

Patient Name: **SAMPLE, REPORT**  
 Accession Number: **512100**  
 Date of Birth: **1/1/1980** Age: **45** Sex: **F**  
 External ID: **ABC123**  
 Comments:

DoctorID: **6206 Dr. Sample Report ND**  
**The Healing Centre**  
 6839 Fort Dent Way  
 Green City, WA 98188  
 United States  
 Phone: 206-209-4200 Fax: 206-209-4211

Test: **2100**  
 Date Collected: **12/31/2025**  
 Date Received: **12/31/2025**  
 Date Reported: **12/31/2025**  
 Tech:

Category	Allergen	Result	Class	Class					
				0/1	I	II	III	IV	V
<b>Animals</b>	Cat Dander	5.71	III						
	<i>Dermatophagoides farinae</i>	0.11	0/1						
	Dog Dander	0.52	I						
<b>Grasses</b>	Bahia Grass	3.58	III						
	Meadow Grass, Kentucky Blue	11.98	III						
	Sweet Vernal Grass*	19.79	IV						
	Timothy Grass	14.80	III						
<b>Molds</b>	<i>Alternaria alternata</i>	< 0.1	Negative						
	<i>Aspergillus fumigatus*</i>	< 0.1	Negative						
	<i>Candida albicans*</i>	< 0.1	Negative						
	<i>Cladosporium herbarum</i>	< 0.1	Negative						
<b>Trees</b>	Cedar, Mountain	< 0.1	Negative						
	Cottonwood Tree	0.33	0/1						
	Elm, American	0.56	I						
	Hickory Pecan Tree	1.64	II						
	Maple/Box Elder	0.76	II						
	Walnut	3.71	III						
	White Ash	6.15	III						
	White Birch*	0.90	II						
	White Oak	3.17	II						
<b>Weeds</b>	Cocklebur, Common*	0.10	0/1						
	Common Pigweed	3.10	II						
	Common Ragweed	0.26	0/1						
	English Plantain	1.80	II						
	Lamb's Quarter*	1.25	II						
	Mugwort	0.26	0/1						
	Russian Thistle	0.84	II						

Analyte		Scale						
Analyte	Value	<5	5	10	31	77	298	>1000
<b>Total IgE</b>	Total IgE*							102.71

Class	Ranges	Interpretation	Class	Ranges	Interpretation
Negative	< 0.1 KU/L	Negative	III	= 3.5 to 17.5 KU/L	Positive with increasing antibody concentration
0/1	= 0.2 to 0.35 KU/L	Equivocal	IV	= 17.5 to 50 KU/L	
I	= 0.35 to 0.7 KU/L	Positive with increasing antibody concentration	V	= 50 to 100 KU/L	
II	= 0.7 to 3.5 KU/L		VI	> 100 KU/L	

\* This test was developed, and its performance characteristics determined by Meridian Valley Lab. It has not been cleared or approved by the US FDA. This test was performed in a CLIA (50D0630590) certified laboratory and is intended for clinical purposes.

