



PATIENT: **Sample Report**

TEST REF: **TST-12345**

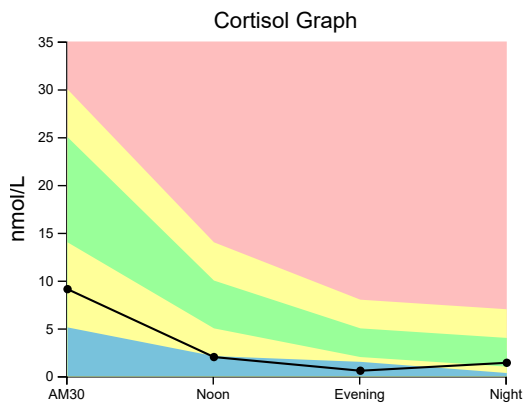
TEST NUMBER: N/A  
 PATIENT NUMBER: N/A  
 GENDER: Female  
 AGE: 58  
 DATE OF BIRTH: MM/DD/YYYY

COLLECTED: MM/DD/YYYY 0000  
 MM/DD/YYYY 0000  
 MM/DD/YYYY 0000  
 RECEIVED: MM/DD/YYYY  
 TESTED: MM/DD/YYYY

PRACTITIONER: **Nordic Laboratories**

**TEST NAME: Adrenal Function Report**

Analyte	Result	Unit	L	WRI	H	Optimal Range	Reference Interval
<b>Cortisol AM30</b>	9.1	nmol/L		◆		14.0 - 25.0	5.1 - 30.0
<b>Cortisol Noon</b>	2.0	nmol/L	↓			5.0 - 10.0	2.1 - 14.0
<b>Cortisol Evening</b>	0.57	nmol/L	↓			2.0 - 5.0	1.5 - 8.0
<b>Cortisol Night</b>	1.4	nmol/L		◆		1.0 - 4.0	0.33 - 7.0
<b>DHEA*</b>	110	pg/mL		◆			106 - 300



**Hormone Comments:**

- Diurnal cortisol pattern and reported symptoms are consistent with evolving (Phase 2) HPA axis (adrenal gland) dysfunction.

**Adrenal Phase: 2**



**Notes:**

RI= Reference Interval, L (blue)= Low (below RI), WRI (green)= Within RI (optimal), WRI (yellow)= Within RI (not optimal), H (red)= High (above RI)  
 The current samples are routinely held three weeks from receipt for additional testing.

\*This test was developed and its performance characteristics determined by Doctor's Data, Inc. The FDA 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.

Methodology: Enzyme Immunoassay

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