family/resources/join-mailing-list/ to receive the magazine. Presentations from previous Alpha-1 educational events are available via E-Education at alf.org/education-videos.

Programs and Services

The Foundation's Public Policy Program advocates for the Alpha-1 community by monitoring and influencing legislative and regulatory issues. Primary concerns include stimulating medical research, blood product safety, developing new therapies, screening, and detection, access to care and reimbursement, federal and state funding, education, awareness, and the recognition of the special needs of people with Alpha-1. For more information, contact Jeanne Kushner at jkushner@alpha1.org or call 1-877-228-7321 ext. 204 or visit alf.org/action-alert.

Grants and Awards: The Foundation's peer-reviewed grants program is intended to promote research that will lead to improved health for Alphas and ultimately, find a cure. We offer grants for both basic science and clinical research. For more information, contact Vanessa Valencia at vvalencia@alpha1.org or call 1-877-228-7321 ext. 242.

The Alpha-1 Foundation DNA & Tissue Bank at the University of Florida is the central storage site for DNA and tissue samples from Alphas and other donors. The Bank is a resource for researchers investigating Alpha-1 and other conditions. Researchers should contact Vanessa Valencia at vvalencia@alpha1.org or call 1-877-228-7321 ext. 242. For other information, call toll-free 1-866-284-2708.

The Targeted Detection Program promotes worldwide awareness among medical professionals, the media, and the public, and the identification of Alphas in population groups at risk. For further information, please call 1-877-228-7321 ext. 306.

The Alpha-1 Clinical Resource Center Network is a steadily growing group of centers throughout North America that specialize in patient care and education for those with Alpha-1. Centers also offer other resources for Alphas such as support groups, transplant centers, and pulmonary rehabilitation. Alphas and their physicians are encouraged to contact their regional Clinical Resource Center for information and guidance. To find an Alpha-1 specialist near you, visit alf.org/alphadoc. For more information, contact Vanessa Valencia at vvalencia@alpha1.org or call 1-877-228-7321 ext. 242.

Scientific Meetings, Conferences, Workshops, Working Groups, and Symposia bring scientists together to focus on special topics related to Alpha-1, advance knowledge of the genetic condition, and work toward new therapies and a cure. For more information, contact Adriana De Arce at adearce@alpha1.org or call 1-877-228-7321 ext. 269.

Programs and Services

The Alpha-1 Global Initiative works on a global, national, and local level to enhance care and support for all those affected by Alpha-1. The Foundation is committed to providing reliable resources and information worldwide. As part of the Global Alpha-1 Initiative, the Foundation is creating new tools focusing on connecting Alphas all over the world with resources – and with each other. For more information, visit alpha-1global.org, contact Randel Plant at rplant@alpha1.org, or call 1-877-228- 7321 ext. 306.

Access and Reimbursement: The Alpha-1 Foundation is committed to providing the tools and resources necessary to assist Alphas in making informed decisions about their healthcare plan and assistance program selection. Patients with questions or issues concerning access or reimbursement should contact Cathey Henderson at 1-800-245-6809 or chenderson@alpha1.org. <https://www.alpha1.org/alphas-friends-family/resources/assistance-organizations/>

The Clinical Trial Outreach and Education Program is an online tool used to educate Alphas about clinical trials with curated educational materials for Alpha-1. For more information, visit <https://www.alpha1.org/alphas-friends-family/resources/clinical-trials/> or contact Alison Keaveny at akeaveny@alpha1.org or call 1-877-228-7321 ext. 206.

Honor Gallery

The Alpha-1 Foundation's Honor Gallery salutes the individuals, corporations, and foundations that share our commitment to finding a cure for Alpha-1 Antitrypsin Deficiency and to improving the lives of people affected by Alpha-1 worldwide. The Foundation gratefully welcomes several new members who have supported our mission for the first time through their generous contributions.

The Foundation acknowledges the continued support of the many long-time donors who have invested in our mission through contributions, events, and workplace giving. It is with sincere gratitude and appreciation that we recognize our Honor Gallery. The following is a list of donors who contributed \$500 or more between July 1, 2021 and June 30, 2022.

\$2.5 Million or More

AlphaNet, Inc.

\$1 Million - \$2,499,999

CSL Behring, L.L.C.

Grifols

Takeda Pharmaceutical Company

\$500,000 - \$999,999

Estate of James L. Fermoile

\$250,000 - \$499,999

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Honor Gallery

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Helen E. Bueker
Anthony Callea
Erin L. Carr
John M. Carter
Estate of Donald J. Ceppi
Mary M. Chilton
Rosemary B. Chiodo
Jan and Michael Chnupa
Donna A. Clark, R.N.
Kathryn W. and Daniel C. Coffin
Jane-Ellen and Joseph Collins
Joan and *Curts* Cooke
Timothy J. Craig, D.O.
Creating Healthier Communities
Barbara E. and Don E. Davis, Jr.
Connie A. DeWald
James F. Donohue, M.D.
Sally D. and John H. Dorf
EchoStor Technologies- Michael McMillan
Traci and Steven Egly
Erickson Solutions Group, Inc. - Peter R.
Latessa
Evolve Biologics
Susan W. and Donald C. Ferro
Janet and *Kenneth N.* Findley
Louis G. Fox
Anne and *Paul* Friedlander
FrontStream Workplace Platform
Gain Therapeutics
Donna A. and James A. Geurink
Barbara A. Glenn
Monica P. Goldklang, MD
Google, Inc.
Goose the Market- Mollie and Chris Eley
Anne and Gary Greenwood
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Melissa Johnson
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Andrew Kwatinetz
Norma I. Leising
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