PRADAXA AND YOUR OTHER MEDICATIONS

Some medicines may affect the way PRADAXA works and certain medicines may increase your risk of bleeding

Here’s what you need to know:

Tell your doctor about all of your prescription and nonprescription (over-the-counter) medicines, vitamins, and herbal supplements.

Your doctor needs to know if you take rifampin (Rifater®, Rifamate®, Rimactane®, Rifadin®).

Ask your doctor or pharmacist if you are not sure if any medicine you are taking or plan to take may increase your risk of bleeding. Medicines that increase your risk of bleeding include:

• aspirin or aspirin-containing products
• long-term (chronic) use of non-steroidal anti-inflammatory drugs (NSAIDs)
• warfarin sodium (Coumadin®, Jantoven®)
• a medicine that contains heparin
• clopidogrel bisulfate (Plavix®)
• prasugrel (Effient®)
• if you have kidney problems and take dronedarone (Multaq®) or ketoconazole tablets (Nizoral®)

Know the medicines you take. Keep a list of them and show it to your doctor and pharmacist when you get a new medicine or any time you go to the doctor or pick up a prescription. Download a Medical History and Medications List.

Stopping PRADAXA may increase your risk of having a stroke or a blood clot forming in your body. Do not stop taking PRADAXA without talking to the doctor who prescribes it for you.

Tell your doctor if you are allergic to any medicines.

Please see Important Safety Information on following pages. Please visit PRADAXA.com for full Prescribing Information and Medication Guide.
What is PRADAXA?

PRADAXA is a prescription blood thinner medicine that lowers the chance of blood clots forming in your body.

PRADAXA is used to:

- reduce the risk of stroke and blood clots in people who have a medical condition called atrial fibrillation not caused by a heart valve problem. With atrial fibrillation, part of the heart does not beat the way it should. This can lead to blood clots forming and increase your risk of a stroke.

- treat blood clots in the veins of your legs (deep vein thrombosis) or lungs (pulmonary embolism) and reduce the risk of them occurring again.

PRADAXA is not for use in people with artificial (prosthetic) heart valves.

IMPORTANT SAFETY INFORMATION ABOUT PRADAXA

For people taking PRADAXA for atrial fibrillation: Do not stop taking PRADAXA without talking to the doctor who prescribes it for you. Stopping PRADAXA increases your risk of having a stroke. PRADAXA may need to be stopped prior to surgery or a medical or dental procedure. Your doctor will tell you when you should stop taking PRADAXA and when you may start taking it again. If you have to stop taking PRADAXA, your doctor may prescribe another medicine to help prevent a blood clot from forming.

PRADAXA can cause bleeding which can be serious and sometimes lead to death. Don’t take PRADAXA if you:

- currently have abnormal bleeding;
- have ever had an allergic reaction to it;
- have had or plan to have a valve in your heart replaced

Your risk of bleeding with PRADAXA may be higher if you:

- are 75 years old or older
- have kidney problems
- have stomach or intestine bleeding that is recent or keeps coming back or you have a stomach ulcer
- take other medicines that increase your risk of bleeding, like aspirin products, non-steroidal anti-inflammatory drugs (NSAIDs) and blood thinners
- have kidney problems and take dronedarone (Multaq®) or ketoconazole tablets (Nizoral®)

Please visit PRADAXA.com for full Prescribing Information and Medication Guide.
Call your doctor or seek immediate medical care if you have any of the following signs or symptoms of bleeding:

- any unexpected, severe, or uncontrollable bleeding; or bleeding that lasts a long time
- unusual or unexpected bruising
- coughing up or vomiting blood; or vomit that looks like coffee grounds
- pink or brown urine; red or black stools (looks like tar)
- unexpected pain, swelling, or joint pain
- headaches and feeling dizzy or weak

Spinal or epidural blood clots (hematoma). People who take PRADAXA and have medicine injected into their spinal and epidural area, or have a spinal puncture have a risk of forming a blood clot that can cause long-term or permanent loss of the ability to move (paralysis). Your risk of developing a spinal or epidural blood clot is higher if:

- a thin tube called an epidural catheter is placed in your back to give you certain medicine
- you take NSAIDs or a medicine to prevent blood from clotting
- you have a history of difficult or repeated epidural or spinal punctures
- you have a history of problems with your spine or have had surgery on your spine.

If you take PRADAXA and receive spinal anesthesia or have a spinal puncture, your doctor should watch you closely for symptoms of spinal or epidural blood clots. Tell your doctor right away if you have back pain, tingling, numbness, muscle weakness (especially in your legs and feet), loss of control of the bowels or bladder (incontinence).

Tell your doctor if you are pregnant or plan to become pregnant. It is not known if PRADAXA will harm your unborn baby. Tell your doctor right away if you become pregnant during treatment with PRADAXA.

Tell your doctor if you are breastfeeding or plan to breastfeed. It is not known if PRADAXA passes into your breast milk. You and your doctor should decide if you will take PRADAXA or breastfeed.

Take PRADAXA exactly as prescribed. It is important to tell your doctors about all medicines (prescription and over-the-counter), vitamins, and supplements you take. Some medicines may affect the way PRADAXA works.

PRADAXA can cause indigestion, stomach upset or burning, and stomach pain.

You are encouraged to report negative side effects of prescription drugs to the FDA. Visit www.fda.gov/medwatch or call 1-800-FDA-1088.

Please see full Prescribing Information and Medication Guide.