Measurement of the Thickness of the Urethrovaginal Space in Women with or without Vaginal Orgasm

Giovanni Luca Gravina, MD, PhD,* Fulvia Brandetti, MD,* Paolo Martini, MD, PhD,* Eleonora Carosa, MD, PhD,* Savino M. Di Stasi, MD,‡ Susanna Morano, MD,‡ Andrea Lenzi, MD,§ and Emmanuele A. Jannini, MD*

*Department of Experimental Medicine, University of L’Aquila, L’Aquila, Italy; †Department of Surgery-Urology, “Tor Vergata” University, Rome, Italy; ‡Department of Clinical Sciences, University of Rome “La Sapienza,” Rome, Italy; §Department of Medical Physiopathology, University of Rome “La Sapienza,” Rome, Italy

ABSTRACT

Introduction. The physiology and anatomy of female sexual function are poorly understood. The differences in sexual function among women may be partly attributed to anatomical factors.

Aim. The purpose of this study was to use ultrasonography to evaluate the anatomical variability of the urethrovaginal space in women with and without vaginal orgasm.

Methods. Twenty healthy, neurologically intact volunteers were recruited from a population of women who were a part of a previous published study. All women underwent a complete urodynamic evaluation and those with clinical and urodynamic urinary incontinence, idiopathic detrusor overactivity, or micturition disorders, as well as postmenopausal women and those with sexual dysfunction were excluded. The reported experience of vaginal orgasm was investigated.

Main Outcome Measure. The urethrovaginal space thickness as measured by ultrasound was chosen as the indicator of urogenital anatomical variability. Designated evaluators carried out the measurements in a blinded fashion.

Results. The urethrovaginal space and distal, middle, and proximal urethrovaginal segments were thinner in women without vaginal orgasm. A direct correlation between the presence of vaginal orgasm and the thickness of urethrovaginal space was found. Women with a thicker urethrovaginal space were more likely to experience vaginal orgasm (r = 0.884; P = 0.015). A direct and significant correlation between the thickness of each urethrovaginal segment and the presence of vaginal orgasm was found, with the best correlation observed for the distal segment (r = 0.863; P < 0.0001). Interobserver agreement between the designated evaluators was excellent (r = 0.87; P < 0.001).

Conclusions. The measurement of the space within the anterior vaginal wall by ultrasonography is a simple tool to explore anatomical variability of the human clitoris-urethrovaginal complex, also known as the G-spot, which can be correlated to the ability to experience the vaginally activated orgasm. Gravina GL, Brandetti F, Martini P, Carosa E, Di Stasi SM, Morano S, Lenzi A, and Jannini EA. Measurement of the thickness of the urethrovaginal space in women with or without vaginal orgasm. J Sex Med 2008;5:610–618.

Key Words. Echography; Female Ejaculation; G-Spot; Skene’s Glands; Female Prostate

Introduction

Female genital anatomy and the physiology of female sexual function have been scientifically neglected in the past [1]. This is particularly true for orgasm, which has been described much more from a political or philosophical perspective [2] than by using scientific tools. Female orgasm is a complex function not perfectly understood where intrapsychic, cognitive, relational, neurohormonal, vascular and anatomical factors play roles. Although literature abounds with descriptions and discussions of vaginal as opposed to clitoral orgasm [3,4] and only few studies map genital erotic sensitivity to tactile stimulation in healthy females [5], it is evident that
some women need direct, external, clitoral stimulation whereas others may reach orgasm also by penetration and thrusting that directly stimulate the internal clitoris and vaginal wall structures and indirectly the external part of the same organ [5]. Interestingly, a study of 200 women revealed that external clitoris length varies by more than 25%, demonstrating that individual differences are macranoanatomic [6]. Furthermore, in cadaver studies, the internal clitoris may have individual differences bigger than 100% [7]. Whether these differences are correlated with ability to reach orgasm is not yet established.

Other areas have been involved in the mechanism of orgasm (urethra, labia minora, Halban’s fascia, periurethral glands; see [4] and references therein). On the basis of its supposedly low presence of sensory receptors, the vagina was considered as poorly responsive by Kinsey [8], and Masters and Johnson [9]. However, Gräfenberg [10], as subsequently reviewed and popularized by Ladas, Whipple, and Perry [11] (but disputed by others [12,13]), suggested a variable area of increased sensitivity over the urethra on the anterior vaginal wall. The urethrovaginal space (where the Halban’s fascia runs [14]) seems critical, being constituted of fibro-connective tissue and large numbers of blood vessels, glands, muscular fibers, and nerve endings. The close physical proximity of the urethra and the clitoris to the anterior vaginal wall suggests an association between these anatomical structures and sexual function [15,16]. In fact, the anterior vaginal wall is an active organ, transmitting, during intercourse, the effect of penile thrusting in the vagina to the clitoris, by stretching the two ligaments that insert around its base [17]. As for the clitoris, microscopic examination reveals that the human vagina’s anterosuperior wall differs from one subject to another [18]. The presence of pseudocavernous tissue (clitoral bulb) in the anterior vaginal mucosa is a frequent but not universal finding (86%) [19]. Around the urethra, the existence of the prostatic embryological remnant—Skene’s glands [20]—is also an anatomical variant and not a constant.

Aims

Differences in women’s sexual function obviously exist and although they have been largely attributed to cultural, religious, intrapsychic and, above all, relational factors [21], it is possible that anatomical factors might be partly responsible. Thus, if “anatomy is destiny” [22], physical differences should be taken into account as a source of physiological variability in female sexual response. With these concepts in mind, the purpose of this study was to use introital ultrasonography to evaluate anatomical variability, measured as the difference in thickness of the urethrovaginal space and to correlate this variability with the presence vaginally activated orgasm.

Methods

Patient Recruitment

A cohort of 37 healthy, neurologically intact, consecutive volunteers were recruited from the normal controls of a population of women who were a part of a previously published study [23]. Approval for this study was obtained from the Internal Review Board. The women underwent physical and neurological examination (including perineal/genital inspection, vaginal, pelvic floor muscle strength, assessment of reflexes such as anal wink, bulbocavernous and perineal-perianal sensation) and urodynamic study (UDS). All women underwent a nonstructured clinical interview at our Medical Sexology Service. The sexual history interview was conducted with each woman in a private room alone with a sexologist. Exclusion criteria consisted of drug or alcohol abuse, medications or medical conditions that might alter sexual function (e.g. diabetes), previous anti-incontinence surgery or any stage of vaginal prolapse (Pelvic Organ Prolapse Quantification System, POP-Q). Subjects with sexual dysfunction were also excluded. For this purpose, a female sexual function index (FSFI) [24] total score of less than 26.55 was considered suggestive of female sexual dysfunction [25]. Specifically, only women with high scores (4 or 5) for Q.11 (how often did you reach orgasm), Q.12 (how difficult was it for you to reach orgasm), and Q.13 (how satisfied were you with your ability to reach orgasm) were selected. It should be noted that these questions do not distinguish vaginal from clitoral orgasm. All subjects were exclusively heterosexual, had stable relationships (median value of 19 months [interquartile range [IQR] 13.5–23 months]) and reported at least two acts of sexual intercourse per week, a regular menstrual cycle and that they had been sexually active within the past 6 months.

The presence of vaginal orgasm was then investigated by the same male investigator (E.A.J.) who collected the sexological history, in a separate setting. Vaginal orgasm was ascertained by the following question: “Have you ever experienced a
vaginal orgasm?" Responses were categorized as “yes” (at least once in the past month) or “no” (never). By vaginal orgasm we mean the orgasm experienced after direct stimulation of the anterior vaginal wall by penetration, without concomitant stimulation of the external clitoris. Other investigators were blinded of these results before echo-graphic studies. All subjects were scheduled for examination during the mid-follicular phase of their menstrual cycle.

**Urodynamic Evaluation**

All enrolled patients underwent uroflowmetry and urodynamic evaluation (Urobenchmark 2000/3, S.I.E.M., Milan, Italy) performed in duplicate. Urodynamic assessment followed the International Continence Society standards and involved water cystometry with 37°C normal saline solution at a filling rate of 30 mL per minute. A 6Fr double lumen transurethral catheter (Bel Bioengineering Lab, Cantù, Co, Italy) was used for infusion and recording of intravesical pressure, and 9Fr intrarectal balloon catheter (Bel Bioengineering Lab, Cantù, Co, Italy) was used for recording abdominal pressure. Electromyographic activity of pelvic floor muscle was recorded by means of surface anal skin electrodes placed at the 3 and 9 o’clock position, to check dysfunctional voiding. Pressure-flow study was carried out in sitting position. For our purposes, the following measurements were detected: detrusor pressure at maximum flow (pDetQmax), maximum flow rate (Qmax) and from the ultrasound images in random order. All investigators (G.L.G. and P.M.) interpreted and measured the thickness of the urethrovaginal space measured the thickness of the urethrovaginal space. The ultrasound evaluation was obtained by an introital approach with the transducer placed over the external urethral orifice and the transducer axis corresponding to the body axis. Care was taken not to distort the anatomy during the procedure. Jet-type print photography, Polaroid photography, and video recording were used to collect all ultrasound images.

**Main Outcome Measures**

Total urethral length and vaginal lumen were viewed in the midsagittal plane. The image was then frozen and filed digitally. Two blinded male investigators (G.L.G. and P.M.) interpreted and measured the thickness of the urethrovaginal space from the ultrasound images in random order. All ultrasound images were displayed through an image processing and analysis program (Scion Image Software version alpha 4.0.3.2, Scion Co., Frederick, MD, USA). The anatomical border between the inner smooth muscle and mucosa-submucosa layer of the urethral wall can be distinguished by ultrasonography, as can the border between the vaginal wall and its lumen, seen as a strip of low echogenicity (Figure 1). To
To standardize the procedures, all measurements were expressed in millimeters and were obtained along a line drawn between the border of the smooth muscle and mucosa-submucosa layer of the urethral wall and the border of the vaginal wall and its lumen. Measurements were taken at various percentiles of the urethra length. The internal urethral meatus was considered as the zero point and the external meatus as the 100th percentile. In the midsagittal plane, we measured the thickness of the urethrovaginal space at the 10th (proximal segment), 50th (middle segment), and 90th percentile (distal segment) of the urethra (Figure 1C). The thickness was measured three times at each location and the median value was considered for statistical purposes.

**Results**

**Clinical and Urodynamic Parameters**

Thirty-seven healthy women were considered for this study. Thirteen were excluded, six because of sexual dysfunction and seven because of menopause with or without sexual dysfunctions. Of the 24 recruitable women, four declined to participate and a total of 20 subjects were finally enrolled (Figure 2). Of these, nine reported experience of vaginal orgasm. No significant difference was noted in age, parity, hormonal pattern, menstrual cycle duration, or FSFI total score among women with and without vaginal orgasm (Table 1). The analysis of urodynamic parameters indicated that all women had both normal cystometry and pressure-flow study. None of them had clinical and urodynamic urinary incontinence or disorders of the emptying phase (i.e., bladder outlet obstruction, impaired detrusor contractility, and electromyographic evidence of dysfunctional voiding). No significant difference between the two groups was found in any urodynamic parameters (Table 1).

**Ultrasound Assessment**

Introital ultrasound examination was well accepted by every volunteer and no adverse events occurred during or after each session. The examination took 20 minutes (IQR 12.5–32 minutes) on average.

Figure 1 (panels A and B) shows the schematic representation and introital ultrasound image of the urogenital organs. The vagina is seen as a hypoanechoic strip, adjacent to the posterior wall of the urethra for a short distance, and running from the probe to the cervical fornix. The symphysis pubis appears as a hyperechoic shell, with a shaded cone. The rectum generates acoustic artifacts with loss of borders between the rectal and posterior vaginal wall. The urethral lumen was seen as an anechoic cylindrical structure. The striated and smooth muscle layers of the urethra cannot always be distinguished in the midsagittal plane. The mucosa and submucosa were uniformly depicted as hypoechoic structures mimicking an open lumen.

Figure 3 shows a box plot diagram of urethrovaginal space thickness, stratified by group and segment. Comparison of ultrasound images in women with and without vaginal orgasm demon-
strated a statistically significant difference in the median thickness of the urethrovaginal space ($P < 0.0001$) (Table 2), which was thinner in women without (median value 10.4 mm; IQR 9.8 to 10.7 mm) than in those with vaginal orgasm (median value 12.4 mm; [IQR12 to 14 mm]). There was no overlapping in the IQR values, suggesting that the two groups had an independent distribution for the measured parameter.

The differences in the thickness of urethrovaginal space segments were also assessed. The median thickness of distal, middle, and proximal segments was less in women without than those with vaginal orgasm. A significant difference was seen for each segment in the two groups (Table 2). A completely independent distribution was found for these variables, as evidenced by the lack of overlapping in the IQR values between the two groups. Although

### Table 1 Clinical and urodynamical parameters

<table>
<thead>
<tr>
<th></th>
<th>Women with vaginal orgasm (N = 9)</th>
<th>Women without vaginal orgasm (N = 11)</th>
<th>$P$ value*</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Clinical parameters†</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age (year)</td>
<td>34 (29.3–36)</td>
<td>32 (30.2–35)</td>
<td>0.67</td>
</tr>
<tr>
<td>Parity (N. of births)</td>
<td>1 (0–1.3)</td>
<td>1 (0–1.5)</td>
<td>0.92</td>
</tr>
<tr>
<td>Menstrual cycle duration (days)</td>
<td>29 (26.5–30)</td>
<td>29 (27.5–31)</td>
<td>0.5</td>
</tr>
<tr>
<td>Body mass index (Kg/m²)</td>
<td>21 (20.6–22.9)</td>
<td>22 (2.4–23.1)</td>
<td>0.65</td>
</tr>
<tr>
<td>17 β estradiol (pmol/L)</td>
<td>198 (145–225)</td>
<td>187 (115–219)</td>
<td>0.79</td>
</tr>
<tr>
<td>Testosterone (nmol/L)</td>
<td>2.22 (1.75–2.7)</td>
<td>2.10 (1.8–2.82)</td>
<td>0.82</td>
</tr>
<tr>
<td>Thyrotropin (mU/L)</td>
<td>1.9 (1.7–2.2)</td>
<td>2.1 (2.2–1.6)</td>
<td>0.76</td>
</tr>
<tr>
<td>FSFI (total score)</td>
<td>29.5 (27.5–32.5)</td>
<td>29 (27–33)</td>
<td>0.77</td>
</tr>
<tr>
<td><strong>Urodynamic parameters on pressure-flow study†</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Qmax (mL/s)</td>
<td>20.5 (19–22.5)</td>
<td>21 (20–23.5)</td>
<td>0.18</td>
</tr>
<tr>
<td>PdetQmax (cmH₂O)</td>
<td>23.5 (18.2–32.5)</td>
<td>25 (22.5–31.5)</td>
<td>0.74</td>
</tr>
<tr>
<td>BOOI</td>
<td>–21 (–32–14.2)</td>
<td>–20 (–28/–15)</td>
<td>0.93</td>
</tr>
<tr>
<td>PVR (mL)</td>
<td>21 (17–33)</td>
<td>23 (18–34)</td>
<td>0.45</td>
</tr>
</tbody>
</table>

*Wilcoxon-Mann Whitney Rank Sum Test; †Median and interquartile range (25th and 75th percentile).
FSFI = female sexual function index; PdetQmax = detrusor pressure at maximum flow; PVR = post-void residual; BOOI = bladder outlet obstruction index.
in both groups proximal segments were thicker than the middle and distal segments, the median thickness of the different segments did not differ significantly within each group for women with \( (P = 0.39) \) and without vaginal orgasm, \( (P = 0.32) \).

Interobserver agreement between the two investigators performing the urethrovaginal space measurement was excellent \( (r = 0.87; P < 0.001) \).

Correlation analysis was estimated for those variables that significantly differed between two groups. This analysis, performed to determine interdependencies between the reported presence of a vaginal orgasm and the thickness of the urethrovaginal space, demonstrated a significant correlation between the presence of vaginal orgasm and the thickness of the urethrovaginal space. Women with a thicker urethrovaginal space were more likely to experience vaginal orgasm \( (r = 0.884; P = 0.015) \) (Table 3).

Correlation analysis, performed by stratifying the urethrovaginal space by segment, showed a direct and significant correlation between the thickness of each segment and the presence of vaginal orgasm. The best correlation was found with the distal segment of the urethrovaginal space \( (r = 0.863; P < 0.0001) \).

**Table 2** Thickness of the urethrovaginal space in women with or without vaginal orgasm

<table>
<thead>
<tr>
<th>Segment</th>
<th>Women without vaginal orgasm (N = 11)</th>
<th>Women with vaginal orgasm (N = 9)</th>
<th>( P ) value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Median thickness of urethrovaginal space*</td>
<td>10.4 (9.8–10.7)</td>
<td>12.4 (12.0–14.0)</td>
<td>&lt;0.0001†</td>
</tr>
<tr>
<td>Median thickness of urethrovaginal space segments*</td>
<td>Distal 9.9 (9.7–10.6)</td>
<td>12.2 (11.9–13.5)</td>
<td>&lt;0.0001†</td>
</tr>
<tr>
<td></td>
<td>Middle 10.4 (9.8–10.6)</td>
<td>12.5 (12.0–13.9)</td>
<td>&lt;0.0001†</td>
</tr>
<tr>
<td></td>
<td>Proximal 10.6 (10.1–11.9)</td>
<td>12.9 (12.3–14.4)</td>
<td>&lt;0.0001†</td>
</tr>
</tbody>
</table>

*Median and interquartile range (25th and 75th percentile).
†Wilcoxon–Mann Whitney Rank Sum Test.
‡Kruskal–Wallis test with Tukey HSD post hoc analysis.

**Discussion**

For cultural and historical reasons, scientific knowledge of female sexual function is still in its infancy. Here, we describe for the first time the use of a simple tool—introital ultrasonography—to characterize the thickness of the urethrovaginal space and correlate this with the presence or absence of vaginal orgasm. Although the number of analyzed subjects was small, the strict selection criteria render this population unique for a study of this kind. To eliminate possible confounders and improve the reproducibility of urethrovaginal space measurements, we excluded patients with clinical and urodynamic urinary incontinence [32],

![Figure 3](image-url)
Table 3  Correlation between thickness of urethrovaginal space and vaginal orgasm

<table>
<thead>
<tr>
<th>Correlation</th>
<th>Correlation coefficient</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thickness of urethrovaginal space vs. vaginal orgasm</td>
<td>0.884</td>
<td>0.015</td>
</tr>
<tr>
<td>Distal segment vs. vaginal orgasm</td>
<td>0.863</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Middle segment vs. vaginal orgasm</td>
<td>0.801</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Proximal segment vs. vaginal orgasm</td>
<td>0.810</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>

*Eta correlation.

Table 3: Correlation between thickness of urethrovaginal space and vaginal orgasm

- The most interesting finding of our study is the variability in female orgasm [38] could be at least partially attributed to differences in the presence, extension and reciprocal relationship of these anatomical structures.
- As the anatomy of the external clitoris changes [6], it is possible that some women may have more extensive clitoral bulbar tissue, exocrine glands, (highly variable embryological remnants), and nerves embedded in the anterior vaginal wall than others, such that they can achieve orgasm by direct stimulation of the urethrovaginal space and not only by stimulation of the external clitoris. Further well-controlled studies, based on the method here described, on much larger samples are needed to confirm the possible correlation between the anatomy of the anterior vaginal wall and vaginal orgasm. As the clitoris [39] and the Skene’s glands are under androgen control, we are currently measuring differences in the urethrovaginal space in hirsute women before and after antianrogen therapy, in women taking various estroprogestins, and in women reporting the much-debated phenomenon of female ejaculation (FE) [40,41]. Interestingly, women who experience vaginal orgasm have an urethrovaginal space thicker than those who do not. However, our data cannot directly demonstrate that the thickness of an apparent correlation

- Intramural detrusor overactivity, and micturition disorders [33], as well as postmenopausal patients and those with sexual dysfunction. As female genital anatomy is affected by estrogen and androgen activity [34], we also standardized the day of menstrual cycle. The measurement sites were also strictly uniform.

- These results raise several questions. What anatomical structures did we measure? Does urethrovaginal space thickness differ in all women with vaginal orgasm, or have we identified a subset of patients? Have we measured the controversial G-spot in vivo?

- Ultrasonography is a widely used tool for the assessment of the female urogenital tract, but its spatial and structural resolution remains much lower [35] than other imaging techniques [34]. Consequently, the anatomical structures within the urethrovaginal space cannot be fully resolved by ultrasonography. Gräfenberg described an erogenous zone located in the anterior vaginal wall [10] and subsequent studies have correlated the focus of female sensitivity with the external urethral sphincter [11]. Effectively, we made our measurements where the urethra is surrounded by the corpora cavernosa of the bulb of the clitoris—rich in nitric oxide synthase and type V phosphodiesterase activity [19], [36]—and by the exocrine, the Prostatic Specific Antigen-expressing Skene’s glands and a plethora of nerve endings [18,37]. However, histological studies have demonstrated clear interindividual differences [18,19]. The variability in female orgasm [38] could be at least partially attributed to differences in the presence, extension and reciprocal relationship of these anatomical structures. As the anatomy of the external clitoris changes [6], it is possible that some women may have more extensive clitoral bulbar tissue, exocrine glands, (highly variable embryological remnants), and nerves embedded in the anterior vaginal wall than others, such that they can achieve orgasm by direct stimulation of the urethrovaginal space and not only by stimulation of the external clitoris. Further well-controlled studies, based on the method here described, on much larger samples are needed to confirm the possible correlation between the anatomy of the anterior vaginal wall and vaginal orgasm. As the clitoris [39] and the Skene’s glands are under androgen control, we are currently measuring differences in the urethrovaginal space in hirsute women before and after antianrogen therapy, in women taking various estroprogestins, and in women reporting the much-debated phenomenon of female ejaculation (FE) [40,41]. Interestingly, women who experience vaginal orgasm have an urethrovaginal space thicker than those who do not. However, our data cannot directly demonstrate that the thickness of an apparent correlation.
anatomical “space” may generate a mechanism that can be related to the creation of an orgasm. But, in conclusion, the results here presented allow us to speculate that there may be a functional correlation between the thickness of urethrovaginal space, or G-spot, and the ability to experience the vaginal orgasm.

Acknowledgments
Our compliments and gratitude to Ms. Marie-Hélène Hayles and Dr. Rosaria Caruso for adapting their English expertise to our needs. This paper was partially supported by Italian Ministry of Research and Education PRIN grants and by an unrestricted grant from Pfizer Italia.

Corresponding Author: Emmanuele Jannini, MD, School of Sexology, Department of Experimental Medicine, University of L’Aquila, Via Vetoio, Coppito, L’Aquila, 67100, Italy. Tel: (+39) (0) 862 433530; Fax: (+39) (0) 862 433523; E-mail: jannini@univaq.it

Conflict of Interest: None declared.

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