ALBERTA PRECISION LABORATORIES

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| DATE: | 2021 April 06 |
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| TO: | All Zones |
| FROM: | Genetics and Genomics, Cytogenetics North & South Laboratories |
| | Discontinuation of Sodium Heparin (NaHep) blood specimen collection and processing for Chromosomal Microarray (CMA) follow-up testing |

PLEASE POST OR DISTRIBUTE AS WIDELY AS APPROPRIATE

Key Message:

- Effective April 5, 2021, only one EDTA blood specimen will be collected for postnatal Chromosomal Microarray (CMA) testing. The Sodium Heparin (NaHep) blood specimen will no longer be required.
- Effective April 5, 2021, any NaHep blood specimens inadvertently collected and received by the Cytogenetic laboratories will not be processed.
- For cases requiring follow-up testing, recommendations will be made in the CMA report.

Background:

- Current practice is to collect a blood specimen in both an EDTA and a NaHep tube for patients requiring postnatal CMA testing. CMA testing is performed on DNA extracted from the EDTA blood specimen, while the NaHep blood specimen is processed for potential follow-up Fluorescence In Situ Hybridization (FISH) and/or Karyotype testing.
- A recent review of cases revealed that follow-up FISH and/or Karyotype testing is performed in less than 5% of cases. This change is a quality improvement and cost savings initiative, as a significant number of specimens are being collected and processed at significant cost to the healthcare system.

How this will impact you:

- The test directory changes reflect that only EDTA blood specimens are required for postnatal CMA testing.
- Occassionally, test orders and subsequent sample collection for follow-up testing will be required, indicated in the CMA report.

Action Required:

- For palliative patients, infants, cases where an aneuploidy is suspected, or in situations where sample recollection would be extraordinarily difficult, a test orderable for constitutional FISH and/or Karyotype testing, indicating "<u>hold for follow-up</u>", should be used so that a blood in a NaHep tube is collected and processed, supporting potential follow-up testing (see below for links).
- For test requests submitted using paper requisitions, please use the most recent version; for orders submitted electronically through Connect Care the electronic requisition has not be altered.
- The most recent version of Cytogenetic requisitions are available: Chromosomal Microarray (CMA) requisition - <u>https://www.albertahealthservices.ca/frm-09591.pdf</u> Constitutional Cytogenetics requisition (FISH and/or Karyotype testing) <u>https://www.albertahealthservices.ca/frm-21248.pdf</u>
- Follow APL test directory instructions for CMA sample collection: https://www.albertahealthservices.ca/webapps/labservices/indexAPL.asp



Effective: April 5, 2021

Questions/Concerns:

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Approved by:

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