



# POSTPARTUM DEPRESSION

FLAME LECTURE: 134

TOOHEY / BURNS 5.6.15

# Learning Objectives



- ▶ Identify risk factors for postpartum blues and depression
- ▶ Differentiate between postpartum blues and depression
- ▶ Describe treatment options for postpartum blues and depression
- ▶ Recognize appropriate treatment options for mood disorders during lactation
- ▶ Prerequisites:
  - ▶ NONE
- ▶ See also – for closely related topics
  - ▶ FLAME LECTURE 39 – Depression in Pregnancy
  - ▶ FLAME LECTURE 235 – Depression in Women
  - ▶ FLAME LECTURE 135 – Postpartum Psychosis

# Postpartum Psychiatric Disease

- ▶ **Postpartum Blues**
  - ▶ Accounts for 60-80% of patient's with depressive symptoms
    - ▶ Self limiting, no medication required (typically regress within 2 weeks)
    - ▶ Supportive care
    - ▶ 20% will progress to major depression
- ▶ **Postpartum Depression (formally called MDD with peripartum onset)**
  - ▶ The single most common complication of pregnancy and post partum!
    - ▶ 10-15% of postpartum women
  - ▶ DSM V criteria
    - ▶ Symptoms present for most of the day almost every day for > 2 weeks which causes clinically significant distress or impairment in social, occupation, or other areas of functioning
    - ▶ Depressed mood, loss of interest or pleasure AND 4 of the following:
      - ▶ Changes in appetite/weight, sleep and psychomotor activity, decreased energy, feelings of worthlessness/guilt, difficulty thinking, concentrating, or making decisions, recurrent thoughts of death or suicide, plans or attempts.
- ▶ **Post Partum Psychosis (FLAME LECTURE 135)**
  - ▶ Most commonly, an untreated bipolar disorder

# Risk Factors<sup>1,2</sup>

## MAJOR RISK FACTORS:

- ▶ History of prior psychiatric or psychological problems
  - ▶ The strongest predictor of postpartum depression is the presence of depression and anxiety during pregnancy
- ▶ History of postpartum depression
  - ▶ Recurrence risk of 30%
- ▶ Life events
  - ▶ Death of a loved one, divorce, loss of job, moving
- ▶ Social support
  - ▶ Informational, instructional, and emotional
  - ▶ History of physical or sexual abuse, or current IPV
- ▶ Teenagers – incidence of PP Depression can be as high as 26%

## MODERATE RISK FACTORS

- ▶ Personality characteristics- nervous, shy, self conscious, low self esteem, pessimistic, poor coping skills
- ▶ Marital problems
- ▶ Family History of Depression

## MINOR RISK FACTORS

- ▶ Obstetric Factors-pregnancy related complications, prematurity, hemorrhage

### Thyroid Hormones & Depression risk:

- Conflicting studies regarding relationship
- 4-10% of postpartum pts have transitory thyroid dysfunction
  - Most of these women do NOT have depression & most depressed women do NOT have thyroid dysfunction
- However, Hypo/Hyperthyroid states can mimic depression, mania, & psychosis
  - In individual cases it is reasonable to screen for thyroid function

# Screening – *Edinburgh Postnatal Depression Scale (EPDS)*

In the past 7 days:

1. I have been able to laugh and see the funny side of things	5. I have felt scared or panicky for no very good reason	*9. I have been so unhappy that I have been crying
<input type="checkbox"/> As much as I always could <input type="checkbox"/> Not quite so much now <input type="checkbox"/> Definitely not so much now <input type="checkbox"/> Not at all	<input type="checkbox"/> Yes, quite a lot <input type="checkbox"/> Yes, sometimes <input type="checkbox"/> No, not much <input type="checkbox"/> No, not at all	<input type="checkbox"/> Yes, most of the time <input type="checkbox"/> Yes, quite often <input type="checkbox"/> Only occasionally <input type="checkbox"/> No, never
2. I have looked forward with enjoyment to things	*6. Things have been getting on top of me	*10 The thought of harming myself has occurred to me
<input type="checkbox"/> As much as I ever did <input type="checkbox"/> Rather less than I used to <input type="checkbox"/> Definitely less than I used to <input type="checkbox"/> Hardly at all	<input type="checkbox"/> Yes, most of the time I haven't been able to cope at all <input type="checkbox"/> Yes, sometimes I haven't been coping as well as usual <input type="checkbox"/> No, most of the time I have coped quite well <input type="checkbox"/> No, I have been coping as well as ever	<input type="checkbox"/> Yes, quite often <input type="checkbox"/> Sometimes <input type="checkbox"/> Hardly ever <input type="checkbox"/> Never
*3. I have blamed myself unnecessarily when things went wrong	*7. I have been so unhappy that I have had difficulty sleeping	Scoring: QUESTIONS 1, 2, & 4 (without an *) Are 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 *) Are reverse scored, with the top box scored as a 3 and the bottom box scored as 0.  Maximum score: 30 <b>Possible Depression: 10 or greater</b> Always look at <u>item 10</u> (suicidal thoughts)
<input type="checkbox"/> Yes, most of the time <input type="checkbox"/> Yes, some of the time <input type="checkbox"/> Not very often <input type="checkbox"/> No, never	<input type="checkbox"/> Yes, most of the time <input type="checkbox"/> Yes, sometimes <input type="checkbox"/> Not very often <input type="checkbox"/> No, not at all	
4. I have been anxious or worried for no good reason	*8. I have felt sad or miserable	
<input type="checkbox"/> No, not at all <input type="checkbox"/> Hardly ever <input type="checkbox"/> Yes, sometimes <input type="checkbox"/> Yes, very often	<input type="checkbox"/> Yes, most of the time <input type="checkbox"/> Yes, quite often <input type="checkbox"/> Not very often <input type="checkbox"/> No, not at all	

# Treatment<sup>1</sup>

## ▶ Medications:

- ▶ Tricyclic Antidepressants
- ▶ Monoamine Oxidase Inhibitors
- ▶ Serotonin Reuptake Inhibitors

## ▶ Interpersonal Psychotherapy:

- ▶ Currently thought to be one of the most effective models for PPD given the importance of interpersonal relationships and role transitions in the peripartum period
- ▶ Based on the premise that there are 4 problem areas
  - ▶ Grief, Role transitions, Interpersonal disputes, Interpersonal deficits

## ▶ Support Groups

## ▶ ECT (Electroconvulsive Therapy)

- ▶ Safety has been demonstrated for many years in pregnancy

## ▶ Consider hospitalization

## ▶ Duration of treatment:

- ▶ Treat for at least 6-12 months
- ▶ Continue maintenance therapy if:
  - ▶ Pt has a history of 3 or more episodes of major depression
  - ▶ Previous episodes have been severe or prolonged
  - ▶ There is a coexisting anxiety disorder
  - ▶ Depression has an early or late onset

## ▶ Remember to counsel regarding future fertility plans

# Treatment - *medications*

- ▶ Physicians often hesitate to prescribe psychiatric medications during pregnancy & lactation because of unknown effects of the medication on the fetus/infant
- ▶ But stopping or avoiding treatment itself poses serious risk!!!
  - ▶ 68% of pregnant women who stop meds during or just prior to pregnancy relapse<sup>3</sup>
    - ▶ Usually during the 1<sup>st</sup> trimester
    - ▶ Rate is higher than non-pregnant women
  - ▶ Depression itself carries a higher risk for abnormalities in neonatal development
    - ▶ **Preterm delivery**: Children of mothers with MDD born at 35 weeks vs. 39.4 weeks<sup>4</sup>
    - ▶ ↓ motor maturity<sup>4</sup>, ↓ head circumference<sup>5</sup>, ↓ Apgars<sup>5</sup>, ↑ Maternal self-harm<sup>5</sup>
    - ▶ Children of mothers with post-partum depression suffer from lower cognitive and language function
  - ▶ Increased stress and depression *during* pregnancy cause hormonal changes in fetus
    - ▶ Increased cortisol<sup>4</sup>, catecholamines, alter uterine blood flow
    - ▶ Associated with neuronal cell death and abnormal development of brain structures as well as sustained HPA dysfunction in the neonate
  - ▶ Anti-depressants typically take 2-6 weeks to become effective

# Treatment - *medications*



## Tricyclic Antidepressants

- Imipramine, nortryptiline, amitriptyline
- Are relatively non selective
- Until recently were 1<sup>st</sup> line
- Have many side effects due to affinity to histamine, muscarinic &  $\alpha$ -adrenergic receptors
- High overdose toxicity
- No increased risk for birth defects
- Good for women with frequent headaches
- Some neonatal withdrawal

## Monamine Oxidase Inhibitors

- Parnate, Nardil, and Marplan
- Require restricting tyramine containing foods
  - No ETOH, aged cheese, liver, orange pulp, smoked fish, packaged soups amongst many other food limitations
  - If not followed, can result in hypertensive crisis OR intracranial bleed
- Dangerous to use with a TCA
- High overdose toxicity
- Concern about ↓ uterine blood flow
- Possible ↑ risk for birth defects
- **AVOID IN PREGNANCY**

## Serotonin Reuptake Inhibitors

- SSRI's are not a major risk factor for infant malformations
  - Only paroxetine found to have increased risk for cardiac malformations
- In 2006 Public Health Advisory warning released regarding SSRI's increasing risk for Persistent Pulmonary HTN, but since then there have been conflicting studies and FDA advises not to change practice in prescribing SRI's

## Neonatal Withdrawal Syndrome<sup>6</sup>:

- Most often associated with TCA's, most commonly after 3<sup>rd</sup> trimester exposure
- Symptoms: neonatal tachycardia, cyanosis, tachypnea, clonus, irritability, feeding difficulties, temperature instability
- Usually self-limited, encourage breastfeeding

# Treatment – Medications<sup>6</sup>

Antidepressant Medications			
Generic Name	Brand Name	Pregnancy Risk Category	Lactation Risk Category
<i>TCA's</i>			
Amitriptyline	Elavil, Endep	C	L2
Amoxapine	Asendin	C	L2
Clomipramine	Anafranil	C	L2
Desipramine	Norpramin	C	L2
Doxepin	Sinequan/Adaptin	C	L5
Imipramine	Tofranil	C	L2
Maprotiline	Ludiomil	B	L3
Nortriptyline	Pamelor/Aventyl	C	L2
Protriptyline	Vivactil	C	N/A
<i>SSRI's</i>			
Citalopram	Celexa	C	L3
Escitalopram	Lexapro	C	L3*
Fluoxetine	Prozac	C	L2*
Fluvoxamine	Luvox	C	L2
Paroxetine	Paxil	D	L2
Sertraline	Zoloft	C	L2

**Remember, physiologic changes during pregnancy affect pharmacology:**

- 50% increase in blood volume by 24-26 weeks
- Glomerular filtration rate increases by 50% in the second trimester
- Therefore, adjustments in doses of medications during pregnancy are often necessary

## Treatment – *during lactation*

- ▶ Tricyclics or metabolites have not been detectable in breast milk
- ▶ Fluoxetine - low levels detected
- ▶ Sertraline - no levels detected, no data on metabolites
- ▶ Bupropion - no levels detected
- ▶ Dose infant is exposed to is much less than during pregnancy
  - ▶ An infant would need to breastfeed exclusively for 2 years to accrue the level of exposure during one month of pregnancy
  - ▶ Infants have comparable level of metabolic enzymes with adults
- ▶ Metoclopramide: often prescribed to ↑ milk production but has a risk of exacerbating depression in patients with a history of mental illness

# IMPORTANT LINKS

- ▶ [ACOG Practice Bulletin 92](#), April 2008 (“Use of Psychiatric Medications during Pregnancy and Lactation”)
  1. UpToDate.com
  2. Robertson, et al, *Gen Hospital Psychiatry*, 2004
  3. Cohen et al, *JAMA*, 2006
  4. Marcus, et al, *Psychiatry*, 2006
  5. Kurki, et al, *ObGyn*, 2000