

# Psychopharmacology for Collaborative Care Managers

## Keep a Perspective

Treating patients with depression and anxiety:

- Three main tools
  - Biology which may mean a psychiatric med.
  - Psychology therapy of various kinds
  - Patient behavior and self management
- Which of these is more effective?
- Which is easier for the patient to do?
- The goal is to return a patient to functioning and normal emotional range – use <u>all</u> the tools



## **Scenario**

Dr. Kim brings a patient to you and says, "Mr. M. here is an 70 yo man with hypertension and coronary artery disease. He recently moved to an assisted care facility; his family says he isn't like himself anymore as he doesn't want to do anything.

They think he's depressed. Could you ask your psychiatrist if we should start an antidepressant?

What more information is needed?

# Does the patient have an indicated condition to use an antidepressant?

#### Clinical history

- Does the chief complaint and history suggest a primary depressive or anxiety disorder according to DSM5 criteria?
- Is there evidence that might suggest another reason for the way the patient is presenting?
- Symptom measures like your blood pressure measure.
   A high level = look closer, not diagnose hypertension.
- Depression
  - PHQ-9 ≥ 10
- Anxiety
  - GAD7 ≥ 10

# Depression

#### Common

- Lifetime prevalence of 16% (>20% in women)
- 10-20 percent of primary care patients are depressed.

## Dangerous

- Depression history = 2 X risk of CAD
- Increases risk of HTN and stroke by 50%
- Depression post MI = 6 X risk of death in 18 mos\*

<sup>»</sup> Frasure-Smith N, Lesperance F, Talajic M. Depression and 18-month prognosis after miocardial infarction Circulation 1995; 15;91;999-1005.

#### PATIENT HEALTH QUESTIONNAIRE (PHQ-9)

## Screening, Monitoring Tool - PHQ-9

- Quick, many languages
- First 2 questions must have a positive score
- Score ≥ 10
  - For Major Depression
    - Sensitivity 88%
    - Specificity 88%
- Mild (5),mod (10), mod severe
   (15) severe depression (20)

NAME: John Q. Sample			DATE:			
Over the last 2 weeks, how often have you been bothered by any of the following problems? (use "\scriv" to indicate your answer)	M. Hall	grant tori	her the their	he art treet for		
1. Little interest or pleasure in doing things	0	1	✓	3		
2. Feeling down, depressed, or hopeless	0	√.	2	3		
Trouble falling or staying asleep, or sleeping too much	0	1	V.	3		
4. Feeling tired or having little energy	0	1	2	<b>✓</b>		
5. Poor appetite or overeating	0	✓	2	3		
6. Feeling bad about yourself—or that you are a failure or have let yourself or your family down	0	1	V	3		
7. Trouble concentrating on things, such as reading the newspaper or watching television	0	1	¥	3		
Moving or speaking so slowly that other people could have noticed. Or the opposite—being so fidgety or restless that you have been moving around a lot more than usual	0	1	1	3		
Thoughts that you would be better off dead, or of hurting yourself in some way	€	1	2	3		
	add columns:	2	+ 10 +	3		
(Healthcare professional: For interpretation of TOTAL, please refer to accompanying scoring card).			15			
10. If you checked off any problems, how difficult have these problems made it for you to do your work, take care of things at home, or get along with other people?		Si	ot difficult at all omewhat difficul ery difficult tremely difficult			

<sup>\*</sup>Patient Health Questionnaire (PHQ-9). Spitzer et al. Copyright Pfizer Inc. 1999

# Could this be something else?

Bereavement – feelings tied to loss

Thyroid disease – weight changes, energy changes

**Cancer** – weight and energy changes, pain

**Substance induced mood disorder** – tied to use patterns Bipolar disorder – rapid improvement, mania/hypomania (earlier onset (mean age 24)

Attention deficit – concentration and irritability more than sadness or loss of interest

**Dementia** – lack of interest or initiative

Pain disorders – related to opiate use

**Persistent Depressive disorder** – last 2 years or more with no break

# Maybe more than one problem?

- Among patients diagnosed with Major Depression
  - Anxiety (75% w features, 37% w diagnosis lifetime prev.)
  - Bipolar features (mixed features in 16%)
  - Personality Disorder (32% w diagnosis)
  - Substance abuse (58% w diagnosis)
- Insomnia/sleep apnea
- Pain, Thyroid disorder
- Social determinant abuse, housing, finances, etc.

# Information to gather as we decide on a medication

#### Medical

• Hypothyroid, pain, sleep issues, evidence of cognitive decline, fall risk, pregnancy, other medications...

#### Psychological/Social

- Pattern when did this start? What was going on?
- Drinking or drug use?
- Life stressors and timing of mood changes
- Past history of depression what happened?
  - Past medication trials dose, duration, response?
- Other mental health problems
- Current life stressors, level of functioning & supports

# **Comorbid anxiety? GAD7**

Over the <u>last 2 weeks</u>, how often have you been bothered by any of the following problems?

Please circle your answers.

GA	ND-7	Not at all sure	Several days	Over half the days	Nearly every day
1.	Feeling nervous, anxious, or on edge.	0	1	2	3
2.	Not being able to stop or control worrying.	0	1	2	3
3.	Worrying too much about different things.	0	1	2	3
4.	Trouble relaxing.	0	1	2	3
5.	Being so restless that it's hard to sit still.	0	1	2	3
6.	Becoming easily annoyed or irritable.	0	1	2	3
7.	Feeling afraid as if something awful might happen.	0	1	2	3

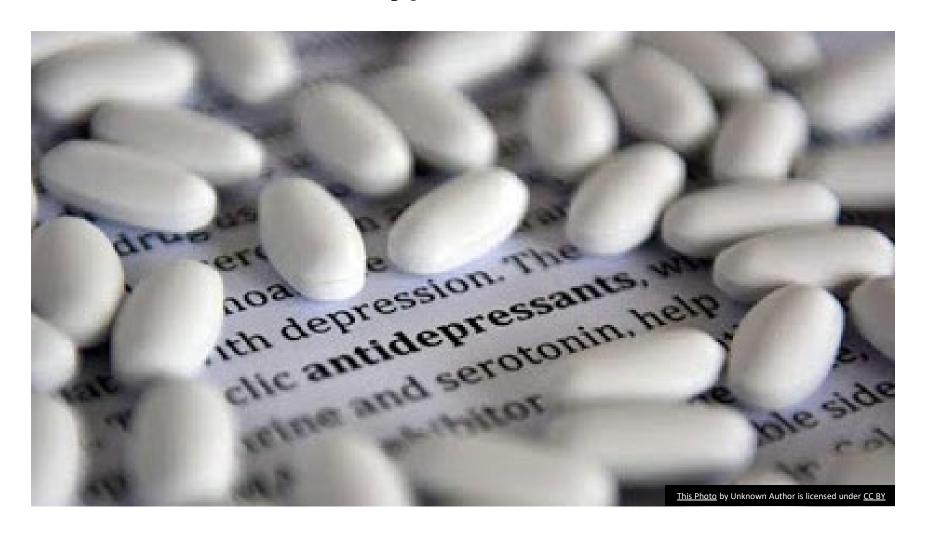
- GAD should not be diagnosed when only occurring with a mood disorder or better explained by other anxiety disorder
- Other symptoms of GAD: muscle tension, fatigue, insomnia, poor concentration
- Several types of anxiety panic, OCD, social anxiety, PTSD

# How does the data gathering impact the decision?

### Examples:

- If no history of mood problems and symptoms came up after a sad event
  - Might want to work on coping first
- If also has pain
  - Might look at an antidepressant that also helps pain
- If on a drug that could cause drug interactions
  - Think about that drug when choosing a med
- Failed responding to several antidepressants
  - Is there a pattern? Maybe try a different class?

## Evaluation for Antidepressant (or Antianxiety) Medication



# Is there a reason not to use an antidepressant (contraindication)?

Only in rare cases are SSRIs absolutely contraindicated

#### Depression/anxiety secondary to another condition

- We address that condition and check back
- Substance use disorders are not contraindications to antidepressant treatment

#### Bipolar disorder

- Antidepressants can lead to mood swings and/or mania if bipolar is the real issue
- May still be effective for comorbid anxiety disorder

Medical condition or medications that would interact with antidepressant treatment

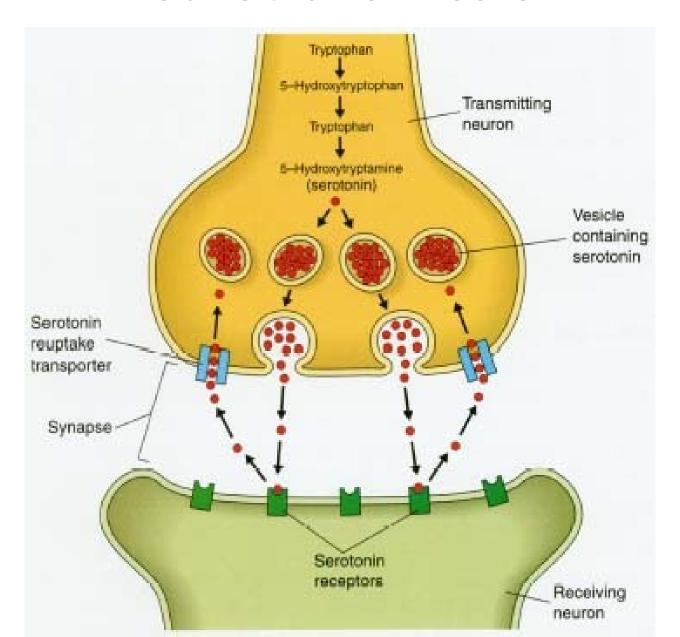
We also may find patients who have had bad reactions to antidepressants

# **Mechanisms of Psychopharmacology**

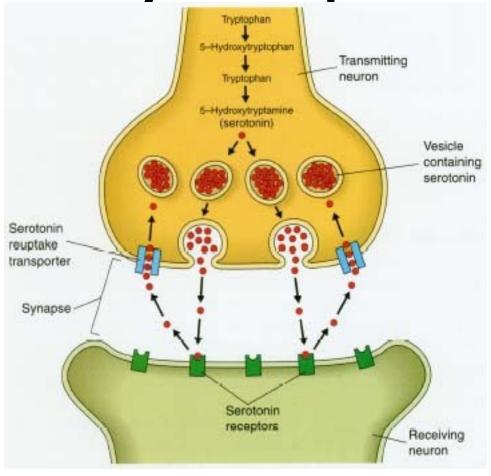


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## Neurotransmission



**Psychotropic Sites of Action** 



Increase release

Decrease reuptake

Decrease breakdown

Direct receptor activation

Receptor modulation

Direct receptor blockade

## Things to think about with medications

#### Half life – how long does it take medicine to build up or exit?

- Impacts how long to feel a benefit
- Impacts how likely we will see withdrawal

#### Does the medication impact the cytochrome p450 system?

- Enzymes that help break down medications
- Basis for some drug interactions

#### Can the medication do more than one thing?

Sleep and mood for example

#### Is this medication covered by insurance?

#### Makes my head spin, what do I tell patients?



Antidepressants impact brain chemicals in a way that is meant to return you to a normal mood range

- Not to make you into someone else.
- They are NOT addictive

The initial effects on your body from the medicine are to increase those chemicals

Side effects generally come early and go away

#### Improvement comes after your body adjusts

So we need to stay on the medicine to reach benefit

Therapeutic effects Effects of antidepressant treatment Side effects

Time in weeks

# How long to wait?

### Get to a minimal therapeutic dose

- Good sign if seeing some improvement in 2 weeks
- Leveling off of further improvement in 6-8 weeks

# Care Manager should have basic information available on these medicines

- What is the minimal effective dose of each medicine?
- What is the usual maximum dose?

## **Neurotransmitters: Monoamines**

### <u>Serotonin</u>: CALM – Reduce strong negative emotions

- mood, anxiety, sleep, anger/aggression
- sexual functioning, gastrointestinal functioning

#### Norepinephrine: Can help with focus and pain

- mood, anxiety
- heart rate, blood pressure, "fight or flight"

### **Dopamine**: enhancing versus blocking

- motivation, mood, psychosis, attention, cognition, reward
- motor activity, inhibits lactation

#### Histamine & Melatonin: sleep

## **Neurotransmitters: Other**

#### GABA: Think of the Valium drugs here as well as gabapentin

- major inhibitory role, anxiety
- sedation, cognition

#### Glutamate: This is a work in progress but think ketamine

- major excitatory role, cognition, mood
- psychosis

#### Acetylcholine: blocking versus enhancing

- cognition and memory
- heart rate, bladder, gastrointestinal: "rest & digest"
- "anticholinergic" side effects

## How to use this information

#### **Explaining it to clients:**

 Example: "This medication affects the level of a chemical called serotonin in the brain"

Helps to understand what are expected side effects

## Establishes classes of medications (e.g. SSRIs or SNRIs)

Easier to remember than learning each individually

#### Helps understand new medications

Really new or "me too"?



# **Antidepressant Medications**



# **Old-school Antidepressants**

#### **Monoamine Oxidase Inhibitors (MAOIs)**

 Require strict dietary restrictions to avoid dangerous side effects, rarely used anymore

#### **Tricyclic antidepressants (TCAs)**

- Significant anticholinergic side effects
- Dangerous in overdose (cardiac arrhythmias)
- Still used for migraine headaches, nerve pain, sleep
  - Amitriptyline (Elavil), Nortriptyline (Pamelor), Doxepin (Sinequan)
- Generally, not first choice for depression/anxiety
- Often see low dose at night added to another antidepressant but watch for drug interactions

## **Commonly used Antidepressants**

SSRIs – serotonin recycling blocker

SNRIs – impacts serotonin AND norepinephrine

Bupropion (Wellbutrin) – serotonin not involved – impacts norepinephrine and dopamine

Mirtazapine (Remeron)

Trazodone – also serotonin in another way but is so sedating that used mostly for sleep

Others – newer options such as vilazodone

### **SSRIs**

- Fluoxetine (Prozac)
- Sertraline (Zoloft)
- Paroxetine (Paxil)
- Citalopram (Celexa) & Escitalopram (Lexapro)
- Fluvoxamine (Luvox)
  - FDA approved for major depressive disorder
  - This group is often picked when also having anxiety
  - Some also approved for:
    - Posttraumatic stress disorder
    - Generalized anxiety disorder
    - Obsessive compulsive disorder
    - Social anxiety disorder

## **SSRIs: Common Side Effects**

Gastrointestinal upset (nausea, diarrhea), usually transient over the first few days

Sexual side effects – difficulty with libido, erection, orgasm, reversible upon stopping medication

"Early activation" – transient period of increased anxiety, restlessness upon initiating treatment

Discontinuation syndrome – "Brain zaps", electric shock-like sensations in the neck and head

Insomnia or somnolence

Weight gain, average about 1% per year

## **SSRIs & Serotonin Syndrome**

- Serotonin Syndrome: uncommon but dangerous consequence of excessive serotonin activity
  - Symptoms: muscle rigidity, fever, agitation
- Causes: overdose of SSRI antidepressants or combination of medications that affect serotonin
- Other pro-serotonin drugs include:
  - Tramadol and other opiates
  - Triptans for migraine headaches
  - Stimulants and drugs of abuse: cocaine, ecstasy (MDMA)
  - Anti-nausea medications, some antibiotics
  - St. John's Wort, some herbal supplements

## **SSRIs: Differences within class**

- Citalopram, escitalopram, and sertraline have the fewest interactions with other medications
  - Good for older patients on lots of medications
- Fluoxetine has the longest half-life
  - Possible better for patients apt to miss doses
  - Also most weight neutral
- Paroxetine may have greater anticholinergic side effects and worse discontinuation syndrome
  - Also more concerns in pregnancy

### **SNRIs**

Venlafaxine (Effexor) & Desvenlafaxine (Pristiq)

**Duloxetine (Cymbalta)** 

Levomilnacipran (Fetzima) – generic just becoming available may be costly

Block reuptake of serotonin and norepinephrine

Efficacy and side effects are generally similar to SSRIs

#### Advantage vs. SSRIs:

also effective for neuropathic pain (e.g. from diabetes, fibromyalgia)

**Disadvantage vs. SSRIs:** 

greater hypertensive effects

# **Bupropion (Wellbutrin)**

- Mechanism: Inhibits norepinephrine <u>and</u> dopamine reuptake
- Effective for major depression and smoking cessation
- Common side effects: headache, insomnia
- Advantages vs. SSRIs: Less weight gain or sexual dysfunction
- Disadvantage vs. SSRIs: not effective for anxiety disorders
- Avoid in patient with a seizure history

# Mirtazapine (Remeron)

- Complex mechanism: works not on reuptake but on receptors that in turn, impact release of serotonin and norepinephrine. Also works on histamine receptors
- Effective for major depression
- Common side effects: sedation and weight gain
- Advantage vs. SSRIs: useful if insomnia and weight loss are present, less sexual side effects
- <u>Disadvantage vs. SSRIs</u>: weight gain, not proven effective for comorbid anxiety disorders

## **SRI plus Serotonin Modulator**

Vilazodone (Viibryd) – (2011) Vortioxetine (Trintellix) – (2013)

Serotonin reuptake inhibitor and partial serotonin receptor activator

Might not be covered by insurance

Vilazodone may have less sexual side effects

Vortioxetine may help with cognitive issues in depression

No clear reason to expect these are better by being newer.

## **Trazodone**

Weak serotonin reuptake inhibitor, blocks and partially activates some serotonin receptors

Used most often for its primary side effect in low doses: sleep

Rare side effect: priapism (erection that won't go away)

Other common side effect: hangover

# **Choice of Antidepressant**

- 38 year-old woman with depression and anxiety
- A) Sertraline
- B) Venlafaxine
- C) Bupropion
- D) Mirtazapine
- What if she also has ADHD symptoms but no anxiety?
- What if she also has chronic neuropathic pain?
- Remember, the patient has to keep taking the med (50% stop)
- There is no best antidepressant choice is based on side effects, preferences, cost, comorbidities
  - Mayo antidepressant shared decision aid for a first medication choice
  - https://depressiondecisionaid.mayoclinic.org/index

#### What if initial treatment fails?

Up to 2/3<sup>rds</sup> of patients fail to reach remission on 1<sup>st</sup> treatment

#### Options for the next step include:

- Increasing dose
- Adding a second "augmenting" antidepressant from other class
  - SSRI + bupropion or mirtazapine are common choices
- Switching to another antidepressant (< 60% improvement)</li>
  - SSRI to other SSRI is as good as switching to bupropion
- Augmenting with an antipsychotic or other medication
  - VA trial found augmentation with aripiprazole (Abilify) was more effective than switch to bupropion.

After 2 failures: Right diagnosis? Intensify treatment?

## Other Common Psychotropic Medications



#### Benzodiazepines



- Alprazolam (Xanax)
- Clonazepam (Klonopin)
- Lorazepam (Ativan)
- Diazepam (Valium)
- Mechanism: act on GABA receptors to enhance GABA effects
- Indicated for panic disorder, generalized anxiety disorder
  - Also used to treat alcohol withdrawal
- Best if used short-term (in primary care)
- Not effective for depression or PTSD
- Potential for abuse and dependence
- Caution with driving, not to be mixed with alcohol
- Sudden withdrawal syndrome: anxiety, shakes, insomnia, seizures
- Can worsen cognition in elderly and may increase fall risk

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### **Controlled Substance Prescribing**

Benzo use doubles risk of opiate overdose

Michigan MAPS report required prior to prescribing

Only 30-day supply at a time (+/- refills)

#### Consider also:

- Urine drug screen for other substance use
- Patient contract
  - Only one doctor at a time
  - No early refills or replacement for lost medications
  - Attend all appointments

### Other (non-addictive) anti-anxiety

#### Buspirone (Buspar) – indication is GAD

- Serotonin agonist, not effective for depression but <u>can augment</u>
- Takes weeks to work, significant GI side effects

#### Hydroxyzine (Atarax, Vistaril)

- Anti-histamine (like Benadryl), can be taken PRN, works immediately
- Avoid in elderly (confusion, falls). Can impact EKG (QTc)

#### Gabapentin (Neurontin), Pregabalin (Lyrica) – also GABA system

- Works immediately, safe
- Good for alcohol withdrawal and related anxiety & neuropathic pain

#### Prazosin (Minipress)

- Anti-hypertensive medication, increased dose gradually
- Evidence primarily for PTSD-related nightmares
- Effectiveness has been questioned recently

### 'Z drugs' – safer? Maybe not

- Also act on GABA site
- May have additive effect with other sedative medications
- Short term best
- High dose addiction
- Unusual effects
  - Odd night behavior
  - Falls in elderly
  - amnesia

# Zolpidem (Ambien)

## Zaleplon (Sonata)

## Eszopiclone (Lunesta)

### Other Hypnotics ("Sleep Aids")

## CBT for Insomnia recommended 1<sup>st</sup> line for chronic insomnia

- Diphenhydramine (Benadryl, other OTCs)
  - Stops working quickly, anticholinergic side effects
- Melatonin
  - Generally safe, not very effective long term
- Sedating antidepressants: Doxepin, Trazodone, Mirtazapine
- Ramelteon (Rozerem): melatonin agonist
  - Limited effectiveness
- Belsomra (suvorexant): (2015)
  - Orexin antagonist

## Talking with Patients about Antidepressants



#### The nuts and bolts

- Antidepressants need to be <u>taken</u> <u>daily; NOT as needed</u>
- All antidepressant take 2-4 weeks to see a benefit
- Most side effects resolve in a few days, serious side effects are rare
- Antidepressant should be <u>continued for at least 6</u>
   <u>months.</u> Longer if recurrent serious episodes
- If the first antidepressant doesn't work out, there are many other options – generally 60% rule (change/add)

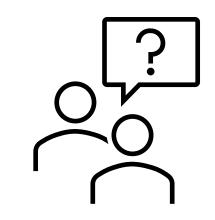
Q: Are antidepressants just a placebo?

A: Antidepressants trials consistently show superiority to placebo: about 30% will get better with a placebo compared to 40% with an antidepressant

 Placebo response is high with depression, some consider this part of antidepressant treatment

Q: Do antidepressants cause suicide?

A: Although the FDA warns against an increase in suicidal thoughts and behaviors in those up to 24 years old, there is no convincing evidence antidepressants result in an increase in suicide death. Epidemiologic studies suggest antidepressant use is associated with fewer suicides



Q: Are antidepressants addictive?

A: Antidepressants are very rarely abused (no real street value) and have no dangerous withdrawal syndromes. Withdrawal occurs in some patients with short acting drugs – more uncomfortable than dangerous



Q: Do antidepressants turn people into zombies?

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A: Most antidepressants are not sedating nor cause problematic slowing of cognition. Some people report feeling overall less emotional on antidepressants. This may be a dose issue or a need to try another medicine. The goal is not lacking emotions but having normal range.



Q: Am I going to be on this medication forever?

A: Recommend at least 6 months after achieving remission if first episode, indefinitely if multiple episodes. Message to patients is, "It's up to you how long you take this medication, and whether you find the benefits outweigh the costs"

#### Conclusion

Antidepressants are effective, generally safe, and preferred by many patients

# Keys are:

Rule out other causes of depression, including bipolar disorder and medical conditions

Provide education to patients about antidepressant treatment, expected response time, and side effects

Follow-up with patients to assess treatment response and to ensure changes are made when response is inadequate

#### **Resources Related to Medications**

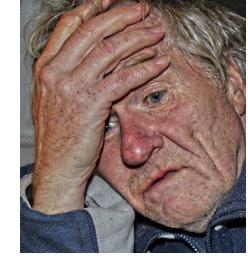
ICSI (Institute for Clinical Systems Improvement), Depression, Adult in primary care depression	https://www.icsi.org/guideline/depression/
APA (American Psychiatric Association) Practice Guidelines	https://psychiatryonline.org/guidelines
American Geriatrics Society 2019 Updated AGS Beers Criteria® for Potentially Inappropriate Medication Use in Older Adults	https://onlinelibrary.wiley.com/doi/full/10.111 1/jgs.15767
Mayo antidepressant shared decision aid	https://depressiondecisionaid.mayoclinic.org/index
Psychopharmacology and Psychiatry Updates Psychopharmacology Institute (Podcasts)	https://podcasts.apple.com/us/podcast/psycho pharmacology-and-psychiatry- updates/id1425185370 (free access to short and preview podcasts)

#### Case Follow-up

You assess Mr. M. and his medical record His PHQ9 is 18, loss of interest started 1 month ago after moving

- No history of mania, substance use, or trauma.
   No past depression.
- Medical history positive for a heart attack 10 years ago with bypass surgery, has hypertension and high cholesterol, treated with medications (beta blocker, ace inhibitor, statin, and aspirin)
- All lab work is normal

Should we start an antidepressant?
What else might help in SCR to decide on next steps?
Cognitive functioning
Recent change in medications, medical condition
Problems related to environment, socializing



## Questions?