

MEMO

To: BCW sonographers, reporting MDs and Obstetrical Care providers
From: Chantal Mayer, Medical lead BCW Ultrasound
Date: January 25, 2021
Re: **NT assessment in pregnancies with NIPT or PGT-A**

Due to limited capacity, BCW offers only a limited number of appointments for NT (nuchal translucency) assessment and is currently unable to offer routine 11-14 week examinations unless referred on the basis of a suspected anomaly.

Referrals for NT assessment are triaged and booked according to the Perinatal Services BC (PSBC) criteria for [eligibility for IPS](#). NT measurement for the purpose of calculation of aneuploidy risk is not currently indicated where NIPT (Non Invasive Prenatal Testing) or PGT- A (Pre-implantation Genetic Testing for Aneuploidy) was performed. This is because, although not diagnostic, the detection rates and accuracy of these tests far exceed that of the NT/SIPS.

Patients with either NIPT or PGT-A are occasionally inadvertently booked for NT assessment at BCW. This typically happens when:

1. The requisition did not specify that PGT-A was performed or that NIPT was planned or performed in a patient who is otherwise a candidate for NT.
2. A booking error occurred.

When a patient who is considered ineligible for NT assessment is examined at 11-14 weeks GA at in the BCW Main ultrasound Department, the following procedure is followed:

1. An ultrasound examination is performed as per PSBC imaging standards which includes a limited fetal anatomical survey.
2. The nuchal region is assessed as part of the anatomical survey; however, the NT measurement is not reported for the purpose of Trisomy 21 risk calculation.
3. If a fetal anomaly is identified, *including an NT measurement greater than 3.5 mm* or a cystic hygroma, the fetal anomaly is reported along with recommendation for referral to Medical Genetics.

References:

1. [BC prenatal Genetic Program: reference cards of key screening table and flow diagrams \(April 2019\) table 3](#)
2. [SOGC Committee Opinion No. 406](#): Prenatal testing after IVF with preimplantation genetic testing for aneuploidy. Nov 2020