

2015 ANNUAL REPORT



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The mission of the Retina Foundation of the Southwest is to prevent vision loss and restore sight through innovative research and treatment. The Retina Foundation is a leader in research to develop treatment and cures for age-related macular degeneration, pediatric eye disorders, and inherited eye diseases. The Retina Foundation is led by five Scientific Directors.



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There are many things that we value in this world. Many of us value our families and our profession. We value our health and our resources to take care of ourselves and one another. Our values have a way of guiding our actions and when our values are firm and strong, it makes it easier to make a decision.

In 2015, our Board and staff completed a strategic plan to reinforce our mission, vision and values. Today, we are deeply focused on preventing vision loss and restoring sight through innovative research and treatment. The values that we find critical are leadership, integrity, compassion, patient education and stewardship. We believe that if we stay true to our values, there is no doubt that we will stand the course of our vision, keeping research at the forefront of our goals by making a difference for all people whose lives are impacted by vision loss.

Our journey began more than 34 years ago, and we have so much to be proud of in 2015. Last year marked the establishment of the Clinical Center of Innovation for Age-Related Macular Degeneration. This collaboration with the Bobby B. Lyle School of Engineering at Southern Methodist University would not have been possible without the generous \$2.5 million grant provided by the W. W. Caruth, Jr. Foundation at Communities Foundation of Texas. By empowering this diverse, unique innovation team, this collaboration places Dallas in the epi-center of a completely transformative process in accelerating medical research to save funds and precious time.

In the pages of this report you will learn about our successes in research in age-related macular degeneration, pediatric eye disorders, and inherited eye diseases. We are exploring the benefits that stem cells may have in slowing age-related macular degeneration. We are closer than ever to establishing our binocular treatment approach as superior to patching for treating amblyopia. We are also exploring a gene therapy approach to treating advanced retinitis pigmentosa that involves the first clinical trial of optogenetics.

Without your generous support, none of this would be possible. Together, our impact is significant and we are grateful to you for helping us enter new territory in vision research and leading the way to save sight for generations to come.

Sincerely,

James H. Merritt, M.D.

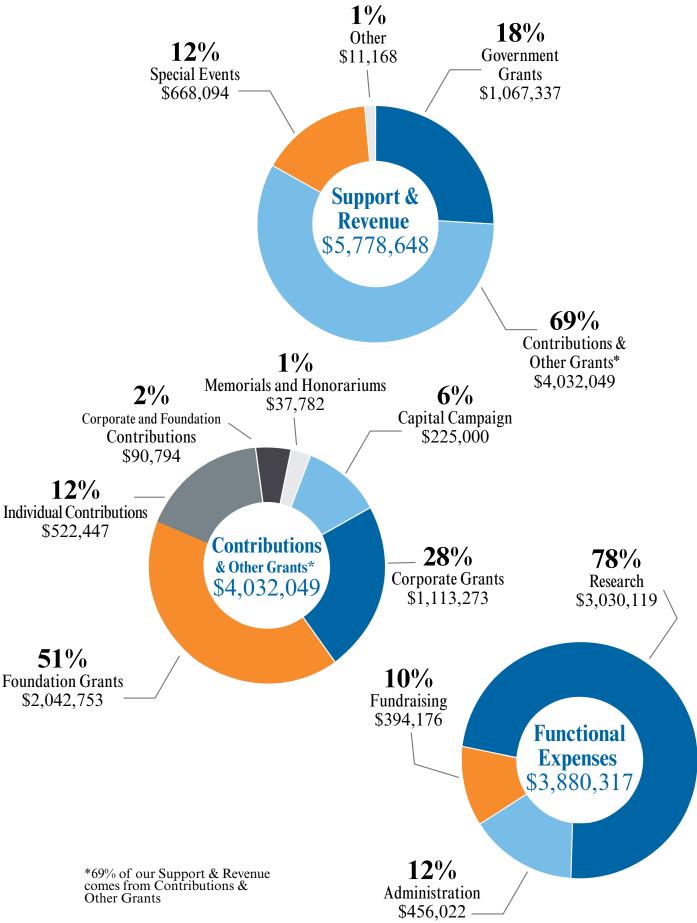
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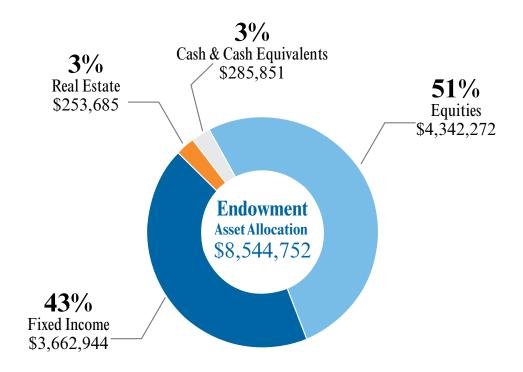




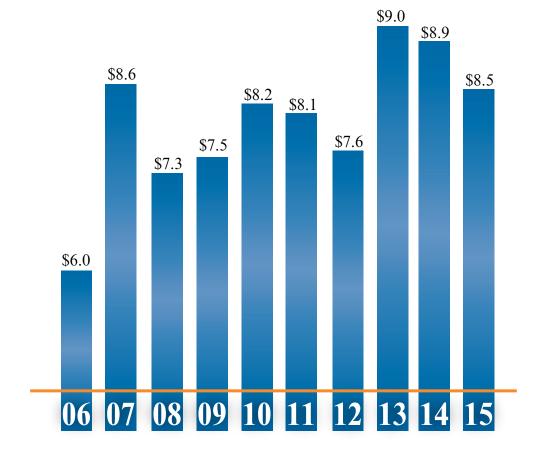
2015 FINANCIALS



2015 FINANCIALS



Endowment Balances2006-2015





New Donors

2,326

RESEARCH STUDIES & **CLINICAL TRIALS**

Enrolled in our



olunteers

2\$4,032,049

Raised

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Phyllis G. & William B. Snyder Legacy Society

Our Legacy Society is named in honor of Phyllis G. & William B. Snyder who have left a legacy to the Foundation through their dedication, leadership and financial support that dates back more than 30 years.

A legacy gift to the Retina Foundation will help ensure that breakthroughs in the treatment and restoration of vision loss continue beyond one's lifetime. It is a simple way to create a lasting personal legacy that will impact lives for future generations to come.

Making a legacy gift to the Retina Foundation can be as simple as making an amendment to your existing will or trust. You can designate a percentage of your estate, a specific dollar amount, specific assets or property for us in establishing your legacy gift. To learn more about the advantages of making a legacy gift to the Retina Foundation please call 214.363.3911.



"I chose, after doing research, to leave a legacy gift to the Retina Foundation of the Southwest. The Foundation, highly rated by Charity Navigator, is a leader in finding a cure for macular degeneration. Both of my parents, in their later years, were afflicted with the "dry" form of this disease. Seeing how difficult it was as they got older to be visually compromised, in their memory I want to help advance research."

-Karla Cross Arundel, *Raleigh, North Carolina,* Member of the Phyllis G. and William B. Snyder Legacy Society



innovative process might be applied to other diseases for which no cure exists."

ONE VISION



To pioneer a new method of medical research to find tools to better diagnose and treat AMD, and help patients live more independent, quality-filled lives.



ONE TEAM

The Retina Foundation of the Southwest (RFSW) and the Bobby B. Lyle School of Engineering at Southern Methodist University (SMU) have joined forces to create the Clinical Center of Innovation for Age-Related Macular Degeneration.

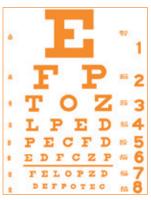
RFSW SCIENTISTS

The Retina Foundation of the Southwest is a world-class leader among its peers in the pursuit of more advanced treatments for AMD with a proud history of groundbreaking discoveries in eye health. Our scientists bring years of clinical experience and medical knowledge to the table.

SMU ENGINEERS

The Bobby B. Lyle School of Engineering at SMU is equipped with a faculty and staff dedicated to understanding the science of innovation, fostering it in their students, and using it to impact the community. The university's commitment to applying its resources to solve real world problems and its expertise in the science and art of innovation make the school a natural partner for RFSW.

18,000,000+ AMD PATIENTS



As the most common cause of vision loss for individuals 50 years and older in the United States, age-related macular degeneration (AMD) is a progressive disease of the retina. AMD gradually destroys the macula, the center part of the retina, which has the highest amount of cells that function to provide fine vision required for activities such as reading and driving. An estimated

18 million Americans suffer from some form of AMD, and the age 50+ population is projected to double by 2030. The disease is complex and there is no known cure.

INVESTMENT OPPORTUNITY

Our goal is to create a self-sustainable program to drive innovation in AMD detection and treatment. An initial grant from the innovation-seeking W. W. Caruth, Jr. Foundation at Communities Foundation of Texas provided our seed funding. These funds have allowed us to begin working on several innovations, however, we still need additional support from forward-thinking donors and investors.



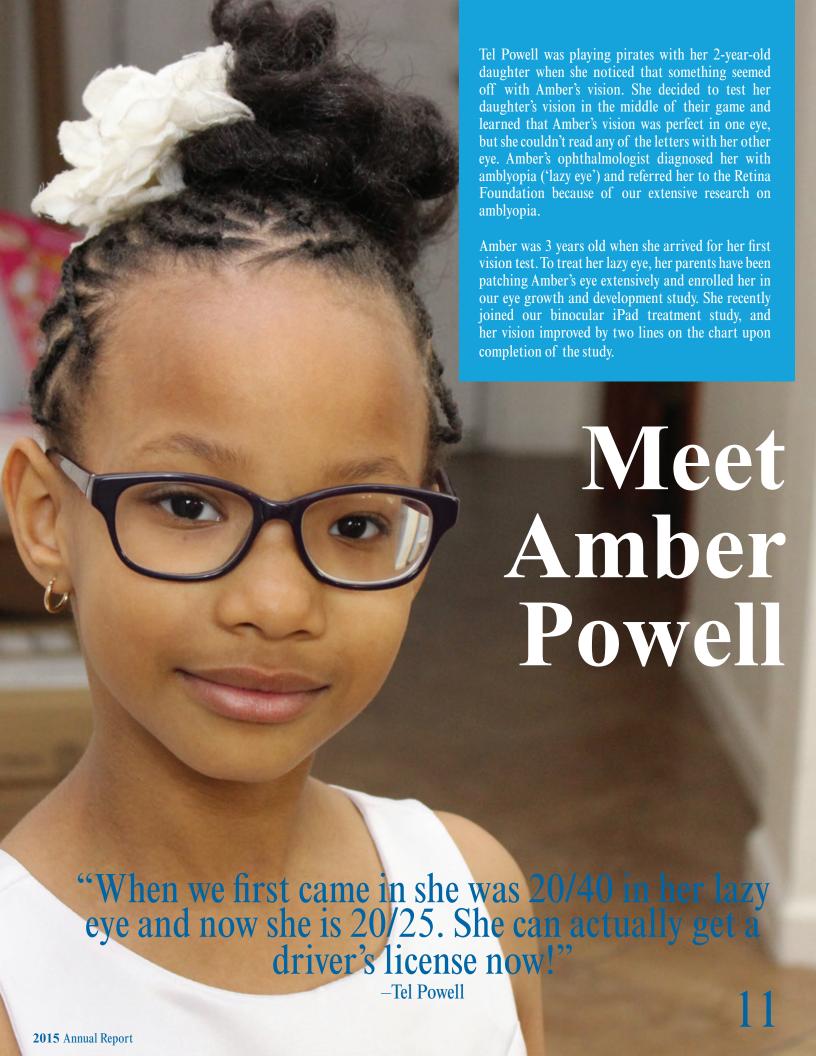
Mrs. Israel believes, "there is an impairment about vision that it almost exceeds not being able to walk." Both her mother and father were diagnosed with age-related macular degeneration (AMD), so she was always very aware of the possibility that she too would be affected. One day, Mrs. Israel was teaching a class at Baylor University when she realized that the faces of her students weren't clear. Mrs. Israel recalls, "I couldn't distinguish if they had brown or blue eyes, and their faces were all a little blurry."

After being officially diagnosed with dry AMD at age 65, Mrs. Israel told her eye doctor that she wanted a solution. Several years later, her eye doctor heard Dr. Karl Csaky speak at a seminar and believed that his new trial might be the answer for Mrs. Israel. It turned out that she was the perfect candidate for a study for patients with Geographic Atrophy Secondary to Age-Related Macular Degeneration. The purpose of this study is to determine the effectiveness of drug delivery to the eye in slowing or stopping the progression of dry AMD. Mrs. Israel has been treated several times now, and she is hopeful that the drug delivery method works.

Meet Donna Israel

"I think <mark>it's magnificent what you are doing here, and it just has to be done" and it just has to be done" and it just has to be done."</mark>

-Donna Isreal





It has been three years since the Retina Foundation diagnosed Wilson Hunt with achromatopsia, a rare inherited eye disease characterized by decreased vision (20/200), extreme light sensitivity and the absence of color vision. Since his initial diagnosis, 5-year-old Wilson has completed weekly visual and mobility therapy and a special education Pre-K class. Three years ago he could not walk up a flight of stairs, slide down a slide or walk from the sidewalk onto the grass due to his eye condition.

Today, Wilson is enjoying playing soccer, swimming in his pool and reading books (with large font). His school has been very helpful in making accommodations such as a slant board to help with Wilson's writing and an iPad to allow Wilson to view what other kids see on the classroom projector screen. Wilson is set to start Kindergarten next year, and his parents couldn't be more proud of him.

Recently, Dr. David Birch successfully identified the CNGA3 gene as the source of Wilson's eye disease. The Hunts are hopeful that their son may one day be part of a gene therapy trial that will help to activate his cones and contribute to meaningful vision improvement. They are very optimistic that "one day he will be able to see some of the beautiful colors in our world."

"We are forever grateful for the Retina Foundation. Without their care I feel like we would still be spinning our wheels trying to figure out Wilson's visual issues."

-Elizabeth & Taylor Hunt

Age-Related Macular Degeneration

Achieving Integrity

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Success in medicine depends on research. We would not have the modern accommodations that we have for several eye conditions if it weren't for the motivation to learn more, to understand more and to overcome more of the diseases that impinge on our health. In 2015, we delved deeper into drug delivery research for treating age-related macular degeneration. We are currently screening a promising new non-invasive drug delivery device. Integrity is our value, and we are *achieving integrity* through our unwavering commitment to find better ways to treat macular degeneration.

RESEARCH HIGHLIGHTS

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Pediatrics

Leading Research

At the Retina Foundation of the Southwest, we are honored to be in a position to combine our expertise and resources to help children whose sight is impacted at birth and during their school-aged years. Each year, we surpass our prior accomplishments, taking our research to new heights and discovering better treatments to improve vision for children with pediatric eye disorders. In 2015, we continued our work on a binocular iPad game approach for treating amblyopia, "lazy eye." The results of the study signify that the treatment was more effective than patching with two weeks of treatment and there is high compliance, suggesting that it offers an engaging approach for children. Leadership is our value, and we are *leading research* in a way that no other organization can do for children with vision loss.

Inherited Eye Diseases

Demonstrating Compassion

No one is more devoted and focused on researching treatments for inherited eye diseases than we are. At the Retina Foundation, we have a combined total of 8 research studies to prevent vision loss for people with Retinitis Pigmentosa, X-linked Retinoschisis, Choroideremia and Stargardt Disease. In 2015, we were the first site in the nation to conduct a Phase I/II clinical trial to evaluate the efficacy of a new, first-in-class gene therapy application of optogenetics for the treatment of retinitis pigmentosa. We are using gene therapy to insert a light sensitive pigment into retinal ganglion cells. Our goal is to provide low vision to patients who are otherwise blind from retinitis pigmentosa. Compassion is our value, and we are *demonstrating compassion* for those who are losing their vision to rare genetic conditions.

13

Capital Campaign

Thanks to you we have raised \$2,762,807 in support of our \$3.6 million building campaign. Our new facility is equipped with seven laboratories and 25 testing rooms, allowing us to see more patients and accelerate our research.

\$500,000+

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A Legacy Gift to the Retina Foundation of the Southwest ensures that our sight saving research continues. A properly structured Legacy Gift may make it possible for you to achieve one or more of the following benefits:

- Reduced current and future taxes
- Conservation of assets for heirs
- Additional current and future income
- A lasting statement of your personal values
- The ability to retain control of your assets during your lifetime

Gifts to Retina Foundation of the Southwest from your estate are exempt from federal estate taxes. For help with specific language for your estate planning, needs please call 214.363.3911. We can assist you in determining what type of Legacy Gift will best fit your individual and family objectives.

Please let us know if you have already included the Retina Foundation in your will or trust. We would like to thank you for your generosity and recognize you as a member of the Phyllis G. and William B. Snyder Legacy Society. Please call us at 214.363.3911.



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2015 Publications

- 1. Aref, A.A., Scott, I.U., Oden, N.L., Ip, M.S., Blodi, B.A., Van Veldhuisen, P.C., SCORE Investigator Group. Incidence, Risk Factors, and Timing of Elevated Intraocular Pressure After Intravitreal Triamcinolone Acetonide Injection for Macular Edema Secondary to Retinal Vein Occlusion: SCORE Study Report 15. JAMA Ophthalmol. 133:1022-9. (2015)
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- 4. Birch DG. Assessing Rod function: ERGs, fields and fundus perimetry. 2nd Casey Gene Therapy Meeting, Denver, CO, May, 2015.
- 5. Birch DG. Structure-Function Relationships in Inherited Retinal Degenerations. Vision Science Distinguished Lecture, Medical College of Wisconsin, Milwaukee, WI, March 2015.
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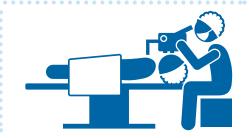
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