Dear Friends

On April 6, 2018, the AЯRE foundation was born to support research that will increase our understanding of the ASXL genes and to improve the treatment of individuals with congenital ASXL mutations.

Families have struggled with extraordinary challenges for decades due to the lack of evidence-based research and significant unknowns about ASXL syndromes.

These syndromes – Bohring-Opitz, Shashi-Pena & Bainbridge-Ropers – are under-diagnosed due to the challenges recognizing the diagnosis, shortage of genetic services, and the cost of genetic testing.

Through research AЯRE aims to improve the longevity and quality of life for these affected individuals and their families.

ARRE key tenets include good stewardship of our resources and transparency.

Our donor summary includes the core focus areas where donations benefit ASXL syndrome patients.

Thank you for making a difference in the lives of these families.

Laura Badmaev
The amount of achievements this organization has had in less than a year since its beginning is nothing short of outstanding! I am so proud to be part of its journey! – anonymous
PRIORITIES

ASXL Registry & Biobank - These are foundational tools based on patient data & samples, which researchers will utilize to study and increase our knowledge of these syndromes.

Research grants - Effective research will bring an understanding of the underlying mechanisms and potential treatments for ASXL syndromes.

Conferences - Bring together professionals and families to deepen our understanding and exchange ideas that will move our mission forward.

ASXL CoE - Establish hub hospitals around the world where many ASXL patients will see specialists who will build their clinical expertise and improve care plans.
“ARRE does so much to help people affected by the ultra-rare ASXL syndromes! They help families connect, help caregivers be better advocates and help bring much-needed awareness to medical professionals.” – Colleen Racciopi
ADVOCACY & COMMUNITY

31 ASXL Symposium Attendees

338 Social Media Followers

104 Newsletter Subscribers
**ADVOCACY & COMMUNITY**

- 8 Research Spotlight Symposium Series
- 2 Campaign Fundraisers
- 6 Family Stories
- 19 Education/Health/Tips
The A ЯRE Foundation means so much to our family. When my granddaughter was born, no one knew anything about Bohring-Opitz Syndrome. The A ЯRE Foundation has given us hope! It funds true research to improve the lives of our children. – Vonnie Iverson S.
Public donations: 8.47%
Private donations: 91.53%

Number of public donations: 64
Average public donation amount: $59.32
Total donations: $44,796.21
We went to the conference sponsored by ARRE and it was wonderful. For the first time we got to hear doctors that are studying these rare diseases and connect with other parents and families. They are doing a great job! – Julie Chandler L.
$31,272.85

- Marketing & Fundraising Expenses (website and credit card donation fees) $934
- Scientific Conference (UCLA, July 2018)* $11,060
- Biobanking $10,944
- Professional Services (lawyer, accountant, incorporation & license fees)** $8,025
- General Expenses $308

COST PER $ RAISED $0.09

* UCLA subsidized the cost of the conference and contributed an additional $13,413 toward the expenses.
** One-time expenses for the set up of the Foundation and required for private donation eligibility
We are thankful for non-profits like ASXL, we hope that with continued research we will know more about how we can help our little ones! Rare diseases, like Shashi-Pena Syndrome-ASXL2 need LOTS of TLC! And of course "ALL" rare disease need TLC! – Michele P.
ARRE will develop a diverse research portfolio that will build on successful studies as well as support innovative investigations with high potential rewards. Key focus areas include:

- Gain or loss of function
- Transcripts expressed postnatally
- Protein expression
- Pathogenic variants
LOOKING AHEAD

Reach for a Cure

The acceleration of technology provides optimism for our future. Whole Exome Sequencing and genetic testing are becoming more available and more families are receiving a diagnosis at an earlier age.

Precision medicine models existing drugs to determine which are effective and optimize them for rare diseases.

Patient cell lines can be converted to organoids — brain, stomach, blood, and lung tissues — that can express the same defects as those seen in patients and can be leveraged for trial therapeutics.

AYRE will strategically invest in our core priorities to bring our vision to fruition.

We look forward to a successful 2019 and invite you to join us on this incredible journey!

Laura Badmaev
Thank you!
The ARRE Foundation is so important for families with children with an ASXL mutation, because they bring researchers and so knowledge together which is very important to understand more about these syndromes! The first conference in July 2018 was a very big success and brought families from all over the world together.” – Suzanne Driessen
LEAVE YOUR LEGACY

Planned giving is an opportunity to make a significant difference in the lives of these children and their families. To learn more and discuss planning your gift to ARRE Foundation, please contact info@arrefoundation.org.

- Securities
- Real Estate
- Gifts by Will
- Life Insurance Policy
- Charitable Gift Annuity
- Charitable Remainder and Lead Trusts
- IRA Distributions
ARRE needs assistance from family, friends, caregivers, and neighbors to make our vision possible. We are a volunteer based organization and your support is critical.

VOLUNTEER

Sign up to receive news and updates about ASXL Rare Research Endowment activities.

SUBSCRIBE

You can help the ASXL Rare Research Endowment remain at the forefront ASXL research: arrefoundation.org/donate

DONATE