

ARE YOU:

- ✓ A professional (medical doctor, psychologist, nurse, or researcher) who has experience with patients with congenital naevi >10 cm on the body (or >1.5 cm on the face)?
- ✓ OR a patient (or parent of a patient) with congenital naevi >10 cm on your body (or >1.5 cm on the face)?
- ✓ Interested in improving research and care delivery for patients with congenital naevi?
- ✓ Available for online surveys in October - November 2018?

YOU CAN HELP PARTICIPATE IN OUR PROJECT!

AIM OF THE PROJECT

The Outcome measures for Congenital Melanocytic Naevi (OCOMEN) project aims to develop an international consensus on the important outcomes to measure (the so-called Core Outcome Set) for patients with Congenital Melanocytic Naevi (CMN).

The results can be used to:

- Harmonize outcomes in future CMN research
- Enable comparison of effectiveness of treatments
- Set up treatment guidelines for CMN



PARTICIPATE by JOINING:

Our **Online Delphi surveys** in **October-November 2018**. There will be 3 survey rounds in which you can evaluate and rate the importance of a list of outcome measures. Each round will take 20-30 minutes and you will have 1-2 weeks to complete it. If you complete the first round, you will be invited to participate in the subsequent rounds.

If you want to participate, kindly send us:

- Your name
- Background (professional, patient/parent)
- to Welling Oei before 30th Sept 2018 at w.oei@erasmusmc.nl

Your participation is highly valuable to us!
We would like to thank you in advance for your kind support.

On behalf of the OCOMEN working group

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