Despite improvements in the tolerability of antipsychotic medications, the development of tardive dyskinesia (TD) still is a significant area of concern; however, clinicians have had few treatment options. Valbenazine, a vesicular monoamine transport type 2 (VMAT2) inhibitor, is the only FDA-approved medication for TD (Table 1).1 By modulating dopamine transport into presynaptic vesicles, synaptic dopamine release is decreased, thereby reducing the postsynaptic stimulation of D2 receptors and the severity of dyskinetic movements.

In the pivotal 6-week clinical trial, valbenazine significantly reduced TD severity as measured by Abnormal Involuntary Movement Scale (AIMS) ratings.2 Study completion rates were high (87.6%), with only 2 dropouts because of adverse events in each of the placebo (n = 78) and 40-mg (n = 76) arms, and 3 in the 80-mg group (n = 80).

Before the development of valbenazine, tetrabenazine was the only effective option for treating TD. Despite tetrabenazine’s known efficacy for TD, it was not available in the United States until 2008 with the sole indication for movements related to Huntington’s disease. U.S. patients often were subjected to a litany of ineffective medications for TD, often at great expense. Moreover, tetrabenazine involved multiple daily dosing, required cytochrome P450 (CYP) 2D6 genotyping for doses >50 mg/d, had significant tolerability issues, and a monthly cost of $8,000 to $10,000. The availability of an agent that is effective for TD and does not have tetrabenazine’s kinetic limitations, adverse effect profile, or CYP2D6 monitoring requirements represents an enormous advance in the treatment of TD.

Clinical implications
Tardive dyskinesia remains a significant public health concern because of the increasing use of antipsychotics for disorders beyond the core indication for schizophrenia. Although exposure to dopamine D2 antagonism could result in postsynaptic receptor upregulation and supersensitivity, this process best explains what underlies withdrawal dyskinesia.3 The persistence of TD symptoms in 66% to

Table 1

<table>
<thead>
<tr>
<th>Fast facts about valbenazine</th>
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<tr>
<td><strong>Brand name:</strong> Ingrezza</td>
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<tr>
<td><strong>Class:</strong> VMAT2 antagonist</td>
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<tr>
<td><strong>Indication:</strong> Tardive dyskinesia</td>
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<tr>
<td><strong>Approval date:</strong> April 11, 2017</td>
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<td><strong>Availability date:</strong> May 1, 2017</td>
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<tr>
<td><strong>Manufacturer:</strong> Neurocrine Biosciences</td>
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<td><strong>Dosing forms:</strong> 40-mg and 80-mg capsules</td>
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<td><strong>Recommended dosage:</strong> 40 mg in the morning, increasing to 80 mg/d after 1 week</td>
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Disclosure
Dr. Meyer is a consultant to Acadia Pharmaceuticals, Neurocrine Biosciences, Inc., Teva Pharmaceutical Industries; and a speaker for Acadia Pharmaceuticals, Alkermes, Allergan, Merck, Otsuka America, Inc., and Sunovion Pharmaceuticals.
80% of patients after discontinuing offending agents has led to hypotheses that the underlying pathophysiology of TD might best be conceptualized as a problem with neuroplasticity. As with many disorders, environmental contributions (e.g., oxidative stress) and genetic predisposition might play a role beyond that related to exposure to D2 antagonism.3

There have been trials of numerous agents, but no medication has been FDA-approved for treating TD, and limited data support the efficacy of a few existing medications (clonazepam, amantadine, and ginkgo biloba extract [EGb-761]),4 albeit with small effect sizes. A medical food, consisting of branched-chain amino acids, received FDA approval for the dietary management of TD in males, but is no longer commercially available except from compounding pharmacies.5

Tetrabenazine, a molecule developed in the mid-1950s to improve on the tolerability of reserpine, was associated with significant adverse effects such as orthostasis.6 Like reserpine, tetrabenazine subsequently was found to be effective for TD7 but without the peripheral adverse effects of reserpine. However, the kinetics of tetrabenazine necessitated multiple daily doses, and required CYP2D6 genotyping for doses >50 mg/d.8

Receptor blocking. The mechanism that differentiated reserpine’s and tetrabenazine’s clinical properties became clearer in the 1980s when researchers discovered that transporters were necessary to package neurotransmitters into the synaptic vesicles of presynaptic neurons.9 The vesicular monoamine transporter (VMAT) exists in 2 isoforms (VMAT1 and VMAT2) that vary in distribution, with VMAT1 expressed mainly in the peripheral nervous system and VMAT2 expressed mainly in monoaminergic cells of the central nervous system.10

Tetrabenazine’s improved tolerability profile was related to the fact that it is a specific and reversible VMAT2 inhibitor, while reserpine is an irreversible and nonselective antagonist of both VMAT isoforms. Investigation of tetrabenazine’s metabolism revealed that it is rapidly and extensively converted into 2 isomers, α-dihydrotetrabenazine (DH-TBZ) and β-DH-TBZ. The isomeric forms of DH-TBZ have multiple chiral centers, and therefore numerous forms of which only 2 are significantly active at VMAT2.3 The α-DH-TBZ isomer is metabolized via CYP2D6 and 3A4 into inactive metabolites, while β-DH-TBZ is metabolized solely via 2D6.3 Because of the short half-life of DH-TBZ when generated from oral tetrabenazine, the existence of 2D6 polymorphisms, and the predominant activity deriving from only 2 isomers, a molecule was synthesized (valbenazine), that when metabolized would slowly be converted into the most active isomer of α-DH-TBZ designated as NBI-98782 (Table 2, page 42). This slower conversion to NBI-98782 from valbenazine (compared with its formation from oral tetrabenazine) yielded improved kinetics and permitted once-daily dosing; moreover, because the metabolism of NBI-98782 is not solely dependent on CYP2D6, the need for genotyping was removed. Neither of the 2 metabolites from valbenazine NBI-98782 and NB-136110 have significant affinity for targets other than VMAT2.11

Use in tardive dyskinesia. Recommended starting dosage is 40 mg once daily with or without food, increased to 80 mg after 1 week, based on the design and results from the phase-III clinical trial.12 The FDA granted breakthrough therapy designation for this compound, and only 1 phase-III trial was performed. Valbenazine produced significant improvement on the AIMS, with a mean 30% reduction in AIMS scores at the Week 6 endpoint from baseline of 10.4 ± 3.6.2 The effect size was large (Cohen’s d = 0.90) for the 80-mg dosage. Continuation of 40 mg/d may be considered for some patients based on tolerability, including those who are known CYP2D6 poor metabolizers, and those tak-
out of the pipeline

Valbenazine and its 2 metabolites lack affinity for receptors other than VMAT2, leading to an absence of orthostasis in clinical trials.\(^1,2\) In the phase-II trial, 76% of participants receiving valbenazine (n = 51) were titrated to the maximum dosage of 75 mg/d. Common adverse reactions (incidence ≥5% and at least twice the rate of placebo) were headache (9.8% vs 4.1% placebo), fatigue (9.8% vs 4.1% placebo), and somnolence (5.9% vs 2% placebo).\(^1\) In the phase-III trial, participants were randomized 1:1:1 to valbenazine, 40 mg (n = 72), valbenazine, 80 mg (n = 79), or placebo (n = 76). In the clinical studies the most common diagnosis was schizophrenia or schizoaffective disorder, and 40% and 85% of participants in the phase-II and phase-III studies, respectively, remained on antipsychotics.\(^1,2\)

There were no adverse effects with an incidence ≥5% and at least twice the rate of placebo in the phase-III trial.\(^2\)

When data from all placebo-controlled studies were pooled, only 1 adverse effect occurred with an incidence ≥5% and twice that of placebo, somnolence with a rate of 10.9% for valbenazine vs 4.2% for placebo. The incidence of akathisia in the pooled analysis was 2.7% for valbenazine vs 0.5% for placebo. Importantly, in neither study was there a safety signal related to depression, suicidal ideation and behavior, or parkinsonism. There also were no clinically significant changes in measures of schizophrenia symptoms.

The mean QT prolongation for valbenazine in healthy participants was 6.7 milliseconds, with the upper bound of the double-sided 90% confidence interval reaching 8.4 milliseconds. For those taking strong 2D6 or 3A4 inhibitors, or known 2D6 poor metabolizers, the mean QT prolongation was 11.7 milliseconds (14.7 milliseconds upper bound of double-sided 90% CI). In the controlled trials, there was a dose-related increase in prolactin, alkaline phosphatase, and bilirubin. Overall, 3% of valbenazine-treated patients and 2%
of placebo-treated patients discontinued because of adverse reactions.

As noted above, there were no adverse effects with an incidence ≥5% and at least twice the rate of placebo in the phase-III valbenazine trial. Aggregate data across all placebo-controlled studies found that somnolence was the only adverse effect that occurred with an incidence ≥5% and twice that of placebo (10.9% for valbenazine vs 4.2% for placebo). As a comparison, rates of sedation and akathisia for tetrabenazine were higher in the pivotal Huntington’s disease trial: sedation/somnolence 31% vs 3% for placebo, and akathisia 19% vs 0% for placebo.

How it works
Tetrabenazine, a selective VMAT2 inhibitor, is the only agent that has demonstrated significant efficacy and tolerability for TD management; however, its complex metabolism generates numerous isomers of the metabolites α-DH-TBZ and β-DH-TBZ, of which only 2 are significantly active (Table 3). By choosing an active isomer (NBI-98782) as the metabolite of interest because of its selective and potent activity at VMAT2 and having a metabolism not solely dependent on CYP2D6, a compound was generated (valbenazine) that when metabolized slowly converts into NBI-98782.

Pharmacokinetics
Valbenazine demonstrates dose-proportional pharmacokinetics after single oral dosages from 40 to 300 mg with no impact of food or fasting status on levels of the active metabolite. Valbenazine has a Tmax of 0.5 to 1.0 hours, with 49% oral bioavailability. The plasma half-life for valbenazine and for NBI-98782 ranges from 15 to 22 hours. The Tmax for NBI-98782 when formed from valbenazine occurs between 4 and 8 hours, with a Cmax of approximately 30 ng/mL. It should be noted that when NBI-98782 is generated from oral tetrabenazine, the mean half-life and Tmax are considerably shorter (6 hours and 1.5 hours, respectively), while the Cmax is much higher (approximately 77 ng/mL) (Table 4, page 44).

Valbenazine is metabolized through endogenous esterases to NBI-98782 and NBI-136110. NBI-98782, the active metabolite, is further metabolized through multiple CYP pathways, predominantly 3A4 and 2D6. Neither valbenazine nor its metabolites are inhibitors or inducers of major CYP enzymes. Aside from VMAT2, the results of in vitro studies suggest that valbenazine and its active metabolite are unlikely to inhibit most major drug transporters at clinically relevant concentrations. However, valbenazine increased digoxin levels because of inhibition of intestinal P-glycoprotein; therefore plasma digoxin level monitoring is recommended when these 2 are co-administered.

Efficacy
Efficacy was established in a 6-week, fixed-dosage, double-blind, placebo-controlled trial of adult patients with TD. Eligible participants had:
• DSM-IV diagnosis of antipsychotic-induced TD for ≥3 months before screening and moderate or severe TD, as indicated by AIMS item 8 (severity of abnormal

### Clinical Point

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### Table 3

VMAT2 Binding affinities of valbenazine and its 2 metabolites

<table>
<thead>
<tr>
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<th>VMAT2 binding affinity</th>
<th>Binding profile</th>
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<tbody>
<tr>
<td>Valbenazine (NBI-98854)</td>
<td>150 nM</td>
<td>Antagonist</td>
</tr>
<tr>
<td>NBI-98782</td>
<td>3 nM</td>
<td>Antagonist</td>
</tr>
<tr>
<td>NBI-136110</td>
<td>220 nM</td>
<td>Antagonist</td>
</tr>
</tbody>
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VMAT2: vesicular monoamine transport type 2
Source: Reference 2

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Clinical Point

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Exclusion criteria included clinically significant and unstable medical conditions within 1 month before screening; comorbid movement disorder (eg, parkinsonism, akathisia, truncal dystonia) that was more prominent than TD; and significant risk for active suicidal ideation, suicidal behavior, or violent behavior. Participants had a mean age of 56, 52% were male, and 65.7% of participants in the valbenazine 40-mg group had a schizophrenia spectrum disorder diagnosis, as did 65.8% in both the placebo and valbenazine 80-mg arms.

Antipsychotic treatments were permitted during the trial and >85% of participants continued taking these medications during the study. Participants (N = 234) were randomly allocated in a 1:1:1 manner to valbenazine 40 mg, 80 mg, or matched placebo. The primary outcome was change in AIMS total score (items 1 to 7) assessed by central, independent raters. Baseline AIMS scores were 9.9 ± 4.3 in the placebo group, and 9.8 ± 4.1 and 10.4 ± 3.6 in the valbenazine 40-mg and 80-mg arms, respectively.

Outcome. A fixed-sequence testing procedure to control for family-wise error rate and multiplicity was employed, and the primary endpoint was change from baseline to Week 6 in AIMS total score (items 1 to 7) for valbenazine 80 mg vs placebo. Valbenazine, 40 mg, was associated with a 1.9 point decrease in AIMS score, while valbenazine, 80 mg, was associated with a 3.2 point decrease in AIMS score, compared with 0.1 point decrease for placebo (P < .05 for valbenazine, 40 mg, P < .001 for valbenazine, 80 mg). This difference for the 40-mg dosage did not meet the prespecified analysis endpoints; however, for the 80-mg valbenazine dosage, the effect size for this difference (Cohen’s d) was large 0.90. There also were statistically significant differences between 40 mg and 80 mg at weeks 2, 4, and 6 in the intent-to-treat population. Of the 79 participants, 43 taking the 80-mg dosage completed a 48-week extension. Efficacy was sustained in this group; however, when valbenazine was discontinued at Week 48, AIMS scores returned to baseline after 4 weeks.

Tolerability

Of the 234 randomized patients, 205 (87.6%) completed the 6-week trial. Discontinuations due to adverse events were low across all treatment groups: 2.6% and 2.8% in the placebo and valbenazine 40-mg arms, respectively, and 3.8% in valbenazine 80-mg cohort. There was no safety signal based on changes in depression, suicidality, parkinsonism rating, or changes in schizophrenia symptoms. Because valbenazine can cause somnolence, patients should not perform activities requiring mental alertness (eg, operating a vehicle or hazardous machinery) until they know how they will be affected by valbenazine.

Valbenazine should be avoided in patients with congenital long QT syn...
drome or with arrhythmias associated with a prolonged QT interval. For patients at increased risk of a prolonged QT interval, assess the QT interval before increasing the dosage.

Clinical considerations
Unique properties. Valbenazine is metabolized slowly to a potent, selective VMAT2 antagonist (NBI-98782) in a manner that permits once daily dosing, removes the need for CYP2D6 genotyping, and provides significant efficacy.

Why Rx? The reasons to prescribe valbenazine for TD patients include:

- currently the only agent with FDA approval for TD
- fewer tolerability issues seen with the only other effective agent, tetrabenazine
- no signal for effects on mood parameters or rates of parkinsonism
- lack of multiple daily dosing and possible need for 2D6 genotyping involved with TBZ prescribing.

Dosing
The recommended dosage of valbenazine is 80 mg/d administered as a single dose with or without food, starting at 40 mg once daily for 1 week. There is no dosage adjustment required in those with mild to moderate renal impairment; however, valbenazine is not recommended in those with severe renal impairment. The maximum dose is 40 mg/d for those with moderate or severe hepatic impairment (Child-Pugh score, 7 to 15) however, valbenazine is not recommended for patients with severe renal impairment (creatinine clearance <30 mL/min) because the exposure to the active metabolite is reduced by approximately 75%. The combined efficacy and tolerability of dosages >80 mg/d has not been evaluated. Adverse effects seen with tetrabenazine at higher dosages include akathisia, anxiety, insomnia, parkinsonism, fatigue, and depression.

A daily dose of 40 mg may be considered for some patients based on tolerability, including those who are known CYP 2D6 poor metabolizers, and those taking strong CYP2D6 inhibitors. For those taking strong 3A4 inhibitors, the maximum daily dose is 40 mg. Concomitant use of valbenazine with strong 3A4 inducers is not recommended as the exposure to the active metabolite is reduced by approximately 75%. Lastly, because VMAT2 inhibition may alter synaptic levels of other monoamines, it is recommended that valbenazine not be administered with monoamine

Bottom Line
Valbenazine is the only medication, other than tetrabenazine, to demonstrate efficacy for tardive dyskinesia (TD) in placebo-controlled studies. Valbenazine is well tolerated, and at the recommended dosages is not associated with development of depression or suicidality, insomnia, or parkinsonism. The binding profile of the active metabolite NBI-98782 is marked by high affinity and selectivity for VMAT2 receptors, a feature that appears to be effective for the treatment of TD, with no appreciable affinity for adrenergic, dopaminergic, or serotonergic receptors.
oxidase inhibitors, such as isocarboxazid, phenelzine, or selegiline.

**Contraindications**

There are no reported contraindications for valbenazine. As with most medications, there is limited available data on valbenazine use in pregnant women; however, administration of valbenazine to pregnant rats during organogenesis through lactation produced an increase in the number of stillborn pups and postnatal pup mortalities at doses under the maximum recommended human dose (MRHD) using body surface area based dosing (mg/m²). Pregnant women should be advised of the potential risk to a fetus. Valbenazine and its metabolites have been detected in rat milk at concentrations higher than in plasma after oral administration of valbenazine at doses 0.1 to 1.2 times the MRHD (based on mg/m²). Based on animal findings of increased perinatal mortality in exposed fetuses and pups, women are advised not to breastfeed during valbenazine treatment and for 5 days after the final dose. No dosage adjustment is required for geriatric patients.

**References**