Children's Mercy Kansas City

SHARE @ Children's Mercy

Presentations

4-23-2020

Missouri Department of Social Services Rare Disease Advisory Committee Update On Ivacaftor/Tezacafto/Elexacaftor

Christopher M. Oermann Children's Mercy Hospital

Let us know how access to this publication benefits you

Follow this and additional works at: https://scholarlyexchange.childrensmercy.org/presentations

Recommended Citation

Oermann, Christopher M., "Missouri Department of Social Services Rare Disease Advisory Committee Update On Ivacaftor/Tezacafto/Elexacaftor" (2020). *Presentations*. 62. https://scholarlyexchange.childrensmercy.org/presentations/62

This Presentation is brought to you for free and open access by SHARE @ Children's Mercy. It has been accepted for inclusion in Presentations by an authorized administrator of SHARE @ Children's Mercy. For more information, please contact histeel@cmh.edu.

Missouri Department of Social Services Rare Disease Advisory Committee Update on Ivacaftor/Tezacaftor/Elexacaftor

Christopher M Oermann, MD

Division Director, Allergy, Immunology, Pulmonary, and Sleep Medicine

Director, Cystic Fibrosis Care Center



Rare Disease Advisory Committee

- Established 2019 to advise Drug Utilization Review Board
- Multidisciplinary committee comprised of allied healthcare professionals (physicians, PharmD, RN, PhDs) and MO HealthNet staff
- Quarterly meetings
- Appointment based on experience researching, diagnosing, and treating rare diseases
 - Provide expert recommendations or determinations regarding access to drugs and/or biological products for rare disease treatment



Ivacaftor/Tezacaftor/Elexacaftor

- FDA Approval: 21 October, 2019
- Indication: treatment of cystic fibrosis (CF) in patients aged 12 years and older who have at least one F508del mutation in the CFTR gene
- Mechanism of Action:
 - Ivacaftor is a CFTR potentiator
 - Tezacaftor and Elexacaftor are CFTR correctors

Dose/Administration:

- 2 tablets (elexacaftor 100 mg, tezacaftor 50 mg, and ivacaftor 75 mg) in AM
- 1 tablet (ivacaftor 150 mg) in PM



Drug Clinical Highlights:

- First CFTR modulator therapy for F508del heterozygotes (90% of CF population)
- FDA approval based on two phase 3 clinical trials including 510 people with CF
 - First trial demonstrated an increase in ppFEV₁ of 13.8% and improvements in sweat chloride, pulmonary exacerbation rate, and BMI
 - Second trial demonstrated an increase in ppFEV₁ of 10%
- Warnings regarding: liver function, concomitant use of CYP3A inhibitors, and cataracts
- Disease Clinical Highlights:
 - Pathophysiology
 - Epidemiology
 - Genetics



Cost:

- \$310,648 per patient annually
- Estimated at \$28,000,000-55,000,000 for MO HealthNet
- No therapeutic alternatives
- Guidelines:
 - Initial Therapy
 - Documented diagnosis of cystic fibrosis
 - Genetic testing documenting F508del
 - Prescribed by or in consultation with "appropriate specialist" at a CF Care Center
 - Age \geq 12 years
 - Screening tests: LFTs, Pulm Function, eye exam, no severe liver disease
 - Continuation
 - Annual eye exam
 - LFTs every 3 months for a year then annually
 - Annual documentation of improvement in measurable goal: ppFEV1, pulmonary exacerbation, BMI

