IS IT TIME TO TAKE A DIFFERENT TREATMENT APPROACH TO TARDIVE DYSKINESIA (TD)?

ANTICHOLINERGICS ARE NOT RECOMMENDED AND MAY AGGRAVATE EXISTING TD^{1,2}

According to the benztropine package insert2:

- Benztropine is **not recommended** for use in patients with TD
- Antiparkinsonism agents **do not alleviate** the symptoms of TD, and in some instances **may aggravate** them
- Benztropine is indicated as an adjunct in the treatment of parkinsonism and is useful in the control of extrapyramidal disorders (other than TD) due to neuroleptic drugs



2020 American Psychiatric Association (APA) guidelines¹

 Anticholinergic medications do not improve and may even worsen TD

2013 American Academy of Neurology (AAN) guidelines³

 There are insufficient data to recommend anticholinergics for the treatment of TD

IT IS IMPORTANT TO DIFFERENTIATE TD FROM ACUTE EXTRAPYRAMIDAL SYMPTOMS (EPS), AS EACH REQUIRES UNIQUE MANAGEMENT^{1,2,4}

	CONSIDER TD	CONSIDER ACUTE EPS
WHEN did onset occur?	Delayed Generally emerges 3 months to years after initiating antipsychotics ⁵⁻⁷	Acute Generally emerges hours, days, or weeks after initiating antipsychotics ⁵⁻⁷
WHAT does it look like?	 Athetoid – Slow, snake-like, and writhing movements^{5,8} Choreiform – Rapid and jerky movements^{5,8} Involuntary movements often seen in the tongue, lower face, jaw, trunk, and upper and lower limbs^{5,9} Movements persist for at least a few weeks⁵ 	Akathisia – Restlessness, an inner urge to move, fidgety movements of the legs, rocking from foot to foot, pacing, and inability to sit or stand still ⁵ Dystonia – Abnormal and prolonged contraction of the muscles of the eyes, head, neck, limbs, or trunk ⁵ Parkinsonism – Parkinsonian tremor, muscular rigidity, disturbed posture, difficulty moving and walking, and slowing of movement ^{5,6}
HOW may it change?	 May be associated with discontinuation of antipsychotics^{5,6} May be temporarily masked by antipsychotic dose increase^{6,10} May be worsened by anticholinergics^{8,11} 	 May resolve days after discontinuation of antipsychotics⁶ May be improved by anticholinergics⁷

Clinical guidelines and recommendations for optimizing the management of TD



Screen regularly for TD

2020 APA guidelines¹

- Screen for TD before starting or changing patients' DRBA treatment
- Monitor for signs of TD at each visit
- Conduct structured TD assessment every 6 to 12 months, depending on patient's risk, and if new or worsening movements are detected at any visit
- 4 Consider a diagnostic evaluation

2020 Delphi panel consensus recommendations12

- A clinical assessment for TD should be performed at every clinical encounter in all patients taking antipsychotics or DRBAs, regardless of the degree or risk for TD
- Consider possible TD in any patient with even mild movements (≥2 on AIMS) in one body area

DRBA, dopamine receptor blocking agent; AIMS, Abnormal Involuntary Movement Scale.



Preserve stable antipsychotic regimens

2013 AAN guidelines³

The 2013 AAN guidelines indicate that there is a lack of clear evidence to support or refute withdrawing causative agents or switching from first-generation to second-generation antipsychotics to treat TD.



Consider first-line treatment with a VMAT2 inhibitor

2020 APA guidelines¹

- Treatment with a VMAT2 inhibitor is recommended in patients with moderate to severe or disabling TD
- VMAT2 inhibitors can also be considered in patients with mild TD

Systematic review of evidence through 201813

 New-generation VMAT2 inhibitors should be recommended as first-line therapy

2020 Delphi panel consensus recommendations¹²

• Treatment with a VMAT2 inhibitor should be considered as part of a comprehensive treatment plan

VMAT2, vesicular monoamine transporter 2.

VMAT2 inhibitors are considered first-line treatment for TD13



Talk to your patients about managing their TD

There are FDA-approved treatment options. Learn about one at TDtreatmentoption.com.

REFERENCES: 1. Keepers GA, Fochtmann LJ, Anzia JM, et al. *The American Psychiatric Association Practice Guideline for the Treatment of Patients With Schizophrenia*. 3rd ed. American Psychiatric Association Publishing, 2020. https://psychiatryonline.org/doi/pdf/10.1176/appi.books.9780890424841.

Accessed September 1, 2020. 2. Benztropine mesylate [package insert]. Lake Forest, IL: Akorn; 2017. 3. Summary of evidence-based guidelines for clinicians: treatment of tardive syndromes. American Academy of Neurology website. https://www.aan.com/Guidelines/Home/GetGuidelineContent/613. Published 2013. Accessed August 22, 2018. 4. Ward KM, Citrome L. Antipsychotic-related movement disorders: drug-induced parkinsonism vs. tardive dyskinesia-key differences in pathophysiology and clinical management. *Neurol Ther*. 2018;7(2):233-248. 5. American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders*. 5th ed. Arlington, VA: American Psychiatric Association; 2013. 6. Caroff SN, Hurford I, Lybrand J, et al. Movement disorders induced by antipsychotic drugs: implications of the CATIE schizophrenia trial. *Neurol Clin*. 2011;29(1):127-148. 7. Pierre JM. Extrapyramidal symptoms with atypical antipsychotics: incidence, prevention and management. *Drug Saf*. 2005;28(3):191-208. 8. Task Force on Tardive Dyskinesia. *Tardive Dyskinesia: A Task Force Report of the American Psychiatric Association*. Washington, DC: American Psychiatric Association; 1992. 9. Guy W. *ECDEU Assessment Manual for Psychopharmacology: Revised 1976*. Rockville, MD: National Institute of Mental Health; 1976. 10. Egan MF, Apud J, Wyatt RJ. Treatment of tardive dyskinesia. *Schizophr Bels*. 2005;80(1):33-43. 12. Caroff SN, Citrome L, Meyer J, et al. A modified Delphi consensus study of the screening, diagnosis, and treatment of tardive dyskinesia. *J Clin Psychiatry*. 2020;81(2):19cs12983. 13. Bhidayasiri R, Jitkritsadakul O, Friedman JH, Fahn OS. Updating the recommendations for treatment of tardive syndromes: a systematic review of

