



Department of Health and Human Services  
 MaineCare Services  
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DUR COMMITTEE AGENDA

Date: Tuesday, June 12, 2018  
 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
- 6) Closed Session
  - Drug Financial Information Review
- 7) New Business (open session)
  - Retro-DUR
    - Introduction: Use of Naloxone Intolerance
  - Retro-DUR
    - Data presentation: Statin Use in ASCVD
  - Updated Suboxone PA form
  - Therapeutic Drug Class Review  
Phosphate Binders, Non-Calcium
  - New Drug Review
 

Aimovig	Lonhala	Steglujan
Carospir	Luxturna	Symdeko
Cimduo	Lyrice CR	Symfi
Crysvita	Noctiva	Symfi Lo
Daxbia	Osmolex ER	Trogarzo
Erleada	Rhopressa	Xhance
Gocovri	Segluromet	Zilretta
  - FDA Safety Alerts
 

Risk of serious and potentially fatal blood disorder prompts FDA action on oral over-the-counter benzocaine products used for teething and mouth pain and prescription local anesthetics  
<https://www.fda.gov/Drugs/DrugSafety/ucm608265.htm>

Lamictal (lamotrigine): Drug Safety Communication - Serious Immune System Reaction  
<https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm605628.htm>

Juluca, Tivicay, Triumeq (dolutegravir): FDA to Evaluate - Potential Risk of Neural Tube Birth Defects  
[https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm608168.htm?utm\\_campaign=FDA%20MedWatch%20-%20Juluca%2C%20Tivicay%2C%20Triumeq&utm\\_medium=email&utm\\_source=Eloqua](https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm608168.htm?utm_campaign=FDA%20MedWatch%20-%20Juluca%2C%20Tivicay%2C%20Triumeq&utm_medium=email&utm_source=Eloqua)

Keytruda (pembrolizumab) or Tecentriq (atezolizumab): FDA Alerts Health Care Professionals and Investigators: FDA Statement - Decreased Survival in Some Patients in Clinical Trials Associated with Monotherapy  
<https://www.fda.gov/Safety/MedWatch/SafetyInformation/SafetyAlertsforHumanMedicalProducts/ucm608253.htm>

- Next Meeting (Tuesday, September 11, 2018 from 5:30pm to 8:30pm)
- Adjournment: 8:30PM