# **Geriatric Optimization**

Tricyclic Antidepressants (TCA) and Paroxetine (Paxil®)

# Selective Serotonin Reuptake Inhibitors (SSRIs) Paroxetine (Paxil®)

## **Tricyclic Antidepressants (TCAs)**

Amitriptyline (Elavil®)
Clomipramine (Anafranil®)
Doxepin (Silenor®)
Imipramine (Torfanil®)
Trimipramine (Surmontil®)

Amoxapine (Asendin®)
Desipramine (Norpramin®)
Nortriptyline (Pamelor®)
Protriptyline (Vivactil®)

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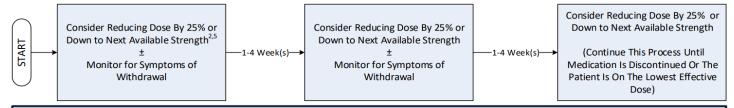


### WHY ARE THESE MEDICATIONS INAPPROPRIATE?

- Older adults are at an increased risk of negative outcomes when taking TCAs and/or paroxetine due to the strong anticholinergic side effects and risk of hypotension. 1,2 Other side effects may include:
  - Sedation
  - o Falls and fractures
  - Visual impairment
  - Hyponatremia
- One meta-analysis that included 248 studies showed that those taking TCAs had higher odds of falling compared to placebo, especially during the first few weeks of therapy.<sup>2</sup>
- Paroxetine is a major concern due to it having the highest risk of anticholinergic side effects and hyponatremia compared to all other SSRIs.
- These medications should be assessed for deprescribing and potentially replaced with alternatives.

### **APPROACH TO OPTIMIZATION**

- Determine the indication(s) of the TCA or paroxetine including any off-label uses.
- Determine if an alternative medication is needed for the condition or if lifestyle changes can be made instead (see appendix for recommended alternatives if using to treat an off-label indication).
- When deprescribing the TCA or paroxetine, providers can consider the following methods<sup>3,4</sup>:
  - 1. Taper-off over several weeks with no alternative antidepressant to replace it
  - 2. Taper-off and add new antidepressant after taper is complete or after a washout period
  - 3. Cross-taper between both the TCA or paroxetine and the new antidepressant
- Deprescribing should be individualized based on treatment goals, side effects, fall risk, and severity of depression/anxiety. The diagram below provides guidance to deprescribing the TCA or paroxetine if any of the above methods are used.<sup>3-5</sup>



- Most tapering schedules last four weeks but can be shorter if member has been on antidepressant short term (≤3 weeks)<sup>5</sup>
  - o Paroxetine has the highest likelihood of withdrawal effects and usually requires a longer taper (i.e., >6 weeks)<sup>3-5</sup>
- A taper can be extended if withdrawal symptoms occur (e.g., reduce dose every 2-3 weeks rather than every 1 week)
- Use caution if cross-tapering between antidepressants as there is an increased risk of adverse reactions (i.e., fractures, serotonin syndrome)<sup>6</sup>

**GOAL:** Full Discontinuation

• Providers may also choose to directly switch between the TCA or paroxetine to the new antidepressant as an option (see appendix for more details on all discontinuation/taper methods).<sup>3-5</sup>

### ADDITIONAL BEST PRACTICES AND CONSIDERATIONS

# Considerations If Switching to Other Antidepressants<sup>1,2,7</sup>

- All SSRIs and serotonin-norepinephrine reuptake inhibitors (SNRIs) are associated with falls and require monitoring. Use minimum effective dose.
- SSRIs and SNRIs also increase risk of hyponatremia and therefore Na+ should be monitored closely especially during initiation or dosage changes.
- •The preferred SSRIs to use in older adults include citalopram (at 20 mg or less), escitalopram, and sertraline.

# Antidepressant Discontinuation Syndrome: FINISH<sup>4,8</sup>

- Flu-like symptoms
- Imbalance
- Nausea
- Insomnia
- Sensory disturbances (blurred vision)
- Hyperarousal (increased anxiety/agitation)



### **APPENDIX**

## **ALTERNATIVE TREATMENT FOR OFF-LABEL USES**

| Off Label Uses   | Alternative Therapies  |
|--|--|
| Fibromyalgia/Neuropathic<br>Pain <sup>9,10</sup>                     | <ul> <li>Cognitive behavioral therapy</li> <li>Duloxetine (Cymbalta®)</li> <li>Milnacipran (Savella®) (only FDA approved for fibromyalgia)</li> <li>Gabapentin (Neurontin®)</li> <li>Pregabalin (Lyrica®)</li> </ul>   |
| Abdominal Pain in<br>Irritable Bowel Syndrome<br>(IBS) <sup>11</sup> | <ul> <li>Gut-directed psychotherapy</li> <li>Peppermint Oil</li> <li>SSRIs may be used in IBS but do not demonstrate as much benefit in improving symptoms.</li> <li>Other agents tailored to the specific symptoms (i.e., diarrhea or constipation) may also help with abdominal pain.</li> </ul> |
| Sleep<br>Disorders/Insomnia <sup>12</sup>                            | <ul><li>Cognitive behavioral therapy</li><li>Melatonin</li></ul>   |

### **OVERVIEW OF ANTIDEPRESSANT TAPER METHODS**

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|---|---|
| Taper Method  | Additional Details <sup>3-5</sup>   |
| Taper-Off (With No<br>New Antidepressant)                     | This method is preferred if member does not need an alternative antidepressant.   |
| Taper Off and Add<br>New Antidepressant<br>After Taper        | <ul> <li>This method is preferred when there is a concern of drug interactions between both antidepressants and if member wants to avoid side effects from taking both antidepressants at once.</li> <li>Providers can consider adding new antidepressant immediately after taper is complete or after a washout period.</li> </ul> |
| Cross-Taper<br>(Between Old<br>Antidepressant and<br>New One) | <ul> <li>This method may be preferred in members who are at increased risk of relapse in depression or anxiety.</li> <li>This method may have higher risk of side effects.</li> </ul>   |
| Direct Switch<br>(Between Both<br>Antidepressants)            | <ul> <li>This method may be preferred if the member is having significant side effects from the TCA or paroxetine and prefers to be switched immediately.</li> <li>This method may have higher risk discontinuation syndrome.</li> </ul>  |



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