

## Repeated IVF/ICSI-ETs failures and impact of hysteroscopy

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### Abstract

**Background:** Despite numerous developments in the field of assisted reproduction, the implantation rate remains low. Among the various reasons of implantation failure, endometrial regularity has an important role. Hysteroscopy is an accurate method for evaluating the endometrial characteristics, with the ability to treat uterine pathology.

**Objective:** The aim of the present study was to evaluate the findings on hysteroscopy and thereafter the result of subsequent IVF/ICSI in infertile women with the history of frequent unexplained and unsuccessful IVF/ICSI attempts.

**Materials and Methods:** In this observational study, the hysteroscopy findings and the outcomes of subsequent IVF/ICSI were evaluated in 89 infertile women admitted in Avicenna Infertility Clinic, with previous repeated (more than two) failed IVF/ICSI-ETs, including the patients with normal Hysterosalpinography (HSG) and excluding severe male factor infertility and also thrombophilia, genetic and immunologic problems. The data were analyzed with SPSS software and Fisher exact, chi-square, and MC-Nemar tests.

**Results:** In 53 (59.5%) cases, hysteroscopy revealed abnormal intrauterine findings including adhesions 7 (13.7%), single polyp 11 (20.7%), endometrial polyposis 10 (18.8%), endometrial hyperplasia 10 (18.8%), uterine cavity hypoplasia 4 (7.8%) and myoma 5 (9.8%). These abnormalities were significantly higher in women with more than 8 years of infertility (chi-square=4.7, p-value=0.03). After hysteroscopy and subsequent IVF/ICSI-ET attempt using standard long protocol, pregnancy rate were significantly higher compared with the previous repeated IVF/ICSI attempts (35.8% versus 0%).

**Conclusion:** According to this study, we strongly suggest evaluation of endometrial integrity by hysteroscopy in patients with repeated IVF/ICSI-ETs failure, before entering any other fertilization procedures.

**Key words:** Hysteroscopy, Repeated IVF/ICSI -ET failure.

### Introduction

Although there was gradual increase in the

success of reproduction, over the years, but many couples also had been left frustrated following repeated failed attempts. Clinicians who treated unsuccessful couples often face a challenge.

The probable causes of repeated IVF failures were classified as: reduced endometrial

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receptivity, embryonic defects or multifactorial causes (1). Intrauterine and endometrial integrity abnormalities such as thin endometrium, altered expression of adhesive molecules and immunological factors like Anti Sperm (ASA), Anticardiolipin (aCL), Lupus anticoagulant (LA), Anti-Phosphatidylserine (aPS), Anti-Phosphatidylethanolamine (aPE), and Antinuclear antibody (ANA), Anti DNA, Anti Zona and Anti ovarian (AOA) antibodies (2), Thrombophilia (3), decrease expression of endometrial integrins, increase of natural killer cells activities and imbalance of cytokine networks (balance between IL-12 and IL-18) (4), may decrease endometrial receptivity, whereas chromosomal and genetic abnormalities of the male sperm or female ovarian defects, embryonic aneuploidy or zona hardening are embryonic reasons for the failure of implantation (5,6). Among the various etiologies that were described, endometrial regularity played an important role in infertility and success of IVF programs.

Hysterosalpingography (HSG) is sensitive but its specificity for detection of intrauterine abnormalities is low (23%), with a false positive rate of 44% and false negative rate of 10% (7). Transvaginal sonography (TVS) is more specific (96.3%) and sensitive (81.8%) than HSG with negative (97.6%) and positive (73.8%) predictive values for the detection of any intrauterine abnormalities (8, 9). Accordingly, saline infusion sonohysterography (SIS), is an accurate and safe method in the evaluation of the uterine cavity (10). Kelekci et al showed equal accuracy between hysteroscopy and SIS but lower pain score in saline infusion sonography (11). Although SIS showed high failure rate in Rogerson study (12), however hysteroscopy showed to be more accurate in the evaluation of intracavity pathology in comparison with other clinical studies (8). Treatment of these abnormalities prior to the subsequent IVF/ICSI-ET raises the pregnancy rate to 22% in Golan and Schiano et al studies (7, 13).

Although the previous trials showed the usefulness of uterine reassessment by hysteroscopy in women with two IVF failures, (14-16) but they didn't exclude the other probable contributory factors like thrombophilia and immunological disorders.

In the present observational study we evaluated hysteroscopy abnormalities; and also the subsequent IVF/ICSI-ET outcome in patients with unexplained repeated IVF/ICSI-ET failure after excluding all other possible ethological factors.

## Materials and methods

### Ethics of experimentation

Investigations have been approved and the trial has been authorized under the decision of Ethical Committee of Avicenna Research Institute.

### Patients

This study was performed from July 2003 to October 2006, at the Avicenna Infertility Clinic on infertile women (25-38 years old) who referred to the clinic with the history of more than two previous IVF/ICSI-ET failures despite transfer of a minimum three good-quality embryos in each attempt. After taking history and physical exam; HSG, and routine hematological, biochemical and hormonal tests, semen analysis, and also flowcytometry, autoantibodies profile like Anti-Cardiolipin (aCL), Lupus Anticoagulant (LA), Anti-Phosphatidylserine (aPS), Anti-Phosphatidylethanolamine (aPE), and Antinuclear Antibodies (ANA), Anti DNA, and thrombophilia profile like MTHFR gene, prothrombin gene, Factor V Leiden gene, serum homocystein, protein C, Protein S, anti-thrombin 3 and karyotype have accomplished in all women referred to our clinic with more than two IVF/ICSI-ET failures. All infertile women underwent standard TVS (Pie medical, model 260 Corvus) with convex 7.5 MHz transvaginal ultrasound probe. We excluded untreated HSG shown abnormalities, thrombophilia, immunological and, genetic problems, and also severe male factor infertility in selected patients of our study.

### Measurements

We informed all cases about the technique, therapeutic effect and potential risks (informed consent) of hysteroscopy and obtained informed consent. The selected patients underwent a diagnostic and/or operative hysteroscopy in early follicular phase of the cycle and all procedures were done between the 7th and 11th day of the cycle. The interventions were performed under general anesthesia in normal lithotomic position. After cervical dilatation of 5-9 mm, operative rigid hysteroscope (Olympus) was entered under visual control into the uterine cavity. A continuous flow instrument with separate in- and out-flow channel was used which was connected to a video camera system. Dextrose 5% was used for distention medium, keeping the distention pressure between

100-150 mmHg. Duration of the procedure was kept as short as possible with continuous surveillance of the fluid balance. Intrauterine adhesions, polyps, and submucosal myomas were treated by scissor and resectoscope during hysteroscopic evaluation procedure; and by curettage after hysteroscopy in the cases with endometrial hyperplasia. Endometrial biopsy has been done in the cases of uterine cavity hypoplasia during hysteroscopy. The given findings in hysteroscopy have been analyzed.

### Protocol of stimulation in subsequent IVF/ICSI attempt

Two months after hysteroscopy, the patients underwent ovarian stimulation with standard long protocol. GnRH-agonist (Buserlin: superfact, Aventis Pharma, Germany 0.5 mg/day) was administered from the day 21 of the cycle. Then all the patients were treated with human menopausal gonadotropin (hMG: Merional, IBSA, Switzerland, 150-300 IU/day) from day 2-3 of the next cycle, while continuing superfact 0.25 mg/day, with the control of follicular growth under sonography every 3-4 days. HCG was administered 10000 IU when the minimum of 3 leading follicles reached 16-18 mm, and 36 hours later oocyte collection was performed.

### Statistical analysis

Data were expressed as mean values  $\pm$  SD for numerical variables and in percentage for categorized variables. The data were analyzed with

SPSS software and Fisher exact, chi-square, and MC-Nemar tests.

### Results

The etiologies of infertility in the study group have been shown in table I. The study population consisted of 89 infertile women, with the mean age of ( $31.02 \pm 3.28$ ) years, mean duration of infertility ( $8.56 \pm 2.91$ ) years and mean numbers of previous ET attempts were ( $2.78 \pm 0.74$ ).

**Table I.** The characteristics of the subject group. (n=89)

Characteristics	mean $\pm$ SD
Age (years)	31.02 $\pm$ 3.28
Duration of infertility (years)	8.56 $\pm$ 2.91
Mean no of previous ET trial (no)	2.78 $\pm$ 0.74
<b>Etiology of infertility No (%)</b>	
Male factor	55(61.7)
Ovarian factor	13(14.6)
Tubal and peritoneal factor	6(6.7)
Unexplained	14(15.7)

Abnormal TVS findings were observed in 47 cases (52.8%) in which endometrial irregularity 17(36.1%), hyperplasia 14(29.7%), polyps 11(23.4%) and myoma 6(6.4%) were detected. Abnormal hysteroscopy findings were observed in 53 cases (59.5%) in which intrauterine adhesions, endometrial hyperplasia, polyps, were the most common abnormalities rather than submucosal myoma, uterine cavity hypoplasia, and endocervical polyps (Table II). According to our results, using Mc-Nemar test ( $\chi^2=4$ ,  $p<0.5$ ), sonography is more specific (100%) but not sensitive (88.6%) compared to the hysteroscopy, with false negative rate of 19.4%.

**Table II.** Findings on hysteroscopy in the study group.

Findings on hysteroscopy, No (%)	Total	Age $\geq 30$ years (n=54)	Age $< 30$ years (n=35)	infertility $\geq 8$ years (n=55)	Infertility $< 8$ years (n=34)
	53 (59.5)	34 (69.3)	19 (65.51)	37 (77.08)	16 (53.3)
Submucosal Myoma	5 (9.8)	5	0	4	1
Single large polyp	11 (20.7)	8	3	9	2
Polypoid endometrium	10 (18.8)	6	4	6	4
Endometrial hyperplasia	10 (18.8)	8	2	7	3
Uterine cavity hypoplasia	4 (7.8)	3	1	2	2
Adhesions	7 (13.7)	3	4	5	2
Endocervical polyp	3 (5.8)	1	2	0	3
Multiple lesions	3 (5.8)	2	1	3	0
Without any finding	36(40.44)	20	16	18	18

**Table III.** Comparison of characteristics of subsequent IVF/ICSI –ET cycle between patients with treated abnormal findings and patients with normal findings on hysteroscopy.

Cycle characteristics after hysteroscopy	With abnormal findings (n=53)	No finding (n=31)	p-value
No of retrieved oocyte ( mean $\pm$ SD)	12.01 $\pm$ 2.1	10.4 $\pm$ 2.8	NS*
No of embryo transferred (mean $\pm$ SD)	3.5 $\pm$ 2.8	3.3 $\pm$ 0.6	NS
Clinical pregnancy rate [no (%)]	19(35.8)**	8(22.2)	NS

\*NS = not significant

\*\* = significant, compared with previous repeated failed IVF/ICSI-attempts (0% versus 35.8%)

However hysteroscopy showed to be more sensitive and specific in the evaluation of intracavity pathology in comparison with TVS in patients with the history of repeated IVF/ICSI-ET failures after exclusion of other possible caus.

Single polyp and endometrial hyperplasia have been found more in patients more than 30 years old and with more than 8 years infertility, but it was not significant by using Fisher exact test (one tail) ( $p=0.27$ ). Abnormal findings in patients with  $\geq 30$  years were not significant compared with younger patients, using chi-square=0.44 ( $p=0.51$ ). More hysteroscopy abnormalities were found in the infertile patients with more than 8 years infertility, (chi-square=4.7,  $p=0.03$ ), which was significant.

Structural abnormalities correlated with the presence of pathological abnormalities were seen in 94.3% ( $n=50$ ) of cases. Also in 13.2% ( $n=7$ ) of cases, nonspecific endometritis was reported. We had no early or late, major or even minor complications in our series.

In patients with abnormal findings in hysteroscopy, subsequent IVF/ICSI-ET cycles after the procedure resulted significantly higher pregnancy rate, compared with previous failed attempts (35.8% versus 0%,  $p<0.001$ ), although, there were not statistically significant difference in pregnancy rate between the patients with abnormal findings compared with whom with normal findings in hysteroscopy (35.8% versus 22.2,  $p=0.17$ ,  $k2=1.88$ ).

## Discussion

TVS is the first clinical diagnostic test in the investigation of the uterine cavity and is especially important as a noninvasive technique to plan hysteroscopy. The sensitivity, specificity, and positive (PPV) and negative predictive values (NPV) for TVS in detecting abnormal uterine cavities have been shown as 100%, 96.3%, 91.3%, and 100%, respectively (8). However SIS is superior to pelvic ultrasound in detecting intracavitary pathology (17). TVS had sensitivity

of 72%, specificity of 92%, PPV of 94% and NPV of 65%, while SIS had sensitivity of 94.1%, specificity of 95%, PPV of 96% and NPV of 90% (18). Hysteroscopy is a gold standard instrument, offers diagnostic accuracy for evaluating the uterine characteristics with the ability to treat uterine pathology in infertile women especially in patients with repeated IVF failures (19), although; it couldn't show the functional status of the endometrium. Ragni et al compared the sensitivity, specificity, positive predictive value and negative predictive value of TVS and hysteroscopy (20).

The TVS's sensitivity and specificity in comparison with hysteroscopy were 91% and 83% respectively. The false positive rate was 9.2% and a 5.1% false negative rate was detected compared to hysteroscopy. Also Hauge et al showed that hysteroscopy and TVS findings were similar in 90.9% (9). Our study showed TVS is specific, but not sensitive compared to the hysteroscopy. Capability to obtain a direct view of the uterine cavity by hysteroscopy showed to be more accurate in the evaluation of intracavity pathology in patients with unexplained IVF failures. Previous studies reported a high incidence of intrauterine abnormalities in patients undergoing IVF/ICSI-ET.

Faghali et al evaluated the benefits of a diagnostic hysteroscopy prior to IVF which shows the systematic hysteroscopy prior to IVF could improve the pregnancy rate (21). La Sala and Oliveira et al showed relation between IVF-ET failure and unsuspected intrauterine abnormalities (22, 23).

Makris et al performed hysteroscopy in patients with history of abortions, infertility and repeated failure of IVF. They showed that abnormal hysteroscopic findings were observed in 40.5% of cases in which intrauterine adhesions, endometrial hyperplasia and polyps were the most common (16). According to the results of our study, abnormal hysteroscopic findings were observed in 59.5% of the cases with unexplained repeated IVF failures which is higher than the result of previous

studies due to exclusion of the other possible reasons of repeated ET failures like, thrombophilia, chromosomal and immunologic factors. Dicker et al showed, uterine abnormalities were detected in about 18% women with normal initial hysteroscopy who had three or more IVF-ET failures and underwent repeated hysteroscopy (15).

Also Schiano et al showed abnormalities in half of the cases like cervical abnormalities (synechia, polyp, and false passage) and hormonal-dependent abnormalities (polyp, hyperplasia, submucous myoma) in repeated uterine hysteroscopy after two implantation failures in IVF (13). Operative hysteroscopy showed to be effective in the treatment of intra-uterine adhesions with infertility in Kdous study with good results in achieving pregnancy (24).

Pabuccu et al observed quite an improvement in access to the endometrial cavity during ET procedure after the hysteroscopic shaving procedure in patients with difficult ETs and failed IVFs attempts due to cervical stenosis (25). We treated the structural abnormalities at the time of hysteroscopy and we had no early or late, major or even minor complications in our series. The pathological reports confirmed the observed lesions in 94.7% comparable to the results of Woolcott study (26).

Dicker study showed, that in elderly women, age-related uterine pathology such as submucous myoma, endometrial hyperplasia, and polyps were more prominent, while in younger patients other uterine lesions such as adhesions and tubal ostia occlusion were more common (15). In our experiences, we have seen single polyp and endometrial hyperplasia more in the patients with  $\geq 30$  years olds rather than the younger group ( $p$ -value= 0.78). Also there were more abnormalities especially single polyp, polypoid endometrium and also endometrial hyperplasia in group of the patient with more than 8 years infertility ( $p$ -values: 0.08, 0.6, and 0.12, respectively).

According to the result of our study, treatment of the endometrial pathology resulted 34.8% pregnancy rate in the subsequent IVF/ICSI-ET, which is more than the results of Golan (22%) (7), Mihaila (16.6) (19) and Schiano (22%) (13). This difference seems to be attributed to the exclusion all other factor related to IVF failure, which limited performing hysteroscopy with best result just in patients with endometrial pathology.

## Conclusion

TVS showed as a noninvasive, specific but not sensitive method for detecting intracavitary pathology. Hysteroscopy proved to be a very useful, accurate and safe method of assessing and treatment of uterine and endometrial pathologies in patients with repeated IVF/ICSI-ET failures after excluding other possible reason for implantation failure, although it couldn't evaluate uterine and endometrial functional status.

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