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PATIENT REPORT

500 Chipeta Way, Salt Lake City, Utah 84108-1221 phone: 801-583-2787, toll free: 800-522-2787

Jonathan R. Genzen, MD, PhD, Chief Medical Officer

Patient Age/Sex: 32 years Female

Specimen Collected: 04-Jun-24 13:14

Preeclampsia, sFlt-1/PlGF Ratio, | Received: 04-Jun-24 13:14 Report/Verified: 04-Jun-24 13:17

S/P

Units Reference Interval Procedure Result

10 ⁱ¹ Preeclampsia sFlt-1/PlGF Ratio, [<=39]ratio

S/P

Test Information

Preeclampsia sFlt-1/PlGF Ratio, S/P

REFERENCE INTERVAL: Preeclampsia sFlt-1/PIGF Ratio, S/P

The sFlt-1/PIGF ratio is indicated to be used as an aid in the management of the patient and is a prognostic assay intended to stratify hospitalized patients into two risk groups (low risk and high risk of progression to preeclampsia with severe features within two weeks from presentation).

The clinical cutoff for the sFlt-1/PIGF ratio is 40.

If the result of the ratio is greater than or equal to 40, the pregnant woman would be at high risk for progression to preeclampsia with severe features within 2 weeks.

If the result of the ratio is less than 40, the pregnant woman would be at low risk for progression to preeclampsia with severe features within 2 weeks.

The assay results should only be used in conjunction with information available from clinical evaluations and other standard of care procedures. The test result is not to be used to replace clinical judgment.

*=Abnormal, #=Corrected, C=Critical, f=Result Footnote, H-High, i-Test Information, L-Low, t-Interpretive Text, @=Performing lab

Unless otherwise indicated, testing performed at:

ARUP Laboratories

500 Chipeta Way, Salt Lake City, UT 84108

Laboratory Director: Jonathan R. Genzen, MD, PhD

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