Risk factors for ectopic pregnancy in assisted reproductive technology: a 6-year, single-center study

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Objective: To explore factors affecting the incidence of ectopic pregnancy (EP) in assisted reproductive technology (ART). **Design:** A retrospective cohort study on the incidence of EPs in IVF/intracytoplasmic sperm injection (ICSI) and IUI cycles from June 2009 to August 2015. Age of patients, tubal factor infertility, type of cycle (fresh or thawed), embryo being transferred (cleavage embryo or blastocyst), and number of embryos transferred were analyzed to explore their relationship with the incidence of EP. **Setting:** Teaching hospital.

Patient(s): A total of 18,432 pregnancies resulting from ART treatment were retrospectively analyzed.

Intervention(s): None.

Main Outcome Measure(s): Ectopic pregnancy rate.

Result(s): For IVF/ICSI cycles, the incidence of EP was different between cycles transferred with cleavage embryo and blastocyst (3.45% vs. 2.47%). In multivariate logistic regression analysis, tubal infertility was associated with EP (adjusted odds ratio 1.716, 95% confidence interval 1.444–2.039). For IUI cycles, EP was significantly higher in stimulated cycles compared with natural cycles (2.62% vs. 0.99%). The EP rate in cycles with sperm from donor and husband was 1.08% and 3.54%, respectively. However, when patients were stratified according to tubal infertility, the EP rate increased with level of peak estrogen. In thawed embryo transfer cycles, the EP rate was lower in blastocyst transfer cycles and in cycles transferred with fewer embryos.

Conclusion(s): Irrespective of tubal infertility, for fresh IVF/ICSI cycles the rate of EP is positively associated with ovarian stimulation; for thawed IVF/ICSI cycles, blastocyst transfer or transfer with fewer embryos reduces the EP rate. In IUI cycles, EP is associated with sperm source. (Fertil Steril® 2016; ■ : ■ – ■. ©2016 by American Society for Reproductive Medicine.) **Key Words:** ART, ectopic pregnancy, relative risk, tubal factor



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ctopic pregnancy (EP) after assisted reproductive technology (ART) procedures is a complication with potentially serious consequences for patients. There are multiple reports that describe a higher incidence of EPs after various methods of ART (2.2% to

4.5%) than in spontaneous pregnancy (1–3).

In previous studies, many factors have been described as being associated with EP (3, 4). Of these, the difference of EP between fresh and thawed embryo transfer cycles and blastocyst transfer have attracted much interest from

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clinicians. It has been shown that blastocyst transfer may reduce EP in IVF/intracytoplasmic sperm injection (ICSI) cycles (5, 6), even though this conclusion is inconsistent with other studies (7, 8). Meanwhile, many studies demonstrate a lower incidence of EP in thawed embryo transfer cycles compared with fresh embryo transfer in stimulated cycles (9-12), indicating that the tubal-uterine environment after ovarian stimulation has been different from physiologic status, which contributes to abnormal implantation eventually. This point of view has been supported by a recent study with a large sample size (>100,000 pregnancies) from the

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Society for Assisted Reproductive Technology registry in United States (13).

Because the incidence of EP is relatively low, and many factors are associated with EP, a large sample size is needed in such studies to reach a reasonable conclusion. Moreover, several EP-associated factors should be analyzed at the same time. Thus, to explore factors affecting the incidence of EP in ART, and to explore possible ways to reduce EP in different ART procedures, we retrospectively analyzed nearly 50,000 ART cycles during 6 years at our center. We hypothesize that factors associated with EP may be different in multivariate analysis.

MATERIALS AND METHODS

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134 This study was approved by the institutional review board of 135 First Affiliated Hospital of Zhengzhou University. For patients 136 undergoing ART (IVF/ICSI and related technology, IUI) treatment in our center, all have allowed use of their medical re-138 cord data for research. Written, informed consent was 139 obtained from all patients before IVF treatment. All patients 140 were registered in our data management system, which is used to save all medical information of patients trying to 142 conceive through ART.

143 From June 2009 to August 2015, 18,432 cycles resulted in 144 a reported clinical intrauterine, ectopic, or heterotopic preg-145 nancy, and were included in this study. For the ART follow-146 up procedure in our center, all patients are asked to come 147 back for a blood hCG test 14 and 18 days after ET or IUI. Those 148 with hCG elevation return for ultrasound examination 149 35 days after ET or IUI. Thus, [1] most of the EP cases are diag-150 nosed with ultrasound examination (ultrasound scan, espe-151 cially transvaginal, showing a gestational sac with fetal 152 heart in the adnexa area). [2] Pregnancy of unknown location, 153 which indicates there is a positive pregnancy test result but no 154 pregnancy has been visualized using transvaginal ultrasound 155 examination, is also included. In this case, we usually suggest 156 that patients stay in hospital, and we administer mifepristone 157 or methotrexate when necessary. [3] Sometimes laparoscopy 158 or laparotomy are also used to visually confirm an EP. In 159 this case, EP is also confirmed by pathology. However, most 160 of these patients are treated in local hospitals and inform us 161 by telephone. [4] In rare cases dilation and curettage is also 162 performed to diagnose EP. The clinical intrauterine pregnancy 163 was defined as documentation of one or more gestational sacs 164 visible by ultrasound examination. Heterotopic pregnancy 165 was defined as a pregnancy that met the criteria for both 166 ectopic and clinical intrauterine pregnancy. Heterotopic preg-167 nancy was also grouped into EP in this study. 168

The parameters evaluated were patient age, infertility diagnosis (tubal infertility or not), type of cycle (fresh or frozen-thawed), type of embryos transferred (cleavage embryo or blastocyst), and number of embryos transferred (one, two, or three). For IUI cycles the impact of IUI cycle type (natural or stimulated) and sperm source (husband or donor) on incidence of EP was also evaluated.

Multivariate logistic regression analysis was used to identify the prognostic variables mentioned before and to evaluate the association between the variables and EP. For IVF/ICSI

cycles, all patients were stratified by tubal infertility, and risk factors for EP were re-evaluated. Tubal infertility mainly includes [1] surgery history on fallopian tubes: previous EP, salpingostomy, and tube reconstruction surgery for infertility treatment; [2] hysterosalpingography examination: hydrosalpinx, salpingitis (however, saline salpingogram or clamydia antibody titers were not used to diagnose tubal factor); [3] salpingectomy and ligation are not included in this study. In fact, for patients with hydrosalpinx, we sometimes suggest tube ligation and a microcoil device to perform hysteroscopic tubal occlusion for treating hydrosalpinx. A *P* value of < .05was considered statistically significant.

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RESULT

In the 18,432 pregnancies, the incidence of EP was 3.17% (585 of 18,432). The incidence of EP in IVF/ICSI and IUI cycles was 3.33% (538 of 16,139) and 2.05% (47 of 2,293), respectively.

Overall, EP had no difference in the three age groups (<28, 29-37, and 38 years). For patients diagnosed with secondary infertility, the EP rate was significantly higher than that in patients with primary infertility (3.92% vs. 2.61%, P=.000). In addition, EP in patients with tubal infertility was also higher when compared with that in patients without tubal infertility (4.36% vs. 2.50%, P=.000) (Table 1).

TABLE 1

Ectopic pregnancy rate in different groups.			
Cycles	EP rate, % (n)	χ²	P value
All ART cycles			
Age (y) <28 29–37 >38	2.87 (168/5855) 3.38 (381/11257) 2.73 (36/1320)	0.7647	.3819
Infertility diagnosis Primary infertility Secondary infertility	2.61 (275/10518) 3.92 (310/7914)	24.9325	.000
Tubal factor existed Yes No	4.36 (292/6698) 2.50 (293/11734)	48.1318	.000
Type of ART IVF/ICSI IUI IVF/ICSI cycles	3.33 (538/16139) 2.05 (47/2293)	10.7682	.001
Type of transfer Fresh embryo Thawed embryo	3.22 (327/10149) 3.52 (211/5990)	1.0559	.304
Stage of embryo Cleavage stage Blastocyst stage	3.45 (491/14240) 2.47 (47/1899)	4.9232	.0265
No. of embryos transferred 1 2	3.51 (53/1512) 3.09 (346/11214)	3.3802	.0660
3 IUI cycles	4.07 (139/3413)		
Type of cycle Natural cycle Stimulated cycle	0.99 (8/805) 2.62 (39/1488)	6.8892	.0087
Sperm source Husband Donor	3.54 (32/904) 1.08 (15/1389)	16.5046	.000
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