Mental Health Issues in Women Across the Reproductive Cycle

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Disclosure

I have no financial interests or relationships to disclose.

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Objectives

- At the end of this session, participants will be able to:
 - Describe how to identify and treat peripartum depression.
 - Discuss considerations of medication-based treatment in the postpartum period, particularly if the patient is breastfeeding
 - Assess and manage mood symptoms in menopausal and postmenopausal women.

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Premenstrual Syndrome (PMS) and Premenstrual Dysphoric Disorder (PMDD)



- Longitudinal research indicates some women demonstrate greater sensitivity to gonadal steroid shifts
- h/o severe premenstrual mood sx is associated with increased risk of perimenopause onset or relapse of MDD



Freeman EW. Hormones and menopausal status as predictors of depression in womer in transition to menopause. Arch Gen Psych 2004;61:62-70

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PMS/PMDD

- Characterized by:
 - Presence of physical and/or behavioral symptoms that occur repetitively in the second half of the menstrual cycle and often the first few days of menses.
 - Symptom-free during the follicular phase. Otherwise, would be thinking more Major Depressive Disorder.
- Clinically significant PMS ~3 to 8 %
- PMDD ~ 2%

Treatment:

Mild symptoms (Sx)

- Lifestyle measures
 - Exercise
 - Relaxation techniques



Moderate to severe Sx

- Cognitive behavioral therapy
- Medications
 - Selective serotonin reuptake inhibitors [SSRIs]
 - Combined estrogen-progestin contraceptives (COCs)
 - Vitex agnus castus (chasteberry)?
 - Continuous versus intermittent?

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Medication Treatment

- Antidepressants: Most data is available for sertraline, citalopram, escitalopram and fluoxetine. Effexor
 was effective but withdrawal is more problematic so not as frequently recommended
- Vitex: Two trials in women with PMDD (vitex versus fluoxetine) were also analyzed; fluoxetine was the
 more effective intervention. However, vitex did show some benefit in one of the PMDD trials. The most
 common dosing studied is 20 to 40 mg of vitex extract.
- OCPs: Suppressing the hypothalamic-pituitary-ovarian axis to abolish cyclic changes in gonadal steroids (combined estrogen-progestin oral contraceptives [COCs], gonadotropin-releasing hormone [GnRH] agonists). Monophasic OCP pills recommended, as multiphasic preparations can worsen mood symptoms

Continuous versus Intermittent for Antidepressant Treatment

- In continuous dosing patient takes antidepressant daily vs intermittent dosing where patient takes daily during the luteal phase, i.e., from ovulation to menses only.
- Systematic review concluded that continuous and intermittent dosing had equivalent efficacy

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Perinatal Depression

Pregnancy as a Risk Factor?

- Overall, population prevalence rates of depression for women are similar before and during pregnancy
 - General prevalence post puberty: 10-20%
 - 1st trimester prevalence: 11%
 - 2nd and 3rd trimester prevalence: 8.5%
 - But...there are things that increase risk

T Pearlstein. Depression During Pregnancy. Best Practice & Research Clinical Obstetrics and Gynecology 29 (2015) 754-764.

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Time for Audience Participation!

What do you think are risk factors for perinatal depression?



Risk Factors for Perinatal Depression

- Personal or family hx of major or postpartum depression
- Gestational diabetes
- Difficulty breastfeeding
- Fetal/newborn loss
- Lack of personal or community resources
- Financial challenges
- Current anxiety
- Single

- Substance use/addiction
- Complications of pregnancy, labor/delivery, or infant's health
- Adolescence
- Unplanned pregnancy
- Major life stressors
- Violent or abusive relationship
- Isolation from family or friends

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Pregnancy as a Risk Factor?

- There does appear to be an increased risk for anxiety symptoms worsening during pregnancy
 - Particularly seen in:
 - Generalized anxiety disorder
 - 8.5% prevalence in pregnancy compared to 3% in gen pop
 - Some confluence with adjustment disorder with anxious features
 - Panic disorder
 - Overall prevalence is not higher (~2%) though panic disorder symptoms may worsen in severity
- Pregnancy can be a very anxiety provoking stimulus

Thorsness KR, Watson C, LaRusso EM. Perinatal anxiety: approach to diagnosis and management in the obstetric setting. American Journal of Obstetrics & Gynecology. (2018); October: 326-345.

How Do You Detect Perinatal Depression?



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Screening

Ideally..

- Once per trimester
- At postpartum visit (2 weeks if high risk, or 6-7 weeks routine visit)
- At well-child visit



Stewart DE, Vigod S. Postpartum Depression. N Engl J Med (2016);375;22:2177-2186. T Pearlstein. Depression During Pregnancy. Best Practice & Research Clinical Obstetrics and Gynecology 29 (2015) 754-764.

Matthy S. Using the edinburgh postnatal depression scale to screen for anxiety disorders. Depression and Anxiety (2008) 25:926–931.

What Screening Tools to Use?

EPDS	Sensitivity = 0.86 Specificity = 0.78 For positive screen >10	5–10 minutes, self-administered, could be self-scored	www.dbpeds.org/articles/detail.cfm?TextID=485
PHQ-2	Sensitivity = 0.83 Specificity = 0.92 For positive screen >3	<1 minute, self-administered or can be asked	www.pfizer.com/pfizer/download/do/phq-9.pdf The two questions from the PHQ-9 for mood and anhedonia are used
PHQ-9	Sensitivity = 0.88 Specificity = 0.88 For positive screen >10	5–10 minutes, self-administered and self-scored	www.pfizer.com/pfizer/download/do/phq-9.pdf

Perinatal depression: Implications for child mental health. Mental Health in Family Medicine 2010:7:239-47

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So, They Are Depressed and/or Anxious and Pregnant...Now What?



Please See End of Slide Set for the Data Regarding The Following:

- SSRIs and risk of spontaneous abortion
- SSRIs and birth outcomes
- Congenital malformations
- Persistent Pulmonary
 Hypertension of the Newborn
 (PPHN)



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Poor Neonatal Adaptation Syndrome (PNAS)

- Occurs in up to 30% of newborns exposed to SSRIs
- Symptoms are transient and usually mild
- Resolve with supportive care in days to weeks
- Unclear if this is due to withdrawal from SSRIs, SSRI toxicity, and/or some combination

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Irritability

Weak cry

Poor tone

Decreased feeding

Jitteriness

Tachypnea

Respiratory distress

Temperature instability

Hypoglycemia

T Pearlstein. Depression During Pregnancy. Best Practice & Research Clinical Obstetrics and Gynecology 29 (2015) 754-764. Becker M, Weinberger T, Chandy A, Schmukler S. Depression During Pregnancy and Postpartum. Curr Psychiatry Rep (2016) 18:32.

Ugh... Can I Have a Digest Version Please?

- There are risks associated with taking SSRIs during pregnancy:
 - May be some small increased risk of miscarriage
 - Increased risk of preterm birth
 - Average decreased gestation of 3 days
 - Increased risk of low birth weight
 - ~2% lower weight
 - There may be an increased risk for PPHN
 - Overall risk still < 1%
 - PNAS symptoms are the most likely, concrete outcome, but these are generally mild and transient

Suggests Medication May Not be



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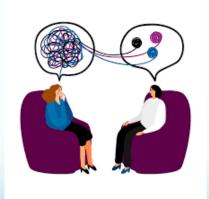
Considerations for Prescribing Medication

Indicated	Should be Strongly Considered
 Mild depression based on clinical assessment No suicidal ideation Engaged in psychotherapy or other non-medication treatment Depression has improved with psychotherapy in the past Able to care for self/baby Strong preference and access to 	 Moderate/severe depression based on clinical assessment Suicidal ideation Difficulty functioning caring for self/baby Psychotic symptoms present History of severe depression and/or suicide ideation/attempts
psychotherapy	 Comorbid anxiety

Suggests Medication Treatment

diagnosis/symptoms

Treatment: Psychotherapy



- First-line treatment for mild to moderate depression
- Indicated for residual symptoms, high risk of relapse
- Encourage in preconception period
- Cognitive behavioral therapy (CBT) or interpersonal psychotherapy(IPT) have been shown to be effective
- Can consider web-based CBT

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Treatment Pearls

- Antidepressant doses may well need to be increased during pregnancy to achieve consistent effect
 - Due to changes in metabolism, GFR, and volume of distribution
 - Use symptoms for guidance
- Be cautious of undertreating illness in the hope of "limiting fetal exposure"
 - Discontinuing medications prior to delivery is not a great idea
- Ensure that initial and ongoing screening includes direct questions on:
 - Suicidal thoughts
 - Thoughts of harming others, including baby
 - Symptoms of psychosis

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T Pearlstein. Depression During Pregnancy. Best Practice & Research Clinical Obstetrics and Gynecology 29 (2015) 754-764.
Thorsness KR, Watson C, LaRusso EM, Perinatal anxiety: approach to diagnosis and management in the obstetric setting. American Journal of Obstetrics & Gynecology. (2018); October: 326-345.

Treatment Pearls

- Screening for patient and familial history of bipolar disorder is very important
 - Average of onset of bipolar disorder (30s) coincides with common childbearing periods
- Patients with bipolar disorder have a 30-40% chance of relapse during the postpartum period
 - This is substantially reduced (~66%) with medication-based treatment
- Patients with a history of postpartum psychosis have a ~30% chance of relapse during subsequent postpartum periods
- Psychiatric consultation is STRONGLY encouraged in these scenarios

Wesseloo R et. al. Risk of Postpartum Relapse in Bipolar Disorder and Postpartum Psychosis: A Systematic Review and Meta-Analysis. Am J Psychiatry (2016) 173;2:117-127.
Hermann A, Gorun A, Benudis, A. Lithium Usa and Non-use for Pregnant and Postpartum Women with Bipolar Disorder. Current Psychiatry Reports (2019) 21:114

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Risks from Benzo Use?

Benzo Use During Pregnancy

- Older data demonstrated a positive association with benzos use and cleft lip and palate
 - Newer, larger, and more well-designed studies have shown no association
- Studies have not consistently demonstrated developmental or cognitive effects in newborns and babies that are present when confounders are controlled
- Higher dosed, consistently dosed benzos in the 3rd trimester can lead to clinically significant withdrawal after delivery

Thorsness KR, Watson C, LaRusso EM. Perinatal anxiety: approach to diagnosis and management in the obstetric setting. American Journal of Obstetrics & Gynecology. (2018); October: 326-345.

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Risk of Untreated Depression and Anxiety?

This Coin Has Two Sides

Risk of Untreated Depression

- Data is much less concrete and difficult to parse from other psychosocial factors and realities
 - Numerous studies have demonstrated significant impact from maternal depression, stress, and anxiety on neonatal development and outcomes
 - Preterm birth
 - Low birthweight
 - Decreased vagal tone and reactivity
 - Altered temperament and increased irritability
 - Altered attention, sleep problems, and delayed neuromotor development

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Risk of Untreated Depression

- Additionally:
 - Maternal depression leads to:
 - Impaired maternal infant bonding
 - Decreased desire to breastfeed
 - Shorter feeding sessions
 - Young children of untreated, depressed mothers are at increased risk for:
 - Externalizing behaviors
 - Fearful temperament and anxiety
 - Delayed motor development
 - Delayed cognitive development

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Risk of Untreated Depression

- Older children of mothers with untreated depression are at increased risk of:
 - ADHD
 - Depression
 - Anxiety disorders
 - Altered stress response

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Stewart DE, Vigod S. Postpartum Depression. N Engl J Med (2016);375;22:2177-2186.

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Risk of Untreated Anxiety?

- Maternal anxiety disorders lead to substantial morbidity including:
 - Sleep disruption
 - Decreased exercise
 - Poor nutrition
 - Substance use
 - Increased use of prenatal services
- Maternal anxiety disorders also impact birth outcomes:
 - Low birth weight
 - Preterm birth
 - Increased risk of preeclampsia

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Risk of Untreated Anxiety?

- Fetal exposure to untreated maternal anxiety has been linked to:
 - Behavioral and emotional problems
 - Attentional disorders
 - Increased risk of psychiatric disorders later in life
 - Decreased gray matter volumes in areas related to learning, memory, and auditory language processing

Thorsness KR, Watson C, LaRusso EM. Perinatal anxiety: approach to diagnosis and management in the obstetric setting. American Journal of Obstetrics & Gynecology. (2018); October: 326-345.

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Putting It All Together



Informed Consent for Antidepressant Use

- No decision regarding whether to use antidepressants during pregnancy is perfect or risk free, including not treating
- SSRIs are among the best studied class of medications during pregnancy
- · Both medication and non-medication options should be considered
- Encourage non-medication treatments (e.g., psychotherapy) in addition to medication treatment or as an alternative when clinically appropriate

Risks of antidepressant use during pregnancy

- Small, but inconsistent increased risk of birth defects when taken in first trimester, particularly with paroxetine
- Possible transient neonatal symptoms
- Low risk for persistent pulmonary hypertension
- Studies do not suggest long-term neurobehavioral effects on children

Risks of under treatment or no treatment of depression during pregnancy

- Association with preterm deliveries
- Increases the risk of postpartum depression
- Can make it harder for moms to take care of themselves and their babies
- Can make it harder for moms to bond with their babies

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So, the Baby Is Delivered...then What?

[·] If pregnant: In your situation, the benefits of taking an antidepressant outweigh the chance of the things we just discussed.

[·] If lactating: SSRIs and other antidepressants are considered a reasonable treatment option during breastfeeding. The benefits of breastfeeding while taking antidepressants generally outweigh the risks.

Na	me			Ad	dre	ss:	
Yo	ur D	Date of Birth:					
Baby's Date of Birth:				Phone:			
As the	you	are pregnant or has	ve recently had a baby, we wo sest to how you have felt IN Th	uld lii	ce to	know how you are feeling. Please check 7 DAYS, not just how you feel today.	
He	re is	an example, alread	ly completed.				
l ha	ave	felt happy:					
		s, all the time					
8	Yes	s, most of the time				most of the time" during the past week.	
		, not very often	Please complete the other qu	uesti	ons	in the same way.	
	No	, not at all					
In t	the p	oast 7 days:					
	Lbe	our been able to laugh	and one the funeu cide of things	*6	This	ngs have been getting on top of me	
		As much as I always	could	0.		Yes, most of the time I haven't been able	
		Not quite so much no	ow			to cope at all	
		Definitely not so muc	th now			Yes, sometimes I haven't been coping as well	
		Not at all				as usual No, most of the time I have coped quite well	
2	Lbi	ave looked forward wit	h enjoyment to things		-	No, I have been coping as well as ever	
-	п	As much as I ever di	d			red, i mare been coping as well as ever	
	п	Rather less than I us	ed to	*7		ave been so unhappy that I have had difficulty sleep	
		Definitely less than I	used to			Yes, most of the time	
		Hardly at all				Yes, sometimes Not very often	
*3.	Lha	ave blamed myself unr	necessarily when things		-	No. not at all	
		nt wrong					
		Yes, most of the time	•	*8		ave felt sad or miserable	
		Yes, some of the tim Not very often	e		0	Yes, most of the time Yes, guite often	
		No. never				Not very often	
					ö	Not very often No, not at all	
4.			orried for no good reason				
	-	No, not at all Hardly ever		•9		eve been so unhappy that I have been crying Yes, most of the time	
		Yes, sometimes			-	Yes, quite often	
		Yes, very often				Only occasionally	
						No, never	
*5			cky for no very good reason				
	8	Yes, quite a lot Yes, sometimes		10	The	e thought of harming myself has occurred to me Yes, quite often	
		No. not much			H	Sometimes	
		No, not at all				Hardly ever	
						Never	
Adr	minis	tered/Reviewed by		Date			
¹So	urce	Cox, J.L., Holden, J.M., a	and Sagovsky, R. 1987. Detection of a Scale. British Journal of Psyci	postn	atal c	depression: Development of the 10-item	
			C. M. Piontek, Postpartum Depressi	,			
194	-199						

- QUESTIONS 1, 2, & 4 (with out an*) → scored 0, 1, 2 or 3 with top box scored as 0 and the bottom box scored as 3.
- QUESTIONS 3, 5-10 (marked with an*) → reverse scored, with the top box scored as a 3 and the bottom box scored as
 - Maximum score: 30
- Possible Depression: score of 10 or greater
- Items 3,4,&5 investigate anxiety symptoms
 - Score 6 or more on these items screens + for anxiety

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Prevalence

- Postpartum psychosis (0.1-0.2%)
- Postpartum blues (50-85%)
- Postpartum depression (10-15%)



Postpartum Blues

- Onset 3-5 days after delivery
- Sx: weepiness, sadness, fatigue, irritability
- Tx: partner, community support
- Should resolve in 10 days. If not consider Peripartum depression



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Postpartum Depression

- Usually develops slowly over first 3 months
- Sx- MDD sx
- Often unable to sleep
- Often see anxiety
- Up to 50% will have intrusive obsessional ruminations/images usually focused on baby. Often violent in nature but egodystonic and no problem with reality testing.

Postpartum Depression

- Studies vary, but the prevalence of postpartum depression is roughly 15-20%
- Distinctly different from "baby blues"
 - Mild symptoms that peak between 2-5 days post delivery and do not meet criteria for a MDE and are not severe enough to impair function
- Exact pathophysiology is not known, but suspected to be a combination of:
 - Rapid decline of hormone levels post delivery
 - Genetic factors
 - Social factors

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Zhao XH, Zhang ZH. Risk factors for postpartum depression: An evidence-based systemic review of systematic reviews and meta-analyses. Asian Journal of Psychiatry 53 (2020) 102353.
Pinheiro, E. et. al. Sertraline and breastfeeding: review and meta-analysis. Arch Womens Ment Health (2015) 18:139-146.

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Postpartum Depression

- Risk factors:
 - Hx of depression***
 - Women with untreated depression during pregnancy have a 7 times greater likelihood of developing postpartum depression
 - AND...

Violence and Abuse	Low birthweight infant	Immigration status
Gestational diabetes	Vit D deficiency	Obesity
Sleep disruption	Lack of social supports	Multiple births
Postpartum anemia	Negative birth experience	

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Zhao XH, Zhang ZH. Risk factors for postpartum depression: An evidence-based systemic review of systematic reviews and meta-analyses. Asian Journal of Psychiatry 53 (2020) 102353.

Postpartum Depression

- Preventative/Protective Factors:
 - Treatment of peripartum depression
 - In-home supportive care programs
 - Skin-to-skin contact and care
 - Healthy diet
 - Adequate sleep*
 - Adequate exercise*
 - *- Limited data for outcomes

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Zhao XH, Zhang ZH. Risk factors for postpartum depression: An evidence-based systemic review of systematic reviews and meta-analyses. Asian Journal of Psychiatry 53 (2020) 102353.

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Postpartum Depression

- Treatment:
 - Mild to moderate symptoms:
 - Psychotherapy often recommended as the treatment of choice
 - SSRIs are considered first line medication-based treatments
 - Severe symptoms:
 - Medication based treatment is encouraged
 - Consultation with psychiatric care if possible

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Zhao XH, Zhang ZH. Risk factors for postpartum depression: An evidence-based systemic review of systematic reviews and meta-analyses. Asian Journal of Psychiatry 53 (2020) 10:2353.
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Psychotropics and Breastfeeding

- Most antidepressants are transmitted in breastmilk at levels that are 10% or lower than the maternal dose
 - This is 5-10 X lower than gestational exposure
- Sertraline has substantial data demonstrating that it has some of the lowest transmission rates (often undetectable in milk)
 - · Likely due to the high level of protein binding
- The impact of low dose SSRIs in breastmilk on newborns has limited data
 - Case reports are conflicting, though most common findings are irritability, restlessness, and/or colic

Becker M, Weinberger T, Chandy A, Schmukler S. Depression During Pregnancy and Postpartum. Curr Psychiatry Rep (2016) 18:32.
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Psychotropics and Breastfeeding

- TCA use is generally discouraged while breastfeeding due to cardiac side effects
- Cases of newborn seizures in mothers who were breastfeeding while taking bupropion have been reported
- Expert consensus recommends that individual decisions be made based on relative risk ratios, but SSRI use is generally considered to be safe while breastfeeding and that the risks of discontinuation are more concerning

Becker M, Weinberger T, Chandy A, Schmukler S. Depression During Pregnancy and Postpartum. Curr Psychiatry Rep (2016) 18:32.

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Psychotropics and Breastfeeding

- Benzodiazepines are also transmitted in breastmilk
 - Effective doses are low, but can lead to infant sedation
 - Occurs in < 2% of cases
 - Case reports of
 - Apnea
 - Irritability
 - Poor weight gain
- Lithium passes into breastmilk in substantial doses
 - Breastfeeding on lithium should only be attempted with close, expert consultation and support

Becker M, Weinberger T, Chandy A, Schmukler S. Depression During Pregnancy and Postpartum. Curr Psychiatry Rep (2016) 18:32. Stewart DE, Vigod S. Postpartum Depression. N Engl J Med (2016);375;22:2177-2186. Thorsness KR, Watson C, LaRusso EM, Perinatal anxiety: approach to diagnosis and management in the obstetric setting. American Journal of Obstetrics & Gynecology. (2018); October: 326-345.

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Informed Consent for Antidepressants when Breastfeeding

 SSRIs and other antidepressants are considered a reasonable treatment option during breastfeeding. The benefits of breastfeeding while taking antidepressants generally outweigh the risks.



Resources

- Center for Women's Mental Health at MGH
 - https://womensmentalhealth.org/
 - This is my go to resource for thoughtful commentary on new studies as well as great provider and patient resources to make informed decisions about mental health treatment, especially during pregnancy and breastfeeding.
- LactMed: A TOXNET DATABASE
 - https://toxnet.nlm.nih.gov/newtoxnet/lactmed.htm
 - This database allows you to look up any medication and get thoughtful guidance about the use in breastfeeding mother.
- MCPAP For Moms
 - https://www.mcpapformoms.org/Toolkits/Toolkit.aspx
 - This is a great toolkit with information and decision aids to support assessing and treating perinatal depression.

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A Little Further Down the Road...

Perimenopause and Menopause



Depression in Women

- Overall, woman experience depression roughly twice as frequently as men
 - This increased is generally confined to reproductive years
- The perimenopausal period is one of particularly vulnerable time with 2-5X û risk for depression
- Increased risk is predominantly seen in women with a history of MDD

Bromberger JT, Epperson, CN. Depression During and After the Perimenopause. Obstet Gynecol Clin N Am. 2018. 45:663-678. Maki PM et. al. Guidelines for the Evaluation and Treatment of Perimenopausal Depression: Summary and Recommendations. J of Women's Health. 2019. 28:2:117-134.

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Can We Articulate Depression Risk in Terms of Hormonal Changes?

- ... Not really
- Studies vary in their association of specific sex hormones with specific depression outcomes throughout the menopausal transition
 - This is complicated by the complex relationship between sex hormones, neurotransmitters, and dx
- There appear to be more consistent associations with overall change in estradiol levels rather than absolute levels
 - More about the change and lack of stability than anything else

Bromberger JT, Epperson, CN. Depression During and After the Perimenopause. Obstet Gynecol Clin N Am. 2018. 45:663-678. Maki PM et. al. Guidelines for the Evaluation and Treatment of Perimenopausal Depression: Summary and Recommendations. J of Women's Health. 2019. 28;2:117-134.

Menopause and Mood Disorders: Risk Factors

- Lower income
- Lower education
- Younger age
- Unpartnered
- Poor social supports
- Childhood and lifetime adversity

- Smoking
- Physical inactivity
- Sleep disturbance
- Vasomotor symptoms
- h/o depression
- h/o PMS
- Anxiety

Maki PM et. al. Guidelines for the Evaluation and Treatment of Perimenopausal Depression: Summary and Recommendations. J of Women's Health. 2019. 28;2:117 134.

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Treatments During Perimenopause



- Antidepressants
- Psychotherapy
- Exercise
- Estrogen

Which Antidepressant?

Consider...

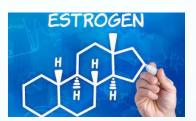
- Prior antidepressant trials and responses
- Adverse side effects
- Safety (drug-drug interactions)
- SSRIs and SNRIs at typical doses
- Added benefit of improving menopause-related sx such as hot flashes and pain



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Does Estrogen Treat Depression?

 Several small, double blinded studies have demonstrated estrogen therapy to be effective in improving depressive symptom rating scales in perimenopause



Bromberger JT, Epperson, CN. Depression During and After the Perimenopause. Obstet Gynecol Clin N Am. 2018. 45:663-678. Maki PM et. al. Guidelines for the Evaluation and Treatment of Perimenopausal Depression: Summary and Recommendations. J of Women's Health. 2019. 28;2:117-134.

Does Estrogen Treat Depression?

- Data indicates estrogen may be an augmentation agent for antidepressant treatment in midlife and older women but should be considered with caution
- Estrogen monotherapy is ineffective in treating symptoms of depression in postmenopausal women
- Estrogen is NOT an FDA approved treatment for depression

Bromberger JT, Epperson, CN. Depression During and After the Perimenopause. Obstet Gynecol Clin N Am. 2018. 45:663-678. Maki PM et. al. Guidelines for the Evaluation and Treatment of Perimenopausal Depression: Summary and Recommendations. J of Women's Health. 2019. 28:2:117-134.

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Key Take Home Points

- Perinatal depression is common and often undetected and untreated!
- Screening tools really help with identification
- Treatment is critical for both baby and mom
- The perimenopausal period is a high-risk time for depression so look for it and treat it!
- Risk vs benefit assessment requires multifactorial consideration
- A collaborative approach is key



SSRIs and Risk of Spontaneous Abortion

- Kjaersgaard et al. 2013: Population based study of pregnancies in Denmark reported 12% rate of spontaneous abortion with exposure to antidepressants
 - RR 1.14 (95% CI 1.10-1.18)
 - RR lowered to 1.0 when compared to spontaneous abortion rates in women with depression who did not take an antidepressant
- Andersen et al. 2014: Large study of pregnancies in Denmark
 - Adjusted hazard ratio of spontaneous abortion with exposure to an SSRI of 1.27 (95% CI 1.22-1.33)
 - Adjusted hazard ratio of spontaneous abortion with discontinuation of an SSRI one year prior to pregnancy of 1.24 (95% CI 1.18-1.30)

T Pearlstein. Depression During Pregnancy. Best Practice & Research Clinical Obstetrics and Gynecology 29 (2015) 754-764. Kjaersgaard MIS, Parner ET, Vestergaard M. et al. Prenatal antidepressant exposure and risk of spontaneous abortion – a population-based study. PLoS One 2013;8:e72095.
Andersen JT, Andersen NL, Howitz H, et al. Exposure to selective serotonin reuptake inhibitors in early pregnancy and the risk of miscarriage. Obstet Gynecol 2014; 124:655-61.

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SSRIs and Birth Outcomes

- Numerous studies have examined outcomes related to preterm birth and low birth weight
 - 2013 systemic review and metanalysis found:
 - Pooled OR 1.55 (95% 1.38-1.74) of preterm birth in mothers taking antidepressants
 - Averaged 3 days shorter gestational age
 - Same analysis found a mean birthweight difference of -74 grams in newborns exposed to SSRIs during gestation (95% CI -117 to -31)
 - ~2% birthweight of a 3.5 kg newborn

T Pearlstein. Depression During Pregnancy. Best Practice & Research Clinical Obstetrics and Gynecology 29 (2015) 754-764. Ross LE, Grigoriadis S. Mamisashvili L, et. al. Selected pregnancy and delivery outcomes after exposure to antidepressant medication: a systemic review and meta-analysis. JAMA Psychiatry 2013;70:436-43.

Congenital Malformations

- Studies have varied somewhat in findings
 - Huybrechts KF et. al. 2014 NEJM found no substantial increase in the risk of cardiac malformation with first-trimester exposure to antidepressants
 - Other, contemporary studies have found an increased risk of malformations with SSRIs, particularly paroxetine, though the clinical significance of findings has been inconsistent
 - Reports of increased risk of malformations usually report OR of 1.2 to 2.0

T Pearlstein. Depression During Pregnancy. Best Practice & Research Clinical Obstetrics and Gynecology 29 (2015) 754-764. Becker M, Weinberger T, Chandy A, Schmukler S. Depression During Pregnancy and Postpartum. Curr Psychiatry Rep (2016) 18-32

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Persistent Pulmonary Hypertension of the Newborn (PPHN)

- PPHN occurs in 1.9 infants per 1000 live births
 - Range of complications and outcomes, but mortality can be 10-20%
- Studies have been somewhat inconclusive in that some demonstrate and association with SSRI use in late pregnancy and others do not
 - Caused the FDA to revise their 2006 warning in 2011
- Studies that do show an association demonstrate an absolute risk of 2.9-3.5/1000 with SSRI use in late pregnancy

T Pearlstein. Depression During Pregnancy. Best Practice & Research Clinical Obstetrics and Gynecology 29 (2015) 754-764 Becker M, Weinberger T, Chandy A, Schmukler S. Depression During Pregnancy and Postpartum. Curr Psychiatry Rep (2016) 18-32