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|---|---|---|--------------------|------------------|
| Patient ID<br><b>SA00043162</b>                       | Patient Name<br><b>SAMPLEREPORT, PT11</b> | Birth Date<br><b>1966-06-10</b>             | Gender<br><b>F</b> | Age<br><b>45</b> |
| Order Number<br><b>SA00043162</b>                     | Client Order Number<br><b>SA00043162</b>  | Ordering Physician<br><b>Client, Client</b> | Report Notes       |                  |
| Account Information<br><b>C7028846 DLMP Rochester</b> |   | Collected<br><b>07 Feb 2012 13:00</b>       |                    |                  |

## DNA Extraction, NPL

**Specimen**

Blood

MCR

**Comment**

Genomic DNA has been extracted.

MCR

**Specimen ID**

1037723

MCR

**Reviewed By**

Melody Elizabeth Kimball

MCR

**Order Date**

09 Feb 2012 07:23

MCR

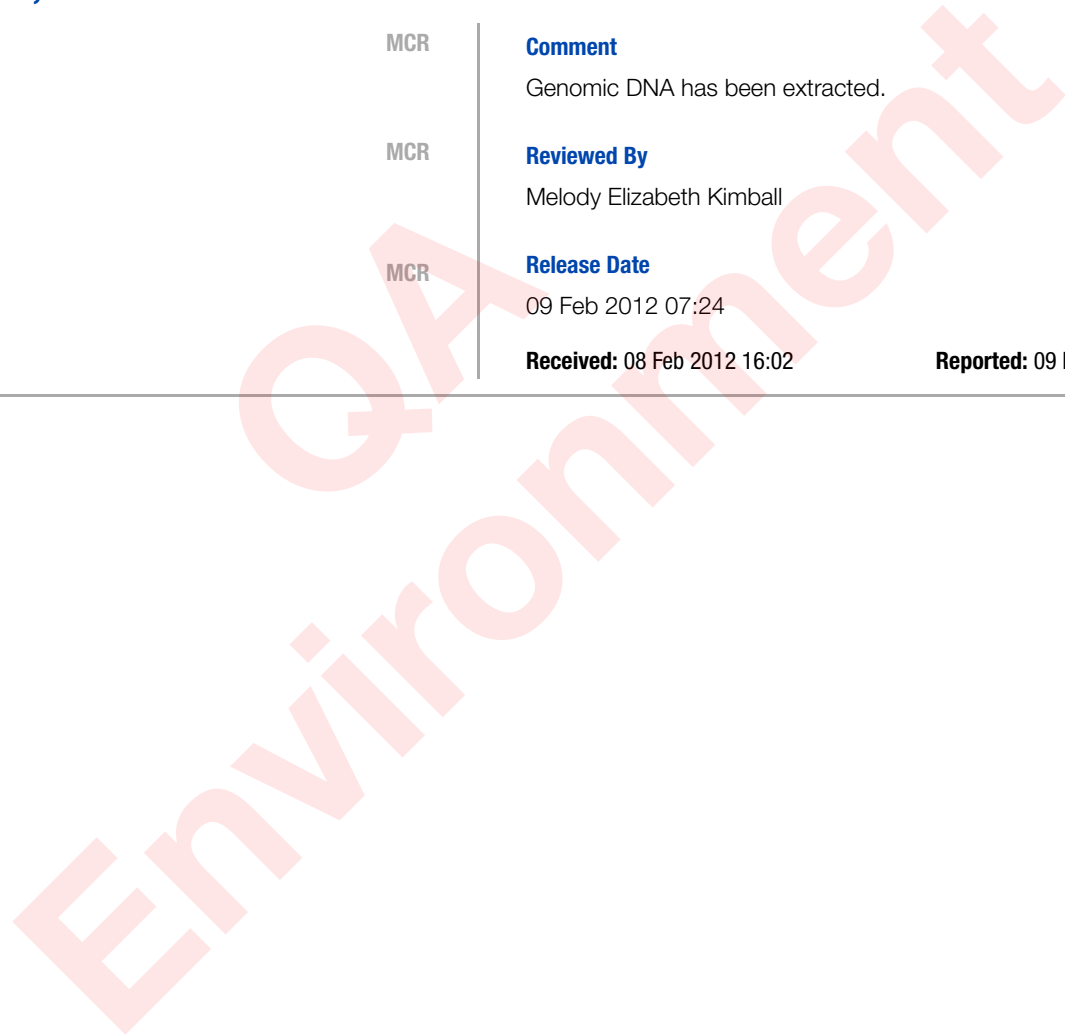
**Release Date**

09 Feb 2012 07:24

MCR

**Received:** 08 Feb 2012 16:02

**Reported:** 09 Feb 2012 07:25



**Performing Site Legend**

| Code | Laboratory                                 | Address                                  |
|------|--|--|
| MCR  | Mayo Clinic Dept. of Lab Med and Pathology | 200 First Street SW, Rochester, MN 55905 |

|   |  |   |                    |                  |
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| Account Information<br><b>C7028846 DLMP Rochester</b> |  | Collected<br><b>07 Feb 2012 13:00</b>       |                    |                  |

## PTPN11, Full Gene Sequence, B

### PTPN11, Full Gene Sequence

#### PTPN11 Full Gene Result

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A mutation was not detected in PTPN11.

#### PTPN11 Full Gene Interpretation

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This result does not rule out the diagnosis of Noonan syndrome or other PTPN11-associated syndrome. Some individuals with features of Noonan syndrome or other PTPN11-associated syndrome and involvement of the PTPN11 gene may have a mutation that is not identified by the described testing methodology. In addition, some individuals with this phenotype may have a mutation in a gene other than PTPN11, including RAF1, SOS1, and KRAS.

#### ADDITIONAL INFORMATION

Fluorescent DNA sequence analysis was used to test for the presence of mutations in all coding exons (1–15) and corresponding exon-intron boundaries of the PTPN11 gene (GenBank accession number NM\_002834.3).

A genetic consultation may be of benefit.

A list of common polymorphisms identified for this patient is available from the lab upon request.

#### CAUTIONS:

Rare polymorphisms exist that could lead to false negative or positive results. If results obtained do not match the clinical findings, additional testing should be considered.

Test results should be interpreted in the context of clinical findings, family history, and other laboratory data. Misinterpretation of results may occur if the information provided is inaccurate or incomplete.

If the patient has had an allogeneic blood or marrow transplant or a recent (i.e. less than 6 weeks from time of sample collection) heterologous blood transfusion these results may be inaccurate due to the presence of donor DNA. Laboratory developed test.

#### Reviewed By

MCR

Jamie Bruflat

### PTPN11, Full Gene Sequencing

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Performed

**Received:** 08 Feb 2012 16:02

**Reported:** 08 Feb 2012 16:06

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