

Hologic is your dedicated partner in sexual and vaginal health. We deliver comprehensive and flexible testing with proven performance—enabling accurate diagnosis to help prevent the spread of disease and protect reproductive health.

Our breadth of assays offers the convenience of testing for multiple specimen options, along with the versatility provided by the Panther® Scalable Solutions.

## **Aptima® Sexual and Vaginal Health**

- CT/NG
- Mycoplasma genitalium
- Trichonomas vaginalis
- Bacterial vaginosis
- Candida vaginitis/Trichomonas vaginalis

- HSV 1&2
- HPV
- HPV 16 18/45
- HIV-1
- O HIV-1 Qual Claim<sup>†</sup> ‡
- Zika Virus\*









## **Sample-to-Result Automation Meets Versatile Testing from a Single Sample**

Collection Kit Sa		Sample Type	Aptima Combo 2® Assay (CT/NG)	Aptima® Mycoplasma genitalium Assay	Aptima® Trichomonas vaginalis Assay	Aptima® HSV 1 & 2 Assay	Aptima® BV Assay	Aptima® CV/TV Assay	Aptima® HPV Assay	Aptima® HPV 16 18/45 Genotype Assay
The charge of th	Aptima® Specimen Transfer Kit	ThinPrep® (PreservCyt®)	<b>√</b>		<b>√</b>				<b>√</b>	1
	- Aptima® Multitest Swab Collection Kit	Clinician Collected Vaginal Swab	<b>✓</b>	<b>✓</b>	<b>✓</b>		1	1		
		Patient Collected Vaginal Swab	1	1			1	1		
		Patient Collected Penile Meatal Swab		1						
		Clinician Collected Throat Swab	1							
		Clinician Collected Rectal Swab	1							
		Clinician Collected Anogenital Lesion Swab				<b>√</b>				
) and	Aptima® Unisex Swab Collection Kit	Clinician Collected Endocervical	<b>√</b>	<b>✓</b>	<b>√</b>					
		Clinician Collected Male Urethral Swab	<b>✓</b>	<b>✓</b>						
	Aptima® Urine Specimen Collection Kit	Female Urine	1	<b>✓</b>						
		Male Urine	1	1						
		✓ = US-FDA Clearance or Approval								

MISC-06950-001 Rev. 001 © 2020 Hologic, Inc. All rights reserved. Hologic, Panther, Aptima, Aptima Combo 2, ThinPrep, PreservCyt, and associated logos are trademarks and/or registered trademarks of Hologic, Inc. and/or its subsidiaries in the United States and/or other countries. All other trademarks, registered trademarks, and product names are the property of their respective owner. This information is intended for medical professionals in the U.S. and is not intended as a product solicitation or promotion where such activities are prohibited. Because Hologic materials are distributed through websites, eBroadcasts and tradeshows, it is not always possible to control where such materials appear. For specific information on what products are available for sale in a particular country, please contact your local Hologic representative or write to diagnostic solutions@hologic.com. representative or write to diagnostic.solutions@hologic.com.

<sup>‡</sup> Seeking dual claim for the HIV-1 Quant assay.

<sup>\*</sup> The Aptima® Zika Virus:

The Aptima® Zika Virus:

This 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 detection of RNA from Zika and the diagnosis of Zika virus infection, not for any other viruses or pathogens; and

This test is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of in vitro diagnostic tests for detection of Zika virus and/or diagnosis of Zika virus infection under Section 564(b)(f) of the Act, 21 U.S.C.§ 360bbb-3(b)(f), unless the authorization is terminated or revoked sooner.