



2018 ANNUAL REPORT

FISCAL YEAR ENDING DECEMBER 31, 2018





Dear friends of NEA,

In the 30 years since the National Eczema Association (NEA) was founded, we have seen the experience of living with eczema move out of the shadows into the light. Our shared accomplishments during 2018 highlight this transformation, and I'm honored to share them with you here.

During Eczema Awareness Month people from around the world came out of the shadows on social media to help #unhideECZEMA. You shared your stories and showed the strength and courage not only to persevere for your own sake and the sake of your families, but to share yourselves with our community and prop each other up when the going gets tough.

Another bright spot — this year NEA celebrated a million-dollar milestone in eczema research. As of the end of 2018, our research program had awarded more than \$1.1 million in grant funding for ground-breaking studies of eczema symptoms, comorbidities, therapies, burden of disease, and quality of life.

NEA spread the news about the promising new therapies making their way through the development pipeline. And we engaged you in many of these processes. We continued to learn about alternative and complementary treatment options that help people with eczema live their best possible lives.

We raised our collective voice, and advocates nationwide shared their stories with policy makers. NEA's advocacy efforts ensured the needs of patients and caregivers continue to be meaningfully incorporated into drug development and evaluation. We worked to change state and federal laws that hinder our community's access to therapies — keeping issues like prior-authorization, step therapy, and non-medical switching policies at the fore.

NEA equipped thousands of healthcare professionals nationwide to provide patient-centered care to people with eczema. Last year we formed a dynamic multidisciplinary team of healthcare providers and families living with eczema. Together we produced education for nurses, doctors, and medical students that is patient-focused and responsive to the real-life needs of people with eczema.

Our community inspires me every day and all year long. We've accomplished so much together, and your engagement and generosity made it all possible. Thank you!

Sincerely,

Julie Block President & CEO

2018 Impact



Research



15 YEARS

OF THE NEA RESEARCH PROGRAM



MILLION DOLLAR MILESTONE:

\$1.1 MILLION

DISTRIBUTED IN GRANT FUNDING



26RESEARCHERS FUNDED IN 2018



Advocacy



484GRASSROOTS ADVOCATES
RECRUITED



18
MEETINGS WITH POLICY MAKERS



18
STATE AND FEDERAL ADVOCACY
COALITIONS JOINED

2018 Impact



Healthcare Provider Engagement



2,501
HEALTHCARE PROVIDERS EDUCATED
IN PATIENT-CENTERED ECZEMA CARE



650
DOCTORS REGISTERED WITH
NEA ECZEMA PROVIDER FINDER



11,959
HEALTHCARE PROVIDER
MEMBERS IN NEA NETWORK



Community Support



473
ATTENDEES AT 2018
ECZEMA EXPO IN CHICAGO



5,913 WEBINAR ATTENDEES IN 2018



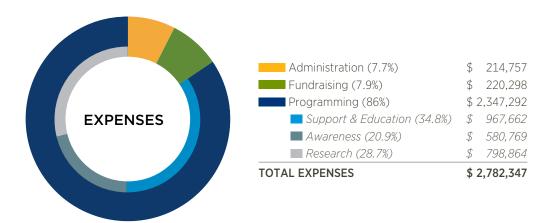
17,302COMMUNITY MEMBERS ACCESSED
THE NEA ECZEMA PROVIDER FINDER



314,179
COMMUNITY MEMBERS ACCESSED
THE SEAL OF ACCEPTANCE
PRODUCT DIRECTORY

Financials

Individual Contributions/Grants \$ 264,994 Grants 30,000 Service Program \$ 997,411 Awareness Events \$ 22.109 **INCOME** Live Events/Conferences \$ 203,011 Corporate Sponsorships \$ 1,619,316 Publications & Brochures \$ 254,651 \$ (27,219) Investment Income TOTAL INCOME \$ 3,391,493



STATEMENT OF ACTIVITIES YEAR ENDED 2018

ASSETS

Current assets:

Cash and cash equivalents	\$ 1,903,505
Investments	\$ 1,373,737
Accounts receivable	\$ 100,015
Prepaid expenses & other current assets	\$ 28,637
TOTAL CURRENT ASSETS	\$ 3,405,894
Fixed assets, net of accumulated depreciation	\$ 54,188
TOTAL ASSETS	\$ 3,460,082

LIABILITIES & NET ASSETS

Current liabilities:

TOTAL LIABILITIES & ASSETS	\$ 3,460,082
TOTAL NET ASSETS	\$ 3,401,364
Temporarily Restricted	\$ 0
Unrestricted	\$ 3,401,364
Net assets:	
TOTAL CURRENT LIABILITIES	\$ 58,718
Accrued vacation payable	\$ 30,723
Accounts payable	\$ 27,995

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NEA Strategic Vision

NEA continues to embrace the new era of eczema, leveraging scientific advancements and increased community engagement to improve the health and quality of life for people with eczema.



IMPACT STATEMENT

Eczema will be better understood as the serious disease it is — driven by expanded research and the contributions of empowered community members. Patients will increasingly be active partners in their own care. People with eczema will be more connected to the community and the support they need and deserve. NEA's commitment to people living with eczema will improve lives.

FIVE TRANSFORMATION KEYS

Our two-year strategic plan is driven by five transformation keys:

1 Erase the Stigma of Eczema

Raise awareness about the damaging impact and cost of eczema on individuals, families, and society to foster understanding and acceptance, and to drive innovations in care.

- 2 Advance and Promote Standards of Eczema Care
 Educate and inspire medical professionals to advance the best standards
 of eczema care, including new treatments.
- 3 Forward Patient-Centered Care
 Develop and coordinate patient-centered, multi-disciplinary tools to improve patient outcomes beyond the physician's office.
- 4 Drive Innovative Research
 Fund, promote, and advance innovative research and researchers to
 expand eczema treatments and drive to a cure.
- (5) Ensure Access and Affordability

 Advocate for and promote accessible and affordable treatments to ensure eczema patients get the care they need.

CORE VALUES

The National Eczema Association embraces the following core values:



VISIONARY

We embrace the future and are passionate leaders.



AUTHENTIC

We are "all in" for the NEA community.



COMPASSIONATE

We have genuine heart and respect for individuals and relationships.



SUPPORTIVE

We empower one another and seek collaboration.



RESPONSIVE

We are organizationally nimble and agile.



ACCOUNTABLE

We are honest, trustworthy, and transparent.



TRANSFORMATIVE

We think big, take risks, and create innovative change.



Thank you to our supporters

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