Janet T. Mills Governor

Jeanne M. Lambrew, Ph.D. Commissioner



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DUR COMMITTEE AGENDA

Date: Tuesday, March 12, 2019

Time: 5:30PM to 8:30PM

Location: Augusta Armory, 179 Western Avenue, Augusta, ME

- 1) Call to Order
- 2) MaineCare Updates
- 3) Public Comments
- 4) Old Business
- 5) Closed Session
 - Drug Financial Information Review
- 6) New Business (open session)
 - Reviews on Biosimilars
 - Retro DUR
 - Data Presentation: Vivitrol Adherence
 - Introduction: Continuous use of antidepressants at 3, 6 and 12 months after initiation
 - New Drug Review

Abilify Mycite Lorbrena Xelpros Arikayce Panzyga Xerava Qbrexza pad Bryhali Xospata Copiktra Siklos **Xyosted** Yupelri Cequa Symjepi Daurismo Sympazan Zemdri

Galafold Tiglutik Invenitys Tolsura

A. FDA Safety Alerts

FDA warns that symptoms of a serious condition affecting the blood cells are not being recognized with the leukemia medicine Idhifa (enasidenib)

https://www.fda.gov/Drugs/DrugSafety/ucm626923.htm?utm_campaign=New%20FDA%20Drug%20Safety%20Communication%20on%20Idhifa%20%28enasidenib%20mesylate%29-%20Drug%20I&utm_medium=email&utm_source=Eloqua

Fluoroquinolone Antibiotics: Safety Communication - Increased Risk of Ruptures or Tears in the Aorta Blood Vessel in Certain Patients

https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm6 28960.htm?utm_campaign=FDA%20MedWatch%20-

%20Fluoroquinolone%20Antibiotics&utm medium=email&utm source=Eloqua

FDA warns about rare but serious risks of stroke and blood vessel wall tears with multiple sclerosis drug Lemtrada (alemtuzumab)

https://www.fda.gov/Drugs/DrugSafety/ucm624247.htm?utm_campaign=FDA%20MedWatch%20-%20Lemtrada%20%28alemtuzumab%29%3A%20Drug%20Safety%20Communication&utm_medium=e mail&utm_source=Elogua

- B. Next Meeting (Tuesday, June 11, 2019 (from 5:30pm to 8:30pm)
- C. Adjournment: 8:30PM