

October 20, 2010

Dear Physician/Midwife/Nurse Practitioner,

Canadian Blood Services will be implementing a new Laboratory Information System (MAK TRACE LINE) on November 22, 2010 for use in its Patient Services Laboratory in Vancouver. The change to the TRACE LINE system provides a stable, state of the art computer environment for perinatal testing.

Here are the changes you should be aware of:

1. Revised Perinatal Results Report

- The appearance of the reports will change. See attached example. Note: date format is YYYY-MM-DD.
- Maternal information can be linked to reports fathers if maternal information including expected date of delivery (EDC) is provided on the requisition.

2. Report Send-outs

- Perinatal reports will now be faxed instead of mailed to the ordering physician. Each patient report can be sent to multiple health care providers if full first and last name, address, phone number, and fax number of each physician requiring a copy is included. Please ensure all information for sending reports is on the requisition and is legible. A copy of all perinatal reports will also be faxed to the delivery hospital.

Examples of the attached reports can be copied and distributed for training purposes within your office. Thank you for your support and cooperation as we make the necessary changes to upgrade our information technology infrastructure. If you have any questions, please contact Tony Dolnik, Diagnostic Services Manager, at 604-707-3481 or tony.dolnik@blood.ca.

Sincerely yours,

Gershon Growe, MD, FRCPC
Medical Director, BC & Yukon

Copy: Tony Dolnik, Diagnostic Services Manager
Janet Unrau, Hospital Liaison Specialist

encl



TRANSFUSION MEDICINE RESULTS REPORT

Sample Number:

Date Received:
2009-10-21
Request Number:

Trace Line Number:
#####



Date Printed: 2009-10-21 10:00

Patient Summary

Blood Group: B Pos
Phenotype: C+ E- c- e+
Transfusion Protocols:
Patient must receive E- c- red cells

SURNAME, FIRST NAME**PHN:** BC 9999 999 999**DOB:** 1970-01-01**Sex:** FEMALE**Other ID:** V10-1234**Ordering Facility:** University Hospital**Physician:** Jones, John**Receiving Facility:** University Hospital**Sample Comments****Date Collected:** 2009-10-21**Tests Performed:**

ABO/Rh

Antibody Screen

Antibody Identification
Detected using Peg IATAntibody Identification
Detected using Peg IAT**Results:**

B Pos

Positive




Anti-E

Anti-c

Remarks

Anti-E and anti-c implicated in hemolytic transfusion reactions. Antibodies to other major blood group antigens were excluded by PEG-IAT test method.

PERINATAL RESULTS REPORT

 <p>Canadian Blood Services Société canadienne du sang</p>		<p>SURNAME, FIRST NAME</p> <p>PHN: AB 99999 9999</p>  <p>DOB: 1970-01-01</p>		<p>PHN is scannable and eye readable, the 2 letters indicate the patient's province of residency and is not included in the bar code</p>																									
<p>Sample Number: #####</p> <p>Date Received: 2009-10-21</p> <p>Request Number: #####</p> <p>Trace Line Number: #####</p> 		<p>Other ID: RCMP748392</p> <p>Medical Record Number: 44568</p> <p>Ordering Facility: General Hospital</p> <p>Physician: Jones, John</p>		<p>Other ID and Medical Record Number will only print if this information is provided</p>																									
<p>Date Printed: 2009-10-21 10:00</p>		<p>Receiving Facility: General Hospital</p>		<p>Hospital for delivery</p>																									
<p>Patient Summary</p> <p>Blood Group: O Neg</p> <p>Phenotype:</p> <p>Known Antibodies:</p>																													
<p>Sample Comments</p>		<p>Date Collected: 2009-10-21</p>																											
<p>Tests Performed:</p> <p>ABO/Rh</p> <p>Test Comment: Antibody Screen</p>		<p>Results:</p> <p>O Neg</p> <p>Negative</p>																											
<p>Sample Comments provide information about problems with sample quality or labeling</p>		<p>Results for current sample</p>																											
<p>Remarks</p> <p>Small chance (<1%) patient has a weak or variant D antigen and may be reported as Rh positive by another laboratory.</p>																													
<p>Remarks will display important information about the management of the patient</p>																													
<p>Guidelines For Perinatal Testing</p> <table border="1"> <thead> <tr> <th></th> <th>Initial Visit</th> <th>Father</th> <th>25 - 28 Weeks</th> <th>As Requested</th> </tr> </thead> <tbody> <tr> <td>First pregnancy</td> <td>X</td> <td></td> <td>X</td> <td></td> </tr> <tr> <td>Rh positive previous reportable</td> <td>X</td> <td></td> <td></td> <td></td> </tr> <tr> <td>Rh negative</td> <td>X</td> <td>If requested</td> <td>X</td> <td></td> </tr> <tr> <td>Clinically significant antibodies detected</td> <td>X</td> <td>X</td> <td></td> <td>X</td> </tr> </tbody> </table>						Initial Visit	Father	25 - 28 Weeks	As Requested	First pregnancy	X		X		Rh positive previous reportable	X				Rh negative	X	If requested	X		Clinically significant antibodies detected	X	X		X
	Initial Visit	Father	25 - 28 Weeks	As Requested																									
First pregnancy	X		X																										
Rh positive previous reportable	X																												
Rh negative	X	If requested	X																										
Clinically significant antibodies detected	X	X		X																									
<p>Additional samples may be submitted for patients at increased risk of allo-immunization (previous transfusion, fetal trauma or procedure, IV drug use).</p>																													