

# **Test Definition: F68KD**

68kD (hsp-70) antibodies by Line Blot

_						
O	VP	r	V	Δ	W	

## **Method Name**

Line Immunoassay

#### **NY State Available**

Yes

### **Specimen**

# **Specimen Type**

Serum

# **Specimen Required**

Draw blood in a plain red-top tube(s), serum gel tube(s) is acceptable. Spin down and send 3 mL of serum frozen in a plastic vial.

#### **Specimen Minimum Volume**

2.0 mL

Reject Due To

## **Specimen Stability Information**

Specimen Type	Temperature	Time	Special Container
Serum	Frozen (preferred)	365 days	
	Ambient	48 hours	
	Refrigerated	5 days	

## **Clinical & Interpretive**

### **Reference Values**

Qualitative test - Positive or Negative

#### Interpretation

Antibodies to inner ear antigen (68kDa) occur in approximately 70% of patients with autoimmune hearing loss. The antibody tests to this 68kDa antigen parallel with disease activity. In addition, a majority of patients positive for antibodies to 68kDa are responsive to corticosteroid treatment. (Hirose et al. The Laryngoscope. 109:1769-1999)



# **Test Definition: F68KD**

68kD (hsp-70) antibodies by Line Blot

#### **Performance**

#### PDF Report

No

# Day(s) Performed

Once per week

#### **Report Available**

3 to 18 days

#### **Performing Laboratory Location**

IMMCO Diagnostics, Inc.

#### **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 has been developed and performance parameters have been validated by IMMCO Diagnostics, Inc. This test has not been approved by the U.S. Food and Drug Administration (FDA); however, US FDA approval is not required for clinical use. It is not intended that clinical diagnosis and patient management decisions be made using these results alone.

This test has been validated using serum samples. The manufacturer has not determined the efficacy of this test when performed on CSF, plasma, joint or pleural fluid specimens. The performance characteristics of this test were determined by IMMCO Diagnostics Inc.

#### **CPT Code Information**

84182

#### **LOINC®** Information

Test ID	Test Order Name	Order LOINC® Value
F68KD	68kD (hsp-70) antibodies	43597-4

Result ID	Test Result Name	Result LOINC® Value
Z0909	68kD (hsp-70) antibodies	43597-4