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DUR COMMITTEE AGENDA

Date: Tuesday, December 13, 2016
Time: 5:30PM to 8:30PM
Location: Augusta Armory, 179 Western Avenue, Augusta, ME

- 1) Call to Order
- 2) Introductions of New Committee Members
- 3) MaineCare Updates
- 4) Public Comments
- 5) Old Business
 - Review of Minutes
- 6) Closed Session
 - Drug Financial Information Review
- 7) New Business (open session)
 - A. PCM Report- Hepatis C
 - B. Use of Antipsychotics in Maine Report
 - C. Retro-DUR
 - Introduce: Co-prescribing of stimulants, benzodiazepines and "Z" drugs.
 - D. New Drug Review
 - Bromsite
 - Byvalson
 - Emverm
 - Exondys
 - Inflectra
 - Invokamet XR
 - Jentadeuto XR
 - Onzetra
 - Otovel
 - Probuphine
 - Qbrexis
 - Relistor Tabs
 - Sustol Inj
 - Zurampic

E. FDA Safety Alerts

FDA analyses conclude that Xarelto clinical trial results were not affected by faulty monitoring device
http://www.fda.gov/Drugs/DrugSafety/ucm524678.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery

Testosterone and Other Anabolic Androgenic Steroids (AAS): FDA Statement - Risks Associated With Abuse and Dependence
http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm526151.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery

FDA launches competition to spur innovative technologies to help reduce opioid overdose deaths
http://www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm520945.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery

Eye Wash/Eye Irrigating Solutions Distributed by Major Pharmaceuticals and Rugby Laboratories: Recall - Microbial Contamination
http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm519570.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery

Hyoscyamine Sulfate 0.125mg by Virtus Pharmaceuticals: Recall - Superpotent and Subpotent Test Results
http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm520868.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery

Lamotrigine Orally Disintegrating Tablet 200 mg by Impax: Recall - Incorrect Labeling of Blister Cards
http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm518486.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery

Opioid Pain or Cough Medicines Combined With Benzodiazepines: Drug Safety Communication - FDA Requiring Boxed Warning About Serious Risks and Death
http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm518710.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery

Ovarian Cancer Screening Tests: Safety Communication - FDA Recommends Against Use
http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm519540.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery

Direct-Acting Antivirals for Hepatitis C: Drug Safety Communication - Risk of Hepatitis B Reactivating
http://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm523690.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery

F. Next Meeting (Tuesday, March 14th, 2017 from 5:30pm to 8:30pm)

G. Adjournment: 8:30PM