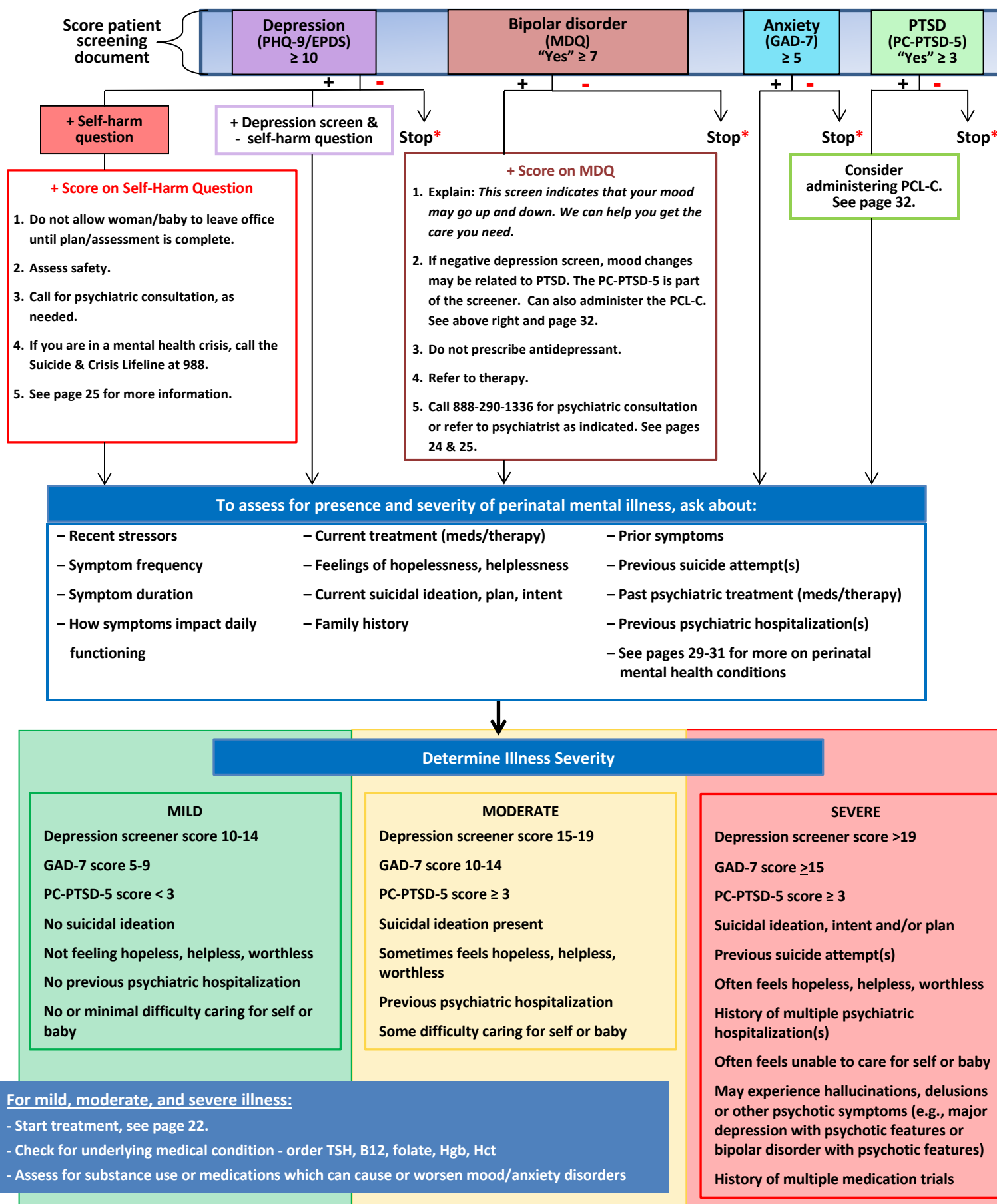


## Assessing Perinatal Mental Health



\*If all screens are negative, tell her they were negative and say, "if something changes, please let us know. We are here."

Continue to other side →

EPDS – Edinburgh Postnatal Depression Scale; GAD – Generalized Anxiety Disorder; MDQ – Mood Disorder Questionnaire; PHQ – Patient Health Questionnaire  
PTSD – Posttraumatic Stress Disorder; PC-PTSD-5 – Primary Care Post Traumatic Stress Disorder; PCL-C – PTSD Check List-Civilian

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Authors: Byatt N., Mittal L., Brenckle L., Logan D., Masters G., Bergman A., Moore Simas T.

### Consider treatment options based on highest level of illness severity

If severity of symptoms overlap, clinical decisions should be based on the assessment, with strong consideration of higher level treatment options.

| MILD   | MODERATE  | SEVERE  |
|--|---|---|
| <p>Therapy referral</p> <p>Consider medication treatment</p> | <p>Therapy referral</p> <p>Strongly consider medication treatment</p> <p>If onset of depression symptoms occurs in 3rd trimester to 4 weeks postpartum and if the patient is &lt;6 months postpartum at screening, consider postpartum brexanolone (IV allopregnanolone infusion over 60 hours in an inpatient setting). See page 20.</p> | <p>Therapy referral</p> <p>Medication treatment</p> <p>If onset of depression symptoms occurs in 3rd trimester to 4 weeks postpartum and if the patient is &lt;6 months postpartum at screening, consider postpartum brexanolone (IV allopregnanolone infusion over 60 hours in an inpatient setting). See page 20.</p> |

- If you are in a mental health crisis, call the Suicide & Crisis Lifeline at 988. If you are experiencing a medical emergency, call 911.
- Call the Arizona Perinatal Psychiatry Access Line at 888-290-1336
- Visit the APAL website at [APAL.arizona.edu](http://APAL.arizona.edu) or [APAL.arizona.edu/moms-families](http://APAL.arizona.edu/moms-families) to find a provider in Arizona.

### Therapy and support options

- All women who screen positive, regardless of illness severity, should be referred to therapy or be advised to continue therapy
- Always discuss and encourage prevention and support options (e.g., peer and social supports and groups, sleep hygiene, self-care, and exercise). See page 27.

### How to educate patients about treatment with antidepressants

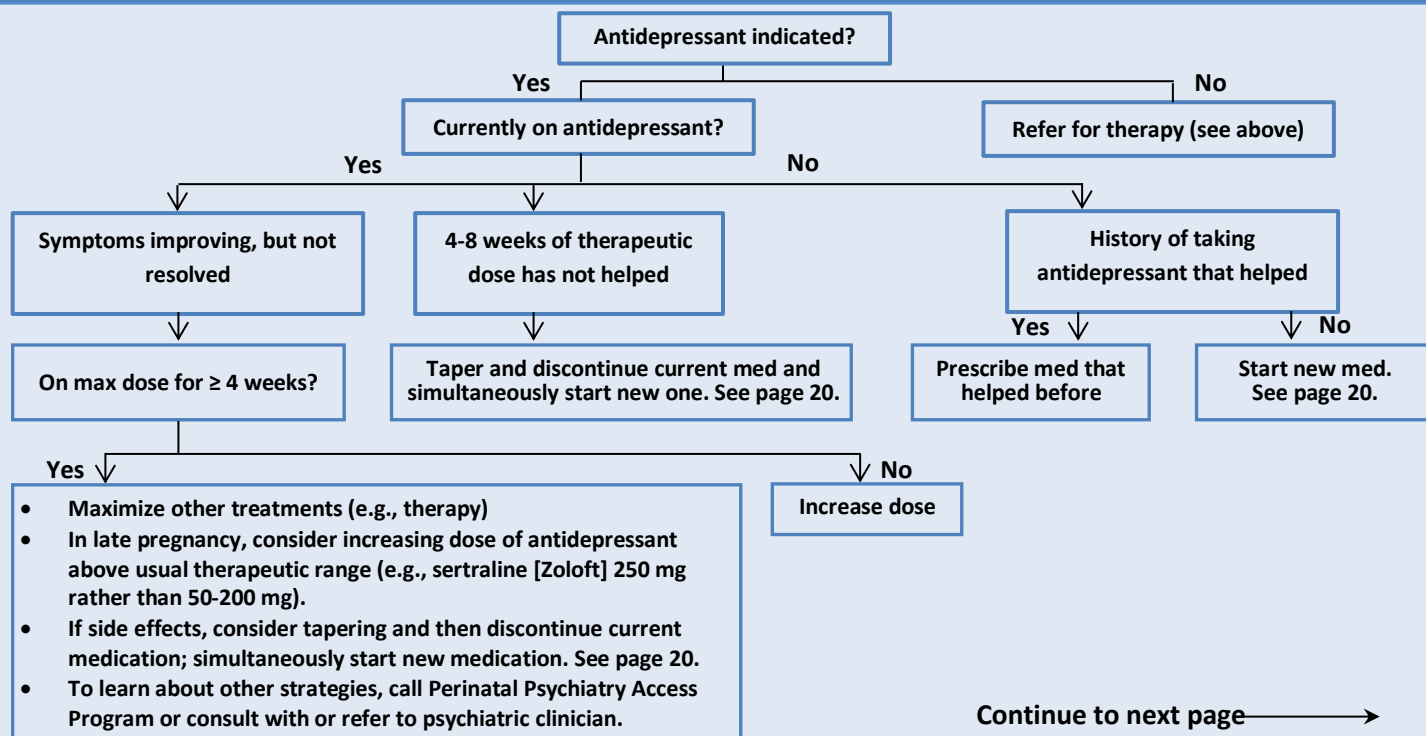
#### Antidepressant use during pregnancy:

- Does not appear to be linked with birth complications
- Has been linked with small but inconsistent risk of birth defects when taken in the first trimester, particularly paroxetine
- Has been linked with transient (days to weeks) neonatal symptoms (tachypnea, irritability, insomnia)
- Has inconsistent, overall reassuring, evidence regarding long-term (months to years) neurobehavioral effects on children

#### Under-treatment or no treatment of perinatal mental health conditions:

- Has been linked with birth complications
- Can increase the risk or severity 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
- Can increase risk of mental illness among offspring
- Has been linked with possible long-term neurobehavioral effects on children

### Medication treatment (when indicated)



Continue to next page →

## Pharmacological Treatment Options for Depression, Anxiety, and PTSD

- Choose antidepressant that has worked before. If antidepressant naïve, choose antidepressant based on table below with patient preference in consideration. Antidepressants are similar in efficacy and side effect profile.
- In late pregnancy, you may need to increase the dose above usual therapeutic range (e.g., sertraline [Zoloft] 250mg rather than 50-200mg).
- If a patient presents with pre-existing mood and/or anxiety disorder and is doing well on an antidepressant, do not switch it during pregnancy or lactation. If patient is not doing well, see page 21.
- Evidence does not support tapering antidepressants in the third trimester.
- Minimize exposure to both illness and medication.
  - Untreated/inadequately treated illness is an exposure
  - Use lowest effective doses with goal of remission of symptoms.
  - Minimize switching of medications
  - Monotherapy preferred, when possible

See page 19 for how to educate patients about treatment with antidepressants

## First-line Treatment Options for Mild, Moderate, or Severe Depression, Anxiety Disorder, and PTSD

| Medication  | sertraline*<br>(Zoloft)  | fluoxetine<br>(Prozac) | citalopram**<br>(Celexa) | escitalopram**<br>(Lexapro) |
|---|--|------------------------|--------------------------|-----------------------------|
| Starting dose and timing                                      | 25 mg<br>qAM (if sedating,<br>change to qHS)   | 10 mg<br>qAM           | 10 mg<br>qAM             | 5 mg<br>qAM                 |
| Initial increase after 4 days                                 | ↑ to 50 mg   | ↑ to 20 mg             | ↑ to 20 mg               | ↑ to 10 mg                  |
| Second increase after 7 more days                             | ↑ to 100 mg  |                        |                          |                             |
| Reassess Monthly (increase as needed<br>until symptoms remit) | ↑ by 50 mg   | ↑ by 20 mg             | ↑ by 10 mg               | ↑ by 10 mg up to 20 mg      |
| Therapeutic range***  | 50-200 mg  | 20-80 mg               | 20-40 mg                 | 10-20 mg                    |
| Individualized approach to titration                          | Slower titration (e.g., every 10-14-days) is often needed for patients who are antidepressant naïve or with anxiety symptoms |                        |                          |                             |

\*Lowest degree of passage into breast milk compared to other first-line antidepressants; \*\*Side effects include QTc prolongation (see below);

\*\*\*May need higher dose in 3<sup>rd</sup> trimester and when treating an anxiety disorder

In general, if an antidepressant has helped during pregnancy, it is best to continue it during lactation.

Prescribe a maximum of two (2) antidepressants at the same time.

## Second-line Treatment Options for Mild, Moderate, or Severe Depression, Anxiety Disorder, and PTSD

| Medication  | duloxetine<br>(Cymbalta)   | venlafaxine<br>(Effexor XR) | fluvoxamine<br>(Luvox) | paroxetine<br>(Paxil)                           | mirtazapine<br>(Remeron) | bupropion HCL<br>(Wellbutrin XL) |
|---|--|-----------------------------|------------------------|---|--------------------------|----------------------------------|
| Starting dose and timing                                      | 30 mg***<br>qAM  | 37.5 mg<br>qAM              | 25 mg<br>qHS           | 10 mg***<br>qAM (if sedating,<br>change to qHS) | 7.5 mg<br>qHS            | 150 mg<br>qAM                    |
| Initial increase after 4 days                                 |  | ↑ to 75 mg                  | ↑ to 50 mg             | ↑ to 20 mg                                      | ↑ to 15 mg               |                                  |
| Second increase after 7 more days                             | ↑ to 60 mg   |                             | ↑ to 100 mg            |   |                          |                                  |
| Reassess Monthly (increase as<br>needed until symptoms remit) | ↑ by 30 mg   | ↑ by 75 mg                  | ↑ by 50 mg             | ↑ by 10 mg                                      | ↑ by 15 mg               | ↑ by 150 mg                      |
| Therapeutic range ***   | 30-120 mg  | 75-300 mg                   | 50-200 mg              | 20-60 mg  | 15-45 mg                 | 300-450 mg                       |
| Individualized approach to<br>titration                       | Slower titration (e.g., every 10-14-days) is often needed for patients who are antidepressant naïve or with anxiety symptoms |                             |                        |   |                          |                                  |

\*\*\*May need higher dose in 3<sup>rd</sup> trimester and when treating an anxiety disorder

### Temporary (days to weeks)

Nausea (most common)

Constipation/diarrhea

Lightheadedness

Headaches

### Long-term (weeks to months)

Increased appetite/weight gain

Sexual side effects

Vivid dreams/insomnia

\*\*QTc prolongation (citalopram & escitalopram)

- Tell women to take medication with food and only increase dose if tolerating; otherwise wait until side effects dissipate before increasing.
- Start medication in morning; if patient finds it sedating recommend that she takes it at bedtime

## Medication Treatment for Moderate/Severe Depression with Onset in Late Pregnancy or Within 4 weeks postpartum – brexanolone (Zulresso)

Brexanolone is an FDA-approved medication that can be considered for treatment of moderate to severe postpartum depression.

Brexanolone:

- is a formulation of intravenous allopregnanolone (a neurosteroid) that acts on GABA-A receptors
- requires an IV infusion over 60 hours
- has a faster onset of action (symptom reduction in 1-2 days) compared to available oral antidepressants, which generally take 4-8 weeks to work
- has been shown to maintain the reduction in depression symptoms at 30 days post-infusion

When is Brexanolone indicated?

If onset of depression occurs in 3<sup>rd</sup> trimester through 4 weeks postpartum and if patient is <6 months postpartum at screening, consider Brexanolone (IV allopregnanolone infusion over 60 hours in an inpatient setting).

More information can be found at Reprotox and LactMed on all pharmacological treatments