

PRACTITIONER

PATIENT

Name: Sample Patient
DOB:
Gender:

TEST	RESULT			
	IN RANGE (Normal)	EQUIVOCAL*	OUT OF RANGE	REFERENCE (ELISA Index)
Array 5 – Multiple Autoimmune Reactivity Screen **				
Parietal Cell + ATPase	<0.50			0.0-2.2
Intrinsic Factor	0.51			0.0-2.5
ASCA + ANCA	0.68			0.4-1.6
Tropomyosin	0.89			0.0-3.0
21-Hydroxylase (Adrenal Cortex)	0.96			0.6-1.9
Myocardial Peptide	0.87			0.0-1.9
Alpha-Myosin	0.79			0.6-2.0
Phospholipid	0.90			0.4-1.7
Platelet Glycoprotein	0.32			0.6-1.8
Ovary/Testis ***	0.73			0.6-1.8
Fibulin	0.37			0.4-1.3
Collagen Complex	0.38			0.2-1.5
Arthritic Peptide	0.49			0.0-1.7
Osteocyte	0.93			0.7-2.0
Cytochrome P450 (Hepatocyte)	1.05			0.8-2.3
Myelin Basic Protein	0.73			0.6-1.7
Asialoganglioside	0.63			0.6-1.6
Alpha-Tubulin + Beta-Tubulin	0.80			0.0-2.7
Cerebellar	0.37			0.4-1.5
Synapsin	1.03			0.0-2.1

* Reference ranges are calculated based on the mean \pm 2 standard deviations (SD). Results > 1 SD, and <2 SDs above the mean are considered to be equivocal. An equivocal result represents the range between negative and suspicious low positive results. Results >2 SDs are considered out of range, and positive.

** All analytes, except for Thyroid Peroxidase, Thyroglobulin, GAD, and IA-2 are tested for IgG and IgA combined. Thyroid Peroxidase (TPO) and Thyroglobulin (Tg) are tested for IgG only. GAD and IA-2 are tested for multiple autoantibodies. TPO results were obtained with the Inova Diagnostics QUANTA Lite TPO ELISA Kit. Thyroglobulin results were obtained with the Inova Diagnostics QUANTA Lite Thyroid T ELISA Kit. GAD65 results were obtained with the Kronus Glutamic Acid Decarboxylase (GAD) Autoantibody ELISA Kit. IA-2 results were obtained with the Kronus IA-2 Autoantibody (IA-2Ab) ELISA Kit. Values obtained with different manufacturers' assay methods may not be used interchangeably. The magnitude of the reported IgG levels cannot be correlated to endpoint titer. Shipping and storage criteria for specimens may vary from Inova Diagnostics' and Kronus' protocol, however, this does not appear to significantly affect patient results.

*** Ovary and Testis are tested together to avoid any confusion arising out of potential cross-reactivity

Mark G. Kartub, M.D., Medical Director

Cyrex Laboratories is certified under the Clinical Laboratory Improvement Amendments of 1988 ("CLIA") as qualified to perform high-complexity clinical testing. Test result data on its own does not constitute a diagnosis of any disease. Only a physician or qualified healthcare professional should interpret the significance of a clinical lab test or make a diagnosis. This test was developed and its performance characteristics determined by Cyrex Laboratories, LLC. This test is a lab developed test and therefore not subject to clearance or approval by the US Food and Drug Administration. The names and titles of tests and arrays are for reference purposes only.

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TEST	RESULT			REFERENCE (WHO Units)
Array 5 – Multiple Autoimmune Reactivity Screen **	Negative	Weak Positive	Moderate to Strong Positive	
Thyroglobulin IgG	<3.92			0.0-200.0

TEST	RESULT			REFERENCE (WHO Units)
Array 5 – Multiple Autoimmune Reactivity Screen **	Negative	Positive		
Thyroid Peroxidase IgG	9.24			0.0-100.0

TEST	RESULT			REFERENCE
Array 5 – Multiple Autoimmune Reactivity Screen **	Negative	Positive		
IA-2 Autoantibody		108.65		0.0-7.49 U/mL
Glutamic Acid Decarboxylase Autoantibody		190.03		0.0-5.0 IU/mL

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