



PATIENT INFORMATION		PHYSICIAN INFORMATION	
<b>Test Patient</b>		John Pathologist, M.D.	
SEX: F		123 Main Street	
DOB: 1/1/1971 (35)		San Jose, CA 95124	
SPECIMEN INFORMATION			
Collected: 02/12/2006		Accession #:	<b>TC06-8</b>
Received: 02/13/2006			
Reported: 09/19/2006			

**\*\*\* THIS IS A TEST REPORT, PLEASE DESTROY \*\*\***

#### CLINICAL INFORMATION

Clinical History: Routine Check-up - 2/12/06  
Abnormal Bleeding - Y  
History of abnormal paps.

**SOURCE:** Vaginal

**ADEQUACY OF SPECIMEN:** SATISFACTORY FOR EVALUATION, ENDOCERVICAL/TRANSFORMATION ZONE COMPONENT PRESENT.  
Scanty cellularity.

**GENERAL CATEGORY:** **EPITHELIAL CELL ABNORMALITY, SEE NARRATIVE DESCRIPTION.**

**NARRATIVE DESCRIPTION:** **ATYPICAL SQUAMOUS CELLS OF UNDETERMINED SIGNIFICANCE. (ASC-US)**  
Abundant polymorphonuclear leukocytes.

**SCREENED BY:** Werner J. Stamm, M.D.

**RECOMMENDATION:** Recommend repeat pap smear and/or colposcopy within 4-6 months.

Werner J. Stamm, M.D.

This Pap test has been evaluated with the assistance of the ThinPrep Pap Test Imaging System, unless otherwise specified.

Education Note: The PAP test is not a diagnostic procedure and should not be used as the sole means to detect cervical cancer. It is only a screening procedure to aid in the detection of cancer and its precursors. Both false-negative and false-positive results have been experienced. Any visible lesion should be biopsied.

#### HPV - HIGH RISK RESULTS

HPV - HIGH RISK RESULTS: **\*\*\*DETECTED\*\*\***

(HPV Types: 16,18,31,33,35,39,45,51,52,56,58,59,68)

Testing was performed using the FDA-approved Digene Hybrid Capture II methodology.

#### HPV - LOW RISK RESULTS

HPV - LOW RISK RESULTS: NOT DETECTED

(HPV Types: 6,11,42,43,44)

Testing was performed using the FDA-approved Digene Hybrid Capture II methodology.

#### NEISSERIA GONORRHEA RESULTS

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PLACE OF TESTING AND SERVICE: Associated Pathology Medical Group, Inc., 105A Cooper Ct., Los Gatos, CA 95032, unless otherwise specified.

#### FINAL REPORT



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NEISSERIA GONORRHEA RESULTS: **POSITIVE**

Testing was performed using the Digene Hybrid Capture II methodology, which is FDA-approved for GC/CT testing. The specimen type (ThinPrep Pap or PreserveCyt solution) is not specified in the kit insert; however, FDA clearance is pending and the performance characteristics have been validated by Digene and APMG.

**CHLAMYDIA TRACHOMATIS RESULTS**

CHLAMYDIA TRACHOMATIS RESULTS: **NEGATIVE**

Testing was performed using the Digene Hybrid Capture II methodology, which is FDA-approved for GC/CT testing. The specimen type (ThinPrep Pap or PreserveCyt solution) is not specified in the kit insert; however, FDA clearance is pending and the performance characteristics have been validated by Digene and APMG.