

# **Test Definition: FADBC**

Adenosine Deaminase, RBC

## **Overview**

#### **Useful For**

As a marker of severe combined immunodeficiency (SCID)

#### **Method Name**

Kinetic Spectrophotometry

#### **NY State Available**

No

# **Specimen**

### **Specimen Type**

Whole Blood EDTA

## **Specimen Required**

Container/Tube: Lavender top (EDTA), pink top (K2 EDTA), or green top (sodium or lithium heparin)

Specimen Volume: 1 mL

**Collection Instructions**: Send 1 mL whole blood refrigerate.

# **Specimen Minimum Volume**

0.5 mL

#### **Reject Due To**

Hemolysis	Reject	
	-	

# **Specimen Stability Information**

Specimen Type	Temperature	Time	Special Container
Whole Blood EDTA	Refrigerated (preferred)	15 days	
	Ambient	15 days	

# Clinical & Interpretive

# **Clinical Information**

Adenosine deaminase (ADA) deficiency is an autosomal recessive disorder of purine metabolism primarily affecting lymphocyte development, viability, and function. Lack of ADA allows deoxyadenosine to accumulate and kill lymphocytes.



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#### **Reference Values**

400 - 900 mU/g Hb

#### Interpretation

Affected individuals have less than 1% of normal adenosine deaminase (ADA) catalytic activity in red cell hemolysates. ADA deficiency is the cause of 20% to 30% of severe combined immunodeficiency (SCID) cases. Heterozygotes cannot be identified by this test.

#### **Cautions**

If the patient has been recently transfused, adenosine deaminase (ADA) deficiency may be masked; interpret results with caution.

#### **Performance**

### **PDF Report**

No

### Day(s) Performed

Sunday, Tuesday, Thursday

#### Report Available

3 to 8 days

#### **Performing Laboratory Location**

**ARUP Laboratories** 

#### **Fees & Codes**

#### **Fees**

- Authorized users can sign in to <u>Test Prices</u> for detailed fee information.
- Clients without access to Test Prices can contact <u>Customer Service</u> 24 hours a day, seven days a week.
- Prospective clients should contact their account representative. For assistance, contact <u>Customer Service</u>.

# **Test Classification**

This test was developed and its performance characteristics determined by ARUP Laboratories. The U.S. Food and Drug Administration has not approved or cleared this test; however, FDA clearance or approval is not currently required for clinical use. The results are not intended to be used as the sole means for clinical diagnosis or patient management decisions.

#### **CPT Code Information**

84311



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# **LOINC®** Information

Test ID	Test Order Name	Order LOINC® Value
FADBC	Adenosine Deaminase RBC	47549-1

Result ID	Test Result Name	Result LOINC® Value
FADBC	Adenosine Deaminase RBC	47549-1