A Guide to Your Pathology Report

To help you understand how to find information on your Pathology Report page, below are descriptions and corresponding annotations of the different sections that may appear.

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### Gyn Cytology Report

<table>
<thead>
<tr>
<th>Patient:</th>
<th>Provider:</th>
<th>Date of Birth:</th>
<th>Received:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Cervical ThinPrep Imaged Liquid Pap Test</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

#### History:
Last Menstrual Period Not given

#### Specimen:
Cervical ThinPrep Imaged Liquid Pap Test

#### Recent Cytology/HPV Results from this lab:

<table>
<thead>
<tr>
<th>Date</th>
<th>Pathology#</th>
<th>Pap Results</th>
<th>HPV Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>06/04/13</td>
<td>XXXX</td>
<td>ASC-US</td>
<td>Positive</td>
</tr>
<tr>
<td>02/10/09</td>
<td>XXXX</td>
<td>LSIL</td>
<td></td>
</tr>
</tbody>
</table>

#### Pap Interpretation:
EPITHELIAL CELL ABNORMALITY, SQUAMOUS.
Atypical squamous cells, cannot exclude high grade squamous intraepithelial lesion.

#### Specimen Adequacy:
Satisfactory for evaluation. Endocervical/transformation zone component present.

#### Molecular Results:
- NEGATIVE for Chlamydia trachomatis (CT).
- NEGATIVE for Neisseria gonorrhoeae (NG).
- POSITIVE for High Risk HPV.
- POSITIVE for HPV 16.
- NEGATIVE for HPV 18.

CT/NG testing and its performance characteristics are determined by the Molecular Pathology Laboratory, 1124 Columbia St, Seattle, WA. Test Method: Nucleic Acid Amplification Test (NAT) using the Gen-Probe Aptima Combo 2 Assay. The laboratory is regulated under the 1988 CLIA amendments as qualified to perform high-complexity clinical testing. This test is FDA-approved for the PANTHER platform.

High Risk HPV testing and its performance characteristics are determined by the Molecular Pathology Laboratory, CellNetix Pathology & Laboratories, 1124 Columbia St, Seattle, WA, using the cobas 4800 assay. This laboratory is regulated under the 1988 CLIA amendments as qualified to perform high-complexity clinical testing. This test is FDA-approved for the Roche cobas platform and ThinPrep specimens.

Initial Evaluation performed by Janet Conte CT(ASCP). Electronically signed

Nisreen Fidda, M.D. Electronically signed

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Find patient and provider information here, as well as the date we received the specimen for testing.

We will provide the past three Pap and HPV results as a way for your physician to track health trends and flag potential risks.

In the Pap interpretation section you will find our diagnosis based on the interpretation of the cells on the prepared slide.

Specimen adequacy is additional information provided to your physician.

Your physician may order additional testing beyond the Pap. These tests are often used to diagnose STDs such as Chlamydia, Gonorrhea, Human papillomavirus (HPV), Herpes (HVS), bacterial vaginosis (BV).

We always include the name of the professionals who diagnosed the case; however, you must contact your physician in case you have any questions or need further explanation on the diagnosis.

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