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PATIENT REPORT

Unknown

Patient Age/Sex:

Specimen Collected: 06-Sep-23 13:	34			
AB Ident Panel-Cold (IRL) Procedure Cold Panel Identification	Received: 06- Result Done	Sep-23 13:37 Units	Report/Verified: Refer	06-Sep-23 14:0 ence Interval
AB Ident Panel-PEG (IRL) Procedure PEG Panel Identification	Received: 06- Result Done ^{f1}	Sep-23 13:37 Units	Report/Verified: Refer	06-Sep-23 14:0 ence Interval
Elution And Antibody Identification, RBC	Received: 06-	Sep-23 13:37	Report/Verified:	06-Sep-23 14:0
Procedure Interp Elution	Result See Below ^t	Units	Refer	ence Interval
Extended Direct Antiglobulin Test	Received: 06-	Sep-23 13:37	Report/Verified:	06-Sep-23 14:0
Procedure	Result	Units	Refer	ence Interval
Int Poly	Positive			
Int IgG	Positive ^{f2}			
Int C3	Positive			
Int LISS IgG	Positive			
Int IgA	Negative			
Int IgM	Positive			
EXT DAT Interp	See Below ^t	2		

Interpretive Text

06-Sep-23 13:34 (Interp Elution) t1:

> The eluate was reactive with all cells tested showing no apparent specificity. This reactivity pattern is consistent with a warm autoantibody.

06-Sep-23 13:34 (EXT DAT Interp) t2:

A 2+ IgM and 3+ IgG autoantibodies were detected on the surface of red blood cells. Serologically, this represents a mixed autoimmune hemolytic anemia. Please correlate with the patient's clinical context.

Result Footnote

f1: PEG Panel Identification

> The patient's plasma was reactive with all cells tested, showing no apparent specificity, using routine test tube methods. This reactivity pattern is most commonly seen with a warm autoantibody however, this reactivity pattern is also consistent with an antibody directed against a high incidence antigen. Please correlate with the patient's clinical context.

If antibody identification is desired, please submit additional samples and order 0013003 IRL-AB PKG. Refer to the test directory for test information and sample requirements. Int IgG

f2:

Spontaneous agglutination was observed with routine testing methods. Patient cells were washed with warm 37C saline to prevent spontaneous agglutination and avoid false positive results.

*=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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