**Mycoplasma genitalium Amplification T7 Oligo**

**Analyte Specific Reagent (ASR).**
Analytical and performance characteristics are not established.

**Intended Use**
Analyte Specific Reagent. Analytical and performance characteristics are not established. This analyte specific reagent is intended to be used as a component of a laboratory developed test (LDT) used exclusively by laboratories. Laboratories can develop their own LDTs using one or more Analyte Specific Reagents and/or general laboratory reagents for *in vitro* diagnostic purposes.

**Materials Provided**
*Mycoplasma genitalium Amplification T7 Oligo* (Cat. No. PRD-03259)

<table>
<thead>
<tr>
<th>Component</th>
<th>Quantity</th>
<th>Concentration</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tube</td>
<td>1 x 0.17 mL</td>
<td>112 nmol/mL</td>
<td><em>T7 amplification primer, complementary to M. genitalium 16S rRNA.</em></td>
</tr>
</tbody>
</table>

**Warnings and Precautions**
- A. For Laboratory Use only. Analytical and performance characteristics are not established.
- B. Use routine laboratory precautions. Wear disposable, powderless gloves, protective eye wear, and laboratory coats when handling specimens and kit reagents. Wash hands thoroughly after handling reagents.
- C. Dispose of all materials that have come in contact with reagents according to local, state, and federal regulations.

**Product Specifications**

**Quality Control**
This analyte specific reagent was manufactured and released in accordance with the Hologic quality control and quality assurance procedures. This reagent has been functionally tested using a system capable of detecting luminescence along with commercially available transcription mediated amplification (TMA) general purpose reagents (GPR). However, it is the responsibility of each laboratory to develop and validate their own tests for clinical diagnostic purposes.

**Storage and Stability**
Please refer to product labels for information on storage and stability.