



Spring CME Event
May 25, 2022

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Referrals and Consults

- Anyone is welcome to staff message me with questions
- If you practice in a BHIP clinic or VFM, you can request that I directly evaluate a patient for a psychiatric consult
- I do NOT have a practice at Valley's Psychiatry & Counseling clinic, but I have 7 awesome colleagues to whom you can refer patients
 - Commercial insurance and older Medicare accepted; REF91



Referrals and Consults

- Other resources: UW Psychiatry Provider Consult Lines
 - Adults: 877-WA-PSYCH (877-927-7924)
 - Pediatric: 866-599-7257 (Seattle Children's)
 - Perinatal: 877-725-4666



Optimizing Depression Management

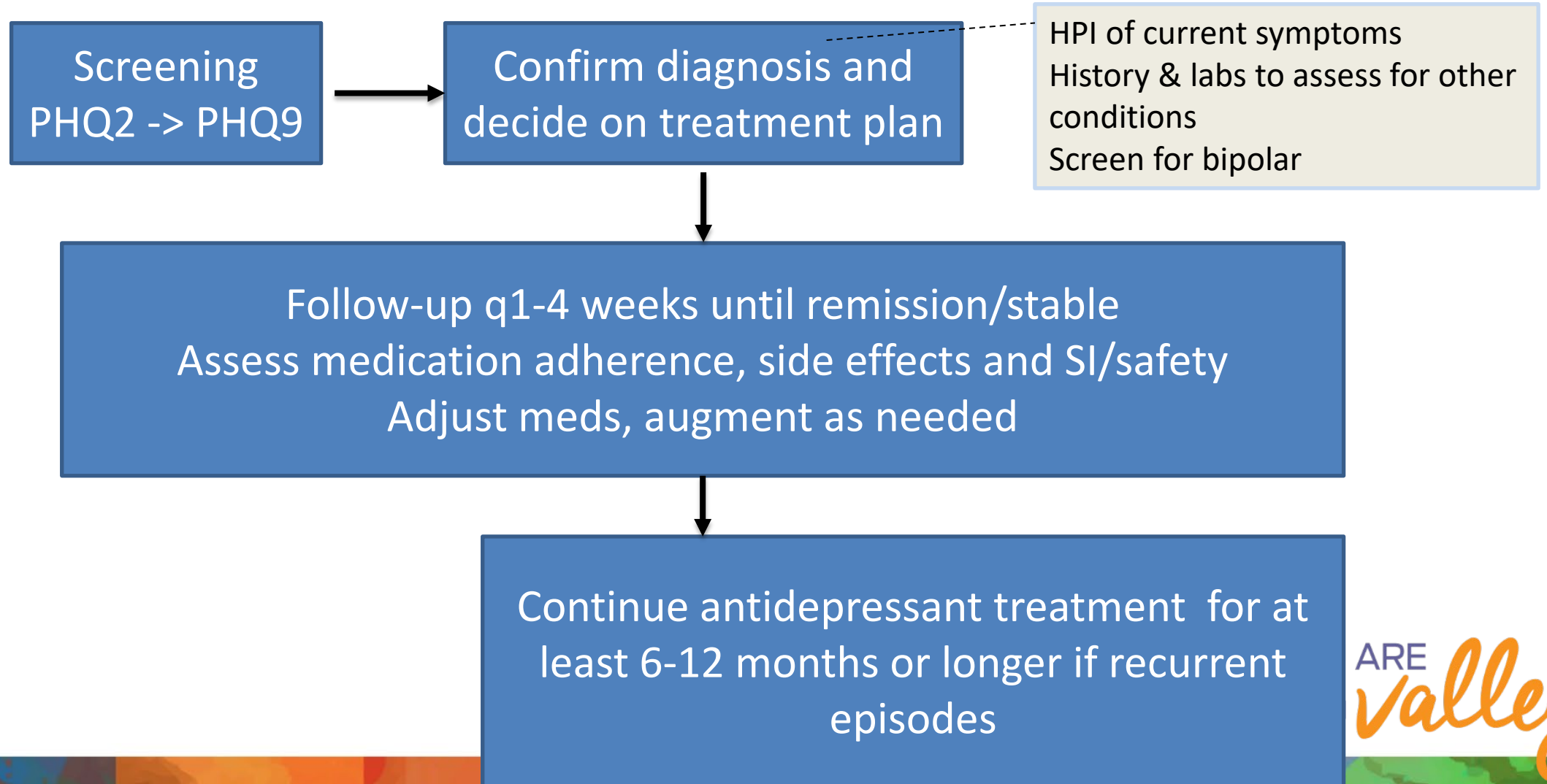
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Objectives

- 1. To discuss the importance of regular follow-up, assessment and treatment adjustment when managing depression.
- 2. To describe evidence-based treatment and augmentation strategies for depression

Brief Overview of Depression CPG



Importance of Screening and Follow-up for Depression

10



5-10% of primary care patients with major depression

5



50% of depression cases detected (w/o universal screening)

2.5



50% adherence to antidepressants

1



33-47% remission with initial treatment

Those with chronic medical conditions are at higher risk for depression and may have more severe or difficult to treat depression than those with depression alone.



Depression Screening & Monitoring

- PHQ9 has been widely validated as a diagnostic measure, and as a responsive and reliable measure of depression treatment outcomes
- Other depression screeners— Edinburgh postnatal, geriatric depression scale, HAM-D
- Also consider patient's objective presentation and functioning (work, school etc...). If discrepancy with PHQ9 score, ask questions to clarify.
- In general, a **5-point reduction** in PHQ9 is considered a significant change



There are so many depression diagnoses in Epic. Which do I use?

Diagnosis	ICD10 Code	When to Use
Major depressive episode (HCC if specifiers included)	F32.x, F33.x	Approx 10+ on PHQ9, and score on question 1 (depressed mood) and/or 2 (anhedonia) =2 or 3]
Adjustment disorder with depressed mood/mixed anxiety and depressed mood	F43.21/ F42.23	Identifiable stressor and pt does not meet criteria for MDE
Depression, unspec Other depression Other recurrent depressive disorders (HCC)	F32.A/F32.89/ F33.8	Pt does not meet criteria for another diagnosis OR You lack sufficient information to make a more specific diagnosis
Major depression, recurrent, chronic (HCC)	F33.9	Major depression for 2+ years w/o full remission

Major Depressive Episode Specifiers

- Single episode vs. recurrent
 - Single -1st time ever meeting criteria for MDE
 - Recurrent – has had at least one prior episode of MDE
- Severity: Mild, moderate, severe
 - Severe, with psychotic features or without psychotic features
- Remission
 - Partial remission (no longer meeting full criteria for a MDE, but still symptomatic)
 - Full remission

- 58 year-old woman with type 2 diabetes mellitus, hyperlipidemia and obesity scores 15 on the PHQ9 and 8 on the GAD7. She is open to trying a medication for depression but is worried about weight gain. Which antidepressant would you start?

Selecting an Antidepressant

- Factors to consider – past history, side effect concerns, co-occurring medical conditions
- Usually start with an SSRI or bupropion
 - Sertraline and escitalopram are good options
- However –
 - If good experience in the past with another AD, try it again
 - If chronic pain (FM/generalized, neuropathy, back), consider duloxetine
 - Avoid bupropion if higher risk for seizures, anorexia/bulimia nervosa
 - For adolescents – Fluoxetine is first line, sertraline/escitalopram 2nd line



Antidepressant Selection: Other Tips

Patient Has	Antidepressant Consideration
Anxiety	Serotonergic antidepressants work best.
Obesity	Weight gain: Bupropion, fluoxetine <<<paroxetine, mirtazapine, TCA's
Cardiovascular disease	Sertraline - best record of CV safety Citalopram, TCA's: Discouraged in patients with or at higher risk for prolonged QTc/TDP
Concern for sexual side effects	Less likely with bupropion or mirtazapine
Pregnancy	Best safety data for fluoxetine, sertraline, citalopram, escitalopram, (tricyclic antidepressants)

Pharmacogenomic testing with decision support tools – insufficient evidence to support routine use



Medication adherence

- Reasons for nonadherence include side effects, no perceived benefit, forgetting, cost
- Educational interventions to improve adherence alone have not been shown to be effective
 - Best strategies involve proactive care management, telephone follow-up, and/or frequent visits, along with identifying and addressing barriers to adherence.

- 58 year-old woman with obesity, diabetes, hyperlipidemia and depression has been on escitalopram 10 mg qday for 1 month. PHQ9 score is 10 and GAD7 score is 3.
- What would you do next?

Depression follow-up



Assess adherence, side effects, response



Partial response?

yes



Increase the dose if able,
if not then augment with
another medication
and/or psychotherapy

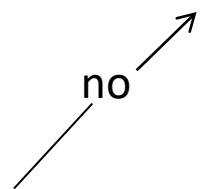
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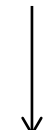
Adequate dose and duration?

Increase dose
and/or wait

no



Yes



Switch medications



Antidepressant Augmentation Strategies

Treatment	Key Symptoms Addressed	Dosing
Buspirone	Anxiety is prominent	10-30mg BID
Bupropion	Depression/fatigue/poor concentration	150mg-450mg qday XL
Psychotherapy	Stressors, coping, negative thoughts	
Lithium*	Depression, anxiety, mood swings	~300-1200mg qday
T3 (Cytomel)*	Depression	25-50mcg/day
Mirtazapine	Depression/anxiety/sleep/appetite	7.5-45mg qbedtime

*Lab monitoring required



Augmenting with Atypical Antipsychotics

1-2 weeks to notice an effect

Typically lower doses than used for psychosis:

Risperidone 0.25-2mg qday

Aripiprazole 2-20mg qday

Quetiapine 50-300mg qday

Olanzapine 5-20mg qday

Ziprasidone 20–80 mg BID

Brexpiprazole 2-3mg qday

Quetiapine also effective as monotherapy for depression (150-300mg qday)



Other Augmentation Strategies

- Light therapy
- Exercise
- Methylfolate (Deplin) – 2 RCT's favoring 15mg qday over placebo
- SAME – 800mg BID, one small RCT supporting
- Stimulants –used off label for residual fatigue, poor concentration; evidence better for late life depression with apathy, fatigue or general medical illness
- Lamotrigine – no evidence for unipolar depression, but good for bipolar and used off label for mood instability/lability



- 58 year-old woman with obesity, diabetes, hyperlipidemia and depression has been on escitalopram for 2 months and is now up to 20mg qday. PHQ9 score is 13 and GAD7 score is 8.
- What would you do next?

Switching Antidepressants

- Switch to another antidepressant if:
 - No significant benefit despite an adequate trial
 - Intolerable adverse effects
 - Loss of effect after years of use (tachyphylaxis) and no benefit with an increase in dose
- A washout period is needed **only** if switching to or from a MAO inhibitor
- If more recent AD/lower dose – can make a direct switch to the other AD
- If longer history on the AD/higher dose – a cross taper may be better tolerated

Switching Antidepressants

- Suggestions re med choice when switching antidepressants
 - If two failed SSRI trials – move on to an SNRI or bupropion
 - If no luck with SSRI, SNRI or bupropion --- Appropriate to seek psychiatric consultation!
 - From a medication standpoint, other options include mirtazapine, TCA, quetiapine, or a newer antidepressant such as Viibryd or Trintellix
- Adolescents – fluoxetine is first line, then sertraline, escitalopram. After that, other SSRI's, bupropion, mirtazapine

Other Strategies for Difficult to Treat Depression

- ECT
 - Electrical current → seizure
 - 60-80% remission rate; relapse 38% at 6 months
 - Cons – General anesthesia, cognitive side effects, stigma
 - Typically given MWF, ~6-12 treatments; for some – monthly maintenance
- rTMS
 - Magnetic field generated below a coil, usually positioned over the patient's dorsolateral prefrontal cortex
 - 50% response rate, 33% remission rate; relapse in ~35%
 - Treatments typically 5 days per week for 4-6 weeks

Other Strategies for Difficult to Treat Depression

- Esketamine
 - NMDA antagonist, opioid receptor agonist, AMPA activation
 - Response typically w/in 2-4 hours in 25-65%
 - Administered onsite: 2x per week for weeks 1-4, tapering down to once every 1-2 weeks

References

- Barkil-Oteo A. Collaborative care for depression in primary care: how psychiatry could “troubleshoot” current treatments and practices. Yale J of Biol and Med 2013; 86: 139-146.
- Chong WW et al. Effectiveness of interventions to improve antidepressant medication adherence: a systematic review. Int J Clin Pract 2011;65(9):954-75.
- Costantini L et al. Screening for depression in primary care with Patient Health Questionnaire-9 (PHQ-9): A systematic review. J Affect Disord 2021; 15(279):473-483.
- Fortney JC et al. Reasons for antidepressant nonadherence among veterans treated in primary care clinics. J Clin Psychiatry 2011; 72(6):827-34.
- Gaines BN et al. What Did STAR* D Teach Us? Results From a Large-Scale, Practical, Clinical Trial for Patients With Depression. Psychiatric Services 2009; 60:1439–1445.
- Holtzheimer P. Unipolar depression in adults: Indications, efficacy, and safety of transcranial magnetic stimulation (TMS). Up to Date, topic last updated May 9, 2019.

References

- Kellner, C. Overview of electroconvulsive therapy (ECT) for adults. Up to Date, topic last updated September 16, 2021.
- Katon W & Shullberg H. Epidemiology of depression in primary care. Gen Hosp Psychiatry 1992; 14(4): 237-47.
- Lowe B et al. Monitoring depression treatment outcomes with the patient health questionnaire-9. Med Care 2004;42(12):1194-201.
- Masan PS. Tolerability and adherence issues in antidepressant therapy. Clin Ther 2003; 25(8):2289-304.
- Nierenberg A. Unipolar major depression in adults: Augmentation of antidepressants with stimulants and stimulant-like drugs. Up to Date, topic last updated March 30, 2021.
- Ontario Health (Quality). Multi-gene Pharmacogenomic Testing That Includes Decision-Support Tools to Guide Medication Selection for Major Depression: A Health Technology Assessment. Ont Health Technol Assess Ser. 2021; 21(13): 1–214.



References

- Seattle Children's. Primary Care Principles for Child Mental Health, version 10 (2021).
- Tao L et al. Light therapy in non-seasonal depression: an update meta-analysis. *Psychiatry Res* 2020; 291:113247
- Thase M & Connolly KR. Ketamine and esketamine for treating unipolar depression in adults: Administration, efficacy, and adverse effects. Up to Date, topic last updated August 26, 2020.
- Thase M & Connolly KR. Unipolar depression in adults: Management of highly resistant (refractory) depression. Up to Date, Topic last updated August 31, 2021.

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