

# Congenital Anomalies SUPPORT YUKON (CASY)

Approximately 420 babies are born to Yukoners every year. The majority are born healthy; however, about 3-5% are born with a birth defect. Over half of birth defects have no known cause.

#### Why does CASY exist?

CASY collects information on birth defects to assist in understanding them and improving maternal and child public health programs. A subset of deidentified information is shared with the Canadian Congenital Anomalies Surveillance System to help understand birth defects Canada wide.

# Who to report/refer to CASY?

All children with a diagnosis or strong suspicion of a congenital anomaly up to 5 years of age should be reported to CASY. When a termination of pregnancy is due to a congenital anomaly, the anomaly should be reported.

If and when a genetic counselling appointment is seen as helpful, parents can be referred to the CASY Coordinator, a certified genetic counsellor.

# How to report/refer to CASY and why?

When a congenital anomaly is identified in a child or fetus, they can be reported and/or referred using the CASY Reporting/Referral form available at the coordinates below, on the following website (www.hss.gov.yk.ca/ casy.php) or on PLEXIA. Confirmatory documentation such as a discharge summary or consultation reports, should accompany the reporting form. They can be sent by fax or mail to the contact information below. Unless documentation is apparent, do not assume a report has already been sent. The coordinator will contact you for further information if necessary.

**Note:** Reporting of birth defects is done indirectly (without consent) to the CASY Coordinator under the authority of the CMOH under the *Public Health and Safety Act* for the purposes of public health surveillance. CASY is governed under the *Health Information Privacy and Management Act (HIPMA).* 

When a referral is made, the CASY Coordinator, a genetic counsellor, will discuss the diagnosis, family history, methods to reduce the risk of congenital anomalies, support services, etc with the family.

# Who should refer?

Physicians / Midwives / Nurses / Any health care provider.

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# Summary of anomalies to refer for

- Congenital anomalies that are confirmed postnatally up to age 5
- Congenital anomalies identified/strongly suspected in any terminated pregnancy or miscarriage
- Chromosome abnormalities
- Prenatally detected club foot, ventriculomegaly hydronephrosis ONLY if confirmed postnatally
- Metabolic disorders
- Confirmed FASD and autism (at any age)

#### Summary of what should NOT be reported

 Prenatal soft markers for aneuploidy (nuchal thickening, echogenic foci, echogenic bowel, choroid plexus cysts, pyelectasis)



#### **CASY Coordinator**

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