

Referral Process for Low-Risk Women with Soft Signs Identified on Prenatal Ultrasound in the Absence of Congenital Anomalies

INDICATION	ACTION REQUIRED
Isolated nuchal fold	Refer to Genetics - increased risk for Down syndrome, cardiac anomalies, and other concerns
Isolated echogenic bowel	Refer to Genetics - increased risk of cystic fibrosis, maternal infection, and other concerns
Isolated short femurs/ humeri (femur measurement that is less than 0.9 of expected for the gestational age, when the gestational age is ascertained by the biparietal diameter)	Refer to Genetics - increased risk of skeletal dysplasia
Mild ventriculomegaly (lateral cerebral ventricles measuring between 10-15mm)	Refer to Genetics - increased risk of maternal infection and other concerns
Mild pyelectasis	Refer to Fetal Assessment Unit - for genitourinary follow-up
Other isolated soft sign	No Action Needed
Echogenic intracardiac focus/foci with choroid plexus cyst/cysts.	No Action Needed
Other cases of multiple soft signs	Refer to Genetics
Major congenital anomalies, irrespective of soft signs	Refer to Genetics
Positive aneuploidy screen (e.g., maternal serum screen, non-invasive prenatal testing) irrespective of soft signs	Refer to Genetics

Call the Program of Genetics & Metabolism (204-787-4631) and ask to speak with the prenatal genetic counsellor on-call if you are uncertain about whether an individual meets referral criteria.