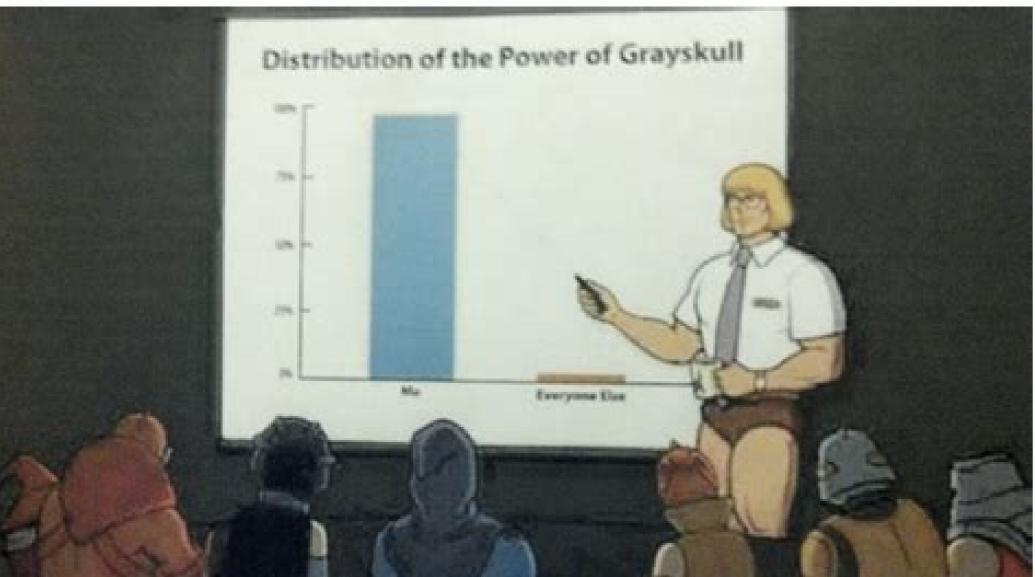
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A Gentle Introduction to Sta



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to give its lenders an idea if a potential borrower will be able to qualify for a new mortgage debt. As a risk-assessment tool, LP is not only the first step in the process of helping lenders underwrite mortgages, but it's also designed to help borrowers find the lowest rates available. Tips LP, which stands for Loan Prospector, is the Freddie Mac automated underwriting service used by third-party loan originators and mortgage wholesale lenders that provides risk assessment for Freddie Mac's credit and pricing terms. The results from running an application through the LP program help a lender to determine if the borrower needs to be reassessed for another loan program or if he simply needs to make some financial changes in order to be qualified for a mortgage. If a borrower receives a result of approve or accept, it means that the loan will close, it simply means that the borrower qualifies for a mortgage. However, it does not mean that the loan will close, it simply means that the proval. LP is a good tool for a mortgage lender to use to weed out successful mortgage applicants from those that will not produce a mortgage closing. Additionally, it tells the lender automatically what changes need to occur in order for the borrower to be able to qualify later. This gives the lender automatically what changes need to occur in order for the borrower to be able to qualify later. Mac's LP is not the only mortgage underwriting software in the industry. Another major program is known as DU, or Desktop Underwriter and is the official loan underwriting software of the other major mortgage underwriting software of the other major mortgage. The lender should review the application and his entry work into the computer to determine if the decline was due to the application to be erroneously denied. LP is a good tool for lenders to use to determine guickly and effectively which clients he should focus on and which ones he should educate for future loan opportunities. It helps the lender to keep from wasting both his and the borrower's time on a mortgage as well. The FDA has approved Astra-Zeneca's Movantik (naloxegol) tablets for the treatment of opioid-induced constipation (OIC) in adults with chronic noncancer pain. 1 Due to its structural similarity to noroxymorphone, Movantik was initially approved as a Schedule II controlled substance; however, its labeling stated that it carried no risk of abuse or dependency. The Drug Enforcement Administration descheduled Movantik in February 2015, and AstraZeneca requested that the FDA remove the controlled substance designation. 1,2 Movantik could be available by June 2015.2Pharmacology and Pharmacokinetics Opioids bind to mu-receptors in the gastrointestinal (GI) tract, resulting in constipation. Movantik is an oral peripherally acting mu-opioid receptor antagonist (PAMORA) that exerts its effect by preventing binding with mu-receptors in the GI tract, with negligible CNS interference.1,2After oral administration, Movantik should be taken on an empty stomach. Its metabolism is primarily through the CYP3A enzyme system.1Dosage and AdministrationMovantik should be given at a dosage of 25 mg once daily; if not tolerated, the dosage may be decreased to 12.5 mg once daily; if this is tolerated, it may be increased to 25 mg once daily. Maintenance laxative therapy should be discontinued prior to initiation of treatment with Movantik, but may be resumed if patients experience OIC symptoms after taking Movantik Movantik has been shown to be efficacious in patients who have been using opioids for at least 4 weeks. It should be taken on an empty stomach at least 1 hour before the first meal of the day or 2 hours after the meal. The tablets should be avoided during treatment with Movantik. Movantik should be discontinued if opioid therapy is discontinued.1Clinical Trials Movantik was evaluated in 2 replicate, randomized, doubleblind placebo-controlled trials in patients with OIC and non—cancerrelated pain. For 12 weeks, 1352 patients received Movantik 12.5 mg, Movantik 25 mg, or placebo. A statistically significant increase in the number of bowel movements per week occurred in the 12.5- and 25-mg groups in study 1 and in the 25-mg group in study 2.1,3Contraindications, Warnings, and Precautions Movantik is contraindicated in patients (1) who are using strong CYP3A4 inhibitors, such as clarithromycin and ketoconazole, or (4) with a known serious or severe hypersensitivity reaction to Movantik or any of its excipients. The overall risks and benefits of Movantik should be monitored for severe, persistent, or worsening abdominal pain, and Movantik should be discontinued if these symptoms develop. The overall risks and benefits should also be considered in patients with disruptions to the blood—brain barrier; these patients should be monitored for symptoms of opioid withdrawal. Concomitant use with moderate CYP3A4 inhibitors, such as diltiazem, erythromycin, and verapamil, should be avoided, as the combination may increase Movantik concentrations. If the combination is unavoidable, the dosage of Movantik should be decreased to 12.5 mg once daily. Concomitant use with strong CYP3A4 inducers, such as rifampin, should be avoided, as the combination may decrease Movantik concentrations. Movantik should not be used in combination with other opioid antagonists. Movantik is a Pregnancy Category C medication. It should not be used during breast-feeding or in patients with severe hepatic impairment. In clinical trials, the most common adverse reactions (>3%) were abdominal pain, diarrhea, nausea, flatulence, vomiting, and headache. 1Dr. Holmberg earned her Pharm D from the University of Connecticut and completed an ambulatory care residency at the Phoenix VA Healthcare System. Her practice has also included pediatrics and inpatient mental health. She resides in Phoenix, Arizona. References 1. Movantik [package insert]. Wilmington, DE: AstraZeneca Pharmaceuticals LP; 2014. www.azpicentral.com/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/movantik/m tablets-cii. Accessed November 2014.3. FDA approves Movantik for opioid-induced constipation. FDA website. www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm414620.htm. Accessed November 2014. I needed a narrow coffee table for that would allow the space I needed for my home office. This coffee table was quick and easy to

make... 1) I used 2 old metal milk crates for the stand that we were using to store old LP records. 2) 1 wall shelf that I dismantled for the top and then place a pane of glass

over the records to protect them. Glass can be purchased from any glass shop and is very affordable. My glass guy to donate scrap glass for different projects. Will post new pics when I get Phil the glass guy to cut me some glass to size. Gail the Pond Plant Girl

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