



**MyCP** is a web portal for community participation in research through discussions and surveys in the CPRN community registry owned and operated by the **Cerebral Palsy Research Network** (https://cprn.org).

## WHY JOIN US:



# What does it mean to participate in the CPRN Community Registry for cerebral palsy research?

·You will participate in online, secure, anonymous surveys about your experience with cerebral palsy.

·You choose which surveys to participate in.

·You can opt out at any time.

·You also have the option to be contacted about possibly participating in clinical research studies.

# Why should you participate in MyCP?



- ·Advance the pace of research in cerebral palsy.
- ·Help determine the most important issues for people with CP.
- ·Enable new research that cannot be done without your direct input.
- ·Allow you to optionally be contacted to participate in clinical research.
- ·Help understand the natural history of cerebral palsy.



#### What about your privacy and security of my medical information?

•Privacy and security of the information you provide for research will be stored on secure servers at the University of Utah.

·Research will be conducted under the ethics approval of the University of Utah Institutional Review Board (IRB\_00107418).

How do you register?

Go to MyCerebralPalsy.org

and click Join MyCP!

### How do you learn more about MyCP?

Visit the MyCP website https://mycerebralpalsy.org for complete information.

If you have questions about the CPRN Community
Registry (IRB\_00107418) and want
to email or talk to someone, please contact: