Postpartum Depression

Clinical Cases Applicability: Antepartum mood disorders, postpartum blues, postpartum depression

Learning Objectives:
1. Understand the neurobiological mechanisms underlying postpartum depression
2. Describe the pharmacology of SSRIs

What are some sex differences in depression?
- Women 2x more likely to develop depression, greatest difference in reproductive years – suggests sex hormones (estrogen, progesterone, oxytocin) and reproductive events play some role in etiology of depression

What is the definition of PDD? prevalence? Risk factors? Natural history?
- DSM 5 defines peri-partum depression as symptom onset during pregnancy or <4 weeks PP – but greatest incidence of new PPD occurs 2-3 months postpartum, and can occur within 1 year
- Prevalence ~10-15%
- Risk factors: antenatal depression (↑ prevalence in 2\textsuperscript{nd}/3\textsuperscript{rd} trimesters) & depression prior to pregnancy; stressful life events (marital conflict, emigration), poor social & financial support
- Often remits on its own within 12 months after giving birth, 6x increased risk for depression later in life

What are proposed neurobiological mechanisms underlying PPD? Pathogenesis unknown
- Genetics: increased risk with family history of PPD
- Abnormal neurotransmitter levels/activity: ↑ brain monoamine oxidase → ↑ metabolism of neurotransmitters such as dopamine, norepinephrine, serotonin → more rapid depletion of these neurotransmitters → depression
- Significant changes in neural activity in brain regions important for self-regulation, empathy & emotion; fMRI studies identify brain regions affected by postpartum depression at rest and in response to infant or non-infant emotional cues
  - Decreased corticortical and corticolimbic connectivity
  - Weaker connectivity among the amygdala (processes emotions), anterior cingulate cortex (connects the “emotional” limbic system and the “cognitive” prefrontal cortex), dorsal lateral prefrontal cortex (working memory) & hippocampus (long term memory, emotional responses)
  - Decreased connectivity between the amygdala and right insular cortex w/ positive infant cues
- Abnormal growth factors: ↓ Brain-derived neurotrophic factor (supports differentiation, maturation and survival of neurons) concentrations in PPD

What hormones may play a role in PPD? Women with a predisposition for depression may be more sensitive to large fluctuations in steroid hormone levels
- Estradiol: estradiol levels continue to ↑ in 3\textsuperscript{rd} trimester, but drop dramatically after birth “estradiol-withdrawal state”
- Cortisol: high levels of cortisol during pregnancy secondary to placentally-derived CRH; cortisol stimulates CRH production in placenta (positive feedback); withdrawal from excess cortisol postpartum (figure 1)
- Oxytocin: hypothesized to have anti-depressant properties; women at risk for PPD have lower oxytocin concentrations in 3\textsuperscript{rd} trimester; BUT administering oxytocin does not improve mood

Why is treatment of PPD important?
Negative outcomes for mothers and infants: ↓ breastfeeding, lack of maternal emotional and behavioral sensitivity to the infant, poor bonding, negative infant temperament, atypical neurodevelopment, and later emotional and behavioral problems for the children

What are pharmacological treatment options for PPD? Mechanism of action? Women with PPD take longer to recover & require more therapies than in non-postpartum MDD women
- SSRIs – paroxetine and sertraline usually undetectable levels in breast-fed infants; SSRIs block the SERT transporter to ↓ presynaptic serotonin reuptake → ↑ serotonin concentration (figure 2)
References:
- Viguera, A. Severe unipolar major depression: Treatment. In: UpToDate, Post TW (Ed), UpToDate, Waltham, MA. (2017)
- Viguera, A. Postpartum unipolar major depression: Epidemiology, clinical features, assessment and diagnosis. In: UpToDate, Post TW (Ed), UpToDate, Waltham, MA. (2016)