



Department of Health  
and Human Services

Maine People Living  
Safe, Healthy and Productive Lives

Paul R. LePage, Governor

Ricker Hamilton, Acting Commissioner

Department of Health and Human Services  
11 State House Station  
Augusta, Maine 04333-0011  
TTY Users: Dial 711 (Maine Relay)

DUR COMMITTEE AGENDA

Date: Tuesday, September 12, 2017  
Time: 5:30PM to 8:30PM  
Location: Augusta Armory, 179 Western Avenue, Augusta, ME

- 1) Call to Order
- 2) Introductions
- 3) MaineCare Updates
- 4) Public Comments
- 5) Old Business
  - Review of Minutes
  - Assessing patients on Stimulants in combination with psych medication seperated into age bans.
- 6) Closed Session
  - Drug Financial Information Review
- 7) New Business (open session)
  - A. Annouce new DUR Chair
  - B. Retro-DUR
    - Data presentation: Co-prescribing of opiate pain medications (including cough syrups), benzodiazepines and "Z" drugs.
  - C. Retro-DUR
    - Introduce: Compliance with GLP-1 Agonists in Type II Diabetes Mellitus
  - D. New Drug Review
    - Imfinzi
    - Kevzara
    - Mavyret
    - Radicava
    - Seebri Neohaler
    - Siliq
    - Syndros
    - Tremfya
    - Tymlos
    - Vosevi

E. FDA Safety Alerts

**Brilinta (ticagrelor) 90 mg tablets, 8-count Physician Sample Bottles: Recall of Lot # JB5047 - Due to Report of Another Medicine in One Bottle**

[https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm560786.htm?source=govdelivery&utm\\_medium=email&utm\\_source=govdelivery](https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm560786.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery)

**Mibela 24 Fe Chewable Tablets by Lupin Pharmaceuticals Inc.: Recall - Out of Sequence Tablets and Missing Expiry/Lot Information**

[https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm560908.htm?source=govdelivery&utm\\_medium=email&utm\\_source=govdelivery](https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm560908.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery)

**Novopen Echo Insulin Delivery Device by Novo Nordisk: Recall - May Crack or Break If Exposed To Certain Chemicals**

[https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm565955.htm?source=govdelivery&utm\\_medium=email&utm\\_source=govdelivery](https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm565955.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery)

**FDA requests removal of Opana ER for risks related to abuse**

[https://www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm562401.htm?source=govdelivery&utm\\_medium=email&utm\\_source=govdelivery](https://www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm562401.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery)

F. Next Meeting (Tuesday, October 10, 2017 from 1:00pm to 4:30pm)

G. Adjournment: 8:30PM