

Anticonvulsant Medications: Use in Adults



The Centers for Medicare & Medicaid Services (CMS) Medicaid Integrity Group (MIG) has identified issues with the utilization of anticonvulsant medications, also known as antiepileptic drugs (AEDs). The U.S. Food and Drug Administration (FDA) approves product labeling for prescription drugs. The MIG has identified that some providers may have prescribed anticonvulsant medications outside of FDA-approved product labeling for indication, age, dosage, or duration of therapy. Therefore, CMS' goal is to improve quality of care and enhance patient safety by educating providers on the proper use of anticonvulsant medications in adults.

This fact sheet summarizes the current FDA-approved product labeling for the use of anticonvulsant medications in adults. After reading this fact sheet, providers should be able to accurately:

- Identify the FDA-approved indications for the use of anticonvulsant medications in adults;
- Identify the available treatment guidelines for the management of seizures in adults; and
- Summarize the adverse reactions and risks of anticonvulsant medications.

FDA-Approved Indications for Anticonvulsant Medications in Adults

Anticonvulsant medications are used for the prevention and treatment of seizures. Epilepsy, one common cause of seizures, is a neurological disorder that is active in approximately 2.4 million adults in the United States.[1] Patients diagnosed with epilepsy experience recurrent seizures; however, not all seizures are a result of epilepsy. A seizure can also be caused by head trauma,[2] low blood sugar, alcohol or drug withdrawal, or high fever and may last from a few seconds to several minutes.[3]



Anticonvulsant medications are FDA approved to treat seizures and many other medical conditions unrelated to seizure disorders. However, not all anticonvulsant medications are FDA approved for every indication. The FDA-approved indications for the use of anticonvulsant medications in adult patients are:

- Bipolar I disorder;
- Epilepsy;
- Mania;
- Postherpetic neuralgia;
- Prevention and treatment of seizures occurring during and following neurosurgery;
- Prophylaxis of migraine headaches;
- Restless leg syndrome (RLS);
- Seizures (absence, myoclonic, partial-onset, and tonic-clonic [grand mal]);
- Seizures associated with Lennox-Gastaut syndrome (LGS); and
- Trigeminal neuralgia.

ACRONYMS

AED	antiepileptic drugs
AHRQ	Agency for Healthcare Research and Quality
CDER	Center for Drug Evaluation and Research
CMS	Centers for Medicare & Medicaid Services
FDA	U.S. Food and Drug Administration
LGS	Lennox-Gastaut syndrome
MIG	Medicaid Integrity Group
MPIE	Medicaid Program Integrity Education
OTC	over the counter
RLS	restless leg syndrome
SHARE	Support, Help, and Resources for Epilepsy
SJS	Stevens-Johnson syndrome
TEN	toxic epidermal necrolysis
WBC	white blood cell

Dosing schedules for anticonvulsant medications are guided by the specific indication for use. Some anticonvulsant medications require dosage adjustments for renal function, hepatic function, and other patient factors. When anticonvulsant medications are used in combination, or when a patient is transitioned from one anticonvulsant medication to another, a dosage adjustment may be required. Specific recommendations for dosage adjustments can be found in the prescribing information for each anticonvulsant medication.

LGS is a rare condition that primarily affects young children. Because seizures associated with LGS are difficult to control with medication and treatment regimens are highly individualized, the FDA-approved anticonvulsant medications for the treatment of LGS are not discussed in this fact sheet.[4] The benzodiazepines and barbiturates are also not discussed in this fact sheet because they are infrequently prescribed for the long-term treatment of seizures.

The indications and dosages for the anticonvulsant medications discussed in this fact sheet are provided in the “Anticonvulsant Medications: U.S. Food and Drug Administration-Approved Indications and Most Common Dosages for Use in Adults” dosing chart, available at <https://www.cms.gov/Medicare-Medicaid-Coordination/Fraud-Prevention/Medicaid-Integrity-Education/Pharmacy-Education-Materials/anticonvulsant-education.html> on the CMS website.

Monitoring Parameters for Select Anticonvulsant Medications

Some of the anticonvulsant medications require monitoring of drug levels to ensure their safe use. The therapeutic drug levels of the anticonvulsant medications that require drug level monitoring are provided in Table 1.

Table 1. Therapeutic Drug Levels for Anticonvulsant Medications

Medication	Therapeutic Level
carbamazepine[5]	4 mcg per ml to 12 mcg per ml
ethosuximide[6]	40 mcg per ml to 100 mcg per ml
phenytoin[7]	10 mcg per ml to 20 mcg per ml
valproic acid[8]	50 mcg per ml to 125 mcg per ml (mania)
valproic acid	50 mcg per ml to 100 mcg per ml (epilepsy)

Treatment Guidelines for the Use of Anticonvulsant Medications in Adults

The Agency for Healthcare Research and Quality (AHRQ) hosts a database of treatment guidelines. The AHRQ is a branch of the U.S. Department of Health and Human Services. Please visit <https://www.guideline.gov> for the AHRQ's National Guideline Clearinghouse. Links to some of the treatment guidelines for the management of seizures and the use of anticonvulsant medications in adults are provided in Table 2.



Table 2. Treatment Guidelines for Anticonvulsant Medications

Sponsoring Organization	Title of Guideline	Link to Guideline
American Association of Neurological Surgeons, Congress of Neurological Surgeons	The role of prophylactic anticonvulsants in the management of brain metastases: a systematic review and evidence-based clinical practice guideline. [2011, November]	https://www.guideline.gov/content.aspx?id=25751
American Academy of Neurology and American Epilepsy Society, Quality Standards Subcommittee and Therapeutics and Technology Assessment Subcommittee	Practice parameter update: management issues for women with epilepsy—focus on pregnancy (an evidence-based review): teratogenesis and perinatal outcomes. Report of the Quality Standards Subcommittee and Therapeutics and Technology Subcommittee of the American Academy of Neurology and American Epilepsy Society. [2009, July; reaffirmed 2013]	https://www.guideline.gov/content.aspx?id=14680
American Academy of Neurology and American Epilepsy Society, Quality Standards Subcommittee and Therapeutics and Technology Assessment Subcommittee	Practice parameter update: management issues for women with epilepsy-focus on pregnancy (an evidence-based review): vitamin K, folic acid, blood levels, and breastfeeding. Report of the Quality Standards Subcommittee and Therapeutics and Technology Subcommittee of the American Academy of Neurology and American Epilepsy Society. [2009, July; reaffirmed 2013]	https://www.guideline.gov/content.aspx?id=14679
Brain Trauma Foundation, American Association of Neurological Surgeons, Congress of Neurological Surgeons	Guidelines for the acute medical management of severe traumatic brain injury in infants, children, and adolescents—second edition. [2012, January]	https://www.guideline.gov/content.aspx?id=36899

Adverse Reactions and Risks of Anticonvulsant Medications

The prescribing information for each anticonvulsant medication provides details on the adverse reactions and risks of that medication. The adverse reactions and risks vary with each medication. Visit <https://www.accessdata.fda.gov/scripts/cder/drugsatfda> on the FDA website or <http://dailymed.nlm.nih.gov> on the DailyMed website and search by medication name for the prescribing information for that drug.

Several anticonvulsant medications have boxed warnings that draw attention to serious and potentially life-threatening adverse reactions. Anticonvulsant medications that have boxed warnings are carbamazepine, lamotrigine, perampanel, valproic acid, and vigabatrin.

Carbamazepine

Carbamazepine may cause dermatologic reactions, including toxic epidermal necrolysis (TEN) and Stevens-Johnson syndrome (SJS). These reactions can be fatal. The medication should be discontinued if a patient presents with a drug-induced rash.[9] If anticonvulsant therapy is required, the patient should be treated with a different anticonvulsant medication.

Aplastic anemia and agranulocytosis have also been associated with carbamazepine therapy. Baseline white blood cell (WBC) and platelet counts should be obtained before starting carbamazepine and should be monitored periodically based on clinical judgment.[10] Boxed warnings have been added to alert patients and providers to these risks.

The boxed warnings for carbamazepine state:[11]

WARNINGS

Serious Dermatologic Reactions and HLA-B*1502 Allele

Serious and sometimes fatal dermatologic reactions, including toxic epidermal necrolysis (TEN) and Stevens-Johnson Syndrome (SJS), have been reported during treatment with TEGRETOL. These reactions are estimated to occur in 1 to 6 per 10,000 new users in countries with mainly Caucasian populations, but the risk in some Asian countries is estimated to be about 10 times higher. Studies in patients of Chinese ancestry have found a strong association between the risk of developing SJS/TEN and the presence of HLA-B*1502, an inherited allelic variant of the HLA-B gene. HLA-B*1502 is found almost exclusively in patients with ancestry across broad areas of Asia. Patients with ancestry in genetically at-risk populations should be screened for the presence of HLA-B*1502 prior to initiating treatment with TEGRETOL. Patients testing positive for the allele should not be treated with TEGRETOL unless the benefit clearly outweighs the risk (see Warnings and Precautions, Laboratory Tests).

Aplastic Anemia and Agranulocytosis

Aplastic anemia and agranulocytosis have been reported in association with the use of TEGRETOL. Data from a population-based case control study demonstrate that the risk of developing these reactions is 5 to 8 times greater than in the general population. However, the overall risk of these reactions in the untreated general population is low, approximately six patients per one million population per year for agranulocytosis and two patients per one million population per year for aplastic anemia.

Although reports of transient or persistent decreased platelet or white blood cell counts are not uncommon in association with the use of TEGRETOL, data are not available to estimate accurately their incidence or outcome. However, the vast majority of the cases of leukopenia have not progressed to the more serious conditions of aplastic anemia or agranulocytosis.

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Because of the very low incidence of agranulocytosis and aplastic anemia, the vast majority of minor hematologic changes observed in monitoring of patients on TEGRETOL are unlikely to signal the occurrence of either abnormality. Nonetheless, complete pretreatment hematological testing should be obtained as a baseline. If a patient in the course of treatment exhibits low or decreased white blood cell or platelet counts, the patient should be monitored closely. Discontinuation of the drug should be considered if any evidence of significant bone marrow depression develops.

Felbamate

Felbamate is used primarily for adjunct therapy for seizures resulting from LGS in pediatric patients and adults and is not indicated for any other type of seizure. The prescribing information allows for monotherapy, but such use has not been systematically studied. Accordingly, the risks of felbamate therapy should be weighed against the potential benefits. Aplastic anemia and hepatotoxicity have been associated with the use of felbamate. A boxed warning has been added to the prescribing information to alert patients and providers to these risks.[12]

The boxed warning for felbamate states:[13]

WARNING

1. Aplastic Anemia

The use of FELBATOL® (felbamate) is associated with a marked increase in the incidence of aplastic anemia. Accordingly, FELBATOL® should only be used in patients whose epilepsy is so severe that the risk of aplastic anemia is deemed acceptable in light of the benefits conferred by its use (see Indications). Ordinarily, a patient should not be placed on and/or continued on FELBATOL® without consideration of appropriate expert hematologic consultation.

Among FELBATOL® treated patients, aplastic anemia (pancytopenia in the presence of a bone marrow largely depleted of hematopoietic precursors) occurs at an incidence that may be more than a 100 fold greater than that seen in the untreated population (i.e., 2 to 5 per million persons per year). The risk of death in patients with aplastic anemia generally varies as a function of its severity and etiology; current estimates of the overall case fatality rate are in the range of 20 to 30%, but rates as high as 70% have been reported in the past.

There are too few FELBATOL® associated cases, and too little known about them to provide a reliable estimate of the syndrome's incidence or its case fatality rate or to identify the factors, if any, that might conceivably be used to predict who is at greater or lesser risk.

In managing patients on FELBATOL®, it should be borne in mind that the clinical manifestation of aplastic anemia may not be seen until after a patient has been on FELBATOL® for several months (e.g., onset of aplastic anemia among FELBATOL® exposed patients for whom data are available has ranged from 5 to 30 weeks). However, the injury to bone marrow stem cells that is held to be ultimately responsible for the anemia may occur weeks to months earlier. Accordingly, patients who are discontinued from FELBATOL® remain at risk for developing anemia for a variable, and unknown, period afterwards.

It is not known whether or not the risk of developing aplastic anemia changes with duration of exposure. Consequently, it is not safe to assume that a patient who has been on FELBATOL® without signs of hematologic abnormality for long periods of time is without risk.

It is not known whether or not the dose of FELBATOL® affects the incidence of aplastic anemia.

It is not known whether or not concomitant use of antiepileptic drugs and/or other drugs affects the incidence of aplastic anemia.

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Aplastic anemia typically develops without premonitory clinical or laboratory signs, the full blown syndrome presenting with signs of infection, bleeding, or anemia. Accordingly, routine blood testing cannot be reliably used to reduce the incidence of aplastic anemia, but, it will, in some cases, allow the detection of the hematologic changes before the syndrome declares itself clinically. FELBATOL® should be discontinued if any evidence of bone marrow depression occurs.

2. Hepatic Failure

Evaluation of postmarketing experience suggests that acute liver failure is associated with the use of FELBATOL®. The reported rate in the U.S. has been about 6 cases of liver failure leading to death or transplant per 75,000 patient years of use. This rate is an underestimate because of under reporting, and the true rate could be considerably greater than this. For example, if the reporting rate is 10%, the true rate would be one case per 1,250 patient years of use.

Of the cases reported, about 67% resulted in death or liver transplantation, usually within 5 weeks of the onset of signs and symptoms of liver failure. The earliest onset of severe hepatic dysfunction followed subsequently by liver failure was 3 weeks after initiation of FELBATOL®. Although some reports described dark urine and nonspecific prodromal symptoms (e.g., anorexia, malaise, and gastrointestinal symptoms), in other reports it was not clear if any prodromal symptoms preceded the onset of jaundice.

It is not known whether or not the risk of developing hepatic failure changes with duration of exposure.

It is not known whether or not the dosage of FELBATOL® affects the incidence of hepatic failure.

It is not known whether concomitant use of other antiepileptic drugs and/or other drugs affect the incidence of hepatic failure.

FELBATOL® should not be prescribed for anyone with a history of hepatic dysfunction.

Treatment with FELBATOL® should be initiated only in individuals without active liver disease and with normal baseline serum transaminases. It has not been proved that periodic serum transaminase testing will prevent serious injury but it is generally believed that early detection of drug-induced hepatic injury along with immediate withdrawal of the suspect drug enhances the likelihood for recovery. There is no information available that documents how rapidly patients can progress from normal liver function to liver failure, but other drugs known to be hepatotoxins can cause liver failure rapidly (e.g., from normal enzymes to liver failure in 2–4 weeks). Accordingly, monitoring of serum transaminase levels (AST and ALT) is recommended at baseline and periodically thereafter. While the more frequent the monitoring the greater the chances of early detection, the precise schedule for monitoring is a matter of clinical judgement.

FELBATOL® should be discontinued if either serum AST or serum ALT levels become increased ≥ 2 times the upper limit of normal, or if clinical signs and symptoms suggest liver failure (see precautions). Patients who develop evidence of hepatocellular injury while on FELBATOL® and are withdrawn from the drug for any reason should be presumed to be at increased risk for liver injury if FELBATOL® is reintroduced. Accordingly, such patients should not be considered for re-treatment.

Lamotrigine

Serious skin rashes have been associated with lamotrigine. The risk of occurrence may be related to concomitant valproate therapy, exceeding the recommended initial dose, or rapidly increasing the dose. The medication should be discontinued if a patient presents with a drug-induced rash.[14]

The boxed warning for lamotrigine states:[15]

WARNING: SERIOUS SKIN RASHES

LAMICTAL® can cause serious rashes requiring hospitalization and discontinuation of treatment. The incidence of these rashes, which have included Stevens-Johnson syndrome, is approximately 0.3% to 0.8% in pediatric patients (aged 2 to 17 years) and 0.08% to 0.3% in adults receiving LAMICTAL. One rash-related death was reported in a prospectively followed cohort of 1,983 pediatric patients (aged 2 to 16 years) with epilepsy taking LAMICTAL as adjunctive therapy. In worldwide postmarketing experience, rare cases of toxic epidermal necrolysis and/or rash-related death have been reported in adult and pediatric patients, but their numbers are too few to permit a precise estimate of the rate. Other than age, there are as yet no factors identified that are known to predict the risk of occurrence or the severity of rash caused by LAMICTAL. There are suggestions, yet to be proven, that the risk of rash may also be increased by (1) coadministration of LAMICTAL with valproate (includes valproic acid and divalproex sodium), (2) exceeding the recommended initial dose of LAMICTAL, or (3) exceeding the recommended dose escalation for LAMICTAL. However, cases have occurred in the absence of these factors.

Nearly all cases of life-threatening rashes caused by LAMICTAL have occurred within 2 to 8 weeks of treatment initiation. However, isolated cases have occurred after prolonged treatment (e.g., 6 months). Accordingly, duration of therapy cannot be relied upon as means to predict the potential risk heralded by the first appearance of a rash.

Although benign rashes are also caused by LAMICTAL, it is not possible to predict reliably which rashes will prove to be serious or life threatening. Accordingly, LAMICTAL should ordinarily be discontinued at the first sign of rash, unless the rash is clearly not drug related. Discontinuation of treatment may not prevent a rash from becoming life threatening or permanently disabling or disfiguring [see Warnings and Precautions (5.1)].

Perampanel

Perampanel has been associated with serious psychiatric and behavioral reactions. Patients should be monitored for signs and symptoms of psychiatric and behavioral adverse reactions including aggression, anger, irritability, hostility, and homicidal ideation and threats. Patients and caregivers should be instructed to contact a health care provider if any of these reactions or changes in behavior are observed.[16]

The boxed warning for perampanel states:[17]

WARNING: SERIOUS PSYCHIATRIC AND BEHAVIORAL REACTIONS

- **Serious or life-threatening psychiatric and behavioral adverse reactions including aggression, hostility, irritability, anger, and homicidal ideation and threats have been reported in patients taking FYCOMPA (5.1).**
- **These reactions occurred in patients with and without prior psychiatric history, prior aggressive behavior, or concomitant use of medications associated with hostility and aggression (5.1).**
- **Advise patients and caregivers to contact a healthcare provider immediately if any of these reactions or changes in mood, behavior, or personality that are not typical for the patient are observed while taking FYCOMPA or after discontinuing FYCOMPA (5.1).**
- **Closely monitor patients particularly during the titration period and at higher doses (5.1).**
- **FYCOMPA should be reduced if these symptoms occur and should be discontinued immediately if symptoms are severe or are worsening (5.1).**

Valproic Acid and Divalproex

Valproic acid and its derivatives, including divalproex, have been associated with hepatotoxicity, teratogenicity, and pancreatitis. Patients requiring valproic acid therapy should be made aware of these risks prior to the initiation of therapy. Hepatotoxicity usually occurs in children younger than 2 years old and within the first 6 months of therapy, although it may occur at any time and may occur in patients older than 2 years old. Serum liver tests should be monitored prior to initiation of therapy and periodically throughout the course of treatment.[18]

The risks of treatment with valproic acid and its derivatives should be discussed with the patient and weighed against the potential benefits. Valproic acid is a known teratogen. Female patients in their childbearing years should also be provided with the medication guide (included in the prescribing information) that describes the teratogenic potential of valproic acid.[19] In addition, studies indicate that exposure to valproic acid in utero causes adverse effects on cognitive function in children; they “have lower cognitive test scores than children exposed in utero to either another antiepileptic drug or to no antiepileptic drugs.”[20] Female patients who become pregnant while taking valproic acid should be encouraged to enroll in the North American Antiepileptic Drug Pregnancy Registry. Information on the registry can be found at <http://www.massgeneral.org/aed> on the Massachusetts General Hospital website.[21]

Pancreatitis may occur in children and adults. Patients should be educated about and evaluated for the warning signs of pancreatitis and encouraged to seek treatment if they occur.[22]

The boxed warning for valproic acid states:[23]

WARNING: LIFE THREATENING ADVERSE REACTIONS

Hepatotoxicity

General Population: Hepatic failure resulting in fatalities has occurred in patients receiving valproate and its derivatives. These incidents usually have occurred during the first six months of treatment. Serious or fatal hepatotoxicity may be preceded by non-specific symptoms such as malaise, weakness, lethargy, facial edema, anorexia, and vomiting. In patients with epilepsy, a loss of seizure control may also occur. Patients should be monitored closely for appearance of these symptoms. Serum liver tests should be performed prior to therapy and at frequent intervals thereafter, especially during the first six months. [see Warnings and Precautions (5.1)]

Children under the age of two years are at a considerably increased risk of developing fatal hepatotoxicity, especially those on multiple anticonvulsants, those with congenital metabolic disorders, those with severe seizure disorders accompanied by mental retardation, and those with organic brain disease. When Depakote is used in this patient group, it should be used with extreme caution and as a sole agent. The benefits of therapy should be weighed against the risks. The incidence of fatal hepatotoxicity decreases considerably in progressively older patient groups.

Patients with Mitochondrial Disease: There is an increased risk of valproate-induced acute liver failure and resultant deaths in patients with hereditary neurometabolic syndromes caused by DNA mutations of the mitochondrial DNA Polymerase γ (POLG) gene (e.g., Alpers Huttenlocher Syndrome). Depakote is contraindicated in patients known to have mitochondrial disorders caused by POLG mutations and children under two years of age who are clinically suspected of having a mitochondrial disorder [see Contraindications (4)]. In patients over two years of age who are clinically suspected of having a hereditary mitochondrial disease, Depakote should only be used after other anticonvulsants have failed. This older group of patients should be closely monitored during treatment with Depakote for the development of acute liver injury with regular clinical assessments and serum liver testing. POLG mutation screening should be performed in accordance with current clinical practice [see Warnings and Precautions (5.1)].

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Fetal risk

Valproate can cause major congenital malformations, particularly neural tube defects (e.g., spina bifida). In addition, valproate can cause decreased IQ scores following *in utero* exposure.

Valproate is therefore contraindicated in pregnant women treated for prophylaxis of migraine [see *Contraindications (4)*]. Valproate should only be used to treat pregnant women with epilepsy or bipolar disorder if other medications have failed to control their symptoms or are otherwise unacceptable.

Valproate should not be administered to a woman of childbearing potential unless the drug is essential to the management of her medical condition. This is especially important when valproate use is considered for a condition not usually associated with permanent injury or death (e.g., migraine). Women should use effective contraception while using valproate [see *Warnings and Precautions (5.2, 5.3, 5.4)*].

A Medication Guide describing the risks of valproate is available for patients [see *Patient Counseling Information (17)*].

Pancreatitis

Cases of life-threatening pancreatitis have been reported in both children and adults receiving valproate. Some of the cases have been described as hemorrhagic with a rapid progression from initial symptoms to death. Cases have been reported shortly after initial use as well as after several years of use. Patients and guardians should be warned that abdominal pain, nausea, vomiting, and/or anorexia can be symptoms of pancreatitis that require prompt medical evaluation. If pancreatitis is diagnosed, valproate should ordinarily be discontinued. Alternative treatment for the underlying medical condition should be initiated as clinically indicated [see *Warnings and Precautions (5.5)*].

Vigabatrin

Vision loss has been associated with vigabatrin therapy. Patients should have their vision tested within 4 weeks of starting vigabatrin and every 3 months during the course of therapy. Because of the risk of vision loss, vigabatrin is only available through Support, Help, and Resources for Epilepsy (SHARE), a restricted distribution program. Vigabatrin is not indicated as an initial therapy for complex partial seizures. It should only be considered after several alternative therapies have not worked and should only be prescribed for patients for whom the benefits outweigh the risk of vision loss.[24]

The boxed warning for vigabatrin, which includes information on the SHARE restricted distribution program, states:[25]

WARNING: VISION LOSS

- SABRIL causes permanent bilateral concentric visual field constriction. Because assessing vision may be difficult in infants and children, the frequency and extent of vision loss is poorly characterized in these patients. For this reason, the risk described below is primarily based on the adult experience.
- Based upon adult studies, 30 percent or more of patients can be affected, ranging in severity from mild to severe, including tunnel vision to within 10 degrees of visual fixation, and can result in disability. In some cases, SABRIL also can damage the central retina and may decrease visual acuity.
- The onset of vision loss from SABRIL is unpredictable, and can occur within weeks of starting treatment or sooner, or at any time after starting treatment, even after months or years.
- Symptoms of vision loss from SABRIL are unlikely to be recognized by patients or caregivers before vision loss is severe. Vision loss of milder severity, while often unrecognized by the patient or caregiver, can still adversely affect function.
- The risk of vision loss increases with increasing dose and cumulative exposure, but there is no dose or exposure known to be free of risk of vision loss.

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- Unless a patient is formally exempted from periodic ophthalmologic assessment as documented in the SHARE program, vision should be assessed to the extent possible at baseline (no later than 4 weeks after starting SABRIL) and at least every 3 months during therapy. Vision assessment is also required about 3 to 6 months after the discontinuation of SABRIL therapy. Once detected, vision loss due to SABRIL is not reversible. It is expected that, even with frequent monitoring, some patients will develop severe vision loss.
- Drug discontinuation should be considered, balancing benefit and risk, if visual loss is documented.
- It is possible that vision loss can worsen despite discontinuation of SABRIL.
- Because of the risk of visual loss, SABRIL should be withdrawn from patients with refractory complex partial seizures who fail to show substantial clinical benefit within 3 months of initiation and within 2-4 weeks of initiation for patients with infantile spasms, or sooner if treatment failure becomes obvious. Patient response to and continued need for SABRIL should be periodically reassessed.
- SABRIL should not be used in patients with, or at high risk of, other types of irreversible vision loss unless the benefits of treatment clearly outweigh the risks. The interaction of other types of irreversible vision damage with vision damage from SABRIL has not been well-characterized, but is likely adverse.
- SABRIL should not be used with other drugs associated with serious adverse ophthalmic effects such as retinopathy or glaucoma unless the benefits clearly outweigh the risks.
- The possibility that vision loss from SABRIL may be more common, more severe or have more severe functional consequences in infants and children than in adults cannot be excluded.
- The lowest dose and shortest exposure to SABRIL consistent with clinical objectives should be used.

Because of the risk of permanent vision loss, SABRIL is available only through a restricted program under a Risk Evaluation and Mitigation Strategy (REMS) called the SHARE Program [see *Warnings and Precautions (5.2)*]. Further information is available at [www.sabril.net or 1-888-45-SHARE].

Risk of Suicidality

Anticonvulsant medications have shown an increased risk of suicidality. In pooled analyses that included 11 anticonvulsant medications, results showed that patients taking an anticonvulsant were almost twice as likely to experience suicidal behavior or ideation as those patients taking a placebo.[26] The FDA requires the manufacturers of anticonvulsant medications to include a warning in the prescribing information.

A Medication Guide has also been developed to alert patients and caregivers to the risk of suicidality. The Medication Guide is dispensed with every new and refilled prescription. Visit <http://www.fda.gov/Drugs/DrugSafety/ucm085729.htm> on the FDA website to find links to the required Medication Guides.

Resources

Visit <http://www.medicaid.gov/Medicaid-CHIP-Program-Information/By-State/By-State.html> for links to State Medicaid program websites.

The Center for Drug Evaluation and Research (CDER) hosts a website providing health professionals with current information on over-the-counter (OTC) and prescription drugs. Visit <http://www.fda.gov/Drugs/ResourcesForYou/HealthProfessionals> to access drug-related databases, information on drug recalls and alerts, current information on new and generic drug approvals, and information on drug safety and availability.

Section 1927(g)(1)(B) of the Social Security Act identifies the predetermined standards that the State's drug use review program must use to assess data on drug use. Visit http://www.ssa.gov/OP_Home/ssact/title19/1927.htm for information on the compendia.

To see the electronic version of this fact sheet and the other products included in the “Anticonvulsants” Toolkit, visit the Medicaid Program Integrity Education page at <https://www.cms.gov/Medicare-Medicaid-Coordination/Fraud-Prevention/Medicaid-Integrity-Education/Pharmacy-Education-Materials/pharmacy-ed-materials.html> on the CMS website.

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References

- 1 Centers for Disease Control and Prevention. (2015, March 18). Epilepsy: Epilepsy Fast Facts. Retrieved July 31, 2015, from <http://www.cdc.gov/epilepsy/basics/fast-facts.htm>
- 2 Barker, G.L. (n.d.) Coding of Seizures and Epilepsy. Acute Symptomatic Seizures vs. Late onset Epilepsy. [Slide 11]. Retrieved July 22, 2015, from www.cdc.gov/nchs/ppt/icd9/att2BarkleySep08.ppt
- 3 Centers for Disease Control and Prevention. (2011, August 3). Epilepsy: Frequently Asked Questions. Retrieved July 22, 2015, from <http://www.cdc.gov/epilepsy/basics/faqs.htm>
- 4 National Institutes of Health. U.S. National Library of Medicine. Genetics Home Reference. (2010, July). Lennox-Gastaut syndrome. Retrieved July 24, 2015, from <http://ghr.nlm.nih.gov/condition/lennox-gastaut-syndrome>
- 5 Tegretol® and Tegretol®-XR (carbamazepine and carbamazepine extended-release) prescribing information. (2014, September 12). Retrieved July 22, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2014/016608s099,018281s047,018927s040,020234s030lbl.pdf
- 6 Zarontin® (ethosuximide) prescribing information. (2012, April 30). Retrieved July 31, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2012/012380s034lbl.pdf
- 7 Dilantin® (phenytoin) prescribing information. (2015, April 16). Retrieved July 30, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/084349Orig1s074lbl.pdf
- 8 Depakote® (divalproex sodium) prescribing information. (2015, March 15). Retrieved July 22, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/018723s054,019680s041lbl.pdf
- 9 Tegretol® and Tegretol®-XR (carbamazepine and carbamazepine extended-release) prescribing information. (2014, September 12). Retrieved July 22, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2014/016608s099,018281s047,018927s040,020234s030lbl.pdf
- 10 Tegretol® and Tegretol®-XR (carbamazepine and carbamazepine extended-release) prescribing information. (2014, September 12). Retrieved July 22, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2014/016608s099,018281s047,018927s040,020234s030lbl.pdf
- 11 Tegretol® and Tegretol®-XR (carbamazepine and carbamazepine extended-release) prescribing information. (2014, September 12). Retrieved July 22, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2014/016608s099,018281s047,018927s040,020234s030lbl.pdf
- 12 Felbatol® (felbamate) prescribing information. (2012, August 31). Retrieved July 23, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2012/020189s027lbl.pdf
- 13 Felbatol® (felbamate) prescribing information. (2012, August 31). Retrieved July 23, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2012/020189s027lbl.pdf

- 14 Lamictal® (lamotrigine) prescribing information. (2015, May 18). Retrieved July 23, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/020241s053,020764s046,022251s0171bl.pdf
- 15 Lamictal® (lamotrigine) prescribing information. (2015, May 18). Retrieved July 22, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/020241s053,020764s046,022251s0171bl.pdf
- 16 Fycompa[™] (perampanel) prescribing information. (2015, June 19). Retrieved July 23, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/202834s0051bl.pdf
- 17 Fycompa[™] (perampanel) prescribing information. (2015, June 19). Retrieved July 23, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/202834s0051bl.pdf
- 18 Depakote® (divalproex sodium) prescribing information. (2015, March 15). Retrieved July 22, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/018723s054,019680s0411bl.pdf
- 19 U.S. Department of Health and Human Services. U.S. Food and Drug Administration. (2011, June 30). FDA Drug Safety Communication: Children Born to Mothers Who Took Valproate Products While Pregnant May Have Impaired Cognitive Development. [FDA Drug Safety Communication]. Retrieved July 30, 2015, from <http://www.fda.gov/Drugs/DrugSafety/ucm261543.htm>
- 20 U.S. Department of Health and Human Services. U.S. Food and Drug Administration. (2015, April 15). Depakene (valproic acid) Capsules and Oral Solution, Depakote (divalproex sodium) Delayed Release and Depakote ER (Extended Release) Tablets, Depakote Sprinkle Capsules (divalproex sodium coated particles in capsules), Depacon (valproate sodium) Injection. [FDA Drug Safety Communication]. Retrieved July 31, 2015, from <http://www.fda.gov/Safety/MedWatch/SafetyInformation/ucm360487.htm>
- 21 U.S. Department of Health and Human Services. U.S. Food and Drug Administration. (2011, June 30). FDA Drug Safety Communication: Children Born to Mothers Who Took Valproate Products While Pregnant May Have Impaired Cognitive Development. [FDA Drug Safety Communication]. Retrieved July 31, 2015, from <http://www.fda.gov/Drugs/DrugSafety/ucm261543.htm>
- 22 Depakote® (divalproex sodium) prescribing information. (2015, March 15). Retrieved July 22, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/018723s054,019680s0411bl.pdf
- 23 Depakote® (divalproex sodium) prescribing information. (2015, March 15). Retrieved July 22, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2015/018723s054,019680s0411bl.pdf
- 24 Sabril® (vigabatrin) prescribing information. (2014, April 3). Retrieved July 24, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2014/020427s013,022006s0141bl.pdf
- 25 Sabril® (vigabatrin) prescribing information. (2014, April 3). Retrieved July 24, 2015, from https://www.accessdata.fda.gov/drugsatfda_docs/label/2014/020427s013,022006s0141bl.pdf
- 26 U.S. Department of Health and Human Services. U.S. Food and Drug Administration. (2008, December 16). Information for Healthcare Professionals: Suicidal Behavior and Ideation and Antiepileptic Drugs. Retrieved July 31, 2015, from <http://www.fda.gov/Drugs/DrugSafety/PostmarketDrugSafetyInformationforPatientsandProviders/ucm100192.htm>

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