What is SWEET?

The SWEET database is an international registry that collects anonymous data from clinics around the world to better understand trends in

Who can participate?

diabetes

Over 40 centers internationally are participating in the SWEET registry, with over 2000 participants from **Canada enrolled.**

At CHEO we invite all patients with diabetes to take part in the study.

How do I participate?

Your participation is voluntary and no extra visits or tests are needed. If you agree to participate, the study team will collect routine information from your chart. (such as diabetes duration, height, weight, bloodwork data, treatment, etc. The SWEET database will not include any information that can directly identify you.



Routine Clinical Data of children and youth with diabetes: Scientific analysis and international comparison among pediatric diabetes centers

The purpose of the SWEET database is to detect outcomes and differences in diabetes care between pediatric diabetes centers internationally, with hopes to improve treatment.

If you are interested in learning more about this study and/or willing to participate, please visit the site below. This site will direct you to more information and give you the opportunity to electronically sign if interested in participating.

If you have any questions, you can contact the study research assistant, Brittney Bosse, at 613-737-7600 ext. 4156 or by email at bbosse@cheo.on.ca.

https://redcap.link/SWEET_diabetes_study







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This research project was approved by the CHEO Research Ethics Board.

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