### Patient Information
- **DOB:**
- **Age:**
- **Gender:**
- **Patient ID:**
- **Health ID:**

### Specimen Information
- **Specimen:**
- **Collected:**
- **Received:**
- **Reported:**

### Client Information

### Endocrinology

<table>
<thead>
<tr>
<th>Test Name</th>
<th>Result</th>
<th>Reference Range</th>
<th>Lab</th>
</tr>
</thead>
<tbody>
<tr>
<td>ESTRADIOL, ULTRASENSITIVE LC/MS/MS</td>
<td>&lt;2</td>
<td>&lt; OR = 29 pg/mL</td>
<td>EZ</td>
</tr>
</tbody>
</table>

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics Nichols Institute San Juan Capistrano. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

### Physician Comments:

### Performing Site:

**EZ** QUEST DIAGNOSTICS/NICHOLS SJIC 33608 ORTEGA HWY, SAN JUAN CAPISTRANO, CA 92675-2042 Laboratory Director: JON NAKAMOTO, MD PHD, CLIA: 05D0643332