Ectopic pregnancy: A 5-step plan for medical management

Two experts describe a multidose methotrexate regimen, the first choice for unruptured, uncomplicated ectopics.

Although ectopic pregnancy remains a leading cause of life-threatening first-trimester morbidity, accounting for about 9% of maternal deaths annually,1 we now are able to diagnose and treat most cases well before rupture occurs—in some cases, as early as 5 weeks’ gestation. As a result, medical therapy with systemic methotrexate has become the first-line treatment, with surgery reserved for hemorrhage, medical failures, neglected cases, and circumstances in which medical therapy is contraindicated.

Early diagnosis not only makes medical therapy possible, it also is cheaper, since it avoids rupture, blood loss, and surgery; preserves fertility; and minimizes lost productivity. This is important because ectopic pregnancy is an expensive condition, with an annual health-care bill exceeding $1 billion.2

Despite this progress, serious challenges remain. Medical management is not for everyone. Success is inversely related to initial serum human chorionic gonadotropin (hCG) levels3 and diminishes sub-
stantially when embryonic cardiac activity is observed during ultrasound imaging.4

In addition, because medical therapy has made outpatient treatment the norm in most cases, it has become virtually impossible to chart the prevalence of ectopic pregnancy. In past years, when hospital records were used, ectopic pregnancy rates were increasing relentlessly, from 4.5 per 1,000 pregnancies in 1970 to 16.8 in 1989 and 19.7 (108,000 cases) in 1992.1,5

Reasons for increasing rates
Today the prevalence of ectopic pregnancy is probably still rising, for several reasons:
- a greater incidence of risk factors such as sexually transmitted and tubal disease,6
- improved diagnostic methods, and
- the use of assisted reproductive technology (ART) to treat infertility (roughly 2% of ART pregnancies are ectopic).7

This article describes a 5-step approach to diagnosis and medical management with multiple-dose methotrexate, as well as fine points of treatment and basic surgical technique. It includes a protocol for multiple-dose methotrexate, a table summarizing treatment outcomes, and several case histories.

Likelihood of ectopic pregnancy
If 100 women present with a positive pregnancy test and pain and bleeding, approximately 60 will have a normal pregnancy, 30 are experiencing spontaneous abortion, and 9 have an ectopic pregnancy.8

### KEY POINTS

- **In properly selected cases**, medical therapy and surgery produce similar outcomes, but medicine is less expensive.
- **Surgery is still the first choice for hemorrhage**, medical failure, rupture or near-rupture, and when medical therapy is contraindicated.
- **Systemic methotrexate** and laparoscopic salpingostomy produce similar success rates and long-term fertility.
- **Single-dose methotrexate** is associated with a higher risk of rupture than multiple doses.
Cigarette smoking increases the likelihood of ectopic pregnancy 2.5 times,12 probably by affecting ciliary action within the fallopian tubes.

Salpingitis isthmic nodosa is anatomic thickening of the proximal portion of the fallopian tubes with multiple lumen diverticula. It increases the risk of ectopic pregnancy 1.5 times, compared with age- and race-matched controls.13

Don’t depend solely on risk factors. Many ectopic pregnancies present without them. Symptoms. Many ectopic pregnancies never produce symptoms; rather, they resolve spontaneously or are timely diagnosed and treated medically. Risk factors should therefore be examined in any woman in early pregnancy and investigated further if ectopic pregnancy is likely.

When symptoms do occur, they usually involve 1 or all of the classic triad: amenorrhea, irregular bleeding, and lower abdominal pain. In addition, syncope, shock, and pain radiating to the
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**STEP 1**

▲

Document the pregnancy and measure ß-hCG

Once you identify the high-risk patient, or a woman comes in complaining of pain and spotting or bleeding, run a pregnancy test to confirm that she is pregnant and, if it is positive, obtain a quantitative ß-hCG.

ß-hCG levels are normally measured using enzyme-linked immunosorbent assays (ELISA), which detect ß-hCG in urine and serum at levels as low as 20 mIU/mL and 10 mIU/mL, respectively. ß-hCG is produced by trophoblastic cells in normal pregnancy, and approximately doubles every 2 days when titers are below 10,000 mIU/mL—although in some normal pregnancies, ß-hCG may increase as slowly as 53% or as rapidly as 230% over 2 days. Eighty-five percent of abnormal pregnancies—whether intrauterine or ectopic—have impaired ß-hCG production with prolonged doubling time. Thus, in failing pregnancies, ß-hCG levels will plateau or fail to rise normally.

A single ß-hCG level fails to predict the risk of rupture, since ectopic pregnancies can rupture at ß-hCG levels as low as 10 mIU/mL or far exceeding 10,000 mIU/mL, or at any level in between.

**STEP 2**

▲ Document the pregnancy and measure ß-hCG

Once you identify the high-risk patient, or a woman comes in complaining of pain and spotting or bleeding, run a pregnancy test to confirm that she is pregnant and, if it is positive, obtain a quantitative ß-hCG.

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**STEP 3**

▲ Obtain an ultrasound scan

Transvaginal ultrasound reliably detects normal intrauterine gestations when ß-hCG passes somewhere between 1,000 mIU/mL and 2,000 mIU/mL (First International Reference Preparation), depending on the expertise of the ultrasoundographer and the particular equipment used. This is known as the “discriminatory zone.” ß-hCG levels reach this zone as early as 1 week after missed menses.

The discriminatory zone is not the lowest ß-hCG concentration at which an intrauterine pregnancy can be visualized via ultrasound. Rather, it is the value at which any intrauterine pregnancy will be apparent. At that value, the absence of an intrauterine pregnancy confirms—by negative conclusion—that the patient has a nonviable gestation.

**TABLE 1**

<table>
<thead>
<tr>
<th>RISK FACTOR</th>
<th>ODDS RATIO*</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>High risk</strong></td>
<td></td>
</tr>
<tr>
<td>Tubal surgery</td>
<td>21.0</td>
</tr>
<tr>
<td>Tubal ligation</td>
<td>9.3</td>
</tr>
<tr>
<td>Previous ectopic pregnancy</td>
<td>8.3</td>
</tr>
<tr>
<td>In utero exposure to diethylstilbestrol</td>
<td>5.6</td>
</tr>
<tr>
<td>Use of intrauterine device</td>
<td>4.2–45.0</td>
</tr>
<tr>
<td>Documented tubal pathology</td>
<td>3.8–21.0</td>
</tr>
<tr>
<td>Assisted reproduction</td>
<td>4.0</td>
</tr>
<tr>
<td>Emergency contraception</td>
<td>High</td>
</tr>
<tr>
<td><strong>Moderate risk</strong></td>
<td></td>
</tr>
<tr>
<td>Infertility</td>
<td>2.5–21.0</td>
</tr>
<tr>
<td>Previous genital infections</td>
<td>2.5–3.7</td>
</tr>
<tr>
<td>Multiple sexual partners</td>
<td>2.1</td>
</tr>
<tr>
<td>Salpingitis (isthmic)</td>
<td>1.5</td>
</tr>
<tr>
<td><strong>Slight risk</strong></td>
<td></td>
</tr>
<tr>
<td>Previous pelvic, abdominal surgery</td>
<td>0.9–3.8</td>
</tr>
<tr>
<td>Cigarette smoking</td>
<td>2.3–2.5</td>
</tr>
<tr>
<td>Vaginal douching</td>
<td>1.1–3.1</td>
</tr>
<tr>
<td>Early age at first intercourse (&lt;18 years)</td>
<td>1.6</td>
</tr>
</tbody>
</table>


* Single values = common odds ratio from homogeneous studies; point estimates = range of values from heterogeneous studies.

patient’s shoulder can result from hemo-peritoneum.
pregnancy improves sensitivity from 21% to 84% at the expense of lower specificity (93% to 99.5%).

Even when an adnexal mass is visualized, cardiac activity is not usually present. If cardiac activity is apparent, proceed to surgery, since methotrexate usually will not resolve these gestations.

Be aware that some adnexal masses suspicious for ectopic pregnancy may turn out to be other entities, such as a corpus luteum, hydrosalpinx, ovarian neoplasm, or endometrioma. Unless a fetal heart rate is detected by ultrasound, the diagnosis is uncertain and curettage is needed to establish a definitive diagnosis.

No intrauterine pregnancy, no extrauterine mass. Despite the high resolution of transvaginal ultrasound, many patients with ectopic pregnancy have no apparent adnexal mass, particularly when diagnosis is early. In these cases, proceed to curettage (step 4).

Don’t interpret ultrasound findings in a vacuum
This is especially unwise when ß-hCG levels are low—even when the ultrasound report points to intrauterine pregnancy. At ß-hCG levels below 1,500 mIU/mL, the sensitivity of ultrasound in diagnosing intrauterine pregnancy drops from 98% to 33% and predictive value is substantially lower. Interpret ultrasound and ß-hCG levels together for greater accuracy.

How size influences management
Ultrasound can detect ectopic pregnancies as small as 2 cm. In general, an ectopic sac size larger than 4 cm should be treated surgically.

STEP 4
Perform uterine curettage
If ultrasound imaging is inconclusive and ß-hCG levels are plateauing or rising subnormally, perform uterine curettage. If ß-hCG levels decrease 15% or more 8 to 12 hours after the procedure, a complete abortion can be strongly suspected. If ß-hCG levels plateau or rise, the trophoblasts were not removed by curettage, and ectopic pregnancy is diagnosed.

Keep in mind these important points:
• Without uterine curettage, roughly 40% of ectopic pregnancy diagnoses are incorrect.
• Because curettage will result in termination of pregnancy, it is vital that it be limited to cases involving abnormal ß-hCG levels, not normally rising values.

STEP 5
Administer methotrexate
Medical management is indicated when the following circumstances are present:
• No viable intrauterine pregnancy is present.

Multiple-dose methotrexate protocol
Discontinue treatment when there is a decline in 2 consecutive ß-hCG titers or after 4 doses, whichever comes first.

<table>
<thead>
<tr>
<th>DAY</th>
<th>INTERVENTION</th>
<th>DOSE (MG)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Baseline studies ß-hCG titer, CBC, and platelets Methotrexate</td>
<td>1.0</td>
</tr>
<tr>
<td>2</td>
<td>Leucovorin ß-hCG titer</td>
<td>0.1</td>
</tr>
<tr>
<td>3</td>
<td>Methotrexate ß-hCG titer</td>
<td>1.0</td>
</tr>
<tr>
<td>4</td>
<td>Leucovorin ß-hCG titer</td>
<td>0.1</td>
</tr>
<tr>
<td>5</td>
<td>Methotrexate ß-hCG titer</td>
<td>1.0</td>
</tr>
<tr>
<td>6</td>
<td>Leucovorin ß-hCG titer</td>
<td>0.1</td>
</tr>
<tr>
<td>7</td>
<td>Methotrexate ß-hCG titer</td>
<td>1.0</td>
</tr>
<tr>
<td>8</td>
<td>Leucovorin ß-hCG titer CBC and platelets Renal and liver function tests</td>
<td>0.1</td>
</tr>
<tr>
<td></td>
<td>Weekly ß-hCG titer until negative</td>
<td></td>
</tr>
</tbody>
</table>

CONTINUED
The greatest determinant of outcomes?
Location, location, location

Fallopian tubes
98.3% of ectopic pregnancies; site most susceptible to rupture

Isthmus
12.3% of ectopics; high risk of rupture

Ampulla
79.6% of ectopics; likely to rupture

Fimbria
6.2% of ectopics; likely to abort

Abdominal, ovarian, and cervical ectopic pregnancies have been reported but are rare

• No rupture has occurred.
• Any adnexal mass is 4 cm in size or smaller.
• β-hCG levels are below 10,000 mIU/mL.

If there is a positive fetal heart beat, surgery is preferred.

Firm diagnosis of ectopic pregnancy is essential prior to methotrexate administration. If the drug is given to a woman carrying a viable pregnancy, it may result in loss of the pregnancy or methotrexate embryoopathy.21

How methotrexate resolves ectopic pregnancy
Systemic methotrexate has been used successfully in the treatment of ectopic pregnancy for more than 20 years. A folic acid antagonist, it inhibits de novo synthesis of purines and pyrimidines. It thus interferes with DNA synthesis and cell multiplication.24 Actively proliferating trophoblasts are particularly vulnerable.25

When methotrexate is administered to normally pregnant women, it blunts the normal β-hCG increment over the next 7 days. Circulating progesterone and 17-α-hydroxyprogesterone concentrations also decline, and abortion occurs.26

Methotrexate directly impairs trophoblastic production of hCG; the decrement in corpus luteum progesterone is a secondary event.

Side effects include abdominal distress, chills and fever, dizziness, immunosuppression, leukopenia, malaise, nausea, ulcerative stomatitis, photosensitivity, and undue fatigue.

Breastfeeding is an absolute contraindication to methotrexate, while relative con-
If treatment is unsuccessful after 4 doses of methotrexate, proceed to surgery.

### TABLE 2

<table>
<thead>
<tr>
<th>Method</th>
<th>Number of Studies</th>
<th>Number of Patients</th>
<th>Number with Successful Resolution</th>
<th>Tubal Patency Rate</th>
<th>Subsequent Fertility Rate</th>
<th>Ectopic Pregnancy</th>
</tr>
</thead>
<tbody>
<tr>
<td>Conservative laparoscopic surgery</td>
<td>32</td>
<td>1,626</td>
<td>1,516 (93%)</td>
<td>170/223 (76%)</td>
<td>366/647 (57%)</td>
<td>87/647 (13%)</td>
</tr>
<tr>
<td>Variable-dose methotrexate</td>
<td>12</td>
<td>338</td>
<td>314 (93%)</td>
<td>136/182 (75%)</td>
<td>55/95 (58%)</td>
<td>7/95 (7%)</td>
</tr>
<tr>
<td>Single-dose methotrexate</td>
<td>7</td>
<td>393</td>
<td>340 (87%)</td>
<td>61/75 (81%)</td>
<td>39/64 (61%)</td>
<td>5/64 (8%)</td>
</tr>
<tr>
<td>Direct-injection methotrexate</td>
<td>21</td>
<td>660</td>
<td>502 (76%)</td>
<td>130/162 (80%)</td>
<td>87/152 (57%)</td>
<td>9/152 (6%)</td>
</tr>
<tr>
<td>Expectant management</td>
<td>14</td>
<td>628</td>
<td>425 (68%)</td>
<td>60/79 (76%)</td>
<td>12/14 (86%)</td>
<td>1/14 (7%)</td>
</tr>
</tbody>
</table>


Trainindications include abnormal liver function tests, blood dyscrasias, excessive alcohol consumption, HIV/AIDS, psoriasis, ongoing radiotherapy, rheumatoid arthritis, and significant pulmonary disease.

**Multiple-dose methotrexate is superior**

In a recent meta-analysis comparing single- and multiple-dose regimens of methotrexate, Barnhart and colleagues found the latter to be more effective, although the single dose was more commonly given. The single-dose regimen was associated with a significantly greater chance of failure in both crude (OR 1.71; 1.04, 2.82) and adjusted (OR 4.74; 1.77, 12.62) analyses. Consequently, we advocate the multiple-dose regimen detailed in [TABLE 2](#): 1 mg of methotrexate per kilogram of body weight on day 1, alternating with 0.1 mg of leucovorin per kilogram on succeeding days. Continue this regimen until β-hCG levels decline in 2 consecutive daily titers, or 4 doses of methotrexate are given, whichever comes first.

If treatment is unsuccessful after 4 doses, additional methotrexate is unlikely to be effective and will involve significant additional cost and morbidity. If β-hCG levels plateau or continue to rise, surgery is indicated.

Although multiple-dose therapy is more effective than a single dose, the optimal number of doses probably falls somewhere between 1 and 4. A 2-dose protocol currently under investigation may provide an optimal compromise.

**Artificially high efficacy rates with "single-dose" therapy.** Unusually high success rates in initial studies of the single-dose regimen may have been due to the inclusion of spontaneously aborting intrauterine pregnancies. Although 6 subsequent studies—1 cohort and 5 case-control studies involving 304 patients—found an overall success rate (no surgical intervention) of 87.2%, 11.5% of participants required more than 1 dose.

**Still the standard.** Despite its lower efficacy rates, single-dose methotrexate remains the “standard” in the United States, as recommended in an American College of Obstetricians and Gynecologists practice bulletin. The usual intramuscular dose is 50 mg per square meter of body surface area, with β-hCG titers on days 4 and 7 and an additional dose if β-hCG levels fail to resolve.

What the evidence shows

**Multidose methotrexate.** Twelve studies measured the success of multiple-dose systemic methotrexate ([TABLE 3](#)); they included 1 randomized, controlled trial, 1 cohort study, and 10 case series. Between 1982 and 1997, 325 cases were treated with multiple-
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Laparoscopy is the preferred surgical approach. Of these, 93.8% had successful resolution with no subsequent therapy, and 78.9% of the 161 women tested had patent oviducts. In addition, of 95 women hoping to conceive, 57.9% had a subsequent intrauterine pregnancy and 7.4% developed a recurrent ectopic pregnancy. These rates compared favorably with numerous laparoscopic surgery series published during the same years.9

**Methotrexate versus laparoscopic salpingostomy.** In 1 randomized clinical comparison,29 100 women with laparoscopy-confirmed ectopic pregnancy were randomized to methotrexate or laparoscopic salpingostomy. Of the 51 patients treated medically, 7 (14%) required surgical intervention for active bleeding and/or tubal rupture. An additional course of methotrexate was required in 2 patients (4%) for persistent trophoblasts.

Of the 49 patients in the salpingostomy group, 4 women (8%) failed therapy and required salpingectomies, and 10 patients (20%) were treated with methotrexate for persistent trophoblasts.

Tubal patency was present in 23 of 42 women (55%) in the methotrexate group, compared with 23 of 39 (59%) in the salpingostomy group.

Overall, this randomized study29 and previous meta-analysis demonstrate that systemic multiple-dose methotrexate is comparable in efficacy to laparoscopic salpingostomy.

**Fine points of treatment**

During methotrexate therapy, examine the patient only once to avoid triggering a rupture, and counsel her to avoid intercourse for the same reason.30 Do not perform repeat vaginal ultrasound examination. Also inform her that transient pain (“separation pain”) from tubal abortion frequently occurs 3 to 7 days after the start of therapy, lasts 4 to 12 hours, and then resolves.31 This is perhaps the most difficult aspect of methotrexate therapy, as it is not always easy to differentiate the pain of tubal abortion from the pain of rupture.

If ß-hCG titers continue to rise rapidly between methotrexate doses, rupture is more likely and surgery should proceed.32 Overall, isthmic ectopic pregnancies (12.3% of ectopics) appear to be at a particularly high risk for rupture and comprise nearly half of methotrexate failures.32 Unfortunately, there is no way to identify isthmic pregnancies without surgery.

Avoid NSAIDs and GI-“unfriendly” foods. Counsel the patient to avoid nonsteroidal anti-inflammatory drugs (NSAIDs) because they may impair natural hemostasis. Gas-forming foods such as leeks, corn, and cabbage can cause distension, which may be mistaken for rupture. Also instruct the patient to avoid folic acid, which impairs the efficacy of methotrexate.

**Ultrasound surveillance is unnecessary.** If an adnexal mass was identified at initial imaging, there is no need to view it again, since these masses tend to enlarge and form hematomas and can cause undue anxiety in both physician and patient. In properly selected patients, multidose methotrexate with monitoring of ß-hCG levels should suffice.

**When surgery is indicated**

Surgical intervention is necessary when pain is severe, persists beyond 12 hours, and is associated with orthostatic hypotension, falling hematocrit, or persistently elevated ß-hCG levels after methotrexate therapy.

Laparoscopy is the preferred approach. Advantages include less blood loss and analgesia,33 shortened postoperative recovery, and lower costs.

**Technique.** For unruptured ampullary ectopic pregnancy, salpingostomy is preferred. Make a linear incision over the bulging antimesenteric border of the fallopian tube using electrocautery, scissors, or laser. Remove the products of conception using forceps or suction, and leave the incision to heal by secondary intention.
For isthmic pregnancies, use segmental excision followed by delayed microsurgical anastomosis. The isthmic tubal lumen is narrower and the muscularis thicker than in the ampulla. Thus, the isthmus is predisposed to greater damage after salpingostomy and greater rates of proximal obstruction.

An increased risk of persistent ectopic pregnancy has been a criticism of salpingostomy. However, when 1 dose of systemic methotrexate is combined with salpingostomy, the risk of persistent pregnancy is virtually eliminated.

When rupture occurs, salpingectomy is the first choice for treatment, as it arrests hemorrhage and shortens the procedure. Either laparoscopy or laparotomy is appropriate. The authors report no financial relationships relevant to this article.