


Patient ID SA00000630	Patient Name SAMPLEREPORT, RPR19	Birth Date 1945-07-10	Gender M	Age 69
Order Number SA00000630	Client Order Number SA00000630	Ordering Physician CLIENT, CLIENT	Report Notes	
Account Information C7028846 DLMP Rochester		Collected 05 Jul 2015 06:00		

Resp Profile, Reg 19, Puerto Rico

Immunoglobulin E (IgE), S

1800 kU/L
High

 SDL
 Reference Value
 ≤ 214

Johnson Grass, IgE
35.0 kU/L
 Class 4 (Strongly Positive 17.5–49.9)

SDL

House Dust Mites/D.P., IgE
1.00 kU/L
 Class 2 (Positive 0.70–3.49)

SDL

Cockroach, IgE
≥100 kU/L
 Class 6 (Strongly Positive ≥100)

SDL

House Dust Mites/D.F., IgE
4.00 kU/L
 Class 3 (Positive 3.50–17.4)

SDL

Penicillium, IgE
3.20 kU/L
 Class 2 (Positive 0.70–3.49)

SDL

Cat Epithelium, IgE
5.00 kU/L
 Class 3 (Positive 3.50–17.4)

SDL

Cladosporium, IgE
<0.35 kU/L
 Class 0 (Negative <0.35)

SDL

Dog Dander, IgE
6.00 kU/L
 Class 3 (Positive 3.50–17.4)

SDL

Aspergillus Fumigatus, IgE
<0.35 kU/L
 Class 0 (Negative <0.35)

SDL

Bermuda Grass, IgE
85.0 kU/L
 Class 5 (Strongly Positive 50.0–99.9)

SDL

Alternaria Tenuis, IgE
<0.35 kU/L
 Class 0 (Negative <0.35)

SDL

Red Top, IgE
25.0 kU/L
 Class 4 (Strongly Positive 17.5–49.9)

SDL

Aureobasidium Pullulans, IgE
<0.35 kU/L
 Class 0 (Negative <0.35)

SDL

Performing Site Legend

Code	Laboratory	Address	Lab Director	CLIA Certificate
SDL	Mayo Clinic Laboratories - Rochester Superior Drive	3050 Superior Drive NW, Rochester MN 55901	William G. Morice M.D. Ph.D	24D1040592

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Oak, IgE	SDL	Parietaria Officinalis, IgE	SDL
1.00 kU/L Class 2 (Positive 0.70–3.49)		<0.35 kU/L Class 0 (Negative <0.35)	
Elm, IgE	SDL	Rough Pigweed, IgE	SDL
2.00 kU/L Class 2 (Positive 0.70–3.49)		1.00 kU/L Class 2 (Positive 0.70–3.49)	
Eucalyptus, IgE	SDL	Australian Pine, IgE	SDL
<0.35 kU/L Class 0 (Negative <0.35)		<0.35 kU/L Class 0 (Negative <0.35)	
Red Sorrel, IgE	SDL	Blomia Tropicalis, IgE	1 SDL
1.00 kU/L Class 2 (Positive 0.70–3.49)		2.00 kU/L Class 2 (Positive 0.70–3.49)	
		Received: 06 Jul 2015 09:40	Reported: 06 Jul 2015 09:42

Laboratory Notes

- 1 Analyte Specific Reagent: This test was developed and its performance characteristics determined by Mayo Clinic. It has not been cleared or approved by the U.S. Food and Drug Administration.

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