'24

ANNUAL REPORT
THE GLOBAL ALBINISM ALLIANCE

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ANNUAL OVERVIEW

↑ On the cover, Charles (50), Africa
← Adam (14), Europe

2024 was a year of both challenge and progress for the *Global Albinism Alliance*. It was marked by the passing of our beloved co-founder, Mike McGowan, whose dedication was instrumental in the development of our organization. Despite this loss, we managed to advance key initiatives that strengthened global advocacy, research, and community support.

We spoke at numerous events, including *World Health Assembly* side events and *United Nations Human Rights Council* sessions, and participated in over 10 major international meetings, such as *EADV*, *the Clinton Global Initiative*, and the *World Congress on Rare Skin Diseases*, raising awareness about albinism and forging new connections.

On the research front, we concentrated our efforts on organizing the *International Scientific Conference on Albinism (ISCA)* 2025, which received a record-breaking 65 abstract submissions. We also continued coordinating an international group of genetic experts that was officially recognized by *ClinGen* in March.

Our Skin Cancer Program significantly expanded, with initiatives such as the launch of a *Global Call to Action* and preparations for the 2025 World Forum in collaboration with the *International League of Dermatological Societies (ILDS)* and *Standing Voice*. We also continued our collaboration with the *International Agency for Research on Cancer* (IARC) and co-submitted, with the *UN Expert on Albinism*, a new application to add sunscreen to the *WHO Essential Medicines List*.

In the spring, we successfully led an advocacy coalition for the renewal of the *UN Mandate on Albinism*.

In June, to mark the 10th anniversary of *International Albinism Awareness Day*, we co-organized an event at the UN in Geneva, spotlighted albinism at key conferences, and joined the Brazilian albinism community in their online celebration. In collaboration with *GLODERM*, we also released a webinar series covering both the medical and psychosocial aspects of albinism.

Throughout the year, we provided direct support to several albinism groups, including mentoring *Albína*, a new association in the Czech Republic, now serving 62 families with two new counseling services. A field visit to Tanzania and Malawi in August further deepened our understanding of local realities.

Finally, 2024 marked our visual transformation with the launch of a full rebrand, signaling our growth from a pilot project to a professional, sustainable organization.

Thank you to all our partners, organizations, and volunteers for making this year possible. We look forward to continuing this journey with you in 2025.

EDITORIAL

Antoine Gliksohn
EXECUTIVE DIRECTOR

2024 was an especially busy year. Despite limited resources, we managed to launch and complete an increasing number of initiatives. This progress was made possible by the many connections we've built over time with organizations that have since become valued partners and supporters. I am deeply grateful for their continued trust and engagement, which have been essential to our achievements.

Although the year brought few major visible milestones, it was marked by steady behind-the-scenes work that lays the groundwork for important developments in 2025. I'm particularly looking forward to our two upcoming conferences: ISCA 2025 in January and the World Forum on Skin Cancer in October.

On November 24, I lost my friend and closest colleague, Mike McGowan. Since the start of our organization, Mike had been my trusted "partner in crime." His deep expertise and experience often helped guide the Alliance's strategic direction. His passing leaves a profound and painful void.

In this turbulent context, we were unable to put in place the conditions needed to hire a second employee, even though it had long been planned. Nevertheless, I am confident we will achieve this over the coming year.

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150+

ALBINISM ORGANIZATIONS UNDER OUR UMBRELLA

80+

COUNTRIES REPRESENTED

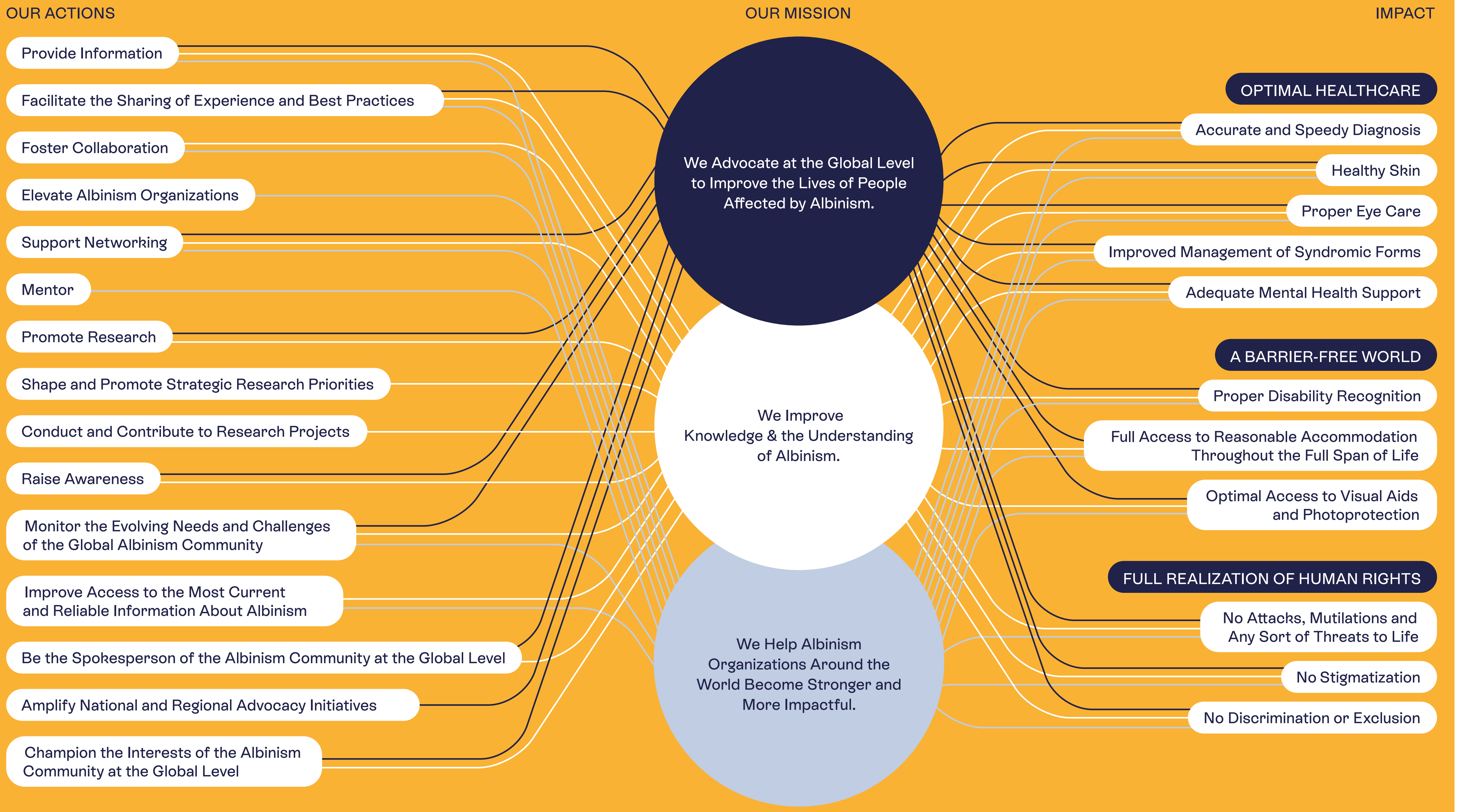
1 in 4,000 to 7,000

PREVALENCE OF ALBINISM IN AFRICA

1 in 12,000 to 15,000

PREVALENCE OF ALBINISM IN EUROPE

HOW WE WORK



OUR ACTIONS

- Provide Information
- Facilitate the Sharing of Experience and Best Practices
- Foster Collaboration
- Elevate Albinism Organizations
- Support Networking
- Mentor
- Promote Research
- Shape and Promote Strategic Research Priorities
- Conduct and Contribute to Research Projects
- Raise Awareness
- Monitor the Evolving Needs and Challenges of the Global Albinism Community
- Improve Access to the Most Current and Reliable Information About Albinism
- Be the Spokesperson of the Albinism Community at the Global Level
- Amplify National and Regional Advocacy Initiatives
- Champion the Interests of the Albinism Community at the Global Level

OUR MISSION

We Advocate at the Global Level to Improve the Lives of People Affected by Albinism.

We Improve Knowledge & the Understanding of Albinism.

We Help Albinism Organizations Around the World Become Stronger and More Impactful.

IMPACT

OPTIMAL HEALTHCARE

- Accurate and Speedy Diagnosis
- Healthy Skin
- Proper Eye Care
- Improved Management of Syndromic Forms
- Adequate Mental Health Support

A BARRIER-FREE WORLD

- Proper Disability Recognition
- Full Access to Reasonable Accommodation Throughout the Full Span of Life
- Optimal Access to Visual Aids and Photoprotection

FULL REALIZATION OF HUMAN RIGHTS

- No Attacks, Mutilations and Any Sort of Threats to Life
- No Stigmatization
- No Discrimination or Exclusion

A World Where People With Albinism Have the Best Possible Quality of Life.

KEY HIGHLIGHTS OF 2024

MARCH 5

Albinism *Variant Curation Expert Panel* Officially Recognized by ClinGen

MARCH 6

Statement at the 55th Session of the *United Nations Human Rights Council*

APRIL 4

Renewal of the United Nations Mandate on Albinism Following Intense Advocacy Work

MAY 8

Meeting with the Research Team at the International Agency for Research on Cancer, Specializing in Skin Cancer Research in People with Albinism

MAY 20

Oral Contribution to Two Side Events of the 77th World Health Assembly

JUNE 1

New Brand Identity & website

JUNE 13

Albinism Highlighted at the World Congress on Rare Skin Diseases in Paris

JUNE 20

International Albinism Awareness Day Celebration in Geneva

JULY 9

Launch of the Call to Action on Skin Cancer in People with Albinism

AUGUST 20

Visit to the *Regional Dermatology Training Center* in Moshi, Tanzania

AUGUST 21

Announcement of *International Scientific Conference on Albinism* 2025

SEPTEMBER 9-12

Annual meeting *of the PanAmerican Society for Pigment Cell Research*

SEPTEMBER 23-24

Clinton Global Initiative Conference

SEPTEMBER 25-28

The European Academy of Dermatology and Venereology Congress

NOVEMBER 1

Submission of an Application to add Sunscreen to the WHO Model Lists of Essential Medicines

NOVEMBER 24

Passing of Mike McGowan

© Elma Okic / UN Photo

WE ADVOCATE AT THE GLOBAL LEVEL

← Human Rights Council
55th Session,
Palais des Nations.

WORLD CANCER DAY EVENT AT THE EUROPEAN COMMISSION

January 31, 2024
Brussels, Belgium

We had the privilege to attend a high-level event at the European Commission, hosted by Stella Kyriakides, EU Health and Food Safety Commissioner, in conjunction with World Cancer Day. This important occasion provided an opportunity to raise awareness about skin cancer among persons with albinism and to engage with key global cancer stakeholders, including the Union for International Cancer Control (UICC).

BUILDING CONNECTIONS

AWARENESS

CALLING FOR EQUAL ACCESS TO EDUCATION FOR PEOPLE WITH ALBINISM AROUND THE WORLD

March 6, 2024
55th Session of the Human Rights Council, United Nations Geneva

For the second time in two years, we had the honor of speaking at the UN in Geneva – in collaboration with International Service for Human Rights – to address people with albinism's right to an education. Our speech focused on their unequal access to education around the globe and some of the reasons for it: some are bullied, some lack any learning or training support whatsoever, and some are afraid to attend school due to violent attacks. Our statement was grounded in the latest report by the UN Independent Expert on Albinism, which included input from fifty-eight different stakeholders: UN Member States, Civil Society Organizations, and individuals with albinism worldwide.

Watch the full session here.

ADVOCACY

COLLABORATION WITH INTERNATIONAL AGENCY FOR RESEARCH ON CANCER

May 8, 2024
📍 IARC Headquarters, Lyon, France

As part of the ongoing collaboration with *the Environmental and Lifestyle Epidemiology Branch of the International Agency for Research on Cancer* (IARC), we visited IARC in Lyon to meet with the research team, review current progress, and explore future opportunities.

Two concrete projects are currently in development: one in partnership with *the South Africa National Cancer Registry*, and another with *Standing Voice*, focused on work in Tanzania and Malawi.

RENEWAL OF THE UN MANDATE ON ALBINISM

Together, with the support of *Standing Voice* and *the Africa Albinism Network*, we led a coalition of civil society organizations to advocate for the renewal of the mandate of *the UN Independent Expert on Albinism.*

In close collaboration with *the Permanent Mission of the Republic of Tanzania*—the penholder of the resolution—we coordinated the drafting process and took part in informal discussions with member states.

ADVOCACY

77th WORLD HEALTH ASSEMBLY SIDE EVENTS

May 29, 2024
📍 Geneva, Switzerland

We were honored to speak at two side events during the 77th World Health Assembly in Geneva, each supporting key projects linked to World Health Assembly resolutions.

The first event, co-hosted by GlobalSkin and the International League of Dermatological Societies (ILDS) with support from the ANESVAD Foundation, focused on a proposed resolution recognizing skin diseases as a global public health priority.

The second event, hosted by Rare Diseases International, centered on a draft resolution aiming to ensure Universal Health Coverage for people living with rare diseases and conditions.

ADVOCACY AWARENESS BUILDING CONNECTIONS

RAISING AWARENESS ABOUT ALBINISM AROUND THE WORLD

June 1, 2024
📍 UN Today

To mark International Albinism Awareness Day, we published an article in *UN Today* highlighting the global challenges faced by persons with albinism—including health risks, vision impairment, discrimination, and limited access to education. The piece also underscores the importance of international cooperation in advancing the rights of people with albinism worldwide.

Read the article here.

AWARENESS

ALBINISM AWARENESS WEBINAR SERIES

June 10–13, 2024
📍 Online

To mark International Albinism Awareness Day 2024, the Global Albinism Alliance and GLODERM co-hosted a special edition webinar series focused on key issues affecting people with albinism. The four sessions featured leading international experts and explored genetics, dermatology, ophthalmology, and the social impacts of albinism.

Watch all sessions here.

KNOWLEDGE ACCESS AWARENESS

INTERNATIONAL ALBINISM AWARENESS DAY 2024 CELEBRATION IN PARIS

June 12, 2024
📍 Paris, France

We co-hosted the Welcome Reception of the 2nd World Congress on Rare Skin Disorders alongside GlobalSkin and ERN Skin. The event provided a valuable opportunity to celebrate International Albinism Awareness Day (IAAD) 2024 and to raise awareness amongst conference attendees about the many challenges faced by people with albinism around the world.

ADVOCACY BUILDING CONNECTIONS

2ND WORLD CONGRESS ON RARE SKIN DISEASES (WCRSD)

June 12–14, 2024
📍 Paris, France

We took part in the 2nd *World Congress on Rare Skin Diseases*, organized in Paris by *ERN-Skin* and *Fondation René Touraine.*

On June 13, Julio Garcia (*Simplemente Amigos, Fundación Nacional de Albinismo – Argentina*) spoke during a patient symposium co-organized by *GlobalSkin*, which focused on the needs and challenges faced by associations and individuals living with rare skin conditions around the world.

The event brought together numerous patient organizations, helping to elevate the global voice of communities that are too often overlooked in public health efforts.

BUILDING CONNECTIONS

AWARENESS

ANNUAL MEETING OF THE INTERNATIONAL ADVISORY COMMITTEE ON NON-IONIZING RADIATIONS

June 13, 2024
📍 World Health Organization, Geneva

For the second year in a row, we had the opportunity to take part in the annual meeting of *the International Advisory Committee on Non-Ionizing Radiations*, hosted by *the World Health Organization's Radiation and Health Unit*. We shared an update on our efforts to address skin cancer in people with albinism—an urgent and often overlooked public health issue.

Our participation reflects our continued commitment to ensuring that the specific health risks faced by people with albinism are recognized and addressed in global discussions on radiation and skin health.

ADVOCACY

BUILDING CONNECTIONS

AWARENESS

→ The 'White Ebony' exhibition was on display during the IAAD celebration at the UN.

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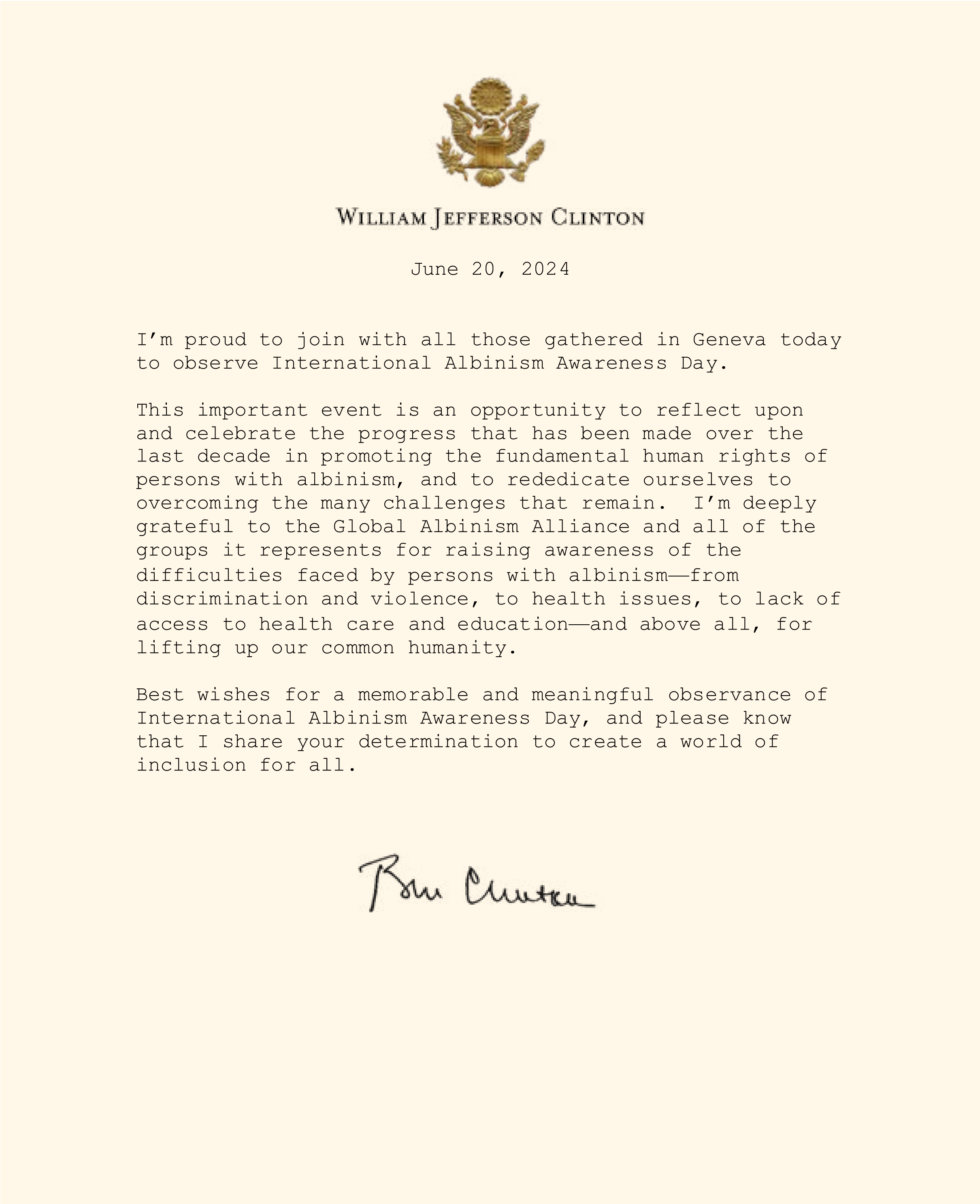

WILLIAM JEFFERSON CLINTON

June 20, 2024

I'm proud to join with all those gathered in Geneva today to observe International Albinism Awareness Day.

This important event is an opportunity to reflect upon and celebrate the progress that has been made over the last decade in promoting the fundamental human rights of persons with albinism, and to rededicate ourselves to overcoming the many challenges that remain. I'm deeply grateful to the Global Albinism Alliance and all of the groups it represents for raising awareness of the difficulties faced by persons with albinism—from discrimination and violence, to health issues, to lack of access to health care and education—and above all, for lifting up our common humanity.

Best wishes for a memorable and meaningful observance of International Albinism Awareness Day, and please know that I share your determination to create a world of inclusion for all.

Bill Clinton

← A letter from Bill Clinton to mark International Albinism Awareness Day.

INTERNATIONAL ALBINISM AWARENESS DAY 2024 CELEBRATION IN GENEVA

June 20, 2024
United Nations Headquarters, Geneva

Hosted by the *United Nations Office at Geneva* and co-organized by *the Global Albinism Alliance*, this International Albinism Awareness Day celebration marked "A Decade of Collective Progress."

The event offered a moment to reflect on the many achievement made over the past ten years, while also acknowledging the ongoing challenges faced by people with albinism worldwide. A highlight of the celebration was the opening of the photo exhibition "White Ebony," showcasing the remarkable exhibit carried out in Congo by Belgian artist *Patricia Willocq*.

The albinism community was well represented, with keynote addresses from our Executive Director, Antoine Gliksohn, Elisabeth Rivaux (*Genespoir*), and Rosalie Malu Muswamba (*Solidarité Albinisme Suisse*), alongside participation Kenneth Skovly Larsen *(Danish Albinism Association)* and Julio Garcia (*Fundación Nacional de Albinismo Simplemente Amigos*).

Watch the video here.

ADVOCACY
AWARENESS
BUILDING CONNECTIONS

WORLD SKIN HEALTH DAY

July 8, 2024
Online

To mark *World Skin Health Day 2024*, we were invited to speak in a webinar hosted by *Resolve Global Health*, focused on the theme "Skin Diseases and the Stigma Around Them." The event brought together voices from across the global skin health community, including Rashmi Sarkar (*International Society of Dermatology; International League of Dermatological Societies*) and David McMahon (*Irish Skin Foundation; GlobalSkin*).

World Skin Health Day is a vital opportunity to raise awareness about the burden of skin conditions and to spotlight the stigma, challenges, and unmet needs faced by people living with these conditions around the world.

AWARENESS

CALL TO ACTION TO END THE EPIDEMIC OF SKIN CANCER IN PEOPLE WITH ALBINISM

July 9, 2024
📍 Online

We joined forces with *the International League of Dermatological Societies*, *the European Academy of Dermatology and Venereology*, *GlobalSkin*, and 32 other national and international organizations to launch a Call to Action. This initiative urges the *World Health Organization*, national Ministries of Health, policymakers, medical professionals, patient groups, and organizations representing people with albinism to implement 10 key actions aimed at preventing premature deaths from skin cancer among people with albinism worldwide.

Our message reached a wide audience across social media, with *LinkedIn* posts generating over 3,300 impressions and more than 560 reactions. To date, the Call to Action has been downloaded at least 1,870 times in multiple languages.

Read the Call to Action here.

ADVOCACY AWARENESS

← Trip to Tanzania and Malawi with Standing Voice.

WORLD HEALTH ORGANIZATION ESSENTIAL MEDICINES LIST APPLICATION

Following the unsuccessful 2022 application co-submitted by the *Global Albinism Alliance* and *the UN Independent Expert on Albinism* to include broad-spectrum sunscreen on the *WHO Model List of Essential Medicines* (EML), a new application was submitted to the 25th WHO EML Expert Committee at the end of 2024. This renewed effort has received the strong support of *Standing Voice*, *Beyond Suncare*, *the Pierre Fabre Foundation*, *the International League of Dermatological Societies* (ILDS), and *the Africa Albinism Network*.

Our application is scheduled to be reviewed by the *WHO Expert Committee* in May 2025.

Learn more about the application here.

BUILDING CONNECTIONS

ADVOCACY

WORLD SKIN HEALTH COALITION

August 2024

We joined the *World Skin Health Coalition*—a patient-led, multi-stakeholder group uniting dermatology patient organizations, healthcare professionals, researchers, and industry partners. Throughout 2024, we actively participated in quarterly meetings focused on shaping a global strategy to support the adoption of a *World Health Assembly* resolution on skin diseases.

As part of our engagement, we contributed to the Coalition's *#NotJustMySkin* online campaign by sharing a series of awareness-raising social media posts in March and April, highlighting the challenges faced by people with albinism in relation to skin health. The campaign reached over 5 million people. Materials featuring data and stories of real people living with skin conditions and diseases were viewed over 17.8 million times. Over 6,800 people in 130 countries across 6 World Health Organization regions signed the open letter urging health policy leaders to act.

BUILDING CONNECTIONS

ADVOCACY

EUROPEAN SOCIETY FOR DERMATOLOGICAL RESEARCH ANNUAL MEETING 2024

September 4–7, 2024
📍 Lisbon, Portugal

We took part in the 2024 Annual Meeting of *the European Society for Dermatological Research* (ESDR), held in Lisbon, Portugal.

Our Executive Director, Antoine Gliksohn, was joined by Dr. Prof. Ncoza DLOVA, head of dermatology, University of Kwazulu-Natal in a special session featuring organizations representing people with various skin conditions. Together, they shared key insights on albinism, focusing on the diverse lived experiences of people with albinism around the world, as well as current research efforts and existing gaps in the field.

Learn more about the event here.

AWARENESS

CLINTON GLOBAL INITIATIVE 2024

September 23–24, 2024
New York City, USA

For the third consecutive year, we joined hundreds of changemakers at the *Clinton Global Initiative* (CGI) in New York City.

We were represented by Antoine Gliksohn, our Executive Director, and Carol Prendergast, our President, who highlighted the multiple challenges faced by people with albinism worldwide, with a special emphasis on skin cancer and the urgent need for increased attention and resources to address this critical health issue.

Learn more about the event here.

BUILDING CONNECTIONS

GLOBALSKIN EUROPE MEMBER MEETING

September 25–28, 2024
Amsterdam, Netherlands

We took part in the *GlobalSkin* Europe Member Meeting, held alongside the 2024 European Academy of Dermatology and Venereology Congress (EADV) in Amsterdam. The gathering offered a valuable opportunity to strengthen connections with *GlobalSkin* staff and fellow member organizations, while contributing to discussions on the future of dermatology patient advocacy across Europe and beyond.

RARE DISEASES INTERNATIONAL (RDI) MEMBERSHIP MEETING

October 21–22, 2024
Barcelona, Spain

This year's gathering in Barcelona was an invaluable opportunity to exchange knowledge, share best practices, and engage in strategic discussions with advocates from the global rare disease community.

A key highlight of the meeting was the discussion on the upcoming *World Health Assembly Resolution* on persons living with a rare disease or condition, which may be adopted by *the World Health Assembly* in May 2025.

ORGANIZATION DEVELOPMENT

BUILDING CONNECTIONS

CONSULTATION WORKSHOP FOR ORGANIZATIONS OF PEOPLE WITH ALBINISM IN EUROPE AND NORTH AMERICA

November 22, 2024
Online

We co-organized an online consultation workshop with the UN Independent Expert on the Rights of Persons with Albinism, Muluka-Anne Miti-Drummond, bringing together organizations from Europe and North America.

Held on the occasion of the 10th anniversary of the UN mandate on Albinism, the workshop offered a moment to reflect on a decade of global progress in advancing the rights of people with albinism—and to recognize the role the mandate has played in that journey. Participants also discussed ongoing challenges and explored strategies to ensure the continued advancement of rights, protection, and inclusion for people with albinism in the decade ahead.

SUPPORT RESEARCH ADVOCACY

ANNUAL CONFERENCE OF THE FRENCH SOCIETY OF DERMATOLOGY

December 3–7, 2024
Paris, France

Our final conference of the year brought us once again to the "*Journées Dermatologiques de Paris*" (JDP), the annual congress of *the French Society of Dermatology*. Since 2021, we have participated in this key event to raise awareness about albinism within the dermatology community.

This year, we were hosted at the booth of *Genespoir*, the French albinism association, where we had the opportunity to present the work of *the Global Albinism Alliance* to participants from several African countries, including Mali, Côte d'Ivoire, Cameroon, the Democratic Republic of Congo, and Guinea.

We also engaged in productive conversations with key partners such as Christophe Przybylski (*Fondation Pierre Fabre*) and Caroline Blondel (*European Society for Dermatological Research*), and had promising exchanges with leading Moroccan dermatologists—Prof. Soumiya Chiheb and Prof. Said Amal (President of *the Moroccan Society of Dermatology*)—on ways to support the albinism community in Morocco.

Learn more about the event here.

BUILDING CONNECTIONS

AWARENESS

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← A young man undergoing treatment for skin cancer.

WE IMPROVE KNOWLEDGE & UNDERSTANDING OF ALBINISM.

ALBINISM VARIANT CURATION EXPERT PANEL

March, 2024
Online

As part of the Clinical Genome Resource (ClinGen) initiative, we continued our coordination of the Albinism Variant Curation Expert Panel throughout 2024.

The panel was officially recognized by the Clinical Genome Resource in March 2024 and is co-chaired by Dr. David Adams (National Institutes of Health, Bethesda, USA) and Dr. Panagiotis Sergouniotis (University of Manchester, United Kingdom).

Nine online meetings were held during the year, focused on reviewing variants in the tyrosinase (TYR) gene, which is associated with oculocutaneous albinism type 1.

Work on the OCA2 gene is expected to begin in 2025, following the publication of rule specifications for TYR.

Learn more about the Project.

BUILDING CONNECTIONS

VISIT TO THE REGIONAL DERMATOLOGICAL TRAINING CENTRE (RDTC)

August 21, 2024
Moshi, Tanzania

During this first visit to Tanzania, we had the opportunity to stop in Moshi to visit *the Regional Dermatological Training Centre* (RDTC). We were warmly welcomed by Professor Daudi Mavura, Principal of the RDTC, and joined by Harry Freeland, *Executive Director of Standing Voice.* Founded in 1990 as a collaboration between *the International League of Dermatological Societies* (ILDS), *the Tanzanian Ministry of Health*, and *Kilimanjaro Christian Medical Center*, the RDTC serves as a supra-regional hub for clinical care, training, and research in dermatology.

Since 1993, the RDTC has run a Comprehensive Care Programme for people with albinism, focused on skin cancer prevention and treatment.

BUILDING CONNECTIONS

ANNUAL MEETING OF THE PANAMERICAN SOCIETY FOR PIGMENT CELL RESEARCH (PASPCR)

September 9–12, 2024
New York, USA

We were pleased to represent the albinism community at the 2024 annual meeting of *the Pan-American Society* for *Pigment Cell Research* (PASPCR), held in New York City. This gathering offered a valuable opportunity to raise awareness among researchers working on melanocytes and melanin production about the needs and expectations of people with albinism—particularly in the field of biomedical research.

We connected with leading scientists contributing to the understanding of albinism through genetics and cell biology, including Jonathan Zippin (*Weill Cornell Medicine*), Mickey Marks (*Children's Hospital of Philadelphia / University of Pennsylvania*), Stacie Loftus (NHGRI), and Elena Oancea (*Brown University*). These experts also serve on the Scientific Advisory Committee for ISCA 2025 and *the ClinGen Albinism Variant Curation Expert Panel.*

The conference featured a powerful keynote by fashion model and advocate Diandra Forrest, who shared her personal story of growing up with albinism in New York City.

RESEARCH

AWARENESS BUILDING CONNECTIONS

RESEARCH LAB VISIT – MEDICAL COLLEGE OF WISCONSIN

September 20, 2024
Milwaukee, USA

We visited the Eye Institute at *the Medical College of Wisconsin*, where we were welcomed by Professor Joseph Carroll and his team. As part of the visit, we participated as a volunteer in an ongoing research study on albinism conducted within *the Advanced Ocular Imaging Program* (AOIP).

This program seeks to deepen the understanding of how retinal development is altered in individuals with albinism, contributing valuable insights to the field of vision science.

RESEARCH

← Antoine Gliksohn, our Executive Director, at the EADV congress

EUROPEAN REFERENCE NETWORK–SKIN BOARD MEETING

October 24–25, 2024
Paris, France

We took part in the *ERN-Skin Board Meeting*, contributing to ongoing efforts to strengthen European collaboration around rare skin conditions, including albinism. Over the past two decades, several European countries have adopted public policies to support people affected by rare conditions. At the regional level, *the European Commission* has established *European Reference Networks* (ERNs) to coordinate care and research.

Since 2021, we have been representing the albinism community within ERN-Skin, working to ensure equitable access to quality healthcare across Europe and to advance the understanding of albinism through data collection and multicenter research studies.

BUILDING CONNECTIONS

RESEARCH

EUROPEAN ACADEMY OF DERMATOLOGY AND VENEREOLOGY 2024 ANNUAL CONGRESS

September 25–28, 2024
Amsterdam, Netherlands

We were proud to participate once again in the annual congress of *the European Academy of Dermatology and Venereology* (EADV)—the largest dermatology conference in Europe. Since 2022, this event has served not only as a key European meeting but as a global platform to connect with dermatologists and partner organizations from around the world.

This year, we had the opportunity to engage with professionals and allies from Argentina, Kenya, Czech Republic, India, USA, South Africa, Dominican Republic, Portugal, Angola, Israel, Guinea, Mexico, France, Brazil, Tanzania, England, Panama, United Arab Emirates, Germany, the WHO—and of course, the Netherlands.

The congress also provided a space to strengthen collaboration with strategic partners such as *GlobalSkin*, the *ILDS*, the *Colegio Ibero Latinoamericano de Dermatologia (CILAD)*, and the organizers of the 2027 *World Congress of Dermatology*.

Learn more about the event here.

BUILDING CONNECTIONS

AWARENESS

EXPLORING THE LINK BETWEEN ALBINISM AND SLEEP DISORDERS

Throughout the year, we engaged in ongoing exchanges with a research team from *the Netherlands Institute of Neuroscience working* on a project exploring the link between albinism and sleep disorders. We look forward to supporting the advancement of this important research.

Learn more about the project here.

RESEARCH

SUPPORTING RESEARCH PUBLICATION ON SKIN CANCER IN ALBINISM

We provided support to Dr. Panawé Kassang for the publication of a long-completed systematic review of the literature on skin cancer in persons with albinism. This study, carried out several years ago and presented at ISCA 2022, had remained unpublished until now, and we were pleased to help ensure that this important research is made accessible to the broader community.

RESEARCH

ISCA 2025 – INTERNATIONAL SCIENTIFIC CONFERENCE ON ALBINISM

ISCA 2025, the second edition of the International Scientific Conference on Albinism, will take place at the end of January 2025. Organized by us with the support of an international scientific committee, the conference builds on the success of ISCA 2022 and reflects our continued commitment to advancing research and global cooperation on albinism.

Much of the organizational work took place throughout 2024, with strong momentum leading into the new year. The Call for Abstracts closed in early December with a notable increase in participation: 65 abstracts were submitted, compared to 50 in the previous edition.

Learn more about the conference here.

RESEARCH

WORLD FORUM ON SKIN CANCER IN PEOPLE WITH ALBINISM

October 27–28, 2025
Cape Town, South Africa

Building on work that began in 2023 with *the International League of Dermatological Societies* (ILDS) and *Standing Voice*, our efforts to organize the first-ever global conference dedicated to skin cancer prevention and management in people with albinism continued throughout 2024. Significant progress was made, including finalizing the location and dates. The Forum is scheduled to take place in Cape Town, South Africa, on October 27–28, 2025. A Save the Date was issued at the end of 2024, with registration planned to open in spring 2025. Fundraising efforts remain ongoing.

The primary goal of the Forum is to develop a global prioritized action plan committed to preventing and managing skin cancer, thereby promoting the health and wellbeing of all people affected by albinism.

BUILDING CONNECTIONS

RESEARCH ADVOCACY

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WE HELP ALBINISM ORGANIZATIONS AROUND THE WORLD BECOME STRONGER AND MORE IMPACTFUL

← Trip to Tanzania and Malawi with Standing Voice.

CELEBRATION OF IAAD 2024 WITH THE BRAZILIAN ALBINISM COMMUNITY

June 14, 2024
Online

We were honored to take part in an online event organized by the Brazilian albinism community to celebrate International *Albinism Awareness Day* 2024, under the theme: "*People with Albinism – 10 Years of Struggle for Inclusion and Accessibility, Without Prejudice and With Equality.*"

Our Executive Director joined *the UN Independent Expert on Albinism*, Muluka-Anne Miti-Drummond, moderator Joselito Luz, and other speakers from Brazil and Argentina to reflect on a decade of collective progress in the global fight for inclusion, accessibility, and equal rights. The event was co-organized by APALBA (*Associação das Pessoas com Albinismo na Bahia*) and *the Coletivo Nacional das Pessoas com Albinismo.*

SUPPORT

VISIT TO TANZANIA AND MALAWI WITH STANDING VOICE

August 14–29, 2024
Tanzania, Malawi, Africa

We spent two intensive weeks in Tanzania and Malawi alongside Harry Freeland, Executive Director of *Standing Voice*, immersing ourselves in grassroots work supporting people with albinism. During our trip, we visited local clinics and witnessed *Standing Voice's* comprehensive, locally led programmes in action.

This visit deepened our understanding of the day-to-day challenges people with albinism face—accessing health services, visual aids, and education—and reinforced the importance of solutions spearheaded by *Standing Voice*. Our shared field experience lays the foundation for stronger collaboration and continued advocacy in the region.

Learn more about *Standing Voice* here.

SUPPORT

MEETING WITH THE TANZANIA ALBINISM SOCIETY

August 25, 2024
Mwanza, Tanzania

As part of our first visit to Tanzania, we had the opportunity to meet with Alfred Kapole, Vice-Chair of *the Tanzania Albinism Society* (TAS). To the best of our knowledge, TAS is the world's first association of persons with albinism—founded in 1977 and officially registered in 1980.

This inaugural meeting between *the Global Albinism Alliance* and TAS offered valuable insight into the situation on the ground. The conversation focused on key priorities for the albinism community in Tanzania, the longstanding work of TAS, and the urgent need for the government to adopt the long-awaited National Action Plan on Albinism.

The exchange marked an important step toward future collaboration in supporting the rights and wellbeing of people with albinism in the country.

SUPPORT

SUPPORT AND MENTORING OF A NEW ORGANIZATION

Online

Throughout the year, we provided ongoing support and mentorship to *Albína*, Czech Republic, a newly established organization dedicated to people with albinism.

In its first year of operation, the organization welcomed 60 members. In partnership with two hospitals, *Albína* launched ophthalmological and dermatological consultation services for individuals with albinism. The organization also facilitated several community meetings and organized its inaugural weekend retreat.

Learn more about Albína here.

SUPPORT

→ Zongyi Lv (12), Asia

© Chinese Organization for Albinism

PEOPLE IN THE GLOBAL ALBINISM ALLIANCE

© Dominik Kučera

Antoine Gliksohn

EXECUTIVE DIRECTOR, FRANCE

Antoine is the Executive Director of the Global Albinism Alliance, where he oversees strategic planning, organizational development, and daily operations. Before joining the Alliance, he worked in Paris as a Project Engineer on a major metro project. A person with albinism himself, Antoine is also an award-winning patient advocate.

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Carol Prendergast

PRESIDENT, USA

Carol is Chair of the Global Albinism Alliance, where she champions the human rights of persons with albinism. Prior to this, she worked as an attorney, program director, consultant, and board member for leading human rights NGOs, and served as a Senior Fellow in Human Rights at the London School of Economics.

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Kelsey Thompson

SECRETARY, USA

Elizabeth Beales

ADMIN VOLUNTEER, AUSTRALIA

© Dominik Kučera

Julio Garcia

LIAISON FOR LATIN AMERICA, ARGENTINA

Mike McGowan

TREASURER

We are sad to share the news of the passing of Mike McGowan, a dedicated advocate for the albinism community, on November 24th, 2024.

Mike was among the initiators of the Global Albinism Alliance. In 2020, he co-organized in Paris an exploratory meeting that marked the launch of our organization. Since then, he remained instrumental in the development of the Alliance, dedicating his energy, expertise, and over three decades of experience as a leader in the albinism community to serving our organization. We would like to share a bit about Mike's life and career to give all of you — those who knew him and those who did not — a sense of the qualities that made him so special to us.

Mike faced the challenges of all people with albinism (PWAs), including low vision that designated him as being legally blind. He often said that coming from a family of seven children who filled their home with "non-stop action" and parents who encouraged them all to always do their best, gave him confidence that he could keep up with anyone despite the challenges of being a PWA. When a high school teacher advised him that he might not be accepted at an academically rigorous college that Mike had applied to, Mike bristled at the suggestion that he could not compete with others who did not have a disability. He enjoyed relating how pleased he was to inform that teacher that he did, in fact, get accepted to that school. Mike wanted everyone to know that just because he was a PWA, he was not to be underestimated. And, indeed, he proved to be right. Mike established a record of professional achievement in a 30 year career in court technology in Cook County Illinois, one of the largest court systems in the United States. Not only did Mike enjoy this demanding position, but it was during this time that Mike's service to the albinism community began. First he assumed a volunteer role in NOAH. Mike began as a local volunteer, then expanded his role nationally by serving as a NOAH board member. He went on to serve as board President and finally, after his retirement from the court system, he served as NOAH's first Executive Director. He held this position for just over ten years, growing the organization in size and scope.

In the past four years Mike was a pivotal figure in the development of the GAA. He used his strategic and organizational skills to help keep us on track. His decades of experience working directly with PWAs, their families and communities always reminded us that while the progress of the broader albinism movement is essential, we can never lose sight of the day-to-day experiences of PWAs. And his deep compassion and commitment to improving the lives of persons with albinism all across the world served as an inspiration to all of us who had the privilege of knowing him. We will continue to be guided by his pure and courageous spirit.

May his memory be a blessing.
Carol Prendergast

> "Mike's vision and leadership were instrumental in shaping the organization, the programs and services that continue to support people with albinism and their families today."
>
> Karen Bly, Executive Director of NOAH

FINANCIAL OVERVIEW

$43,090	RESTRICTED REVENUES
$115,876	UNRESTRICTED REVENUES
$158,965	TOTAL REVENUES
$132,269	TOTAL EXPENDITURES
$105,275	PROGRAMMING
$27,784	SUPPORT
$38,594	AWARENESS & OUTREACH
$27,695	ISCA & RESEARCH
$9,202	IAAD
$3,949	FUNDRAISING
$25,045	MANAGEMENT & GENERAL

$26,696 NET OPERATING INCOME

OUR PARTNERS

ILDS & IFD

GLODERM

WORLD SKIN HEALTH COALITION

ERN SKIN

ISHR

STANDING VOICE

FONDATION PIERRE FABRE

VISION OF CHILDREN FOUNDATION

VISION FOR TOMORROW FOUNDATION

FONDATION VOIR & ENTENDRE
INSTITUT FORESIGHT

EADV

GLOBALSKIN

RARE DISEASES INTERNATIONAL

CLINTON GLOBAL INITIATIVE

NOAH

HPS NETWORK

GENESPOIR

INTERNATIONAL AGENCY FOR RESEARCH ON CANCER

UN MANDATE ON THE ENJOYMENT OF HUMAN RIGHTS BY PERSONS WITH ALBINISM

SCIENTISTS WE PARTNER WITH

ASIA

TAMIO SUZUKI
Yamagata University, Yamagata, Japan

WEI LI
Beijing Children‘s Hospital, Beijing, China

MIDDLE EAST

ANAT BLUMENFELD
Hadassah University Medical Center
Jerusalem, Israel

CLAUDIA YAHALOM
Hadassah University Medical Center
Jerusalem, Israel

AMERICA

BRIAN BROOKS
National Eye Institute, Bethesda, MD, USA

DAVID ADAMS
National Human Genome Research Institute
(NHGRI), Bethesda, MD, USA

STACIE LOFTUS
National Human Genome Research Institute
(NHGRI) Bethesda, MD, USA

ELENA OANCEA
Brown University, Providence, RI, USA

JONATHAN ZIPPIN
Weill Cornell Medicine, New York City, NY, USA

MICHAEL MARKS
Children‘s Hospital of Philadelphia (CHOP)
Philadelphia, PA, USA

JOSEPH CARROLL
Medical College of Wisconsin
Milwaukee, WI, USA

WILLIAM OETTING
University of Minnesota, Minneapolis, MN, USA

EUROPE

BENOIT ARVEILER
Bordeaux University Hospital, Bordeau, France

FANNY MORICE-PICARD
Bordeaux University Hospital, Bordeau, France

MATHIEU FIORE
Bordeaux University Hospital, Bordeau, France

SMAIL HADJ-RABIA
Hopital Necker-Enfants Malades
Institut Imagine, Paris, France

ALEXANDRA REBSAM
Institut de la Vision, Paris, France

KAREN GRØNSKOV
University of Copenhagen
Copenhagen, Denmark

MARIA VAN GENDEREN
University Medical Center Utrecht
Utrecht, Netherlands

PANAGIOTIS SERGOUNIOTIS
Manchester Royal Eye Hospital
Manchester Centre for Genomic Medicine
University of Manchester
Manchester, UK

MERVYN THOMAS
University of Leicester, Leicester, UK

MICHAEL HOFFMAN
University Eye Clinic, Magdeburg, Germany

OVERVIEW OF ALBINISM ORGANIZATIONS

ASIA

- CHINESE ORGANIZATION FOR ALBINISM
- JEEVAN TRUST
- SAHKAR CHARITABLE TRUST
- CENTRAL INDIA ALBINISM SOCIETY
- ALBINO INDONESIA FAMILY
- KOMUNITAS ALBINO INDONESIA / ALBINISM COMMUNITY OF INDONESIA
- JAPANESE ALBINISM NETWORK
- KUALA LUMPUR AND SELANGOR ALBINISM ASSOCIATION
- NATIONAL DISABLED ALBINO NEPAL
- PAKISTAN ALBINISM SOCIETY
- ALBINISM PHILIPPINES
- TAIWAN ALBINO CARING ASSOCIATION
- BẠCH GIA TRANG

MIDDLE EAST

- IRAN ALBINISM ASSOCIATION
- ALBI - ALBINISM ASSOCIATION OF ISRAEL
- ALBINO JORDAN
- ALBINIZM DERNEGI

NORTH AMERICA

- BORN TOO WHITE OTTAWA
- NOAH – NATIONAL ORGANIZATION FOR ALBINISM AND HYPOPIGMENTATION
- HERMANSKY-PUDLAK SYNDROME NETWORK
- THE ASSOCIATION OF AFRICAN AMERICAN ALBINISM AWARENESS LTD
- VISION FOR TOMORROW FOUNDATION
- THE ALBINISM ALLIANCE GROUP
- ALBINISMO EN DOMINICANA
- FONDATION ALBHA
- ALBINO FOUNDATION OF TRINIDAD AND TOBAGO
- VISION OF CHILDREN FOUNDATION

LATIN AMERICA

- ALBINISMO ARGENTINA
- FUNDACIÓN NACIONAL DE ALBINISMO SIMPLEMENTE AMIGOS
- ALBINOS DE MENDOZA
- ALBINOS DE MISIONES
- ALBINOS PATAGONIA
- ALBINOS ZONA SUR
- AMIGOS ALBINOS DE CÓRDOBA
- BUENOS AIRES - SIMPLEMENTE AMIGOS
- ORGANIZACIÓN LATINOAMERICANA DE ALBINISMO
- ALBINISMO SEM FRONTEIRAS
- ALBINOS DO MEU BRASIL E DO MUNDO
- ALBINOS ES
- ALBINOS(AS) DO NOSSO NORDESTE
- ASSOCIAÇÃO DAS PESSOAS COM ALBINISMO EM MATO GROSSO
- ASSOCIAÇAO DOS ALBINOS DE ALAGOAS
- INSTITUTO NÓBREGA
- UNIVERSIDADE FEDERAL DO RIO DE JANEIRO (UFRJ) - PROJETO DE EXTENSÃO UNIVERSITÁRIA
- ASSOCIAÇÃO DAS PESSOAS COM ALBINISMO NA BAHIA
- CORPORACIÓN ALBINOS CHILE
- ALBINOS DE CORAZÓN
- FUNDACIÓN ALBINOS POR COLOMBIA
- FUNDACIÓN VIVE FELIZ
- COMUNIDAD DE PERSONAS CON LA CONDICIÓN DE ALBINISMO EN ORIENTE CUBA
- COMUNIDAD ALBINA SANTO DOMINGO DE LOS TSÀCHILAS
- FUNDACIÓN DE ALBINOS PIEL DE ÁNGEL
- ORGANIZACIÓN NACIONAL DE ALBINISMO DEL ECUADOR
- ALBINOS DE GUATEMALA
- UNION GUATEMALTECA DE ALBINISMO
- ALBIN GIRL
- ALBINO LATINO
- FUNDACIÓN PIEL DE LUNA ALBINOS DE MÉXICO A.C.
- FUNDACIÓN BENÉFICA SOS ALBINOS PANAMÁ
- ALBINOS PARAGUAY
- ALBINOS PERÚ
- ALBINOS DEL URUGUAY
- ALBINOS DE VENEZUELA

EUROPE

- NOAH ALBINISMUS SELBSTHILFEGRUPPE E.V
- LIGUE POUR LA DÉFENSE DES ALBINOS
- ALBINISM EUROPE
- GENESPOIR
- ALBINA
- DANSK FORENING FOR ALBINISME
- SUOMEN ALBINISMIYHDISTYS RY
- ALBINISM FELLOWSHIP UK & IRELAND
- ALBINISMO.EU
- ALBINIT
- ASSOCIATION OMBRE ET LUMIÈRE MONACO
- NORSK FORENING FOR ALBINISME
- STOWARZYSZENIE NA RZECZ OSÓB Z ALBINIZMEM W POLSCE
- COMUNITATEA ALBINISM ROMANIA INTERNATIONALA
- RUSSIAN COORDINATING SUPPORT CENTRE FOR PATIENTS WITH ALBINISM
- ALBA
- OOGVERENIGING ALBINISME
- SVENSKAR MED ALBINISM! | ALBINISM SVERIGE
- SOLIDARITÉ POUR ALBINISME SUISSE

OCEANIA & PACIFIC

- ALBINISM FELLOWSHIP OF AUSTRALIA
- HEALTHY SKIN FIJI
- PARENTS OF CHILDREN WITH ALBINISM FIJI
- FIJI ALBINISM PROJECT
- ALBINISM TRUST
- CLARENCE SEBASTIAN FOUNDATION

OVERVIEW OF ALBINISM ORGANIZATIONS

AFRICA

ALDEIA NISSI

ASSOCIAÇÃO DE APOIO A PESSOAS ALBINAS DO CUNENE

ASSOCIAÇÃO DE APOIO DE ALBINOS DE ANGOLA

FRATERNIDADE ALBINISTA

GRUPO TEATRAL EXCESSO DE COR

GRUPO VOLUNTÁRIO DE APOIO A PESSOAS COM ALBINISMO EM ANGOLA

LIGA ANGOLANA PARA A DEFESA DOS ALBINOS

ASSOCIATION ALGÉRIENNE D'ALBINISME

ASSOCIATION MAROCAINE DES ALBINOS

MOVIMENTO PRO ALBINO EM ANGOLA

TSHIMOLOGO ASSOCIATION FOR ALBINISM

ALBINISM SOCIETY OF BOTSWANA

UNDER THE SAME SUN

ALBINISM SOCIETY OF ESWATINI

MINERVA - ALBINISM IN ESWATINI

STUKIE MOTSA FOUNDATION

SWAZILAND ASSOCIATION OF PERSONS WITH ALBINISM

VITILIGO AND ALBINISM ASSOCIATION

ALBINISM FOUNDATION OF EAST AFRICA

DR CHOKSEY ALBINISM FOUNDATION

KENYA ALBINO CHILD SUPPORT

KISII ALBINISM SUPPORT ORGANIZATION

KWALE ALBINISM GROUP

POSITIVE EXPOSURE KENYA

THE ALBINISM EMPOWERMENT NETWORK

BLACK ALBINISM

EMPOWERED ALBINISM GROUP

ALBINISM SOCIETY OF KENYA

ALBINISME MADAGASCAR

ASSOCIATION OF PERSONS WITH ALBINISM IN MALAWI

STANDING VOICE

KANIMAMBO - ASSOCIAÇÃO DE APOIO AO ALBINISMO

AMOR A VIDA

ASSOCIAÇÃO DE APOIO A ALBINOS DE MOÇAMBIQUE

ASSOCIAÇÃO ZE MANUEL PINTO

THE SHADE TREE PROJECT

NAMIBIAN ALBINO ASSOCIATION

SUPPORT IN NAMIBIA OF ALBINISM SUFFERERS REQUIRING ASSISTANCE

APHAD - ALIKAR CENTER FOR PEACE, HUMAN RIGHTS AND DEMOCRACY

SOMALI ALBINISM EMPOWERMENT ORGANIZATION

ALBINISM SOCIETY IN LIMPOPO

A BEAUX ALBINISM BEAUTY

ALBINISM ADVOCACY FOR ACCESS

ALBINISM RENAISSANCE

ALBINISM SOCIETY OF KWA ZULU NATAL

ALBINISM SOCIETY OF SOUTH AFRICA

BEAUTY BY NATURE ALBINISM SOCIETY

CHILDLINE SA

EASTERN CAPE ALBINISM SOCIETY SOUTH AFRICA

HUMAN RIGHTS MEDIA CENTER

IBUTHO LEZWE FOUNDATION

KWATSADUZA ALBINISM SOCIETY INITIATIVE

WESTERN CAPE ALBINISM AND HYPO-PIGMENTATION FOUNDATION

KHULISA SOCIAL SOLUTIONS

ALBINO PEACEMAKERS

GOLDEN AYA TANZANIA ORGANIZATION

GOOD HOPE STAR FOUNDATION

PROMOTION OF EDUCATION LINK ORGANIZATION

STICHTING INSIDE THE SAME

ALBINOS TANZANIA DEVELOPMENT FOUNDATION

ALL ABOUT ALBINISM CAMPAIGN ASSOCIATION

BRIGITTE ALFRED FOUNDATION

SMA LAY ASSOCIATION ALBINO SUPPORT PROJECT

TANZANIA ALBINO ECONOMIC AND HEALTH ORGANIZATION

TANZANIA ALBINISM SOCIETY

THE KESHO TRUST

SHADE

JOSEPHAT TORNER FOUNDATION EUROPE

ASANTE MARIAMU

AFRICA ALBINO FOUNDATION UGANDA

ALBINISM CRISIS OUTREACH

ASANTE ALBINO ASSOCIATION

LIRA DISTRICT ALBINO ASSOCIATION (LANGO SUB REGION)

LUWERO DISTRICT ALBINO ASSOCIATION

NAZIGO ALBINO ASSOCIATION (KAYUNGA)

NORTHERN UGANDA ALBINO ASSOCATION (NORTHERN UGANDA AND WEST NILE)

SITE FOR COMMUNITY SERVICES PROGRAMME

SOURCE OF THE NILE UNION OF PEOPLE WITH ALBINISM (NATIONWIDE COVERAGE)

TUSIMAME AFRICAN ALBINISM FOUNDATION

UGANDA ALBINISM AID PROJECT

ALBINISM UMBRELLA

EVERY CHILD MINISTRIES

GLOBAL AID MISSION

ALBINO FOUNDATION OF ZAMBIA

NATIONAL ALBINISM INITIATIVE NETWORKING OF ZAMBIA

ZAMBIAN ALBINISM MATTERS ORGANIZATION

ALBINISM MULTIPURPOSE COOPERATIVE

ALBINO GLOBAL FOUNDATION

ALBINO TRUST ZIMBABWE ONLINE

MISS ALBINISM ZIMBABWE TRUST

PRINCESS SAFETY CENTRE FOUNDATION

ALBINISM ALIVE INITIATIVE

ZIMBABWE ALBINO ASSOCIATION

ALBINO CHARITY ORGANISATION OF ZIMBABWE

AFRICA ALBINISM NETWORK

AFRICAN UNION FOR PERSONS WITH ALBINISM

ALBINOS SANS FRONTIERES

ASSOCIATION JHONY CHANCEL POUR LES ALBINOS

ASBL INTERNATIONALE VROUWEN

ASSOCIATION “ALBINOS CRÉATURES HUMAINES À PART ENTIÈRE”

ASSOCIATION POUR LA PROTECTION ET LA PROMOTION DES ALBINOS “ENTRE NOUS”

ASSOCIATION ARTISTIQUE ET CULTURELLE DES PERSONNES ALBINOS - BÉNIN

DIVINE CONNECTION WORLDWIDE

ONG VALEUR ALBINOS

FÉDÉRATION DES ASSOCIATIONS DE PERSONNES VIVANT AVEC L'ALBINISME EN AFRIQUE DE L'OUEST

ASSOCIATION ALBI INTERNATIONAL

ASSOCIATION BURKINABE POUR L'INTEGRATION DES PERSONNES ALBINOS

ASSOCIATION DES FEMMES ALBINOS DU BURKINA

ASSOCIATION POUR LA PROTECTION ET LA PROMOTION DES ALBINOS

ASSOCIATION SIMA

SOLIDARITÉ POUR L'INSERTION DES ALBINOS DU MALI

ASSOCIATION DES FEMMES ALBINOS ESPOIR DU BURUNDI

LIVE TOGETHER AS FAMILY

ORGANISATION DES PERSONNES ALBINOS DU BURUNDI

AFRICAN ALBINISM AMBASSADORS

AMICALE DES FEMMES ALBIBELLES

ASSOCIATION OF HANDICAPPED ALBINO YOUTHS OF CAMEROON AND AFRICA

OVERVIEW OF ALBINISM ORGANIZATIONS

- ASSOCIATION FEMMES ALBINOS DU CAMEROUN
- ASSOCIATION DE PARTAGE ET D'ENTRAIDE DES ALBINOS DU CAMEROUN
- ASSOCIATION NATIONALE DE PROMOTION ET DE PROTECTION DES DROITS DE L'HOMME
- ASSOCIATION FOCUS
- ASSOCIATION POUR LA PROMOTION DES ALBINOS AU CAMEROUN
- MR & MRS ALBINOS CAMEROUN
- VILLAGE D'ENFANTS ALBINOS DU CAMEROUN
- ASSOCIATION MONDIALE POUR LA DEFENSE DES INTERETS ET LA SOLIDARITE DES ALBINOS
- RÉSEAU DES ORGANISATIONS DES PERSONNES VIVANT AVEC ALBINISME D'AFRIQUE CENTRALE
- ALBICARE INTERNATIONAL
- ASSOCIATION NATIONALE DES ALBINOS DE CENTRAFRIQUE
- ASSOCIATION D'APPUI AUX ALBINOS DU TCHAD
- COMITÉ MISS ALBINOS
- FONDATION SALIF KEITA
- ACTION POUR L'ALBINOS (KASAI ORIENTAL)
- ALBISO
- ASBL PLUS DE COULEURS - FIÈREMENT NDUNDU
- ASSOCIATION DES ALBINOS DU SANKURU
- ASSOCIATION DES ALBINOS ET HANDICAPÉS DU KASAÏ CENTRAL
- ASSOCIATION FAMILLE D'ACTION MULTIFORME- COMMUNAUTÉ NATIONALE DES PERSONNES ATTEINTES D'ALBINISME
- ASSOCIATION POUR LA PROTECTION ET LE DEVELOPPEMENT DE LA PERSONNE ALBINOS
- CORBETTA ONG
- FONDATION JULIO LEVOTRE
- FONDATION PAULINE ALBINOS
- LES AMIS DE DEPHILL POUR LA SENSIBILISATION À L'ALBINISME
- ONG DES ALBINOS FONDATION MWINMBA TEXAS
- ORGANISATION DES PERSONNES ENGANGEES POUR LA CAUSE DES ALBINOS
- ORGANISATION POUR LE BIEN ETRE DES ALBINOS AU CONGO
- PARLEMENT DES FEMMES ALBINOS AU SUD-KIVU
- PROMOTION ET PROTECTION DES ALBINOS DU KASAÏ
- SOLIDARITÉ DES ALBINOS DU KASAÏ ORIENTAL
- VOICE OF ALBINO ONG
- HOPE OF ALBINISM
- ASSOCIATION COMPASSION ALBINOS
- AGIR POUR LA CAUSE DES ALBINOS RDC - (GOMA, NORD KIVU)
- ASSOCIATION DE LUTTE POUR LE BIEN-ETRE DES ALBINOS
- ASSOCIATION OF GAMBIAN ALBINOS
- ASSOCIATION NATIONALE DES ALBINOS DU SÉNÉGAL
- ENGAGE NOW AFRICA
- GHANA ASSOCIATION OF PERSONS WITH ALBINISM
- CONFEDERATION NATIONALE DES ALBINOS DE GUINÉE
- UNION POUR LE BIEN ETRE DES ALBINOS DE GUINÉE
- FONDATION POUR LE SECOURS ET L'INTEGRATION SOCIALE DES ALBINOS
- ASSOCIACAO DOS ALBINOS DA GUINEA-BISSAU
- ACTION ET SOLIDARITÉ POUR LES ALBINOS DE CÔTE D'IVOIRE
- ASSOCIATION IVOIRIENNE POUR LA PROMOTION DES FEMMES ALBINOS
- ASSOCIATION NATIONALE DES ALBINOS DE CÔTE D'IVOIRE
- ASSOCIATION POUR LE BIEN-ÊTRE DES ALBINOS DE BOUAKÉ
- ONG BIEN ÊTRE DES ALBINOS DE CÔTE D'IVOIRE
- FÉDÉRATION DES ASSOCIATIONS POUR LE BIEN-ÊTRE DES PERSONNES ALBINOS DE CÔTE D'IVOIRE
- LIBERIA ALBINISM SOCIETY
- STAND-UP STAND-OUT ALBINISM FOUNDATION
- UNITED ALBINOS ASSOCIATION OF LIBERIA
- ASSOCIATION MALIENNE POUR LA DÉFENSE DES DROITS ET LE BIEN-ÊTRE DES ALBINOS
- ASSOCIATION MALIENNE POUR LA PROTECTION DES ALBINOS
- ASSOCIATION MALIENNE POUR LA SENSIBILISATION, LA PROMOTION ET LA PROTECTION DES PERSONNES VIVANTES AVEC ALBINISME - SOS ALBINOS
- COALITION DES ORGANISATIONS DE PERSONNES ATTEINTES D'ALBINISME
- THE SALIF KEITA GLOBAL FOUNDATION
- ORGANISATION MAURITANIENNE POUR L'APPUI ET L'INSERTION DES ALBINOS
- ASSOCIATION NATIONALE DES ALBINOS DU NIGER
- ANAMBRA STATE ALBINISM ASSOCIATION
- THE ALBINISM NETWORK ASSOCIATION
- ONOME AKINLOLU MAJARO FOUNDATION
- WHITE ANGELS FOUNDATION
- THE ALBINO FOUNDATION
- ORGANIZATION FOR INTEGRATION AND PROMOTION OF PEOPLE WITH ALBINISM
- RWANDA ALBINISM SOCIETY
- ALLIANCE NATIONALE POUR LA PROMOTION ET LA RÉINSERTION DES ALBINOS AU SÉNÉGAL
- CLUB ALBINOS SN
- FÉDÉRATION NATIONALE DES ALBINOS SÉNÉGAL
- AROBAZ ALBINISME AU SÉNÉGAL
- CARE ALBINOS
- ALBINISM ROYAL FOUNDATION SIERRA LEONE
- SIERRA LEONE ALBINISM FOUNDATION
- SIERRA LEONE ASSOCIATION FOR PERSONS WITH ALBINISM
- GOLDEN SHADE
- ASSOCIATION INITIATIVE PRO.TO.P.A
- ASSOCIATION NATIONALE DES PERSONNES ATTEINTES D'ALBINISME AU TOGO
- CAS SOCIAL ALBINOS ENFANTS ADULTES
- LES ALBINOS DE HEMA NAYÉLÉ À BANFORA
- AISBL ECRAN TOTAL
- ANIDA ENSEMBLE CHANGEONS LE REGARD SUR L'ALBINISME
- ALBINISM INCLUSION AND EQUALITY ORGANIZATION
- CENTRE CANADIEN DE SENSIBILISATION À L'AMÉLANISME
- BEYOND SUNCARE
- ENGAGE NOW AFRICA
- FONDATION PIERRE FABRE
- SARAH'S VOICE
- VOICE OF ALBINISM

albinismalliance.org