
Please note some of these events are for information only and are not open to the general public. If you would like more information, please contact us. For more information about upcoming events, please visit our website www.isg-uk.org.

The ISG has a focus on tackling stigma and discrimination. However, you can also get good advice from the Action Medical Research, for children, for life. Action Medical Research is a UK charity that supports and represents people who have disfigurements of the face or body from any cause, as well as children with a range of health conditions.

Contact a Family is a UK charity that supports and represents families of children with disabilities and who have been diagnosed with a complex illness, such as ichthyosis. The ISG works closely with other organisations, including the British Association of Dermatologists, the British Skin Foundation, the Genetic Alliance UK, the Specialised Healthcare Alliance, the British Dysmorphology Society and the Specialist Dermatologists in the UK and Eire. The ISG is also a member of the Specialised Healthcare Alliance.

The ISG has a number of initiatives, including a youth forum, which may have some helpful advice. The ISG also has many new initiatives, including a regional contacts for local support and events; lobby for better service provision for patients. The ISG works closely with other organisations, including the British Association of Dermatologists, the British Skin Foundation, the Genetic Alliance UK, the Specialised Healthcare Alliance, the British Dysmorphology Society and the Specialist Dermatologists in the UK and Eire. The ISG is also a member of the Specialised Healthcare Alliance.

The ISG will continue to provide this service by developing and improving upon the leaflets, rewording and design of our literature, producing information that is both informative and user-friendly. The ISG will continue to produce and circulate a regular newsletter, updates, and communications to its members.

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