

Epidermolysis Bullosa

is a rare disease and the DEBRA International Congress 2017 offers you the perfect opportunity to learn from the experts, those who live with EB every day. This is your chance to network with DEBRA Groups and EB researchers and clinicians from around the world.

Increasingly, health care professionals are becoming aware of the importance of developing close relationships with patient-led organisations thus enhancing and progressing their own professional development.

The DEBRA International Congress is acknowledged as a premier event for those working in the EB field to both learn and share knowledge of this condition.

We know and understand the value of networking among a small pool of international health care professionals with experience in EB. This year's Congress will be a notable learning opportunity and will also provide many great occasions to meet and build rapport with your professional peers from a variety of backgrounds. This is your chance to develop lasting relationships enabling comfortable and easy future communication.

The programme for 2017 will be dynamic, covering a wide range of clinical and research projects, DEBRA supported initiatives, and will feature first class presenters from around the world.

Registration

will open in the New Year but now is the time to book your leave, talk to your supervisors and clear your calendar. Now's the time to do whatever you need to do so that you can attend this conference.

Be part of the passionate and committed global team that is making a difference to the lives of those affected by EB.

*Save
the date
24-26 Nov
2017*


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**INTERNATIONAL
CONGRESS**

www.debra.org.nz

24-26 November 2017 | Wellington, New Zealand