The Netherlands CP register will be built up in stages. Visit www.cpregister.nl if you want to participate, and to check if the hospital or rehabilitation center of your child is already working with the register.

If you have any questions, you can always contact your practitioner.

By participating, you and your child are contributing to better care for children with CP!

The Netherlands CP register has been created by the Dutch CP register Consortium:

Amsterdam UMC, CP Nederland, CP-Net, Nederlandse Vereniging van Revalidatieartsen, Adelante, De Hoogstraat Revalidatie, Erasmus MC Rotterdam, Heliomare, Libra revalidatie & audiologie, Maastricht UMC+, Merem Medische Revalidatie, Revant, Rijndam Revalidatie, Roessingh Centrum voor Revalidatie, UMCG, UMC Utrecht.

The Netherlands CP register is made possible by:

ZonMw · STEPTEMBER · Innovatiefonds · HandicapNL · dioraphte

photography: Mladen Pikulić | graphic design: burodeboo

nederlands cp register



The Netherlands CP register collects information from all children with cerebral palsy in the Netherlands. Do you want to participate?

Together we can improve the care for children with cerebral palsy!

CP folder A5 ouders Engels DEF•.indd 1 19/01/2021 16:01

The Netherlands CP Register

Your child has cerebral palsy (CP), a movement and posture disorder caused by a brain lesion during early development. Every child with CP is different. In order to better adapt the care to each individual child, we have developed the Netherlands CP Register. This register collects data from all children with CP in the Netherlands.

You and your child can also participate in the register!

How does participation help?

The register helps the practitioner to monitor the development of your child, to detect problems in time, and to see whether a treatment has worked. The register also provides a great deal of knowledge about the long-term effects of treatments. This allows us to better anticipate which treatment works best for specific children with CP. You can access the data by using the register portal, so that you always remain up to date.

What data do we collect?

We do not include any additional data in the register: only information that the practitioner needs to provide the proper care. This includes data about CP, medical history, physical examination, psychosocial functioning, treatment information, and treatment results.

To be added by you: Patient Reported Outcome Measures (PROMs)

For a good assessment of the treatment results and development of your child, it is important to have information from your side as well. That is why we ask you, together with your child, to enter a number of questionnaires in the register.

What happens to your child's data?

Data from all children with CP in the Netherlands are collected and compared. Your child's data is safely stored in the register. Only you and the person treating your child can view all the data, including personal information. Researchers only have access to anonymized data.

Do you want to participate?

