

## Memo update- Thrombophilia Risk test back in-house

### **Who is this message for:**

New York Presbyterian Hospital/Weill Cornell Medicine healthcare providers and clients

### **What is happening:**

**Effective immediately**, Thrombophilia Risk Test for *Factor V Leiden* and *Factor II Prothrombin mutations* will be routed back to in-house testing using a new platform.

The corresponding tests provided by ARUP are listed below:

EPIC PROCEDURE (ORDERABLE)	Cerner Orderable	EPIC CODE	Emdeon Code
FACTOR V LEIDEN	FV Leiden	LAB6340	21813569
PROTHROMBIN (FACTOR II) MUTATION	FactorII	LAB6103	21811728
Methylenetetrahydrofolate Reductase (MTHFR)	MTHFR Mutation Detection PCR	LAB6170	4262492

### **What you will need to do:**

We will now require collection of 5 mL of peripheral blood (minimum 1 mL) into each of two lavender (EDTA) top tubes, the first for Factor II Prothrombin and Factor V Leiden mutation. Kindly be advised, testing for MTHFR Mutations (Methylenetetrahydrofolate Reductase) will still be *permanately* performed at ARUP Laboratories.

### **Why we are making this change:**

The manufacturer has discontinued production of reagents used for this test. We have completed the validation of a new in-house test for *Factor II Prothrombin Mutation* and *Factor V Leiden Mutation* that will now be performed on the Cepheid platform.

### **For any questions, please contact:**

Hanna Rennert, Director, Molecular Pathology  
[har2006@med.cornell.edu](mailto:har2006@med.cornell.edu), 212-746-6412

Liliana Serrano, Supervisor, Molecular Pathology  
[lis9134@nyp.org](mailto:lis9134@nyp.org), 212-746-2294

Outreach Client Services Department  
212-746-0670