For use in the U.S.

Assay Menu

Infectious Disease

	Atellica® IM Analyzer	ADVIA Centaur® XP Immunoassay System	ADVIA Centaur® XPT Immunoassay System	ADVIA Centaur® CP Immunoassay System
Hepatitis A				
HAV IgM	●	●	●	●
HAV Total	●	●	●	●
Hepatitis B				
Anti-HBe 2	●	●	●	●
HBc IgM	●	●	●	●
HBc Total	●	●	●	●
HBeAg	●	●	●	
Anti-HBs 2	●	●	●	●
HBsAg II	●	●	●	
HBsAg				●
HBsAg Confirmatory	●	●	●	●

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Hepatitis C				
HCV	●	●	●	●
HIV				
CHIV	●**	●**	●**	●**
EHIV	●**	●**	●**	●**
TORCH				
CMV IgG	●	●	●	●
CMV IgM	●*	●*	●*	●*
Herpes-1 IgG	●	●	●	●
Herpes-2 IgG	●	●	●	●
Rubella IgG	●	●	●	●
Rubella IgM	●	●	●	●
Toxoplasma IgG	●	●	●	●
Toxoplasma IgM	●	●	●	●*
Special ID				
EBV-EBNA IgG	●*	●*	●*	●*
EBV-VCA IgG	●*	●*	●*	●*
EBV-VCA IgM	●*	●*	●*	●*
Syphilis	●	●	●	●
Zika	●*,†	●†	●†	

**Under development. Not commercially available. Future availability cannot be guaranteed.*

***Assay developed, manufactured, and sold by Siemens Healthcare Diagnostics Inc. for Ortho Clinical Diagnostics, Inc. and Grifols Diagnostic Solutions Inc.*

†The ADVIA Centaur Zika Test has not been FDA cleared or approved. This test has been authorized by FDA under an EUA for use by authorized laboratories. This test has been authorized only for the diagnosis for Zika virus infection and not for any other viruses or pathogens. This test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of the emergency use of in vitro diagnostic tests for detection of Zika virus and/or diagnosis of Zika virus infection under section 564(b)(1) of the Act, 21 U.S.C. § 360bbb-3(b)(1), unless the authorization is terminated or revoked sooner.

Product availability may vary from country to country and is subject to varying regulatory requirements. Please contact your local representative for availability.

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