



# ISCA 2022

International Scientific Conference on Albinism

*A global forum to improve the health of those with albinism*

Online conference | 16-19 Nov. 2022

## Sponsor Prospectus

**ISCA** (International Scientific Conference on Albinism) is a unique international scientific conference gathering participants from all over the world. This meeting includes **researchers, health care providers and albinism organization leaders** all sharing the same vision: to improve the health of those with albinism. ISCA is the only conference in the world to address all medical aspects of albinism.



**November 16th–19th, 2022**  
**Virtual Conference**



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## About ISCA / Nov. 16–19, 2022

During the four-day virtual event, about 50 albinism experts will address the many health concerns of persons with albinism. This includes dermatology, ophthalmology, psychology, genetics, therapies, syndromic forms of albinism and more, contributing to a rich and educational event. It is anticipated that over 150 scientists will attend in ISCA 2022 along with 80 others including students and albinism group leaders. This is a unique opportunity to reach the most prominent members of the worldwide albinism community.

### The Main Objectives of ISCA are to:

- Improve the clinical and molecular diagnosis of albinism
- Improve medical care for persons with albinism
- Promote and facilitate albinism focused research programs
- Stimulate the development of therapies to treat albinism
- Put medical professionals and researchers in contact at both the local and international levels and foster collaboration in albinism treatment and investigation
- Facilitate the exchange of best practices for treating persons with albinism among medical professionals and patient organizations

### Global Albinism Alliance

#### Scientific Advisory Committee Members:

Lluís Montoliu – Chair, Spain  
Maria Mies van Genderen –  
Vice-Chair, The Netherlands  
Benoit Arveiler, France  
Brian Brooks, USA  
Irene Gottlob, USA  
Karen Gronskov, Denmark  
Michael Hoffmann, Germany

Barbara Kaesmann-Kellner,  
Germany  
Michael Marks, USA  
Fanny Morice-Picard, France  
Mathieu Fiore, France  
Joseph Carroll, USA  
Carolina Reato Marçon, Brazil  
Alexandra Rebsam, France  
Mervyn Thomas, UK

For complete information about the GAA and ISCA see  
<http://isca2022.albinismalliance.org>

Sponsoring ISCA aligns well with the GAA's vision of a world where people with albinism enjoy all their human rights. Access to health is a basic human right. Collaboration between research and clinicians will lead to a better quality of life for hundreds of thousands of persons living with albinism.

### About the Global Albinism Alliance

GAA is a non-profit organization registered in the State of Delaware, USA. It brings together albinism organizations from all regions of the world.

**GAA Vision:** A world where persons with albinism enjoy all of their human rights.

**GAA Mission:** To advance the capacity of albinism organizations around the world, to promote greater knowledge of albinism, and to advocate at the global level to improve the enjoyment of human rights by persons with albinism and their family members.

**GAA Activities** are organized according to three pillars:

- member support,
- advocacy and education
- research and information dissemination

Sponsored by



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### Benefits

	Bronze \$2,500	Silver \$5,000	Gold \$7,500	Platinum \$10,000+
Registration for up to five members of your organization	•	•	•	•
The name and the logo of your company / foundation / society will be displayed on all ISCA 2022 communications	•	•	•	•
Information about your company / foundation / society will be emailed to conference attendees in an electronic packet	•	•	•	•
Your company / foundation / society will be mentioned and acknowledged in the opening session of ISCA 2022. The logo of your company will be included in the official conference slide deck template		•	•	•
A video (up to 90 seconds in length) about your company / foundation / society will be shown during one of the main sessions of ISCA 2022			•	
Your company / foundation / society will be offered the opportunity to give a four minute live presentation at the opening session. The logo of your company will be displayed on the virtual background to be used by all speakers				•

**For more information or to arrange to become a sponsor**

Please contact Mike McGowan at [mmcgowan@albinismalliance.org](mailto:mmcgowan@albinismalliance.org)



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