



Patient Information	Specimen Information	Client Information
<b>DOB:</b> <b>AGE:</b> <b>Gender:</b> <b>Patient ID:</b> <b>Health ID:</b>	<b>Specimen:</b> <b>Collected:</b> <b>Received:</b> <b>Reported:</b>	MOTA, CHRISTINA

## ALLERGEN REPORT

EGG COMPONENT PANEL	CLASS
Performing Lab: EN	0 1 2 3 4 5 6
<b>Test Name</b>	<b>Results kU/L</b>
OVALBUMIN (F232) IGE	<0.10

EGG COMPONENT PANEL	CLASS
Performing Lab: EN	0 1 2 3 4 5 6
<b>Test Name</b>	<b>Results kU/L</b>
OVOMUCOID (F233) IGE	<0.10
	<i>See Endnote 1</i>

### INTERPRETATION

Performing Lab:

See Endnote 2

**Endnote 1** IgE reactivity to whole egg without reactivity to Gal d 1 or Gal d 2 may be explained by IgE reactivity to other egg proteins or non-protein egg constituents. Additional information can be found at <http://www.phadia.com>

**Endnote 2**

Specific IGE Class	kU/L	Level of Allergen Specific IGE Antibody
0	<0.10	Absent/Undetectable
0/1	0.10-0.34	Very Low Level
1	0.35-0.69	Low Level
2	0.70-3.49	Moderate Level
3	3.50-17.4	High Level
4	17.5-49.9	Very High Level
5	50-100	Very High Level
6	>100	Very High Level

The clinical relevance of allergen results of 0.10-0.34 kU/L are undetermined and intended for specialist use.

Allergens denoted with a "\*\*\*" include results using one or more analyte specific reagents. In those cases, the test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by the U.S. Food and Drug Administration. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

### PERFORMING SITE: