

Year in Review 2024



VISION FOR
TOMORROW
FOUNDATION

VFT is committed to funding promising research on aniridia and albinism and supporting families affected by these challenging genetic conditions.

APRIL

Family
Outreach
Fundraising
Campaign

MAY

Launch PFAC

JUNE

Aniridia
Europe
Conference

OCTOBER

Pledged
Support to
International
Scientific
Conference on
Albinism Grant

NOVEMBER

Chicago Event

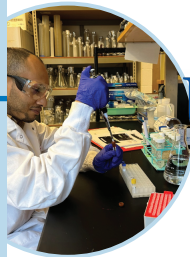
DECEMBER

NOAH
Support

EYE ON RESEARCH



VFT is proud of its ongoing support for the National Organization for Albinism and Hypopigmentation with a grant for its growing Research Program.



UNIVERSITY OF
DELAWARE

Kicked off year one of a new study with Melinda Duncan, Ph.D. to look into specific genes and potential drug therapies to slow or stop aniridic keratopathy.

VFT hosted an event for more than 300 people in Chicago to raise awareness and funds for future research projects.

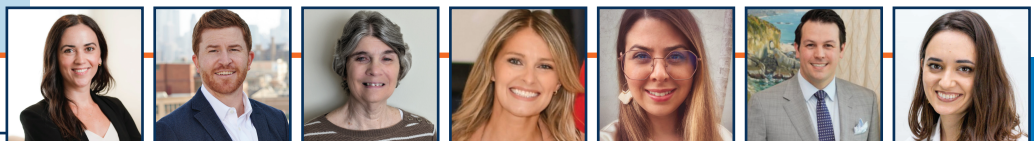


Attended the 7th European Aniridia Conference in Sweden.

VFT networked with European researchers and gained a better understanding of the critical work they are doing.



FAMILY SUPPORT



PATIENT AND FAMILY ADVISORY COMMITTEE • Launched the PFAC to engage with more persons with aniridia and their family members to help VFT identify opportunities for more patient community engagement.

RESOURCE DEVELOPMENT • Created and shared tools to help parents navigate aniridia diagnosis.

NEW PARENT OUTREACH • Provided insights and support to new parents of aniridic children.

LEADERSHIP

- Susan Wolfe, President
- Grayson Chinn, Vice President & Secretary
- Matt Wolfe, Treasurer
- Cheleah Googe, PhD

EMERITUS BOARD

- Susan Ballis, MD
- Wendy Baum, CPA, JD

For ongoing news, please visit: visionfortomorrow.org

or follow us:   

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Gold
Transparency
2024

Candid.