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Eligard package insert pdf

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Any application of this information for any purpose is limited to personal, non-commercial use. Ã, â â â â â â â â â â â «Back to deficiency of drugs Product news Subscribe to AHFS Clinical Drug Information to obtain direct access to integrated content Drug deficiency, more complete and usable drug information. Class: Hormonal therapy Generic name: Leelfolide (Loo-Proe-Lide) Commercial name: EligardÃ‚Â®, LupronÃ‚Â®, Lupron DepotÃ‚Â®, ViadurÃ‚Â® How do you use this medication? Leelfrolide is approved by the FDA for the treatment of advanced prostate carcinoma. It is important for patients to remember that doctors have drug ability to prescribe for conditions other than those for whom the drug has been approved by the FDA. Patients who received a prescription of this drug for a different condition from which it is approved may wish to discuss the question with their doctor. What is the mechanism of action? Leelfolide is classified as a luteinizing hormone Releasing hormone (LHRH) agonist. LHRH is a naturally present hormone in the body that involved in stimulating the production of female and male hormones such as estrogen and testosterone. Leelfrolide inhibits the signaling of LHRH way that the production of estrogen or testosterone is reduced. Some types of cancer, such as prostate cancer, are stimulated to grow from hormones. The reduction in the production of growth-stimulator hormones inhibits the growth of tumor cells. How is Leuprolide Data (administered)? LeUprolide is administered by injection under the skin (subcutaneous), in a muscle (intramuscular), or as an implantable device and the dose depends on several factors, including the condition that is treated, the regime of details used and the general health of the patient. How are monitored patients? Patients are usually scheduled to meet with their health care provider while they are treated with Leelfrolide.ã, Blood can be designed to control hormone levels. Patients can also suffer physical examinations, scans or other measures to evaluate side effects and therapy response. Patients will be monitored for a phenomenon called Ã ¢ FlareÃ ¢ where hormonal levels can raise themselves during initial therapy with Leelfrolide. The high levels of hormones can cause a sudden increase in symptoms, such as bone pain in patients with the spread of bone cancer, or the obstruction of the urinary system with consequent difficulty or incapacity to urinate in patients with prostate cancer. Patients who experience greater bone pain, especially the spine, difficulty to urinate, blood in urine, or numbness of hands or feet should consult immediately doctor. Patients will also have their monitored heart function, such as Leelfolide causes unusually serious side effects for the heart. Furthermore, patients will be monitored for blood clots, as this is another rare but serious condition caused by Leelfolide. If patients experience pain, redness and / or swelling of a limb and none other, the other, breathing, chest pain, or significant changes in heart rate or rhythm, should contact their health care provider immediately.Ã Patients may also be monitored for bone density, such as long-term use of leuprolide may decrease the density of the bones. What are common (occurring in 30% or more of patients) side effects of treatment with leuprolide? Ã Â ¢ Ã Â ¢ Ã flushing ¢ Ã reduced sex drive Ã ¢ Â ¢ Ã Â ¢ Ã ¢ Ã impotence Erectile dysfunction What it is the less common (occurring in 10% to 29% of patients) side effects of treatment with leuprolide? Ã Â ¢ a breast enlargement, tenderness or pain Ã ¢ Â ¢ Ã The increase in cholesterol levels ¢ Ã ¢ Ã fluid retention in the swelling of the feet or ankles Ã Â ¢ Ã generalized weakness or fatigue Ã ¢ Â ¢ a depression Ã ¢ Â ¢ Ã Ã ¢ Ã Ã Ã sweating bone pain increased due to a flareÃ ¢ if patients have cancer spread to the bones ¢ Ã Ã ¢ Ã inability or difficulty urinating due to a Ã ¢ flareÃ ¢ Ã ¢ Â ¢ Ã reactions at the injection site this is not a complete list of side effects. Some patients may have other side effects that are not listed here. Patients may want to discuss with your doctor the other less side effects of this drug, some of which may be severe. Some side effects may require medical attention. Other side effects do not require medical attention and may go away during treatment. Patients should check with their doctor about any side effects that continue or are bothersome. What are the possible late effects of treatment with leuprolide? A rare but serious effect, part of which is associated with treatment with leuprolide is damage to the heart. Patients who experience chest pain, breathing difficulty or significant changes in heart rate or rhythm should contact their health care provider. Treatment with leoprolide can also lead to decreased bone mineral density, which increases the risk of fractures. Patients should talk to their doctor about the risk of late side effects weight carefully the benefits of treatment with these risks. What can patients do to help alleviate or prevent the discomfort and side effects? Ã Â ¢ Ã pay particular attention to the instructions of the physicianÃ ¢ s and tell your doctor about any side effects. Ã Â ¢ Ã Keep rest and proper nutrition. Ã Â ¢ Ã wear sunscreen and protective clothing and try to minimize the exposure to the sun. Ã Ã Ã ¢ Drink plenty of fluids. (Patients should ask their doctor about the amount of liquid to be consumed during the day.) Are there any special precautions patients should be aware of before you start treatment? Ã Â ¢ Ã Patients should inform their doctor if they are pregnant, breastfeeding or planning a family in the near future. This medication can cause birth defects. It 'important to use some type of birth control during treatment. In addition, patients may want to talk with your doctor if you are thinking of having children in the future, since some drugs can cause fertility problems. Ã Ã ¢ Ã It 'important that patients inform their physician of any pre-existing conditions (chickenpox, heart disease, kidney disease, liver disease, lung disease, etc.), since © can worsen with this drug. Ã Â ¢ Ã Patients should inform your doctor of any other medication you are taking (both prescription and over-the-counter medications, including vitamins, herbs, etc.) because they might interfere with treatment. Ã Â ¢ Ã Patients should check with their Before starting any new drug or nutritional supplement. Ãƒ â ¢ Ã, patients must inform your doctor of any known drug or food allergies or any reactions to drugs that have experienced in the past. When patients have to inform his doctor? Ã, â ¢ Ã, â ¢ ¢ breathing difficulty A chest pain Ã ¢ â ¢ A change in heart rate or rhythm Ã ¢ â ¢ Ã Â, numbness or tingling of hands or feet Ãƒ â ¢ Ã, incapacitÃ o The difficulty of urinating â ¢ A severe depression ã, â ¢, memory impairment or confusion Ã ¢ â ¢ Ã Ã Ã Â ¢ ¢ a leg weakness Ã ¢ â ¢ Ã Ã, swelling of feet and ankles Ã ¢ Â ¢ Ã, sudden weight increase Ãƒ Â ¢ Ã, bone pain in the lower back or dock what is an illustrative leaflet? A package insert is By the FDA and contains a synthesis of the essential scientific information necessary for the safe and effective use of the drug for health and consumers.ã, a leaflet generally includes information regarding specific indications, scheduled administration, dosage, side effects , Contraindications, results from some clinical studies, chemical, pharmacokinetics and metabolism of the specific Drug.ã, carefully examining the leaflet, you get the most complete and up-to-date information on how to safely use this medication. If you do not have the paper leaflet of the drug you are using, your pharmacist or doctor can be able to supply with a copy. Copyright Ã © 2016 cancerconnect Last updated 07/10. Important limitations to use the information provided above the drug has been selected is only provided to your information and is not a substitute for consultation with an adequate doctor. 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