1. Background:
Phexxi (lactic acid, citric acid, and potassium bitartrate) vaginal gel is indicated for the prevention of pregnancy in females of reproductive potential for use as an on-demand method of contraception. Phexxi is not effective for the prevention of pregnancy when administered after intercourse.

2. Coverage Criteria:

A. Authorization

1. Phexxi will be approved based on all of the following criteria:
   a. Used for the prevention of pregnancy
   -AND-
   b. Patient is unable to use all of following other methods of contraception due to failure, contraindication, intolerance or refusal (document reason for each method):
      1) Injection (e.g., Depo-Provera)
      2) Oral Contraceptive [e.g., norethindrone (generic Micronor), Yaz]
      3) Transdermal Patch (e.g. Twirla, Xulane)
      4) Vaginal Contraceptive Ring (e.g., Annovera, NuvaRing)
      5) Diaphragm
      6) Sponge (e.g. Today)
      7) Cervical Cap (e.g., FemCap)
      8) Female Condom
   -AND-
   c. History of failure, contraindication, or intolerance to nonoxynol-9 based spermicide
   -AND-
   d. Provider attests they have counseled the patient regarding a higher rate of pregnancy prevention with the use of other methods of contraception (e.g., injection, oral contraception, transdermal patch, vaginal ring) compared to Phexxi

Authorization will be issued for 12 months.
a State mandates may apply. Any federal regulatory requirements and the member specific benefit plan coverage may also impact coverage criteria. Other policies and utilization management programs may apply.

3. Additional Clinical Programs:
   - Notwithstanding Coverage Criteria, UnitedHealthcare may approve initial and re-authorization based solely on previous claim/medication history, diagnosis codes (ICD-10) and/or claim logic. Use of automated approval and re-approval processes varies by program and/or therapeutic class.

4. References:

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<tr>
<th>Program</th>
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<tr>
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<tr>
<td>5/2022</td>
<td>Annual review. Updated references.</td>
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