TECHNICAL BULLETIN

⊠NEW TEST ⊠TEST CHANGE

NOTIFICATION DATE: 03/27/2023 EFFECTIVE DATE: 04/11/2023

Serotonin Releasing Assay (SRA)

On the effective date, Northwell Health Laboratories will transition to a reduced run schedule of 2 days per week for the Serotonin Releasing Assay (SRA) test performed by Versiti Diagnostics Laboratories. This is due to ongoing nationwide supplier constraints. We are unable to confirm when the SRA assay will resume its standard schedule of 6 days per week.

Northwell Health Laboratories recommends the **Heparin-Induced Thrombocytopenia-PEA Assay** performed at Versiti Diagnostic Laboratories as a time-sensitive alternative. The turnaround time for this assay is 48 hours and performed 6 days a week.

Clinical considerations for this assay include:

- The PEA is equivalent to the SRA as a functional assay for the laboratory evaluation of patients with suspected heparin induced thrombocytopenia (HIT). (1)
- The PEA performed at Versiti has an industry-leading turnaround time (48 hours), minimizing the time patients remain on alternative anticoagulation. Parenteral direct thrombin inhibitors have high associated bleeding risk in patients with suspected HIT. (2)

• HIT remains a clinico-pathologic diagnosis and functional assays are best utilized in patients with an intermediate or high pre-test (clinical) probability of HIT when PF4 serology is indeterminate.

Test Requirement/Parameters	Recommended Alternative Assay	Current Assay
	Heparin-Induced Thrombocytopenia-	(Reduced Run Schedule)
	PEA Assay	Serotonin Releasing Assay
Method	Flow Cytometry	Serotonin Release Assay
Specimen Requirements	5 mL serum (1 mL min.)	5 mL serum (1 mL min.)
Test performance	6 days per week	2 days per week
Submission Container/Tube	Red Top Tube	Red Top Tube
Turn Around Time	48 hrs	1-3 days
Specimen Stability	Sample should be spun down and taken off the clot. Plasma is NOT acceptable. Sample must be received within 7 days	Samples should be spun down and taken off the clot. Sample must be received within 7 days of
	of draw date if refrigerated. Send samples refrigerated.	draw date if refrigerated. Send samples refrigerated.
Reference Range	See Report	See Report
Computer Interface Code	PDM #235702	PDM #5910036
Test Order	PEA	SRA

References

- 1. Samuelson Bannow et al A prospective, blinded study of a PF4-dependent assay for HIT diagnosis. Blood 2021; 137 (8): 1082–1089. doi: https://doi.org/10.1182/blood.2020008195
- 2. Rivner H, Parmar R, Cardoso R, et al. a meta-analysis of bivalirudin versus argatroban for the treatment of heparin-induced thrombocytopenia. J Am Coll Cardiol. 2017 Mar, 69 (11_Supplement) 2064.https://doi.org/10.1016/S0735-1097(17)35453-0)

