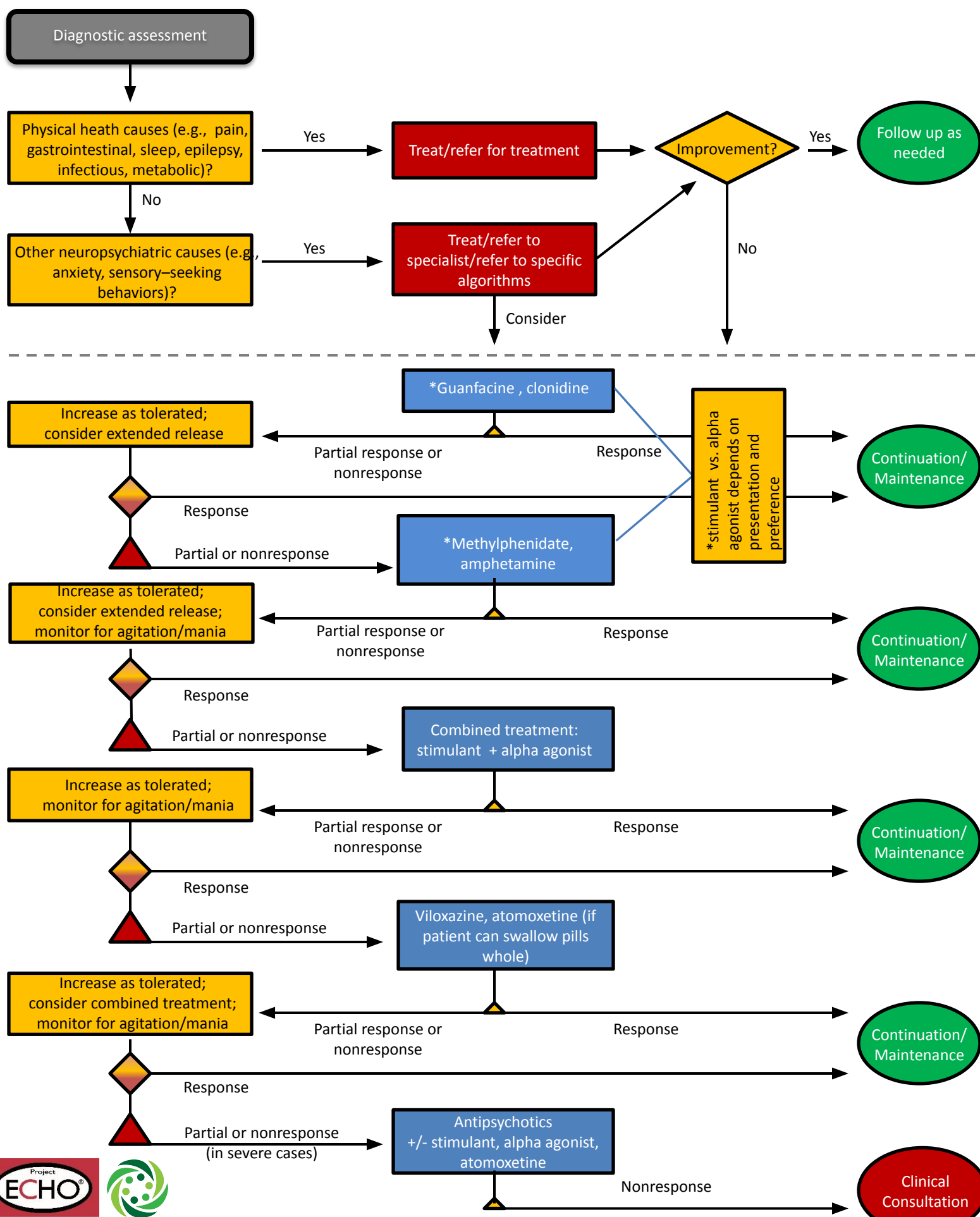
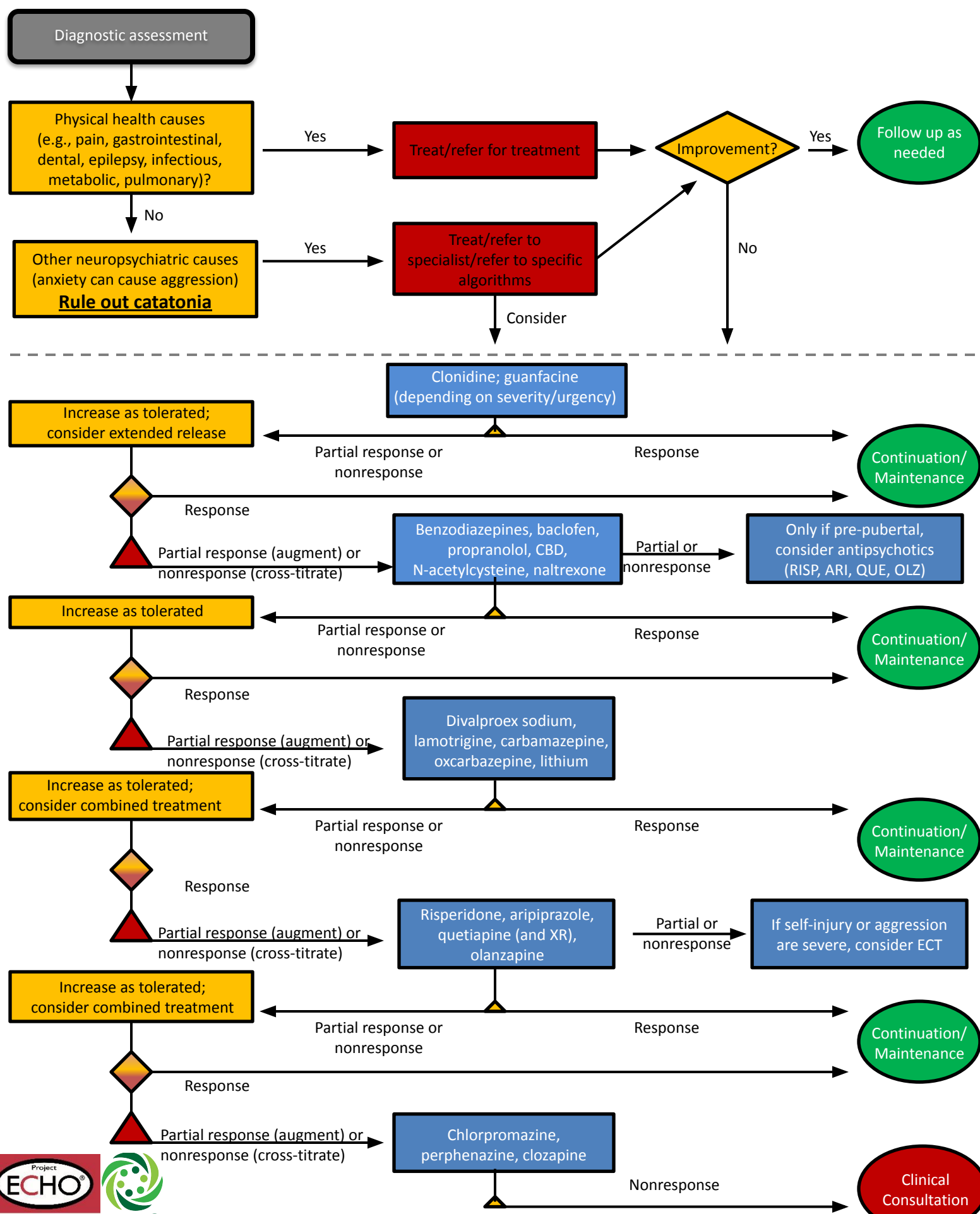


Expert Consensus Recommendations* for the Pharmacological Management of Attention Deficit, Hyperactivity, and Impulsivity in Phelan-McDermid Syndrome



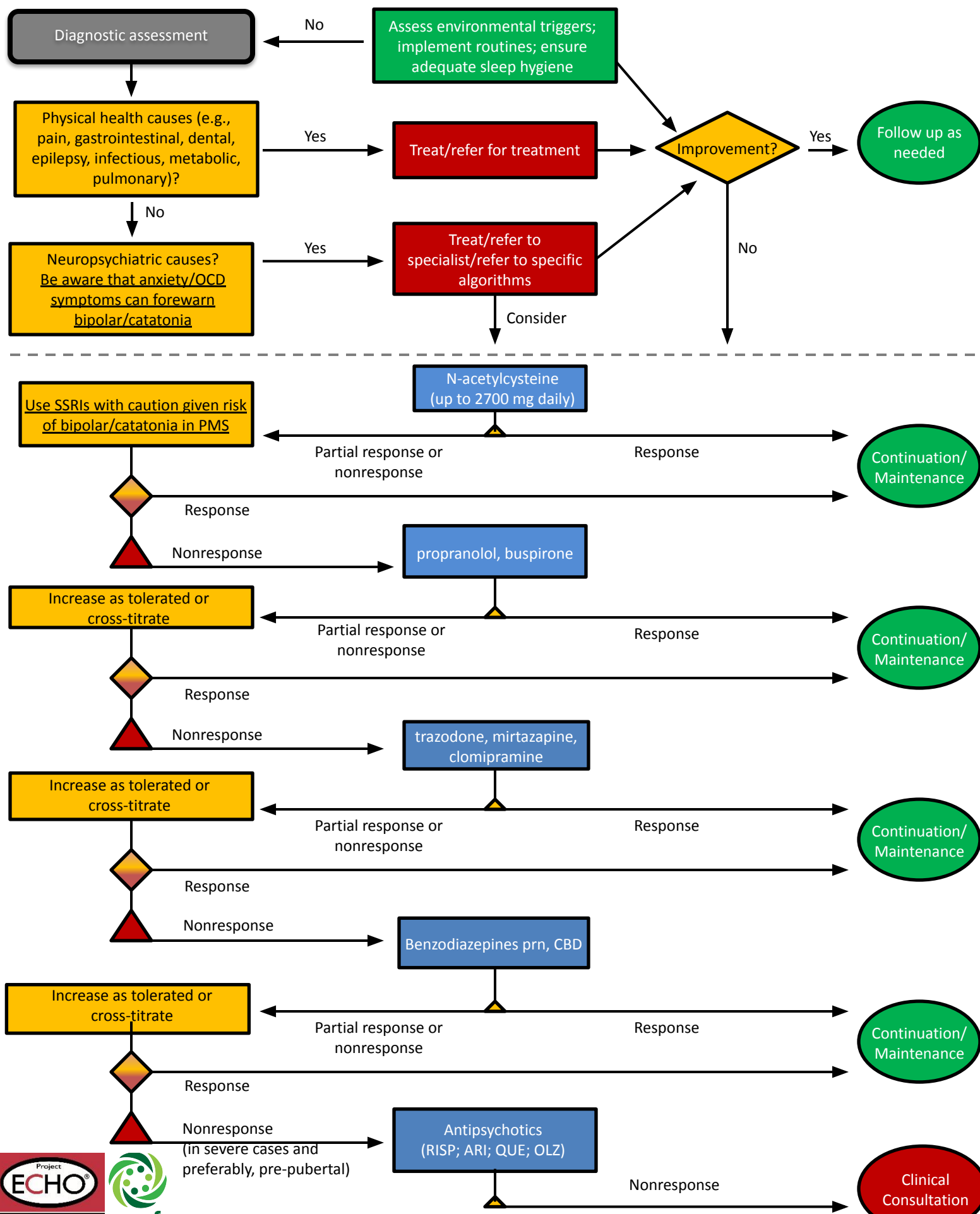
*These recommendations are not established as “evidence-based.” rev 3.1, 12/2024

Expert Consensus Recommendations* for the Pharmacological Management of Irritability, Aggression, and Self-Injury in Phelan-McDermid Syndrome



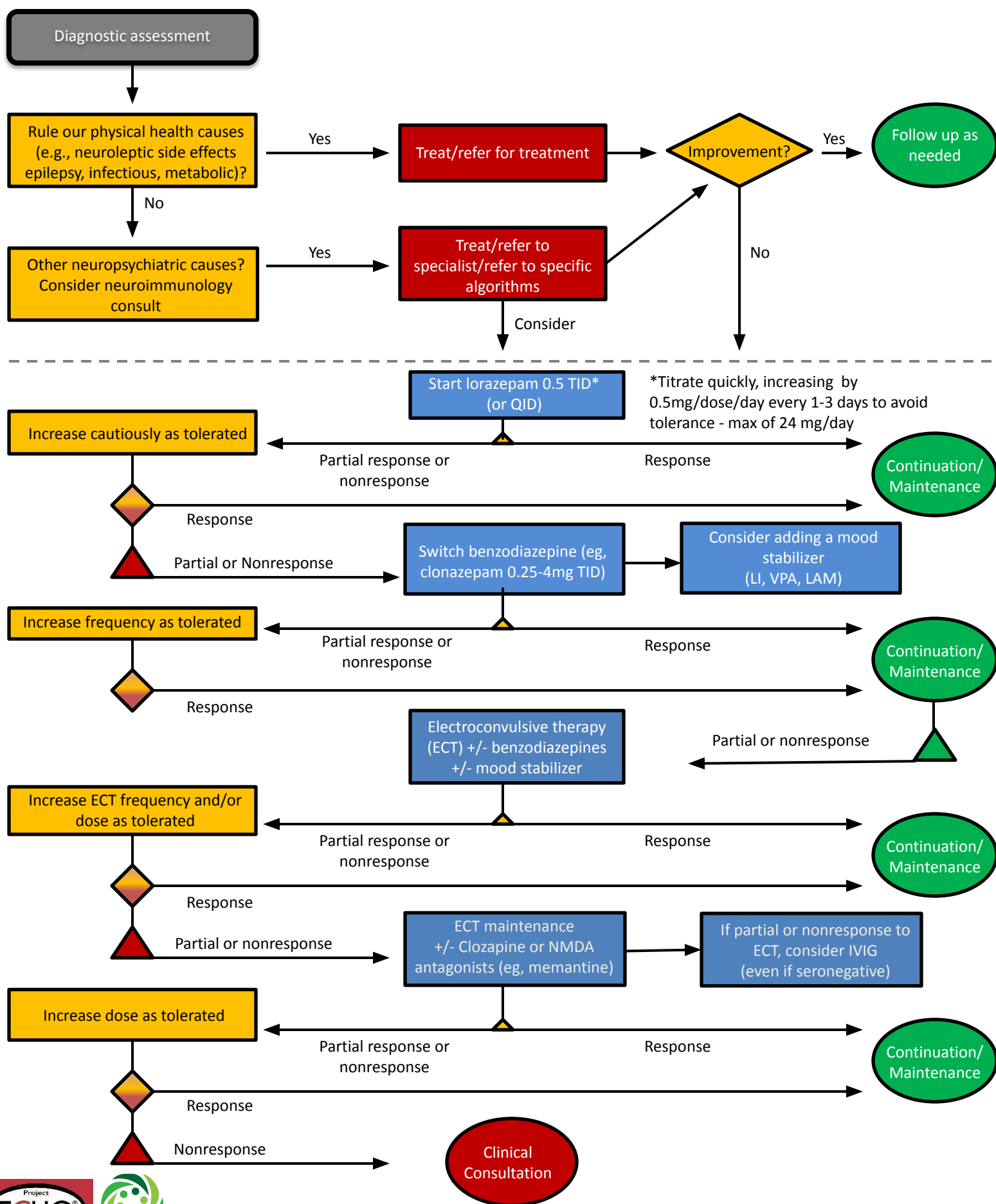
*These recommendations are not established as "evidence-based." rev 3.1, 12/2024

Expert Consensus Recommendations* for the Pharmacological Management of Anxiety in Phelan-McDermid Syndrome



*These recommendations are not established as "evidence-based." rev 3.1, 12/2024

Expert Consensus Recommendations* for the Pharmacological Management of Catatonia** in Phelan-McDermid Syndrome



*These recommendations are not established as “evidence-based.” rev 3.1, 12/2024

**** Notes for the Treatment Algorithm for the Pharmacological Management of Catatonia in Phelan-McDermid Syndrome**

- Would recommend starting lorazepam 0.5-1 mg TID, and increasing by 0.5 mg TID every few days, based on response.
- Track frequency of catatonia symptoms objectively to carefully guide titration; increase lorazepam until symptom improvement plateaus, or until the point of over-sedation.
- Consider QID dosing, especially if there is sedation or “break-through” symptoms.
- Monitor vital signs closely and if unstable, expedite to urgent referral for ECT.
- If no response to benzodiazepines, ECT alone is the next step assuming symptom severity warrants it. If the patient is simply prompt dependent with psychomotor retardation, ECT may not be indicated.
- If only PARTIAL response to benzodiazepines, consider ECT while remaining on the benzodiazepine, and using flumazenil reversal. It is NOT necessary to taper the benzodiazepine.
- If there is PARTIAL response to benzodiazepines, and the remaining symptoms do not warrant ECT, consider adjunctive antidepressant or mood stabilizer, depending on the underlying psychopathology.
- Acute ECT needs to be delivered at least three times weekly with BILATERAL electrode placement and monitoring for seizure quality.
- If inadequate response to bilateral ECT, consult with an expert to address ECT technical parameters and associated medications to improve seizure quality.
- Every patient who responds to ECT requires medication to decrease maintenance ECT frequency while maintaining clinical stability.
- Once the patient is on twice weekly ECT, start lithium and titrate to a therapeutic serum level as you decrease ECT frequency.
- If no improvement with ECT, consider neuroimmunology work-up and IVIG, even if seronegative.
- Note that with stuporous catatonia, as symptoms improve, mania may emerge and should not be considered a worsening of the condition.

****Clinical Guidelines to Initiate Treatment of Catatonia in Phelan-McDermid Syndrome**

Any person with PMS who is in their usual state of health and experiences the onset of at least three of the following symptoms according to the Diagnostic and Statistical Manual for Mental Disorders Fifth Edition (DSM-5, APA, 2013):

- (1) stupor (i.e., no psychomotor activity; not actively relating to environment);
- (2) catalepsy (i.e., passive induction of a posture held against gravity);
- (3) waxy flexibility (i.e., slight, even resistance to positioning by examiner);
- (4) mutism (i.e., no, or very little, verbal response);
- (5) negativism (i.e., opposition or no response to instructions or external stimuli);
- (6) posturing (i.e., spontaneous and active maintenance of a posture against gravity);
- (7) mannerisms (i.e., odd, circumstantial caricature of normal actions);
- (8) stereotypy (i.e., repetitive, abnormally frequent, non-goal-directed movements);
- (9) agitation, not influenced by external stimuli;
- (10) grimacing;
- (11) echolalia (i.e., mimicking another's speech);
- (12) echopraxia (i.e., mimicking another's movements)

*The Diagnostic Manual - Intellectual Disability, Second Edition (DM-ID2; Barnhill et al., 2017) notes that mutism, mannerisms, stereotypies, and grimacing can also be features of intellectual disability, and that echolalia can be a feature of ASD, so the history and time of onset of these symptoms is critical to delineate.

*****Clinical Guidelines to Initiate a Laboratory-Based Assessment to rule out Encephalitis in Phelan-McDermid Syndrome**

Presence of encephalopathy defined as:

- (1) depressed or altered level of consciousness lasting 24 hours OR
- (2) lethargy OR
- (3) personality change

PLUS at least 1 of the following:

- (a) fever;
- (b) seizure;
- (c) focal neurological findings;
- (d) CSF pleocytosis;
- (e) EEG or neuroimaging findings “consistent with encephalitis”

Any person with PMS who is in their usual state of health and experiences the onset of several of the following symptoms accompanied by new focal neurological signs, or in the absence of focal neurological signs, any person whose psychiatric symptoms fail to respond to appropriate trials of psychiatric medications *may* warrant a work-up to exclude encephalitis:

- A marked change in sleep patterns;
- Symptoms characteristic of mania or depression;
- New, intense anxiety (obsessive compulsive symptoms, separation anxiety, phobias);
- Loss of previous abilities; general confusion; disorientation;
- New and unusual motor patterns, such as changes in gait, difficulty making transitions across visual borders, and loss of hand skills;
- New incontinence;
- Note that multiple recurrent episodes may occur.

*****Protocol for the Laboratory-Based Assessment of Catatonia and/or Encephalitis in Phelan-McDermid Syndrome**

Blood Tests:

Comprehensive metabolic panel (CMP)
Complete blood count (CBC) and Differential
Serum Iron, Total Iron Binding Capacity (TIBC), Iron saturation
Erythrocyte sedimentation rate (ESR)
C-reactive protein (CRP)
Vitamin B12 level
Vitamin B6 level
Vitamin D level
Folate level
Free T4 and Thyroid Stimulating Hormone
Serum homocysteine, total
Celiac serology
Fluorescent ANA
Strep titer - if not done in the past three months or history of recurrent strep throats
Anti-thyroid antibodies: Thyroglobulin Ab and Thyroid Peroxidase (TPO) Ab
^aSerum Autoimmune Encephalopathy Evaluation (*profile + reflex tests*)

Cerebral spinal fluid studies using lumbar puncture are recommended if the patient has: (1) seizures; (2) no response to standard treatment of catatonia; (3) new onset movement disorder (e.g., choreiform movements)

Cerebral Spinal Fluid Studies:

^bNeopterin (HPLC Fluorescence)
^aSerum and CSF: IgG, albumin, Oligo bands (Mayo Clinic Multiple Sclerosis Panel)
^cRoutine CSF measures: gram stain, culture cell counts, glucose, and protein
^aSpinal Autoimmune Encephalopathy Evaluation (*profile + reflex tests*)

Other studies:

Brain MRI with or without contrast (must be done prior to lumbar puncture)
Electroencephalogram (EEG) - overnight if possible or extended if indicated. Routine EEG
Is adequate if clinical suspicion is low and there is no evidence of clinical seizures

^aMayo Clinic Laboratories - <https://www.mayocliniclabs.com/>

^bMNG Laboratories - <https://mnglabs.com/>

^cIf any evidence of CNS infection, follow standard of care procedures

Useful References

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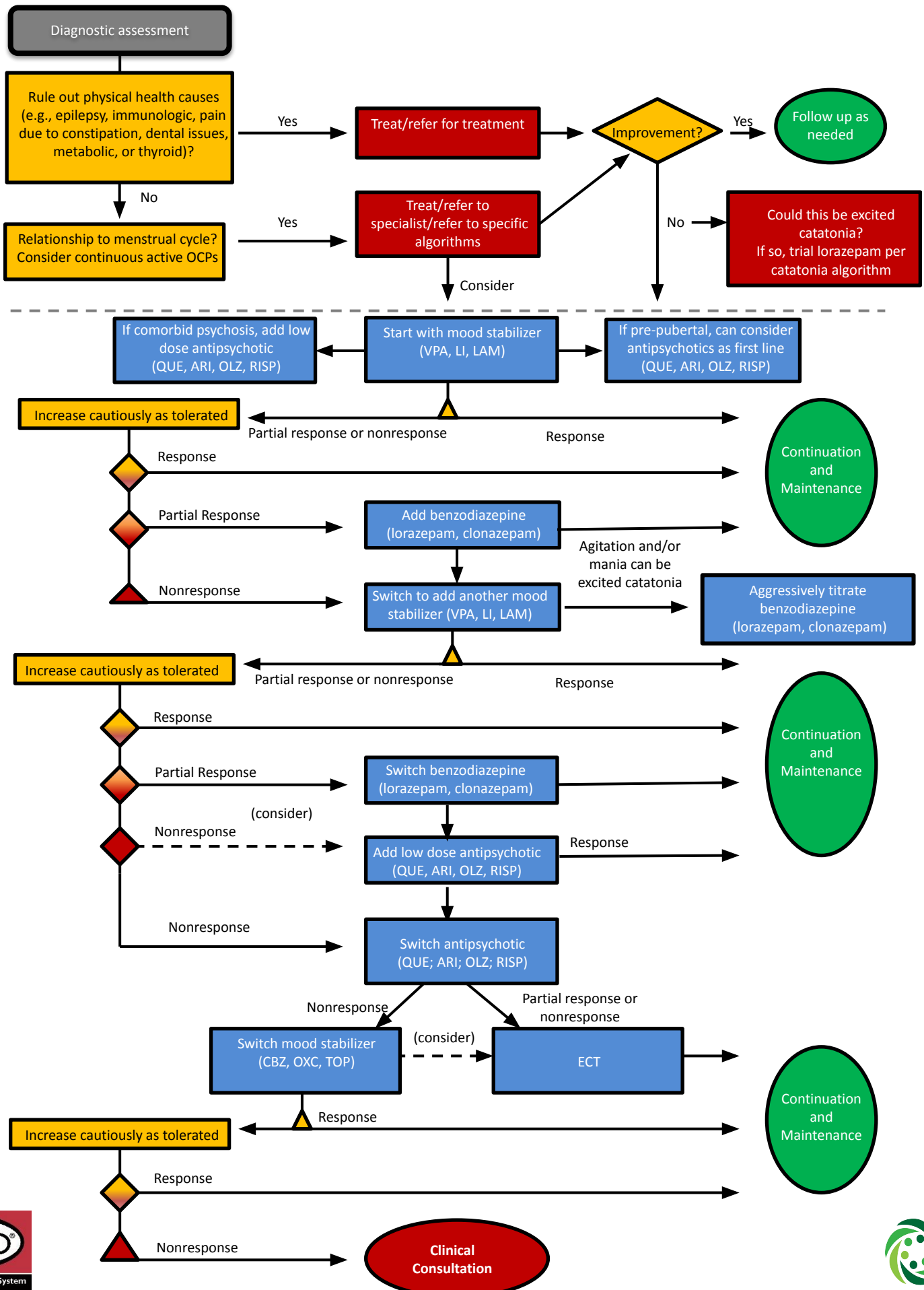
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Expert Consensus Recommendations* for the Pharmacological Management of Mood Cycling in Phelan-McDermid Syndrome



*These recommendations are not established as “evidence-based.” rev 3.1 12/2024

People with PMS can develop repeated episodes of mania which include symptoms of profound insomnia, agitation, aggression, self-injury and new repetitive behaviors:

- Sometimes mania may be accompanied by psychotic symptoms, including paranoia and presumed perceptual disturbances.
- Mania in PMS often begins with the onset of puberty.

People with PMS who experience manic symptoms often have three additional features of concern that require careful monitoring:

1. High rates of catatonia which can be stuporous or hyper-motoric/excited or alternating between the two patterns
2. Severe negative reactions to antipsychotic medications, including new onset or worsening catatonia and neuroleptic malignant syndrome
3. High rates of behavioral activation, including mania, when exposed to antidepressants

Because of accumulating experience with this clinical presentation, our group strongly advises the following:

- Avoid antidepressants
- Avoid antipsychotics when possible; use low doses to address core psychotic symptoms when present
- Treat mood cycling with lithium or anticonvulsants (e.g., divalproex sodium or lamotrigine) and note that some patients may require two agents
- Address sleep disturbance during mania with trazodone up to 100 mg, or with clonidine; see accompanying protocol for more details
- Perform careful evaluations for catatonia and consider an empiric lorazepam trial according to the accompanying protocol if suspected

Expert Consensus Recommendations* for the Pharmacological Management of Sleep Disturbance in Phelan-McDermid Syndrome

