VITROS® 3600 Immunodiagnostic System Assay Menu



The VITROS 3600 immunodiagnostic system provides your lab with the most comprehensive offerings to give your patients the highest-quality results as efficiently as possible.

Anemia	Infectious Disease	Inflammatory	Reproductive Endocrinology
Ferritin ^{MW}	Anti-HAV IgM ™	Procalcitonin MW	Estradiol MW
Folate MW	Anti-HAV Total MW		FSH ^{MW}
Vitamin B12 MW	Anti-HBc ™	Nutritional Assessment	LH ^{MW}
	Anti-HBc IgM ™	Ferritin ^{MW}	Progesterone ^{MW}
Bone Disease	Anti-HBe ™	Folate MW	Prolactin ^{™W}
Intact PTH MW	Anti-HBs ™	Vitamin B12 MW	Testosterone MW
NTx MW	Anti-HCV ™W		Total β -hCG Generation II $^{\mbox{\scriptsize MW}}$
Vitamin D ™	Anti-HIV 1+2 MW	Oncology	TSH ™
	Anti-SARS CoV-2 IgG ™	AFP MW	
Cardiac	Anti-SARS-CoV-2 IgG	CA 15-3 ^{MW}	Sepsis Procalcitonin MW
CK-MB ^{MW}	Quantitative MW,1	CA 19-9 ^{MW}	
hsTroponin (D) ™	Anti-SARS-CoV-2 Total ™	CA 125 II ^{MW}	Thyroid/Metabolic
Myoglobin ^{MW}	Anti-SARS-CoV-2 Total	CEA ^{MW}	aTg (D) ^{MW}
NT-proBNP MW	Nucleocapsid MW,1	PSA ^{MW}	aTPO (D) ^{MW}
Troponin I ES ^{MW}	HBeAg MW		Cortisol MW
·	HBsAg MW	Prenatal	Free T3 MW
Diabetes	HIV Combo MW	Anti-HIV 1+2 MW	Free T4 MW
C-Peptide MW	Rubella IgG ^{MW} SARS-CoV-2 Antigen ^{MW,1}	HBsAg ^{MW}	T3 Uptake MW
Insulin ^{MW}	Syphilis MW	HIV Combo MW	Tg (D) ^{MW}
	бургино	Rubella IgG MW	Total T3 ^{MW}
		Total β-hCG Generation II ^{MW}	Total T4 ^{MW}

1. VITROS SARS-CoV-2 Antigen Assay, VITROS Anti-SARS-CoV-2 Total N and VITROS Anti-SARS-CoV-2 IgG Quant Antibody Tests have not been cleared or approved by the U.S. Food and Drug Administration (FDA). They have been authorized by the FDA under an Emergency Use Authorization (EUA) and testing is limited to laboratories certified under the Clinical Laboratory Improvement Amendments (CLIA) of 1988, 42 U.S.C. §263a, to perform moderate- or high-complexity tests. The VITROS antigen test has been authorized only for the detection of proteins from SARS-CoV-2, not for any other viruses or pathogens. The VITROS antibody tests have been authorized only for the detection of either total or IgG antibodies from SARS-CoV-2, not for any other viruses or pathogens, and results should not be used as the sole basis for diagnosis. These tests are only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostic tests for detection and/or diagnosis of COVID-19 under Section 564(b)(1) of the Act, 21 U.S.C. § 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner.



(D): In development, not currently available.

MW: MicroWell

New QuidelOrtho VITROS system branding may not be available in all markets, subject to country-specific regulatory approval. Please confirm with your local commercial team.

Products are subject to local regulatory requirements and may not be available in all regions.

