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| **Characteristics** | **Healthy Volunteers (n=10)** |
| Age, median (range) | 46.0 (28-70) |
| Race |  |
|  White | 8 (80%) |
|  Asian | 2 (20%) |
| Parity, median (range) | 2 (0-3) |
|  Vaginal Parity, median (range) | 1.5 (0-3) |
| Vaginal Estrogen Use | 2 (20%) |
| History of Hysterectomy | 0 (0%) |
| History of vaginal reconstructive surgery | 0 (0%) |
|  |
| Comfort of use (1= very poor, 5 = very good), median, (range)  |  |
|  Active wearing | 4.5 (1-5) |
|  Device removal | 4.5 (2-5) |
| Time of use daily, median (range) | 2.5 (1-8) hours |
| New symptoms reported after use |  |
|  Vaginal Pain | 2 (20%) |
|  Vaginal Discharge | 1 (10%) |
|  Vaginal Bleeding | 1 (10%) |
|  |
| **Feedback Through Steps of Use, median (range)** | **Insertion** | **Locating Loop for Removal** | **Removal** |
| How intuitive did you find this task? (1= very difficult, 5= very straightforward) | 4 (1-5) | 5 (4-5) | 5 (5) |
| How easy-to-use did you find the device? (1= very difficult, 5= very easy) | 4 (1-5) | 5 (4-5) | 5 (4-5) |
| How safe did you find this task? (1=high risk, 5= very safe) | 5 (4-5) | 5 (3-5) | 5 (4-5) |
| **Sample of free form answer questions asked of the healthy volunteers for feedback:** |
| Did you have any changes in your ability to urinate or defecate? Please explain.* 2 reported slower urine stream, but did not affect ability to empty bladder.
* No changes in defecation reported.
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| Did you have any feeling of possible expulsion with use of the device during daily activity? If yes, please explain.* 1 reported device shifted lower in vagina with full bladder
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| Please provide your thoughts on tolerance of this device by patients who are two weeks post-operative from reconstructive vaginal surgery.* 6 reported insertion/removal of device could be uncomfortable in early post-operative setting.
* 7 reported once device was properly placed, the device was no longer noticeable during active use.
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| Please provide your thoughts on how to improve the device or its application/removal* 6 of responses posed a change to the softness or shape of the retaining feature.
* 3 of responses mentioned more descriptive instructions for use and orientation of device.
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\*over 2 consecutive days total