



## DUR COMMITTEE AGENDA

Date: Tuesday, June 27, 2017  
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
  - Make Elidel and Protopic preferred
  - Retro-DUR: Additional data on Co-prescribing of stimulants, benzodiazepines and Z drugs
- 6) Closed Session
  - Drug Financial Information Review
- 7) New Business (open session)
  - A. Retro-DUR
    - Data presentation: Assessing the pattern of usage of long-acting stimulants, specifically looking at more than once a day dosing patterns and use among different age groups, including pediatrics.
  - B. Retro-DUR
    - Introduce: Co-prescribing of opiate pain medications (including cough syrups), benzodiazepines and “Z” drugs.
  - C. New Drug Review
    - Airduo
    - Alunbrig
    - Arymo ER
    - Austedo
    - Bavencio
    - Dupixent
    - Emflaza
    - Ingrezza
    - Kisqali
    - Lartruvo
    - Morphabond ER

Ocrevus  
Rhofade  
Rydapt  
Spinraza  
Synjardy XR  
Trulance  
Utibron  
Xadago  
Xermelo  
Xultophy  
Zejula

D. FDA Safety Alerts

**FDA Drug Safety Communication: FDA updates warnings for oral and injectable fluoroquinolone antibiotics due to disabling side effects**

[https://www.fda.gov/Drugs/DrugSafety/ucm511530.htm?source=govdelivery&utm\\_medium=email&utm\\_source=govdelivery](https://www.fda.gov/Drugs/DrugSafety/ucm511530.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery)

**Canagliflozin (Invokana, Invokamet): Drug Safety Communication - Increased Risk of Leg and Foot Amputations**

[https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm558605.htm?source=govdelivery&utm\\_medium=email&utm\\_source=govdelivery](https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm558605.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery)

**FDA Drug Safety Communication: FDA identifies no harmful effects to date with brain retention of gadolinium-based contrast agents for MRIs; review to continue**

[https://www.fda.gov/Drugs/DrugSafety/ucm559007.htm?source=govdelivery&utm\\_medium=email&utm\\_source=govdelivery](https://www.fda.gov/Drugs/DrugSafety/ucm559007.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery)

**FDA Drug Safety Communication: FDA restricts use of prescription codeine pain and cough medicines and tramadol pain medicines in children; recommends against use in breastfeeding women**

[https://www.fda.gov/Drugs/DrugSafety/ucm549679.htm?source=govdelivery&utm\\_medium=email&utm\\_source=govdelivery](https://www.fda.gov/Drugs/DrugSafety/ucm549679.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery)

**FDA alerts consumers of nationwide voluntary recall of EpiPen and EpiPen Jr**

[https://www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm550170.htm?source=govdelivery&utm\\_medium=email&utm\\_source=govdelivery](https://www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm550170.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery)

**Information on Erythropoiesis-Stimulating Agents (ESA) Epoetin alfa (marketed as Procrit, Epogen), Darbepoetin alfa (marketed as Aranesp)**

[https://www.fda.gov/Drugs/DrugSafety/PostmarketDrugSafetyInformationforPatientsandProviders/ucm109375.htm?source=govdelivery&utm\\_medium=email&utm\\_source=govdelivery](https://www.fda.gov/Drugs/DrugSafety/PostmarketDrugSafetyInformationforPatientsandProviders/ucm109375.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery)

**FDA approves two hepatitis C drugs for pediatric patients**

[https://www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm551407.htm?source=govdelivery&utm\\_medium=email&utm\\_source=govdelivery](https://www.fda.gov/NewsEvents/Newsroom/PressAnnouncements/ucm551407.htm?source=govdelivery&utm_medium=email&utm_source=govdelivery)

E. Next Meeting (Tuesday, September 12, 2017 from 5:30pm to 8:30pm)

F. Adjournment: 8:30PM