

Clinical comparison of mifepristone and gestrinone for laparoscopic endometriosis

Junhong Song^{1**}, Yan Wang^{1**} and Lili Yu^{2*}

¹Department of Gynecology and Obstetrics, The Affiliated Yantai Yuhuangding Hospital of Medical College, Qingdao University, Yantai, China

²Department of Center for Reproductive Medicine, The Affiliated Yantai Yuhuangding Hospital of Medical College, Qingdao University, Yantai, China

Abstract: Endometriosis is a common disease among women of childbearing age, and it is the main cause of dysmenorrhea and infertility. This article analyzes the efficacy of mifepristone and gestrinone in the treatment of endometriosis. The results showed that the recurrence rate of mifepristone group and gestrinone group were 8.33% and 5%, respectively, which was significantly lower than 23.33% of the control group. Before and after treatment, LH, endocrine test results FSH PRL had no obvious change in mifepristone group and gestrinone group, while E2 decreased, as mifepristone group (141.7±31.2) pmol/L, gestrinone group (64.2±11.7) pmol/L. The incidence of adverse reactions and liver dysfunction in the mifepristone group were significantly lower than those the gestrinone group (P<0.05). Mifepristone and gestrinone can be used for endometriosis postoperative adjuvant treatment, is safe and effective, but using mifepristone has the lower rate of adverse reaction. In conclusion, mifepristone is a current research focus, its mechanism of action in the process of exploration, has broad prospects in the treatment of endometriosis, its long-term application security is paid more and more attention.

Keywords: Mifepristone, endometriosis, clinical efficacy, hormone level, gestrinone.

INTRODUCTION

In recent years, its incidence increased year by year (Rafique *et al.*, 2017). The disease is a benign lesion with metastasis and implantation characteristics, treatment more difficult (Yu *et al.*, 2016). In the past, surgery was the main treatment. But the disease is easy to relapse after surgery. The application of adjuvant therapy may help stabilize postoperative efficacy, improve the prognosis (Rivosecchi *et al.*, 2014). At present, the treatment of endometriosis with infertility is mainly treated with two kinds of medical treatment and surgical treatment, of which surgical treatment can relieve the pain of the patients, but the surgical trauma is larger and the postoperative recurrence is easy (Taylor *et al.*, 2017). Traditional drug therapy can relieve the symptoms of endometriosis and improve the outcome of pregnancy. Mifepristone and gestrinone are the main drugs for the treatment of endometriosis (Salengros *et al.*, 2010).

At present, gestrinone and mifepristone are the most frequently used drugs. Mifepristone is a synthetic 19- to methyl testosterone derivatives, mainly through the estrogen receptor and progesterone receptor down-regulation in ectopic endometrium, the ectopic endometrial degradation to achieve the treatment of endometriosis objective (Tosti *et al.*, 2017). Mifepristone inhibits ovulation, induces corpus luteum dissolution and interferes with endometrial integrity. It is a progesterone antagonist, which can directly act on ectopic endometrium, inhibit its proliferation and differentiation, promote

apoptosis, reduce its growth potential, effectively control endometriosis and reduce its recurrence (Vercellini *et al.*, 2014). Adhesion of endometriosis is common in laparoscopy and it is difficult to remove thoroughly during ectopic cyst separation because of its easy separation (Yu *et al.*, 2016). Surgical treatment can only remove the endometriosis, which can be identified by the naked eye. For the microscopic lesions, atypical lesions that can not be completely removed, the recurrence of the lesion can not be cleared after the operation (Taylor *et al.*, 2017). After the operation, 3~6 months of medication can be given to the atrophy of the unresectable or deep unresectable lesions (Salengros *et al.*, 2010). Therefore, it can prevent or delay the recurrence of the disease. Gestrinone main effect on endometrium and ectopic endometrium may play anti estrogen receptor and progesterone resistance strong, thus reducing the level of estrogen, but also with androgenic activity, can inhibit gonadotropin release, secretion of ovarian suppression, the ectopic endometrial atrophy (Goldstone *et al.*, 2017). To further explore the effective method for the treatment of severe endometriosis, at the same time in our hospital laparoscopic surgery for severe endometriosis patients, we analyzed treatment effect after surgery using mifepristone or gestrinone.

MATERIALS AND METHODS

Research object

180 patients with endometriosis from January 2016 to December 2017 in our hospital were randomly divided

*Corresponding author: e-mail: 1335192968@qq.com

**Junhong Song and Yan Wang contributed equally to this article

into mifepristone group, gestrinone group and control group, each group as 60 cases. The median age of mifepristone group was 21-42 years, with an average age of (31.7 ± 8.4) years, 31 cases in stage III and 29 cases in stage IV. Group gestrinone was 22-39 years old, with an average age of (32.8 ± 7.4) years, 28 cases in stage III and 32 cases in stage IV. The age of the control group was 22-40 years old, with an average age of (33.4 ± 9.5) years, 31 cases of stage III and 29 cases of stage IV. All patients were approved by ethics committee of affiliated Yantai Yuhuangding hospital, ethical approval number as 2015AYYHD12 and all patients signed on the informed consent. All patients had normal liver and kidney function, and no history of steroid therapy and contraindication without laparoscopic operation within six months. There was no significant difference in age, weight, parity, duration and staging between the three groups ($P > 0.05$).

Therapeutic method

The patients in the control group were treated with laparoscope operation. The operation of the patients was to inject the whole-body anesthetic first to the patient, to explore the pelvic abdominal cavity of the patient by laparoscope, and to separate the adhesive tissue in the pelvic cavity. Finally, the ectopic focus in the pelvic cavity was electrocoagulated, and the cyst was eliminated if the cyst appeared in the ovary. If the fallopian tube is closed, the fallopian tube is opened and the methylene blue solution is used as the fluid.

The patients in the control group did not give any drugs; the mifepristone group was treated with laparoscopic surgery as compared with the control group. Mifepristone group was given mifepristone for one week after operation, and 1 times/day, 10mg each time, oral administration for 6 months. Patients in group gestrinone were treated with laparoscopy and control group. In gestrinone group, gestrinone was given a week after operation, orally, two times a week, once 2.5mg, for 6 months. Three groups of patients were reexamined after laparoscopic surgery, and every three months. The review is to understand the patient's symptoms, signs, results of B-ultrasound etc. In mifepristone group and gestrinone group of patients during the treatment, is required for a visit every month, the main purpose of the visit is to understand the adverse drug reactions and follow-up in patients with liver function.

Observation index

The criteria of efficacy: (1) Relief: no symptoms, no signs, and found no pelvic masses; (2) improvement: Although there are primary symptoms, but the degree of symptoms was reduced, compared with before treatment reduced or no positive signs, and found no pelvic masses; (3) recurrence: Ultrasonography again found pelvic endometriosis cysts, or recurrence of symptoms and signs before treatment. And the incidence of liver function

damage records of mifepristone group and gestrinone group of adverse drug reactions.

Adverse reaction

(1) Androgen like effects: Weight gain, hairy, voice changes, acne, seborrheic dermatitis, etc. (2) Low estrogen: bleeding, hot flashes, sweating, irritability, etc. (3) Liver dysfunction: elevated transaminase. (4) Gastrointestinal reaction: Upper abdominal discomfort, nausea and vomiting, fatigue and so on.

STATISTICAL ANALYSIS

Statistical analysis was carried out using SPSS14.0 software, the measurement data using mean and standard deviation ($\bar{x} \pm s$), compared with two groups of data of mean t test, count data using χ^2 test, $P < 0.05$, the difference was statistically significant.

RESULTS

Comparison of the clinical efficacy of three groups of patients after treatment

The recurrence rates of mifepristone group and gestrinone group were 8.33% and 5% respectively, which were significantly lower than the control group 23.33%, the difference was statistically significant ($P < 0.05$); The total effective rate in mifepristone group and gestrinone group was significantly higher than that in the control group ($P < 0.05$). The total effective rate in mifepristone group was 91.66%, and the recurrence rate was 8.33%; while the total effective rate in gestrinone group as 95% and recurrence rate was 5%. The difference was not statistically significant ($P > 0.05$), as shown in table 1.

Hormone level comparison

The two groups before and after treatment, LH, endocrine test results FSH PRL had no obvious change; E2 decreased as the mifepristone group (141.7 ± 31.2) pmol/L, gestrinone group (64.2 ± 11.7) pmol/L, but were in the early levels of follicles, no significant differences between the two groups. The endocrine results of mifepristone group and gestrinone group before and after treatment were shown in table 2.

Serum ovarian cancer associated antigen

Serum levels of ovarian cancer associated antigen (CA125) and liver function were measured. After three groups of treatment, CA125 was lower than before treatment. Compared with the laparoscope group, the difference between mifepristone and Gestrinone group decreased significantly ($P < 0.05$), but there was no statistical difference between group Gestrinone and mifepristone group ($P > 0.05$). The data is shown in table 3.

Adverse reaction

In the mifepristone group, there were 5 patients with

Table 1: Comparison of the clinical efficacy of three groups of patients after treatment

Group	Cases	Relieve	Improve	Recrudescence	Total effective rate
Mifepristone group	60	35(58.33)	20(33.33)	5(8.33)	91.66%
Gestrinone group	60	34(56.66)	23(38.33)	3(5.0)	95.0%
Control group	60	18(30.0)	28(46.66)	14(23.33)	76.66%

Table 2: Endocrine changes before and after medication

Group	FSH (U/L)		LH (U/L)		E2 (pmol/L)		PRL(ng/ml)	
	Before drug use	After drug use	Before drug use	After drug use	Before drug use	After drug use	Before drug use	After drug use
Mifepristone group	5.13±1.35	4.26±1.55	4.35±2.03	3.62±1.45	176.3±35.2	141.7±31.2	17.2±3.8	21.6±5.4
Gestrinone group	4.85±1.26	5.83±2.72	5.12±1.87	4.72±1.41	157.6±18.3	64.2±11.7	22.8±4.7	17.2±5.1
Control group	5.24±1.63	5.26±1.77	4.64±1.56	4.72±1.38	164.1±31.8	168.7±30.5	19.6±3.2	19.1±4.2

Table 3: Difference of CA125 before and after treatment

Group	Cases	Difference before and after treatment	F value	P value
Mifepristone group	60	22.5±1.8	8.13	0.00
Gestrinone group	60	15.4±1.6		
Control group	60	12.1±1.1		

Table 4: Comparison of two groups of symptom scores

group	Observation time	Pelvic symptom score			Syndrome score	
		Dysmenorrhea	Coitus pain	Pelvic pain	Pelvic tenderness	Scleroma
Mifepristone group	Before treatment	7.12±1.52	6.83±1.34	6.75±1.63	5.82±1.36	6.17±1.52
Gestrinone group	After treatment	1.60±0.55	1.34±0.31	1.75±0.88	2.05±0.76	1.38±0.79
Control group	Before treatment	6.91±1.46	6.33±0.92	6.71±1.28	6.04±1.45	5.73±1.50
Mifepristone group	After treatment	2.36±0.84	1.70±0.64	2.48±0.94	2.84±0.63	2.16±0.55
Gestrinone group	Before treatment	6.83±1.53	6.04±1.44	6.13±1.48	6.42±1.59	5.94±1.23
	After treatment	7.12±1.86	5.89±1.23	5.27±1.57	5.31±1.42	5.72±1.15

Table 5: Comparison of adverse reactions

Group	Cases	Abnormal liver function	Gain weight	Acne	Hot flashes	Vaginal bleeding	Intimal thickening
Mifepristone group	60	5	2	1	0	2	5
Gestrinone group	60	17	11	10	8	7	6
Control group	60	1	1	0	0	1	1

abnormal liver function and the liver enzymes were slightly increased. The liver function recovered to normal after the treatment of unstopped medicine and with the treatment of liver protection. In group gestrinone, 17 cases were damaged, and most of the liver enzymes were significantly increased. The liver function recovered to normal after stopping and combining with the treatment of liver protection. The incidence of adverse reactions and liver function damage in mifepristone group were significantly lower than those in group gestrinone ($P < 0.05$). The adverse reactions after medication were compared as shown in table 5.

DISCUSSION

There are many treatments for endometriosis (EM), including drug therapy, surgery or surgery combined with drug therapy (Bhatt *et al.*, 2015). In recent years, with the new understanding and deepening of the pathogenesis of EM, there have been many new developments in the adjuvant regimens before and after operation. Traditional drugs including “tatazol, Gn RH-a” and so on have been severely restricted due to severe side effects and expensive prices (Bergström *et al.*, 1967; Carbonell *et al.*, 2012). In recent years, the application of gestrinone in the

treatment of endometriosis is effective. Gestrinone has strong anti progesterone and moderate anti estrogen, androgen and estrogen also has weak effect, can effectively promote the ectopic endometrial atrophy and absorption (Dindo *et al.*, 2004). However, the long-term use of gestrinone can lead to elevated liver enzymes, androgen effect, and low levels of estrogen, and may also cause osteoporosis and increased blood ester (Emir *et al.*, 2014). Mifepristone is a novel progesterone receptor antagonist, mifepristone has proved: Directly reduce estrogen and progesterone receptor content of endometrial function, and inhibit intimal interleukin secretion of -6 in immune regulation mechanism. Mifepristone can also alleviate the pain symptoms by inhibiting the production of prostaglandins (Goldstone *et al.*, 2017).

Endometriosis can cause the micro environment of the patients with intra-abdominal change, combined with the hinder of the egg and sperm egg cell division and pick up the egg process are the main factors leading to female infertility (Koh *et al.*, 2013). The anatomical structure of severe endometriosis can seriously damage the pelvic, uterus and rectum to cause adhesion and fallopian tube adhesion of uterus backward and pick up the egg function decline situation. In the past, laparoscopy is the first choice for severe endometriosis (Kertmen *et al.*, 2015). It can effectively improve the pregnancy rate of patients after operation. However, the recurrence rate of endometriosis is very high. Although the laparoscopic operation can remove the ectopic focus of the naked eye, the atypical, small and deep focus of the invasive site, the close adhesion, the rupture and the complete lesion can not be removed, and the recurrence of the disease is caused by the effect of hormone after the operation (Liu *et al.*, 2016). If the patient is given medication again after operation, the residual small lesions and naked eye lesions will be inhibited and atrophy and degeneration will occur.

Mifepristone is one of the progesterone receptor antagonists (Mukai *et al.*, 2012). It has many aspects in the treatment of endometriosis. Mifepristone progesterone itself has no activity, but the combination of mifepristone and its receptor, the content of hypothalamus pituitary hormone receptor regulation system ER, PR, to block the endometrial response to estrogen, progesterone, progesterone inhibitory activity (Mellotte *et al.*, 2015). In addition, mifepristone can also have a direct or indirect effect on the ovary, which affects the maturation and development of follicles, delays the ovulation period, induces the dissolution of the corpus luteum, and reduces the levels of progestin and estrogen in the patient's body, in favor of the treatment of endometriosis (Okada *et al.*, 2011). Mifepristone can also directly affect endometrium endometrium, inhibit its increment and differentiation, promote its apoptosis and reduce its growth ability.

In addition, gestrinone can also act directly on the receptor in patients with endometriosis and ectopic

endometrium of the endometrium and ectopic endometrial atrophy, so that the normal ovarian ovulation, improve the pregnancy rate (Carbonell *et al.*, 2013). A lot of information, for severe endometriosis was treated with laparoscopic surgery after treatment; medication can significantly improve the clinical efficacy and pregnancy rate, and reduce the postoperative recurrence rate (Ono *et al.*, 2009). Through this clinical treatment, the recurrence rate of mifepristone group and gestrinone group was significantly lower than that of the control group after treatment (Qi *et al.*, 2015). The difference was statistically significant. The incidence of mifepristone and the incidence of liver function damage were significantly lower than that of the gestrinone group, and the difference was statistically significant. Thus, after laparoscopic surgery patients were given mifepristone can be effective in the treatment of severe endometriosis, alleviate the symptoms, the endometriosis lesions appeared atrophy, and no obvious adverse reactions. Although gestrinone is effective medicine for the treatment of severe endometriosis, can reduce the recurrence rate, but the damage of the drug on liver function is more serious and more adverse reactions, which should be paid attention to.

CONCLUSION

The results showed no significant difference between the two groups of curative effect of mifepristone and gestrinone, the operation will remove the ectopic lesions, combined with drug therapy after operation, can effectively control the symptoms and reduce the recurrence. Mifepristone and gestrinone for postoperative adjuvant therapy of endometriosis are safe and effective. The adverse reaction rate of mifepristone is low. In conclusion, mifepristone is a current research focus, its mechanism of action in the process of exploration, has broad prospects in the treatment of endometriosis, its long-term application security is paid more and more attention. Low dose treatment batch may be an effective solution, without affecting the treatment effect, can effectively reduce the adverse reaction caused by long-term medication.

REFERENCES

- Bhatt A, Abe S, Kumaravel A, Vargo J and Saito Y (2015). Indications and Techniques for Endoscopic Submucosal Dissection. *Am. J. Gastroe.*, **110**(6): 784-791.
- Bergström J, Hermansen L and Hultman E (1967). Diet, muscle glycogen and physical performance. *Acta Physio. Scand.*, **71**(2): 140-150.
- Carbonell JL, Acosta R, Perez Y and Perez R (2012). Safety and effectiveness of different dosage of mifepristone for the treatment of uterine fibroids: A double-blind randomized clinical trial. *Int. J. Wom. Heal.*, **4**(2): 1-10.

- Carbonell JL, Acosta R and Perez Y (2013). 5 mg mifepristone daily versus placebo for 3 months to treat uterine myomas. Double blind randomized clinical trial. *Int. J. Wom. Hea.*, **5**(1): 5-12.
- Dindo D, Demartines N and Clavien PA (2004). Current status of peripheral fatigue research. *Discu. Mode. Econ.*, **240**(2): 205-213.
- Emir S, Sozen S, Bali I, Gürdal SO, Turan BC, Yıldırım O and Yetisyigit T (2014). Outcome analysis of laparoscopic D1 and D2 dissection in patients 70 years and older with gastric cancer. *Int. J. Clin. Exp. Med.*, **7**(10): 3501-3511.
- Goldstone P, Walker C and Hawtin K (2017). Efficacy and safety of mifepristone-buccal misoprostol for early medical abortion in an Australian clinical setting. *Aust. N. Z. J. Obstet. Gynaecol.*, **57**(3): 366-371.
- Koh R, Hirasawa K and Yahara S (2013). Antithrombotic drugs are risk factors for delayed postoperative bleeding after endoscopic submucosal dissection for gastric neoplasms. *Gastr. Endo.*, **78**(3): 476-483.
- Kertmen N, Babacan T, Keskin O, Solak M, Sarici F, Akin S, Arik Z, Aslan A, Ates O, Aksoy S, Ozisik Y and Altundag K (2015). Molecular subtypes in patients with inflammatory breast cancer; A single center experience. *J. BUON.*, **20**(1): 35-39.
- Liu K, Chen XZ, Nakamura I, Ohki S and Eslick GD (2016). Application of automatic data acquisition system in medical logistics system. *Logis. Tech. App.*, **2**(2): 135-142.
- Mellotte G, Maher V, Devitt PG, Shin VY and Leung CP (2015). Pharmaceutical logistics and supply chain management. *Beijing: Peking University Medical Press*, **1**(2): 101-112.
- Mukai S, Cho S and Kotachi T (2012). Analysis of delayed bleeding after endoscopic submucosal dissection for gastric epithelial neoplasms. *Gastro. Res. Pract.*, **12**: 875-883.
- Okada K, Yamamoto Y and Kasuga A (2011). Risk factors for delayed bleeding after endoscopic submucosal dissection for gastric neoplasm. *Surg. Endosc.*, **25**(1): 98-107.
- Ono S, Fujishiro M and Niimi K (2009). Technical feasibility of endoscopic submucosal dissection for early gastric cancer in patients taking anti-coagulants or anti-platelet agents. *Dig. Liver Dis.*, **41**(10): 725-728.
- Qi J and Liu X (2015). Dynamic changes of skeletal muscle ultrastructure, calcium dependent protease and ubiquitin after acute eccentric exercise. *Chi. J. Spo. Med.*, **2**: 13-16.
- Rafique S and Decherney AH (2017). Medical management of endometriosis. *Clin. Obstet. Gynecol.*, **60**(3): 485-496.
- Rivosecchi R, Rice MJ, Smithburger PL, Buckley MS, Coons JC and Kane SL (2014). An evidence based systematic review of remifentanyl associated opioid-induced hyperalgesia. *Expert. Opin. Drug. Saf.*, **13**: 587-603.
- Salengros JC and Huybrechts I (2010). Different anesthetic techniques associated with different incidences of chronic post-thoracotomy pain: Low-dose remifentanyl plus presurgical epidural analgesia is preferable to high-dose remifentanyl with postsurgical epidural analgesia. *J. Cardiothorac Vasc Anesth*, **24**: 608-616.
- Tosti C, Biscione A, Morgante G, Bifulco G and Luisi S (2017). Hormonal therapy for endometriosis: From molecular research to bedside. *Eur. J. Obstet. Gynecol. Reprod. Biol.*, **209**(17): 61-66.
- Taylor HS, Giudice LC, Lessey BA and Abrao MS (2017). Treatment of endometriosis-associated pain with elagolix, an oral GnRH antagonist. *N. Engl. J. Med.*, **377**(2): 28-32.
- Vercellini P, Viganò P, Somigliana E and Fedele L (2014). Endometriosis: Pathogenesis and treatment. *Nat. Rev. Endocr.*, **10**(5): 261-275.
- Yu SH, Yu CC, Yang XT, He SD, He J and Qin CT (2016). Pharmacoproteomic analysis reveals that metapristone (RU486 metabolite) intervenes E-cadherin and vimentin to realize cancer metastasis chemoprevention. *Sci. Rep.*, **6**(1): 17-20.