Year in Review 2022

VISION FOR TOMORROW FOUNDATION

FEBRUARY
Joins Global Genes

JUNE
International Aniridia Day

AUGUST
UVA Research Published

OCTOBER
Great Non-Profits

NOVEMBER
ISCA

DECEMBER
ANA Partnership Agreement

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

NEW BOARD *ONE YEAR IN* • EXCITED ABOUT PROGRESS AND STILL MORE TO DO...

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

EMERITUS BOARD

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

FOCUSED ON RESEARCH

University of Virginia

Funded research that established a first-of-its-kind protocol to examine gene expression in early stages of eye developments in aniridia. The research was published in "Cold Harbor Springs Protocol" medical journal.

ISCA 2022 International Scientific Conference on Albinism
A global forum to improve the health of those with albinism

Sponsored the International Scientific Conference on Albinism. The unique conference included researchers, health care providers and albinism organization leaders from around the globe.

Illinois Eye and Ear Infirmary
University of Georgia

Ongoing research for aniridia cornea repair.

MAKING CONNECTIONS

INTERNATIONAL ANIRIDIA DAY • Hosted meet-ups to bring Aniridia families together in person to share experiences and make new connections.

ANA Aniridia North America

Signed a Memorandum of Understanding to solidify ongoing support and collaboration between VFT and Aniridia North America.

Global Genes® Allies in Rare Disease

Virtually attended the Global Genes Patient Advocacy Summit as part of VFT's new membership in the Global Genes RARE Foundation Alliance.

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The Vision for Tomorrow Foundation
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VFT is a 501(c)(3) charitable organization with 100 percent volunteer support. More than 95 percent of donations directly fund research.