Comparison of uterine and tubal pathology identified by transvaginal sonography, hysterosalpingography, and hysteroscopy in female patients with infertility

The Harvard community has made this article openly available. Please share how this access benefits you. Your story matters

Citation
Phillips, Catherine H., Carol B. Benson, Elizabeth S. Ginsburg, and Mary C. Frates. 2015. “Comparison of uterine and tubal pathology identified by transvaginal sonography, hysterosalpingography, and hysteroscopy in female patients with infertility.” Fertility Research and Practice 1 (1): 20. doi:10.1186/s40738-015-0012-3. http://dx.doi.org/10.1186/s40738-015-0012-3.

Published Version
doi:10.1186/s40738-015-0012-3

Citable link
http://nrs.harvard.edu/urn-3:HUL.InstRepos:33490712

Terms of Use
This article was downloaded from Harvard University’s DASH repository, and is made available under the terms and conditions applicable to Other Posted Material, as set forth at http://nrs.harvard.edu/urn-3:HUL.InstRepos:dash.current.terms-of-use#LAA
Comparison of uterine and tubal pathology identified by transvaginal sonography, hysterosalpingography, and hysteroscopy in female patients with infertility

Catherine H. Phillips1*, Carol B. Benson1, Elizabeth S. Ginsburg2 and Mary C. Frates1

Abstract

Background: The causes of female infertility are multifactorial and necessitate comprehensive evaluation including physical examination, hormonal testing, and imaging. Given the associated psychological and financial stress that imaging can cause, infertility patients benefit from a structured and streamlined evaluation. The goal of such a work up is to evaluate the uterus, endometrium, and fallopian tubes for anomalies or abnormalities potentially preventing normal conception. To date, the standard method for assessing these structures typically involves some combination of transvaginal sonography (TVS), hysterosalpingography (HSG), and hysteroscopy (HSC). The goal of this review is to compare the diagnostic accuracy of TVS, HSG, and HSC for diagnosing abnormalities in infertility patients to determine if all studies are necessary for pre-treatment evaluation.

Results: We identified infertility patients prior to initiation of assisted reproductive technology who had baseline TVS, HSG, and HSC within 180 days of each other. From medical record review, we compared frequencies of each finding between modalities. Of the 1274 patients who received a baseline TVS over 2 years, 327 had TVS and HSG within 180 days and 55 patients had TVS, HSG and HSC. Of the 327, TVS detected fibroids more often than HSG (74 vs. 5, \( p < .0001 \)), and adenomyosis more often than HSG (7 vs. 2, \( p = .02 \)). HSG detected tubal obstruction more often than TVS (56 vs. 8, \( p = .002 \)). Four (1.2 %) patients had endometrial polyps on both HSG and TVS.

In the 55 patients with HSG, TVS, and HSC, HSC identified endometrial polyps more often than TVS (10 vs. 1, \( p = .0001 \)) and HSG (10 vs. 2, \( p = .0007 \)). TVS detected more fibroids than HSC (17 vs. 5, \( p < .0001 \)). Tubal obstruction was identified more often by HSG than HSC (19 vs. 5, \( p < .0001 \)).

Conclusions: TVS is superior for evaluation of myometrial pathology. HSG is superior for evaluation of tubal pathologies. Endometrial pathologies are best identified with HSC.

Keywords: Infertility, Transvaginal sonography, Hysterosalpingography

Background

Infertility is defined as the inability for a couple to conceive a pregnancy following 1 year of unprotected vaginal intercourse [1]. It is estimated that 10–15 % of couples seek treatment for infertility [1–3]. It is generally considered appropriate to evaluate a couple for causes of infertility after 1 year of failed attempts at conception. However, given the inverse relationship of female fertility with age, it is often recommended that women over 35 years of age be evaluated after 6 months of failure to conceive, and women older than 40 be evaluated immediately [1].

A variety of factors may affect normal fertility including patient age, anatomy, ovulatory status, and sperm quality. Potential causes of infertility can be divided into male and female causes and include endocrine, anatomic, genetic, and behavioral conditions [4]. As a result, the evaluation of the infertile couple is multifactorial,
necessitating physical examination, hormonal testing, and imaging. Because the infertility population is under a great deal of psychological and emotional stress, these patients benefit from a structured and streamlined evaluation. In particular, evaluation of the female partner attempting to conceive requires assessment of the uterus, endometrium, and fallopian tubes for anomalies or abnormalities potentially preventing normal conception. The best method for assessing these structures usually involves some combination of transvaginal sonography (TVS), hysterosalpingography (HSG), and hysteroscopy (HSC). Less often, pelvic magnetic resonance imaging (MRI) and saline infusion sonohysterography (SIS) are used.

The objective of this paper is to compare the diagnostic accuracy of TVS, HSG, and HSC for diagnosing uterine and tubal abnormalities in women with infertility to determine if all three modalities are necessary in the work up of these patients.

Methods
We identified all baseline TVS performed on women in our infertility program from October 12, 2011 to October 12, 2013, prior to their initiation of assisted reproductive techniques (ART). From this group, we narrowed our patient population to those who had an HSG within 180 days of the sonogram to maximize the likelihood of concordance between the studies. All TVS and HSG reports were reviewed for tubal, myometrial, or endometrial findings and anatomical variants.

We reviewed patient medical records to identify those patients who also had hysteroscopy (HSC) within 180 days of the baseline TVS, and we recorded the reported findings.

Myometrial abnormalities were categorized as fibroids or adenomyosis; endometrial abnormalities as polyps, cysts, cavity distortion (e.g., synechia, stricture), or nonspecific asymmetry; and tubal abnormalities as obstruction. For each abnormality, the frequency of detection by each modality (TVS, HSG, HSC) was tabulated. For TVS, visualization of a hydrosalpinx was classified as an obstructed fallopian tube. Detection rates of abnormalities were compared among the modalities using the Fisher exact test, with a $p$-value of <0.05 considered significant.

This study was approved by the Brigham and Women’s Hospital Institutional review board, protocol number 2014P000355.

Results
A total of 1274 patients received a baseline TVS as part of a work up for infertility during the study period. Among these patients, 327 underwent a diagnostic HSG within a 180-day interval of the sonogram and comprise our study population. The time between TVS and HSG was 94 ± 49 days (mean ± SD). Among our 327 study patients, 55 also underwent HSC. The time (mean ± SD) between TVS and HSC was 61 ± 41 days and between HSG and HSC was 61 ± 50 days.

Among the study population of 327 patients (Table 1), 74 (23 %) had fibroids and 7 (2 %) had adenomyosis as diagnosed by either modality. Endometrial abnormalities were found in 16 (5 %) patients, based on TVS or HSG. Tubal obstruction was found in 56 (17 %) patients, more commonly unilateral (47 patients) than bilateral (9 patients).

TVS detected myometrial abnormalities significantly more often than did HSG, identifying fibroids in 74 patients while HSG only identified 5 (Fig. 1), and detecting adenomyosis in 7, while HSG detected only 2 ($p < .0001$ for both comparisons). Both HSG and TVS diagnosed endometrial polyps in the same 4 patients. HSG detected 6 patients with cavity distortion, while TVS found none of these ($p < 0.002$). With respect to tubal abnormalities, HSG performed significantly better than TVS, detecting tubal obstruction in all 56 (Fig. 2), while ultrasound only diagnosed 8 ($p < .0001$) (Fig. 3).

Among the subset of 55 patients who were evaluated by all three modalities, TVS, HSG, and HSC (Table 2), 17 (31 %) had fibroids and 6 (11 %) had adenomyosis. Endometrial abnormalities were found in 13 (24 %) patients and tubal obstruction in 19 (35 %).

In this group, TVS detected myometrial abnormalities significantly more often than did HSC, which identified only 4 of the 17 patients with fibroids ($p < .0001$) and none of the patients with adenomyosis. With respect to endometrial abnormalities, HSC outperformed TVS and HSG, identifying 10 polyps, while TVS only detected 1.

Table 1 Myometrial, endometrial, and tubal abnormalities detected by transvaginal ultrasound and/or hysterosalpingography (N = 327)

| Category                | TVS | HSG | Statistical significance |
|-------------------------|-----|-----|--------------------------|
| Myometrium              |     |     |                          |
| Fibroids                | 74  (23 %) | 5   | $p < .0001$              |
| Adenomyosis             | 7   (2 %)  | 2   | $p < .0001$              |
| Cesarean scar           | 1   | 0   |                          |
| Endometrium             |     |     |                          |
| Polyps                  | 4   | 4   |                          |
| Cysts                   | 4   | 0   |                          |
| Cavity distortion       | 0   | 6   | $p < 0.002$              |
| Nonspecific asymmetry   | 0   | 2   |                          |
| Tubes                   |     |     |                          |
| Total obstructed        | 8   | 56  (17 %) | $p < .0001$             |
| Unilateral              | 7   | 47  |                          |
| Bilateral               | 1   | 9   |                          |
| Anomalies               | 4   |     |                          |

Abbreviations: TVS transvaginal ultrasound, HSG hysterosalpingography
Diagnostic imaging plays an important role in the assessment of women with infertility. Although no consensus protocol for work up of these patients exists, the majority of infertility patients undergo a baseline TVS and HSG. TVS is used for evaluating ovaries, fallopian tubes, and the adnexa and is a favored imaging modality in the infertility population because it is readily available, relatively low cost, and does not use ionizing radiation. TVS is the test of choice for diagnosing polycystic ovary syndrome [5], and is helpful for identifying endometriosis and the sequelae of PID. In addition, TVS is invaluable for monitoring ovarian folliculogenesis during treatment with ART [6–8]. In contrast, HSG provides information about tubal patency and uterine cavity abnormalities such as anomalies, polyps, synechiae, and adhesions, any of which could interfere with embryo implantation [9]. However, HSG offers limited evaluation of the cervix and myometrium and does carry the small risks of contrast reaction and of ionizing radiation exposure [10]. Besides TVS and HSG, supplemental evaluation with SIS and hysterosalpingo-contrast sonography (HyCoSy) is sometimes performed. These imaging procedures are becoming more popular because of their ability to combine TVS adnexal evaluation with HSG-like assessment of the uterine cavity, without the risks of contrast reactions and radiation exposure [11–13], but are not yet universally available.

MRI of the pelvis offers multi-planar imaging and does not require the use of ionizing radiation. It is an excellent modality for detecting endometriosis [5] and is helpful for determine the nature of uterine duplication anomalies, leiomyomas, and adenomyosis [14–18]. MRI is also employed for evaluating intracranial causes of infertility, such as pituitary adenomas. However, due to its high cost and limited access, MRI is not typically used in the infertility assessment except for a specific indication requiring such imaging.

\[ p = 0.0001 \] and HSG only 2 \( (p = 0.0007) \). HSG outperformed HSC for tubal obstruction, which only detected 5 of the 19 patients with unilateral or bilateral obstruction \( (<0.0001) \)

**Discussion**

Diagnostic imaging plays an important role in the assessment of women with infertility. Although no consensus protocol for work up of these patients exists, the majority of infertility patients undergo a baseline TVS and HSG. TVS is used for evaluating ovaries, fallopian tubes, and the adnexa and is a favored imaging modality in the infertility population because it is readily available, relatively low cost, and does not use ionizing radiation. TVS is the test of choice for diagnosing polycystic ovary syndrome [5], and is helpful for identifying endometriosis and the sequelae of PID. In addition, TVS is invaluable for monitoring ovarian folliculogenesis during treatment with ART [6–8]. In contrast, HSG provides information about tubal patency and uterine cavity abnormalities such as anomalies, polyps, synechiae, and adhesions, any of which could interfere with embryo implantation [9]. However, HSG offers limited evaluation of the cervix and myometrium and does carry the small risks of contrast reaction and of ionizing radiation exposure [10]. Besides TVS and HSG, supplemental evaluation with SIS and hysterosalpingo-contrast sonography (HyCoSy) is sometimes performed. These imaging procedures are becoming more popular because of their ability to combine TVS adnexal evaluation with HSG-like assessment of the uterine cavity, without the risks of contrast reactions and radiation exposure [11–13], but are not yet universally available.

MRI of the pelvis offers multi-planar imaging and does not require the use of ionizing radiation. It is an excellent modality for detecting endometriosis [5] and is helpful for determine the nature of uterine duplication anomalies, leiomyomas, and adenomyosis [14–18]. MRI is also employed for evaluating intracranial causes of infertility, such as pituitary adenomas. However, due to its high cost and limited access, MRI is not typically used in the infertility assessment except for a specific indication requiring such imaging.
At our institution, we begin the infertility assessment with an HSG. If there is evidence of an abnormal uterine cavity from etiologies such as uterine septa, submucosal fibroids, synchiae, or polyps, HSC is then typically performed [19–21]. The standard practice at our institution is to perform HSC in the office setting, reserving operative HSC and laparoscopy for patients who are not able to tolerate office based procedures and for situations for which surgical correction is required, such as septoplasty for the correction of a subseptate uterus. Hysteroscopy is also performed prior to ART if there is a 6 month or greater delay between the HSG and ART. TVS is obtained when patients begin ART, and continues during folliculogenesis.

Our results indicate that TVS is superior to HSG for detection of myometrial pathology, including fibroids and adenomyosis. These results make intuitive sense, as TVS uses high frequency sound waves to evaluate the 3 dimensional volume and echotexture of the uterine tissue, while HSG uses radiographs and contrast dye to outline the endometrial cavity. By assessing the contour of the contrast-filled cavity, information about the surrounding myometrium can be inferred, but not diagnosed, because the tissue itself is not imaged directly. HSG may detect submucosal fibroids, but other myometrial pathology, such as intramural or subserosal fibroids, are likely to be missed. Similarly, TVS is superior to HSC, which visualizes the walls of the uterine cavity but cannot assess for lesions within the myometrium.

Our results also indicate that HSG is the superior modality for detection of tubal pathology, specifically tubal obstruction. This finding is in keeping with the functional component of HSG, which allows the operator to visualize in real-time contrast medium passing through the tubes and most importantly, spilling into the surrounding peritoneum. TVS can only infer tubal obstruction when a

Table 2 Myometrial, endometrial, and tubal abnormalities detected by each modality (N = 55)

| Category          | TVS  | HSG | HSC | Statistical significance |
|-------------------|------|-----|-----|--------------------------|
| Fibroids          | 17 (31 %) | 6   | 4   | p < .0001 (TVS vs HSC)   |
| Adenomyosis       | 6 (11 %)  | 0   | 0   |                          |
| Endometrium       |      |     |     |                          |
| Polyps            | 1    | 2   | 10  | p = .0001 (HSC vs TVS)   |
|                   |      |     |     | p = .0007 (HSC vs HSG)   |
| Cysts             | 1    | 0   | 0   |                          |
| Cavity distortion | 0    | 4   | 2   |                          |
| Nonspecific asymmetry | 0  | 2   | 1   |                          |
| Tubes             |      |     |     |                          |
| Total Obstructed  | 6    | 19 (35 %) | 5   | <0.0001 (HSG vs HSC)    |
| Unilateral        | 6    | 16  | 5   |                          |
| Bilateral         | 0    | 3   | 0   |                          |
| Anomalies         | 0    | 1   | 31  |                          |

Abbreviations: TVS transvaginal ultrasound, HSG hysterosalpingography, HSC hysteroscopy

Fig. 3 a 38-year-old G2P2 female with history of male factor infertility presenting for baseline assessment prior to initiating ART. Transvaginal grey-scale image of the left adnexa demonstrates an anechoic tubular structure (calipers), measuring 43 × 31 × 19 mm, separate from the left ovary (not shown) and consistent with hydrosalpinx. b HSG demonstrates contrast pooling within a dilated, blind ending fallopian tube (arrows), confirming the presence of a left sided hydrosalpinx. The right fallopian tube is normal in caliber and demonstrates free intraperitoneal spill of contrast, indicating tubal patency.
hydrosalpinx is present, therefore obstructed but nondis-
tended fallopian tubes will be missed with sonography
alone. Endometrial pathologies, specifically endometrial
polyps, were more frequently identified on direct
visualization with HSC than on TVS and HSG combined.
It is possible that, for some of our patients, the HSC pre-
ceded the TVS and/or HSG and, thus, polyps could have
been removed by the time of imaging evaluation. While
TVS and HSG are both potential screening modalities for
endometrial lesions, HSC is required for optimal diagnosis
(Fig. 4), and one reason why flexible office hysteroscopy
remains the gold standard for endometrial assessment.

A weakness in our study is that we did not assess SIS as
a method to evaluate the endometrium. This procedure is
included in some protocols during the work up of women
with infertility, but is not part of the routine assessment at
our institution. SIS has been shown to be superior to TVS
for identifying endometrial abnormalities including polyps
and cavity distortion [11, 13, 22–27]. Some reports have
also shown SIS to be comparable to the gold standard of
HSC for evaluation of intrauterine abnormalities including
polyps, submucosal fibroids, adhesions and uterine anom-
alies, with a sensitivity and specificity for detection of 88
and 94 %, respectively [28, 29]. In addition, none of our
patients were evaluated by HyCoSy, a procedure that uses
aerated saline or contrast to assess tubal patency with
TVS. HyCoSy has been shown to be comparable to HSG
with regards to assessing tubal patency, with sensitiv-
ity ranges from 75–96 % and specificity from 67–100 %
[12, 13, 30, 31]. SIS and HyCoSy can be done in a single
visit and together provide information about the uterine
cavity and the patency of the fallopian tubes, similar to
HSG, but with added information about the myometrium
from the TVS component, all without exposure to ioniz-
ing radiation or iodinated contrast. Despite these advan-
tages, HyCoSy does not provide anatomical information
about the fallopian tubes, which limits its utility.

Given the lack of a single all encompassing imaging
tool for accurately diagnosing endometrial, tubal, and
myometrial causes of infertility, it could be helpful to
outline one step-wise approach for use of the TVS, HSG,
and HSC. Although there is tremendous variability be-
tween practices, at our institution most infertility pa-
tients undergo both a TVS and HSG prior to initiating
ART. Others have found that SIS and HyCoSy provide
comparable information as TVS and HSG combined. If
findings of these tests suggest an abnormality within the
uterine cavity, which could prevent implantation of a vi-
able gestational sac, the patient will be referred for a
HSC for direct inspection and possible treatment. How-
ever, the management of abnormal tubal pathology on
HSG will vary depending on plan for reproductive ther-
apy. If the patient is an In-vitro Fertilization (IVF) candi-
date, tubal obstruction is not of much consequence, as
the embryo is directly implanted into the uterus. However,
if the patient is not a candidate for IVF, tubal obstruction
can be further managed with surgical interventions such
as tuboplasty or salpingostomy.

**Conclusion**

Our study compared the results from TVS, HSG, and
HSC in a cohort of female infertility patients. TVS was
superior for detecting myometrial pathology, HSG was
superior for evaluating tubal patency, and HSC detected

---

**Fig. 4** 44 year-old G0P0 female with inability to conceive for 4 years presents for baseline assessment prior to IVF. TVS demonstrates an 11 x
11 x 10 mm echogenic lesion within the left aspect of the endometrial cavity (calipers). Flow was demonstrated within the lesion with color
Doppler, raising the possibility of endometrial polyp. b Corresponding HSG demonstrates a depended rounded filling defect within the left
aspect of the endometrial cavity, which persisted on multiple projections, suggestive of a polyp. The fallopian tubes are normal in caliber and
patent. The patient went on to HSC, where the lesion proved to be a submucosal fibroid.
more endometrial polyps than HSG and TVS. No single modality provided accurate identification of all different pathologies. Complete work up of women with infertility may include all modalities, given the unique information obtained from each. However, with knowledge of the unique specificity of each imaging test to detect specific pathologies, a combination of HSG, HSC and TVS could be selected based on the clinical presentation of patients.

Competing interests
The authors declare that they have no competing interests.

Authors’ contribution
CP helped to conceive of this study, lead data collection, reviewed the literature, and drafted this manuscript. CB participated in statistical analysis as well as reviewing the manuscript. EG helped to select references and aided in interpretation of hysteroscopic reports. MF conceived of the study and participated in its design and coordination as well as helped to draft the manuscript. All authors read and approved the final manuscript.

Acknowledgements
No other persons, besides those mentioned in the authorship, significantly contributed towards this study. There were no sources of funding for this research paper, all authors dedicated their academic time towards the preparation and design of this study.

Author details
1Department of Radiology, Brigham and Women’s Hospital, Harvard Medical School, 75 Francis Street, Boston, MA 02115, USA. 2Department of Infertility and Reproductive Surgery, Obstetrics and Gynecology, Brigham and Women’s Hospital, Harvard Medical School, 75 Francis Street, Boston, MA 02115, USA.

Received: 9 November 2015 Accepted: 15 December 2015
Published online: 23 December 2015

References
1. McLaren JF. Infertility evaluation. Obstet Gyneco Clin North Am. 2012;39:453–63.
2. Chandra A, Martinez GM, Mosher WD, Abma JC, Jones J. Fertility, family planning, and reproductive health of U.S. women: data from 2002 National Survey of Family Growth. Vital Health Stat. 2012;231–160.
3. Chandran A, Copen CE, Stephen EH. Infertility service use in the United States: Data from the National survey of Family Growth. Vital Health Stat. 2012;23:1.
4. Sadow CA, Sahni VA. Imaging female infertility. Abdom Imaging. 2014;39:92.
5. De Crespiigny LC, O’Herlihy C, Robinson HP. Ultrasound evaluation of the mechanism of human ovulation. Am J Obstet Gynecol. 1981;139:636–9.
6. Simpson WL, Bentia LG, Mester J. Hysterosalpingography: a reemerging study. Radiographics. 2006;26:419–31.
7. Baramki TA. Hysterosalpingography. Fertil Steril. 2005;83:595–606.
8. O’Neill MJ. Sonohysterography. Radio Clin North Am. 2003;41:781–97.
9. Schief R, Deichert U. Hysterosalpingo-contrast sonography of the uterus and fallopian tubes: results of a clinical trial of a new contrast medium in 120 patients. Radiology. 1991;178:213–5.
10. Saunders RD, Shwayder JM, Nakajima ST. Current methods of tubal patency assessment. Fertil Steril. 2001;75:217–9.
11. Behr SC, Courtier JL, Qayyam A. Imaging of Mullerian duct anomalies: MR imaging. Abdom Imaging. 2011;36:756–64.

Submit your next manuscript to BioMed Central and we will help you at every step:
• We accept pre-submission inquiries
• Our selector tool helps you to find the most relevant journal
• We provide round the clock customer support
• Convenient online submission
• Thorough peer review
• Inclusion in PubMed and all major indexing services
• Maximum visibility for your research

Submit your manuscript at www.biomedcentral.com/submit