Women with epilepsy: 5 clinical pearls for contraception and preconception counseling

For women with epilepsy, intrauterine devices are the optimal reversible contraceptive, and, preconception, the use of antiepileptic drugs with the lowest teratogenic potential should be considered

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In 2015, 1.2% of the US population was estimated to have active epilepsy.¹ For neurologists, key goals in the treatment of epilepsy include: controlling seizures, minimizing adverse effects of antiepileptic drugs (AEDs) and optimizing quality of life. For obstetrician-gynecologists, women with epilepsy (WWE) have unique contraceptive, preconception, and obstetric needs that require highly specialized approaches to care. Here, I highlight 5 care points that are important to keep in mind when counseling WWE.

1. Enzyme-inducing AEDs reduce the effectiveness of estrogen-progestin and some progestin contraceptives. AEDs can induce hepatic enzymes that accelerate steroid hormone metabolism, producing clinically important reductions in bioavailable steroid hormone concentration (TABLE 1, page 10). According to Lexicomp, AEDs that are inducers of hepatic enzymes that metabolize steroid hormones include: carbamazepine (Tegretol), eslicarbazepine (Aptiom), felbamate (Felbatol), oxcarbazepine (Trileptal), perampanel (Fycompa), phenobarbital, phenytoin (Dilantin), primidone (Mysoline), rufinamide (Banzel), and topiramate (Topamax) (at dosages >200 mg daily). According to Lexicomp, the following AEDs do not cause clinically significant changes in hepatic enzymes that metabolize steroid hormones: acetazolamide (Diamox), clonazepam (Klonopin), ethosuximide (Zarontin), gabapentin (Neurontin), lacosamide (Vimpat), levetiracetam (Keppra), pregabalin (Lyrica), tiagabine (Gabitril), vigabatrin (Vigadron), and zonisamide (Zonegran).²³ In addition, lamotrigine (Lamictal) and valproate (Depakote) do not significantly influence the metabolism of contraceptive steroids,⁴⁵ but contraceptive steroids significantly influence their metabolism (TABLE 2, page 16).

For WWE taking an AED that accelerates steroid hormone metabolism, estrogen-progestin contraceptive failure is common. In one study 20 healthy women were administered an ethinyl estradiol (20 µg)-levonorgestrel (100 µg) contraceptive, and randomly assigned to either receive carbamazepine 600 mg daily or a placebo pill.⁷ In this study, based on serum progesterone measurements, 5 of 10 women in the carbamazepine group ovulated, compared with 1 of 10 women in the placebo group. Women taking carbamazepine had integrated serum ethinyl estradiol and levonorgestrel concentrations approximately 45% lower than women taking placebo.⁷ Other studies also report that carbamazepine accelerates steroid hormone metabolism and reduces the circulating concentration of ethinyl estradiol, norethindrone, and levonorgestrel by about 50%.⁴⁵
WWE taking an AED that induces hepatic enzymes should be counseled to use a copper or levonorgestrel (LNG) intrauterine device (IUD) or depot medroxyprogesterone acetate (DMPA) for contraception. WWE taking AEDs that do not induce hepatic enzymes can be offered the full array of contraceptive options, as outlined in Table 1. Occasionally, a WWE taking an AED that is an inducer of hepatic enzymes may strongly prefer to use an estrogen-progestin contraceptive and decline the preferred option of using an IUD or DMPA. If an estrogen-progestin contraceptive is to be prescribed, safeguards to reduce the risk of pregnancy include:

- prescribe a contraceptive with ≥35 µg of ethinyl estradiol
- prescribe a contraceptive with the highest dose of progestin with a long half-life (drospirenone, desogestrel, levonorgestrel)
- consider continuous hormonal contraception rather than 4 or 7 days off hormones and
- recommend use of a barrier contraceptive in addition to the hormonal contraceptive.

The effectiveness of levonorgestrel emergency contraception may also be reduced in WWE taking an enzyme-inducing AED. In these cases, some experts recommend a regimen of two doses of levonorgestrel 1.5 mg, separated by 12 hours. The effectiveness of progestin subdermal contraceptives may be reduced in women taking phenytoin. In one study of 9 WWE using a progestin subdermal implant, phenytoin reduced the circulating levonorgestrel level by approximately 40%.

### TABLE 1 Lexicomp risk, severity, and reliability rating of potential interactions between AED and accelerated metabolism of estrogen and progestin contraceptive hormones

<table>
<thead>
<tr>
<th>Antiepileptic medication</th>
<th>Lexicomp risk (letter grade), severity, and reliability rating for AED interaction with estrogen-progestin contraceptive</th>
<th>Antiepileptic medication</th>
<th>Lexicomp rating- interaction with estrogen-progestin contraceptive</th>
</tr>
</thead>
<tbody>
<tr>
<td>Carbamazepine (Tegretol)</td>
<td>D rating-consider therapy modification, Severity Major, Reliability Good</td>
<td>Acetazolamide (Diamox)</td>
<td>A rating. No known interactions with estrogen or progestin hormones</td>
</tr>
<tr>
<td>Eslicarbazepine (Aptiom)</td>
<td>D rating-consider therapy modification, Severity Major, Reliability Good</td>
<td>Clonazepam (Klonopin)</td>
<td>A rating. No known interactions with estrogen or progestin hormones</td>
</tr>
<tr>
<td>Felbamate (Felbatol)</td>
<td>D rating-consider therapy modification, Severity Major, Reliability Good</td>
<td>Ethosuximide (Zarontin)</td>
<td>A rating. No known interactions with estrogen or progestin hormones</td>
</tr>
<tr>
<td>Oxcarbazepine (Trileptal)</td>
<td>D rating-consider therapy modification, Severity Major, Reliability Good</td>
<td>Gabapentin (Neurontin)</td>
<td>A rating. No known interactions with estrogen or progestin hormones</td>
</tr>
<tr>
<td>Perampanel (Fycompa)</td>
<td>D rating-consider therapy modification, Severity Major, Reliability Fair</td>
<td>Lacosamide (Vimpat)</td>
<td>B rating. May increase the serum concentration of ethinyl estradiol by 20%. Severity Minor, Reliability Fair</td>
</tr>
<tr>
<td>Phenobarbital</td>
<td>D rating-consider therapy modification, Severity Major, Reliability Fair</td>
<td>Levetiracetam (Keppra)</td>
<td>A rating. No known interactions with estrogen or progestin hormones</td>
</tr>
<tr>
<td>Phenytoin (Dilantin)</td>
<td>D rating-consider therapy modification, Severity Major, Reliability Fair. Also induces CYP3A4</td>
<td>Pregabalin (Lyrica)</td>
<td>A rating. No known interactions with estrogen or progestin hormones</td>
</tr>
<tr>
<td>Primidone (Mysoline)</td>
<td>D rating-consider therapy modification, Severity Major, Reliability Fair</td>
<td>Tiagabine (Gabitril)</td>
<td>A rating. No known interactions with estrogen or progestin hormones</td>
</tr>
<tr>
<td>Rufinamide (Banzel)</td>
<td>D rating-consider therapy modification, Severity Major, Reliability Fair</td>
<td>Vigabatin (Vigadone)</td>
<td>A rating. No known interactions with estrogen or progestin hormones</td>
</tr>
<tr>
<td>Topiramate (Topamax)</td>
<td>D rating-consider therapy modification, Severity Major, Reliability Good</td>
<td>Zonisamide (Zonegran)</td>
<td>A rating. No known interactions with estrogen or progestin hormones</td>
</tr>
</tbody>
</table>

*aLamotrigine (Lamictal) and valproate (Depakote) are not strong inducers of hepatic enzymes. Hence, they do not accelerate the metabolism of estrogen and progestin contraceptive hormones. However, the metabolism of lamotrigine and valproate is accelerated by estrogen (see Table 2). Abbreviation: AED, antiepileptic drug.*
Do not use lamotrigine with cyclic estrogen-progestin contraceptives.

Estrogens, but not progestins, are known to reduce the serum concentration of lamotrigine by about 50%. This is a clinically significant pharmacologic interaction. Consequently, when a cyclic estrogen-progestin contraceptive is prescribed to a woman taking lamotrigine, oscillation in lamotrigine serum concentration can occur. When the woman is taking estrogen-containing pills, lamotrigine levels decrease, which increases the risk of seizure. When the woman is not taking the estrogen-containing pills, lamotrigine levels increase, possibly causing such adverse effects as nausea and vomiting. If a woman taking lamotrigine insists on using an estrogen-progestin contraceptive, the medication should be prescribed in a continuous regimen and the neurologist alerted so that they can increase the dose of lamotrigine and intensify their monitoring of lamotrigine levels. Lamotrigine does not change the metabolism of ethinyl estradiol and has minimal impact on the metabolism of levonorgestrel.

Preconception counseling: Before conception consider using an AED with low teratogenicity.

Valproate is a potent teratogen, and consideration should be given to discontinuing valproate prior to conception. In a study of 1,788 pregnancies exposed to valproate, the risk of a major congenital malformation was 10% for valproate monotherapy, 11.3% for valproate combined with lamotrigine, and 11.7% for valproate combined with another AED, but not lamotrigine. At a valproate dose of ≥1,500 mg daily, the risk of major malformation was 24% for valproate monotherapy, 31% for valproate plus lamotrigine, and 19% for valproate plus another AED, but not lamotrigine. Valproate is reported to be associated with the following major congenital malformations: spina bifida, ventricular and atrial septal defects, pulmonary valve atresia, hypoplastic left heart syndrome, cleft palate, anorectal atresia, and hypospadias.

In a study of 7,555 pregnancies in women using a single AED, the risk of major congenital anomalies varied greatly among the AEDs, including valproate (10.3%), phenobarbital (6.5%), phenytoin (6.4%), carbamazepine (5.5%), topiramate (3.9%), oxcarbazepine (3.0%), lamotrigine (2.9%), and levetiracetam (2.8%). For WWE considering pregnancy, many experts recommend use of lamotrigine, levetiracetam, or oxcarbazepine to minimize the risk of fetal anomalies.

Folic acid: Although the optimal dose for WWE taking an AED and planning to become pregnant is unknown, a high dose is reasonable.

The American College of Obstetricians and Gynecologists (ACOG) recommends that women planning pregnancy take 0.4 mg of folic acid daily, starting at least 1 month before pregnancy and continuing through at least the 12th week of gestation. ACOG also recommends that women at high risk of a neural tube defect should take 4 mg of folic acid daily. WWE taking a teratogenic AED are known to be at increased risk for fetal malformations, including neural tube defects. Should these women take 4 mg of folic acid daily?
Epilepsy and operation of a motor vehicle

For most women with epilepsy, maintaining a valid driver’s license is important for completion of daily life tasks. Most states require that a patient with seizures be seizure-free for 6 to 12 months to operate a motor vehicle. Estrogen-containing hormonal contraceptives can reduce the concentration of some AEDs, such as lamotrigine. Hence, it is important that the patient be aware of this interaction and that the primary neurologist be alerted if an estrogen-containing contraceptive is prescribed to a woman taking lamotrigine or valproate. Specific state laws related to epilepsy and driving are available at the Epilepsy Foundation website (https://www.epilepsy.com/driving-laws).

ACOG notes that, for women taking valproate, the benefit of high-dose folic acid (4 mg daily) has not been definitively proven, and guidelines from the American Academy of Neurology do not recommend high-dose folic acid for women receiving AEDs. Hence, ACOG does not recommend that WWE taking an AED take high-dose folic acid.

By contrast, the Royal College of Obstetricians and Gynecologists (RCOG) recommends that all WWE planning a pregnancy take folic acid 5 mg daily, initiated 3 months before conception and continued through the first trimester of pregnancy.

The RCOG notes that among WWE taking an AED, intelligence quotient is greater in children whose mothers took folic acid during pregnancy. Given the potential benefit of folic acid on long-term outcomes and the known safety of folic acid, it is reasonable to recommend high-dose folic acid for WWE.

Final takeaways
Surveys consistently report that WWE have a low-level of awareness about the interaction between AEDs and hormonal contraceptives and the teratogenicity of AEDs. For example, in a survey of 2,000 WWE, 45% who were taking an enzyme-inducing AED and an estrogen-progestin oral contraceptive reported that they had not been warned about the potential interaction between the medications. Surprisingly, surveys of neurologists and obstetrician-gynecologists also report that there is a low level of awareness about the interaction between AEDs and hormonal contraceptives. When providing contraceptive counseling for WWE, prioritize the use of a copper or levonorgestrel IUD. When providing preconception counseling for WWE, educate the patient about the high teratogenicity of valproate and the lower risk of malformations associated with the use of lamotrigine, levetiracetam, and oxcarbazepine.

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References


