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Introduction and General Principles

The use of psychotropic medications by children and youth is an issue confronting parents, other caregivers, and health care professionals across the United States. Children and youth in foster care, in particular, have multiple needs, including those related to emotional or psychological stress. They typically have experienced abusive, neglectful, serial or chaotic caretaking environments. Birth family history is often not available. These children often present with a fluidity of different symptoms over time reflective of past traumatic events that may mimic many psychiatric disorders and result in difficulties with attachment, mood regulation, behavioral control, and other areas of functioning.

Because of the complex issues involved in the lives of foster children, it is important that a comprehensive evaluation be performed before beginning treatment for a mental or behavioral disorder. Except in the case of an emergency, a child should receive a thorough health history, psychosocial assessment, mental status exam, and physical exam before prescribing a psychotropic medication. The physical assessment should be performed by a physician or another healthcare professional qualified to perform such an assessment. It is recognized that in some emergency situations, it may be in the best interest of the child to prescribe psychotropic medications before a physical exam can actually be performed. In these situations, a thorough health history should be performed to assess for significant medical disorders and past response to medications, and a physical evaluation should be performed as soon as possible. A thorough psychosocial assessment should be performed by an appropriately qualified mental health clinician (masters or doctoral level), a psychiatrist/child psychiatrist, or a primary care physician with experience in providing mental health care to children and youth. The child’s symptoms and functioning should be assessed across multiple domains, and the assessment should be developmentally age appropriate. It is very important that information about the child’s history, including history of trauma and current functioning be made available to the treating physician in a timely manner, either through an adult who is well-informed about the child or through a comprehensive medical record. It is critical to meet the individual needs of patients and their families in a culturally competent manner. This indicates a need to address communication issues as well as differences in perspective on issues such as behavior and mental functioning.

Interpretation of clinical symptoms and decisions concerning treatment should, whenever possible, be informed by the child’s developmental history of trauma, neglect or abuse and the timing of these stressors. In general, optimal outcomes are achieved with well-coordinated team based care with members of different professions (e.g., child psychiatrist, child psychologist, social worker, primary care physician, etc.) each contributing their particular expertise to the treatment plan and follow-up. Additionally, at present there are no biomarkers to assist with the diagnosis of mental disorders, and imaging (e.g., MRI) and other tests (e.g., EEG) are not generally helpful in making a clinical diagnosis of a mental disorder.

The role of non-pharmacological interventions should be considered before beginning a psychotropic medication, except in urgent situations such as suicidal ideation, psychosis, self-injurious behavior, physical aggression that is acutely dangerous to others, or severe impulsivity endangering the child or others; when there is marked disturbance of psychophysiological functioning (such as profound sleep disturbance), or when the child shows marked anxiety, isolation, or withdrawal. Given the history of trauma, unusual stress and change in environmental circumstances associated with being a child in foster care, psychotherapy should generally begin before or concurrent with prescription of a psychotropic medication. Referral for trauma-informed, evidence-based psychotherapy should be considered when available and appropriate. Equally important, the role of the health care provider and the health care environment’s potential to exacerbate a child’s symptoms, given their respective trauma history, should be considered and minimized. Patient and caregiver education should be provided about the condition to be treated, treatment options (non-pharmacological and pharmacological), treatment expectations, and potential side effects that may occur during the prescription of psychotropic medications.

It is recognized that many psychotropic medications do not have Food and Drug Administration (FDA) approved labeling for use in children. The FDA has a statutory mandate to determine whether pharmaceutical company sponsored research indicates that a medication is safe and effective for those indications that are listed in the approved product labeling. The FDA assures that information in the approved product labeling is accurate, and limits the manufacturer’s marketing to the information contained in the approved labeling. The FDA does not regulate physician and other health provider practice. In fact,
Psychotropic Medication Utilization Parameters

During the prescription of psychotropic medication, certain parameters should be considered to ensure the best possible outcome for the patient. Except in cases of emergency, whenever possible, trauma-informed, actively engaged, and documented in the medical record. To that end, clear documentation of the physician’s rationale in the electronic medical record facilitates continuity of care and minimizes misinterpretation.

Role of Primary Care Providers

Primary care providers play a valuable role in the care of youth with mental disorders. Not only are they the clinicians most likely to initially interact with children who are in need of medication due to an emotional or psychiatric disorder, but also an inadequate number of child psychiatrists are available to meet all children’s mental health needs. Primary care clinicians are in an excellent position to perform screenings of children for potential mental disorders, and they should be able to diagnose and treat relatively straightforward situations such as uncomplicated ADHD, anxiety, or depression. Primary care providers should provide advice to youth in foster care and their care givers about handling feelings and behaviors, recognizing the need for help, making decisions regarding healthy life styles, and the available treatments for childhood mental disorders. As always, consideration should be given regarding the need for referral for counseling, psychotherapy, or behavioral therapy. Primary care providers vary in their training, clinical experience, and confidence to address mental disorders in children. Short courses and intensive skills oriented seminars may be beneficial in assisting primary care clinicians in caring for children with mental disorders. Active liaisons with child psychiatrists who are available for phone consultation or referral can be beneficial in assisting primary care clinicians to meet the mental health needs of children. “The management of common presentations of ADHD, depression and anxiety, psychotherapy referral, psychopharmacology and appropriate child psychiatry referral are within the scope of general pediatric practice” (Southammososane 2015). In addition, the American Academy of Pediatrics has recently provided a policy statement (“Health Care Issues for Children and Adolescents in Foster Care and Kinship Care”) which can be found at: http://pediatrics.aappublications.org/content/136/4/e1131

General principles regarding the use of psychotropic medications in children include:

- A DSM-5 psychiatric diagnosis should be made before the prescribing of psychotropic medications.
- Clearly defined target symptoms and treatment goals for the use of psychotropic medications should be identified and documented in the medical record at the time of or before beginning treatment with a psychotropic medication. These target symptoms and treatment goals should be assessed at each clinic visit with the child and caregiver in a culturally and linguistically appropriate manner. Whenever possible, standardized clinical rating scales (clinician, patient, primary caregiver, teachers, and other care providers) or other measures should be used to quantify the response of the child’s target symptoms to treatment and the progress made toward treatment goals.
- In making a decision regarding whether to prescribe a psychotropic medication in a specific child, the clinician should carefully consider potential side effects, including those that are uncommon but potentially severe, and evaluate the overall benefit to risk ratio of pharmacotherapy.
- Except in the case of an emergency, informed consent should be obtained from the appropriate party(s) before beginning psychotropic medication. Informed consent to treatment with psychotropic medication entails diagnosis, expected benefits and risks of treatment, including common side effects, discussion of laboratory findings, and uncommon but potentially severe adverse events. Alternative treatments, the risks associated with no treatment, and the overall potential benefit to risk ratio of treatment should be discussed.
- Whenever possible, trauma-informed, evidence-based psychotherapy, should begin before or concurrent with the prescription of psychotropic medication.
- Before starting psychopharmacological treatment in preschool-aged children even more emphasis should be placed on treatment with non-psychopharmaceutical interventions. Assessment of parent functioning and mental health needs, in addition to training parents in evidence-based behavior management can also reduce the need for the use of medication.
- Medication management should be collaborative. Youth, as well as caregivers, should be involved in decision-making about treatment, in accordance with their developmental level. Parents providing informed consent should be engaged, and where applicable, other caregivers, family, and child related agencies should be involved.
- During the prescription of psychotropic medication, the presence or absence of medication side effects should be documented in the child's medical record at each visit.
- Appropriate monitoring of indices such as height, weight, blood pressure, or laboratory findings should be documented.
- Monotherapy regimens for a given disorder or specific target symptoms should usually be tried before polypharmacy regimens. While the goal is to use as few psychotropic medications as can be used to appropriately address the child’s clinical status, it is recognized that the presence of psychiatric comorbidities may affect the number of...
psychotropic medications that are pre-
scribed. When polypharmacy regimens are needed, addition of medications should occur in a systematic orderly process, accompanied by on-going monitoring, evaluation, and documentation. The goal remains to minimize polypharmacy while maximizing therapeutic outcomes.

- Medications should be initiated at the lower end of the recommended dose range and titrated carefully as needed.

- Only one medication should be changed at a time, unless a clinically appropriate reason to do otherwise is documented in the medical record. (Note: starting a new medication and beginning the dose taper of a current medication is considered one medication change).

- The use of “prn” or as needed prescriptions is discouraged. If they are used, the situation indicating need for the administration of a prn medication should be clearly indicated as well as the maximum dosage in a 24 hour period and in a week. The frequency of administration should be monitored to assure that these do not become regularly scheduled medications unless clinically indicated.

- The frequency of clinician follow-up should be appropriate for the severity of the child’s condition and adequate to monitor response to treatment, including: symptoms, behavior, function, and potential medication side effects. At a minimum, a child receiving psychotropic medication should be seen by the clinician at least once every ninety days.

- The potential for emergent suicidality should be carefully evaluated and monitored, particularly in depressed children and adolescents as well as those initiating antidepressants, those having a history of suicidal behavior or deliberate self-harm and those with a history of anxiety or substance abuse disorders.

- If the prescribing clinician is not a child psychiatrist, referral to or consultation with a child psychiatrist, or a general psychiatrist with significant experience in treating children, should occur if the child’s clinical status has not shown meaningful improvement within a timeframe that is appropriate for the child’s diagnosis and the medication regimen being used.

- Before adding additional psychotropic medications to a regimen, the child should be assessed for adequate medication adherence, appropriateness of medication daily dosage, appropriateness of the diagnosis, the occurrence of comorbid disorders (including substance abuse and general medical disorders), and the influence of psychosocial stressors.

- If a medication has not resulted in improvement in a child’s target symptoms (or rating scale score), discontinue that medication rather than adding a second medication to it.

- If a medication is being used in a child for a primary target symptom of aggression associated with a DSM-5 non-psychotic diagnosis (e.g., conduct disorder, oppositional defiant disorder, intermittent explosive disorder), and the behavior disturbance has been in remission for six months, then serious consideration should be given to slow tapering and discontinuation of the medication. If the medication is continued in this situation, the necessity for continued treatment should be evaluated and documented in the medical record at a minimum of every six months.

- The clinician should clearly document care provided in the child’s medical record, including history, mental status assessment, physical findings (when relevant), impressions, rationale for medications prescribed, adequate laboratory monitoring specific to the drug(s) prescribed at intervals required specific to the prescribed drug and potential known risks, medication response, presence or absence of side effects, treatment plan, and intended use of prescribed medications.

### Use of Psychotropic Medication in Preschool Age Children

The use of psychotropic medication in young children of preschool ages is a practice that is limited by the lack of evidence available for use of these agents in this age group. The Preschool Psychopharmacology Working Group (PPWG) published guidelines (Gleason 2007) summarizing available evidence for use of psychotropic medications in this age group. The PPWG was established in response to the clinical needs of preschoolers being treated with psychopharmacological agents and the absence of systematic practice guidelines for this age group, with its central purpose to attempt to promote an evidence-based, informed, and clinically sound approach when considering medications in preschool-aged children.

The PPWG guidelines emphasize consideration of multiple different factors when deciding on whether to prescribe psychotropic medications to preschool-aged children. Such factors include the assessment and diagnostic methods utilized in evaluating the child for psychiatric symptoms/illness, the current state of knowledge regarding the impact of psychotropic medication use on childhood neurodevelopmental processes, the regulatory and ethical contexts of use of psychotropic medications in small children (including available safety information and FDA status), and the existing evidence base for use of psychotropic medication in preschool aged children.

The publication includes specific guidelines and algorithm schematics developed by the PPWG to help guide treatment decisions for a number of psychiatric disorders that may present in preschool-aged children, including Attention-Deficit Hyperactivity Disorder, Disruptive Behavioral Disorders, Major Depressive Disorder, Bipolar Disorder, Anxiety Disorders, Post-Traumatic Stress Disorder, Obsessive-Compulsive Disorder, Pervasive Developmental
Disorders, and Primary Sleep Disorders. The working group’s key points and guidelines are similar to the general principles regarding the use of psychotropic medication in children already detailed in this paper. However, the working group’s algorithms put more emphasis on treating preschool-aged children with non-psychopharmacological interventions (for up to 12 weeks) before starting psychopharmacological treatment, in an effort to be very cautious in introducing psychopharmacological interventions to rapidly developing preschoolers.

The working group also emphasizes the need to assess parent functioning and mental health needs, in addition to training parents in evidence-based behavior management, since parent behavior and functioning can have a large impact on behavior and symptoms in preschool-aged children.

**Distinguishing between Levels of Warnings Associated with Medication Adverse Effects**

Psychotropic medications have the potential for adverse effects, some that are treatment limiting. Some adverse effects are detected prior to marketing, and are included in the FDA approved product labeling provided by the manufacturers. When looking at product labeling, these adverse effects will be listed in the “Warnings and Precautions” section. As well, the “Adverse Reactions” section of the product labeling will outline those adverse effects reported during clinical trials, as well as those discovered during post-marketing evaluation. Many tertiary drug information resources also list common adverse effects and precautions for use with psychotropic medications.

At times, post-marketing evaluation may detect critical adverse effects associated with significant morbidity and mortality. The Food and Drug Administration (FDA) may require manufacturers to revise product labeling to indicate these critical adverse effects. If found to be particularly significant, these effects are demarcated by a box outlining the information at the very beginning of the product labeling, and have, in turn, been named boxed warnings. Boxed warnings are the strongest warning required by the FDA. It is important for clinicians to be familiar with all medication adverse effects, including boxed warnings, in order to appropriately monitor patients and minimize the risk of their occurrence. The medication tables include boxed warnings as well as other potential adverse effects. The list of potential adverse effects in the tables should not be considered exhaustive, and the clinician should consult the FDA approved product labeling and other reliable sources for information regarding medication adverse effects.

The FDA has in recent years taken additional measures to try to help patients avoid serious adverse events. New guides called Medication Guides have been developed, and are specific to particular medication and medication classes. Medication Guides advise patients and caregivers regarding possible adverse effects associated with classes of medications, and include precautions that they or healthcare providers may take while taking/prescribing certain classes of medications. The FDA requires that Medication Guides be issued with certain prescribed medications and biological products when the Agency determines that certain information is necessary to prevent serious adverse effects, that patient decision-making should be informed by information about a known serious side effect with a product, or when patient adherence to directions for the use of a product are essential to its effectiveness. During the drug distribution process, if a Medication Guide has been developed for a certain class of medications, then one must be provided with every new prescription and refill of that medication.

Copies of the Medication Guides for psychotropic medications can be accessed on the FDA website at:

http://www.fda.gov/Drugs/DrugSafety/ucm085729.htm

**Usual Recommended Doses of Common Psychotropic Medications**

The attached medication charts are intended to reflect usual doses and brief medication information for commonly used psychotropic medications. The tables contain two columns for maximum recommended doses in children and adolescents — the maximum recommended in the FDA approved product labeling, and the maximum recommended in medical and pharmacological literature sources. The preferred drug list of medications potentially prescribed for foster children is the same as for all other Texas Medicaid recipients.

The tables are intended to serve as a resource for clinicians. The tables are not intended to serve as comprehensive drug information references or a substitute for sound clinical judgment in the care of individual patients. Circumstances may dictate the need for the use of higher doses in specific patients. In these cases, careful documentation of the rationale for the higher dose should occur, and careful monitoring and documentation of response to treatment should be performed. If the use of higher medication doses does not result in improvement in the patient’s clinical status within a reasonable time period (e.g., 2-4 weeks), then the dosage should be decreased and other treatment options considered.

Not all medications prescribed by clinicians for psychiatric diagnoses in children and adolescents are included in the following tables. However, in general, medications not listed do not have adequate efficacy and safety information available to support a usual maximum dose recommendation.

See Psychotropic Medication Tables beginning on page 8.
Criteria Indicating Need for Further Review of a Child’s Clinical Status

The following situations indicate a need for review of a patient’s clinical care. These parameters do not necessarily indicate that treatment is inappropriate, but they do indicate a need for further review.

For a child being prescribed a psychotropic medication, any of the following suggests the need for additional review of a patient’s clinical status:

1. Absence of a thorough assessment for the DSM-5 diagnosis(es) in the child’s medical record

2. Four (4) or more psychotropic medications prescribed concomitantly (side effect medications are not included in this count)

3. Prescribing of:
   - Two (2) or more concomitant stimulants *
   - Two (2) or more concomitant alpha agonists *
   - Two (2) or more concomitant antidepressants
   - Two (2) or more concomitant antipsychotics
   - Three (3) or more concomitant mood stabilizers

   * The prescription of a long-acting and an immediate-release stimulant or alpha agonist of the same chemical entity does not constitute concomitant prescribing.

   Note: When switching psychotropics, medication overlaps and cross taper should occur in a timely fashion, generally within 4 weeks.

4. The prescribed psychotropic medication is not consistent with appropriate care for the patient’s diagnosed mental disorder or with documented target symptoms usually associated with a therapeutic response to the medication prescribed.

5. Psychotropic polypharmacy (2 or more medications) for a given mental disorder is prescribed before utilizing psychotropic monotherapy

6. The psychotropic medication dose exceeds usual recommended doses (literature based maximum dosages in these tables).

7. Psychotropic medications are prescribed for children of very young age, including children receiving the following medications with an age of:
   - Stimulants: Less than three (3) years of age
   - Alpha Agonists: Less than four (4) years of age
   - Antidepressants: Less than four (4) years of age
   - Mood Stabilizers: Less than four (4) years of age
   - Antipsychotics: Less than five (5) years of age

8. Prescribing by a primary care provider who has not documented previous specialty training for a diagnosis other than the following (unless recommended by a psychiatrist consultant):
   - Attention Deficit Hyperactive Disorder (ADHD)
   - Uncomplicated anxiety disorders
   - Uncomplicated depression

9. Antipsychotic medication(s) prescribed continuously without appropriate monitoring of glucose and lipids at least every 6 months.
## Stimulants for treatment of ADHD

<table>
<thead>
<tr>
<th>Drug (generic)</th>
<th>Drug (brand)+</th>
<th>Initial Dosage</th>
<th>Literature Based Maximum Dosage</th>
<th>FDA Approved Maximum Dosage for Children and Adolescents</th>
<th>Schedule</th>
<th>Patient Monitoring Parameters</th>
<th>Black Box Warning**</th>
<th>Warnings and Precautions</th>
</tr>
</thead>
</table>
| Amphetamine mixed salts* | Adderall® | • Age 3-5 years: 2.5 mg/day  
Age ≥ 6 years: 5-10 mg/day | Age 3-5 years: 30 mg/day | Approved for children 3 years and older: 40 mg/day | One to three times daily | | | |
| | Evekeo® | Age 3-5 years: 5mg/day  
Age 6-12 years: 5-10 mg/day  
Age ≥13 years: 10 mg/day | Age ≥ 6 years: >50 kg: 60 mg/day | Approved for children 6 years and older: 30 mg/day | Once daily | | | |
| | Adderall®XR | Age 3-5 years: 30 mg/day | | | | | | |
| Amphetamine base | Adzenys®XR-ODT  
(oral disintegrating tablet) | • Age ≥ 6 years: 6.3 mg/day  
(3.1 mg = 5 mg Adderall®XR) | Age 6-12 years: 18.8 mg/day  
Age 13-17 years: 12.5 mg/day | Approved for children 6 years and older:  
• Ages 6-12 years: 18.8 mg/day  
• Ages 13-17 years: 12.5 mg/day | Once daily | Baseline and ongoing: height, weight, heart rate, and blood pressure  
Baseline: Assessment using a targeted cardiac history of the child and the family, and a physical examination of the child with an EKG and/or a pediatric cardiology consult as indicated | | • Abuse potential  
• Sudden death and serious cardiovascular events (Only boxed warning for amphetamine salts and dextroamphetamine)  
• Hypertension  
• Potential for psychiatric adverse events (hallucinations, delusional thinking, mania, aggression, etc.)  
• Stimulants do not appear to affect ultimate adult height. If mild growth suppression occurs, it is likely reversible upon discontinuation of stimulant  
• Tics  
• Decreased appetite and weight  
• Sleep disturbance |
| | Dyanavel®XR  
(oral suspension) | • Age ≥ 6 years: 2.5-5 mg/day  
(2.5 mg = 4 mg Adderall®XR) | Age ≥ 6 years: 20 mg/day | Approved for children 6 years and older: 20 mg/day | Once daily | | | |
| Dextroamphetamine* | Dexedrine® | • Age 3-5 years: 2.5 mg/day | Age 3-5 years: 30 mg/day | Approved for children 3 years and older: 40 mg/day | Once or twice daily | | | |
| | Zerozed® | Age ≥ 6 years: 5 mg twice daily | Age ≥ 6 years: >50 kg: 80 mg/day | Age ≥ 6 years: 40 mg/day | | | | |
| | Procentra®  
(oral suspension) | | | | | | | |
| | Dexedrine Spansule® | • Age 3-5 years: 5 mg/day  
Age ≥ 6 years: 5 mg/day | Age ≥ 6 years: >50 kg: 80 mg/day | | | | | |
| | | | | | | | | |
| Lisdexamfetamine | Vyvanse® | • Age 3-5 years: 10 mg/day  
Age ≥ 6 years: 30 mg/day | Age 3-5 years: 30 mg/day  
Age ≥ 6 years: 70 mg/day | Approved for children 6 years and older: 70 mg/day | Once daily | | | |

* Generic available  
** See the FDA approved product labeling for each medication for the full black box warnings.  
+ XR, extended-release

(Continued on Page 9)

March 2016  
(Tables Updated July 2016)
<table>
<thead>
<tr>
<th>Drug (generic)</th>
<th>Drug (brand)*</th>
<th>Initial Dosage</th>
<th>Literature Based Maximum Dosage</th>
<th>FDA Approved Maximum Dosage for Children and Adolescents</th>
<th>Schedule</th>
<th>Baseline/ Monitoring</th>
<th>Black Box Warning</th>
<th>Warnings and Precautions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ritalin®</td>
<td>Age 3-5 years: 2.5 mg twice daily</td>
<td>Age 3-5 years: 20 mg/day</td>
<td>Approved for children 6 years and older: 60 mg/day</td>
<td>One to three times daily</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Methylin®</td>
<td>Age ≥ 6 years: 5 mg twice daily</td>
<td>Age ≥ 6 years: ≤50 kg: 60 mg/day</td>
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<td></td>
</tr>
<tr>
<td>Methylin®</td>
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<td>Age ≥ 6 years: &gt;50 kg: 100 mg/day</td>
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<tr>
<td>Methylin®SR</td>
<td>Age ≥ 3 years: 10 mg/day</td>
<td>Age 3-5 years: 20 mg/day</td>
<td>Approved for children 6 years and older: 60 mg/day</td>
<td>Once daily</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Methylin®ER</td>
<td></td>
<td>Age ≥ 6 years: 10-20 mg/day</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Metadate®ER</td>
<td>Age ≥ 3-5 years: 10 mg/day</td>
<td>Age 3-5 years: 20 mg/day</td>
<td>Approved for children 6 years and older: 60 mg/day</td>
<td>Once daily</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Metadate®CD</td>
<td>Age ≥ 6 years: 10 mg/day</td>
<td>Age ≥ 6 years: 108 mg/day</td>
<td>Approved for children 6 years and older: Age 6-12 years: 54 mg/day</td>
<td>Once daily</td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Quillivant®XR (oral suspension)</td>
<td>Age ≥ 3-5 years: 10 mg/day</td>
<td>Age 3-5 years: 36 mg</td>
<td>Approved for children 6 years and older: Age 6-12 years: 54 mg/day</td>
<td>Once daily</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>QuilliChew®ER (chewable)</td>
<td>Age ≥ 6 years: 10-20 mg/day</td>
<td>Age ≥ 6 years: 108 mg/day</td>
<td>Approved for children 6 years and older: Age 6-12 years: 54 mg/day</td>
<td>Once daily</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Concerta®</td>
<td>Age ≥ 3 years: 18 mg/day</td>
<td>Age 3-5 years: 36 mg</td>
<td>Approved for children 6 years and older: Age 6-12 years: 54 mg/day</td>
<td>Once daily</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Daytrana®TD patch*</td>
<td>Age ≥ 3 years: 10 mg/day</td>
<td>Age 3-5 years: 20 mg</td>
<td>Approved for children 6 years and older: Age 6-12 years: 54 mg/day</td>
<td>Once daily</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Dexmethylphenidate*</td>
<td>Age ≥ 3-5 years: 2.5 mg/day</td>
<td>Age 3-5 years: 10 mg/day</td>
<td>Approved for children 6 years and older: Age 6-12 years: 20 mg/day</td>
<td>Twice daily</td>
<td></td>
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<tr>
<td>Focalin®</td>
<td>Age ≥ 6 years: 5 mg twice daily</td>
<td>Age 3-5 years: 10 mg/day</td>
<td>Approved for children 6 years and older: Age 6-12 years: 20 mg/day</td>
<td>Once daily</td>
<td></td>
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<tr>
<td>Focalin®XR</td>
<td>Age ≥ 6 years: 50 mg/day</td>
<td>Age ≥ 6 years: 50 mg/day</td>
<td>Approved for children 6 years and older: Age 6-12 years: 50 mg/day</td>
<td>Once daily</td>
<td></td>
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<td></td>
<td></td>
</tr>
</tbody>
</table>

* Generic available
** See the FDA approved product labeling for each medication for the full black box warnings.
+ IR, immediate release; SR, sustained-release formulation; CD, combined immediate release and extended release; ER and XR, extended-release; LA, long-acting; TD, transdermal
+ Daytrana®TD patch: Post marketing reports of acquired skin depigmentation or hypopigmentation of the skin
## Other ADHD Treatments

<table>
<thead>
<tr>
<th>Drug (generic)</th>
<th>Drug (brand)*</th>
<th>Initial Dosage</th>
<th>Literature Based Maximum Dosage</th>
<th>FDA Approved Maximum Dosage for Children and Adolescents</th>
<th>Schedule</th>
<th>Baseline/Monitoring</th>
<th>Black Box Warning</th>
<th>Warnings and Precautions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Atomoxetine</td>
<td>Strattera®</td>
<td>Age ≥ 6 years and weight ≤70 kg: 0.5 mg/kg/day / Age ≥ 6 years and weight &gt;70 kg: 40 mg/day</td>
<td>Age ≥ 6 years: 1.8 mg/kg/day or 100 mg/day, whichever is less</td>
<td>Approved for treatment of ADHD (age 6-17 years): 1.4 mg/kg/day or 100 mg/day, whichever is less</td>
<td>Once or twice daily</td>
<td>Baseline and ongoing: height, weight, heart rate, and blood pressure / Onset of therapeutic effect typically delayed 3 weeks</td>
<td>Suicidal ideation in children and adolescents being treated for ADHD</td>
<td>Severe liver injury / Contraindicated to use within 14 days of an MAOI / Increased blood pressure and heart rate / Psychiatric adverse events / Priapism (rare)</td>
</tr>
<tr>
<td>Clonidine*</td>
<td>Catapres® (IR)</td>
<td>Age ≥ 6 years and weight &lt;45 kg: 0.05 mg/day / Age ≥ 6 years and weight &gt;45 kg: 0.1 mg/day</td>
<td>Age ≥ 6 years AND Weight 27-40.5 kg: 0.2 mg/day / Weight 40.5-45 kg: 0.3 mg/day / Weight &gt;45 kg: 0.4 mg/day</td>
<td>Not approved for treatment of ADHD in children and adolescents</td>
<td>One to four times daily</td>
<td>Baseline and ongoing: heart rate and blood pressure / Personal and family cardiovascular history</td>
<td>None</td>
<td>Hypotension / Bradycardia / Syncope / Sedation/Somnolence / Taper, do not discontinue abruptly</td>
</tr>
<tr>
<td>Guanfacine*</td>
<td>Kapvay® (ER)</td>
<td>Age ≥ 6 years: 0.1 mg/day</td>
<td>Age ≥ 6 years: 0.4 mg/day</td>
<td>Approved for monotherapy and adjunctive therapy to stimulants for treatment of ADHD (age 6-17 years): 0.4 mg/day</td>
<td>Once or twice daily</td>
<td>Baseline and ongoing: heart rate and blood pressure / Personal and family cardiovascular history</td>
<td>None</td>
<td></td>
</tr>
<tr>
<td>Tenex® (IR)</td>
<td>Intuniv® (ER)</td>
<td>Age ≥ 6 years: 1 mg/day / Age 6-12 years: 4 mg/day / Age 12-17 years: 7 mg/day</td>
<td>Age ≥ 6-12 years: 4 mg/day / Age 12-17 years: 7 mg/day</td>
<td>Approved for monotherapy and adjunctive therapy to stimulants for treatment of ADHD</td>
<td>Once daily</td>
<td>Baseline and ongoing: heart rate and blood pressure / Personal and family cardiovascular history</td>
<td>None</td>
<td></td>
</tr>
<tr>
<td>Wellbutrin®</td>
<td>Wellbutrin®</td>
<td>Age ≥ 6 years: 3 mg/kg/day or 150 mg/day, whichever is less</td>
<td>Age ≥ 6 years: 6 mg/kg/day or 300 mg/day with no single dose &gt;150 mg, whichever is less</td>
<td>Not approved for children and adolescents</td>
<td>One to three times daily</td>
<td>Blood pressure and Pulse / Mental status exam and suicide assessment</td>
<td>Increased risk of suicidal thinking and behavior (suicidality) in short-term studies in children and adolescents with major depressive disorder (MDD) and other psychiatric disorders</td>
<td>Lowers seizure threshold (use caution with other agents that may lower seizure threshold-e.g. antipsychotics, TCA’s, excessive alcohol) / Discontinuation syndrome / Activation of mania/ hypomania / Suicidal ideation / Contraindicated for use within 14 days of an MAOI</td>
</tr>
<tr>
<td>Bupropion*</td>
<td>Wellbutrin®SR</td>
<td>Same as above</td>
<td>400 mg/day</td>
<td>Not approved for children and adolescents</td>
<td>Once or twice daily</td>
<td>Baseline and ongoing: heart rate and blood pressure / Personal and family cardiovascular history</td>
<td>None</td>
<td></td>
</tr>
<tr>
<td>Nortriptyline*</td>
<td>Wellbutrin®XL</td>
<td>Same as above</td>
<td>450 mg/day</td>
<td></td>
<td>Once daily</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Imipramine*</td>
<td>Tofranil®</td>
<td>Age ≥ 6 years: 1 mg/kg/day or 25 mg/day, whichever is less</td>
<td>Age ≥ 6 years: 4 mg/kg/day or 200 mg/day, whichever is less</td>
<td>Approved for treatment of enuresis in children</td>
<td>Twice daily</td>
<td>CBC / Blood pressure and Pulse / EKG / Mental status exam and suicide assessment</td>
<td>Caution with cardiac disease / Cardiac conduction abnormalities / Orthostatic hypotension / Activation of mania/hypomania / Anticholinergic and cognitive adverse effects / Lowers seizure threshold / Discontinuation syndrome / Suicidal ideation / Contraindicated for use within 14 days of an MAOI / Use caution in those with history of suicide attempts; may be cardiotoxic in overdose</td>
<td></td>
</tr>
<tr>
<td>Nortriptyline*</td>
<td>Aventyl®</td>
<td>Age ≥ 6 years: 0.5 mg/kg/day</td>
<td>Age ≥ 6 years: 2 mg/kg/day or 100 mg/day, whichever is less</td>
<td>Not approved for children and adolescents</td>
<td>Twice daily</td>
<td>CBC / Blood pressure and Pulse / EKG / Mental status exam and suicide assessment</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* Generic available
+ IR, immediate release; SR, sustained-release formulation; ER, extended-release; XL, extended-length
## Antidepressants, SSRIs

<table>
<thead>
<tr>
<th>Drug (generic)</th>
<th>Drug (brand)*</th>
<th>Initial Dosage</th>
<th>Literature Based Maximum Dosage</th>
<th>FDA Approved Maximum Dosage for Children and Adolescents</th>
<th>Schedule</th>
<th>Patient Monitoring Parameters</th>
<th>Black Box Warning**</th>
<th>Warnings and Precautions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Citalopram*</td>
<td>Celexa®</td>
<td>• Age 6-11 years: 10 mg/day</td>
<td>• Age ≥ 6 years: 40 mg/day</td>
<td>Not approved for children and adolescents</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Age ≥ 12 years: 20 mg/day</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Esclotram*</td>
<td>Lexapro®</td>
<td>• Age 6-11 years: 5 mg/day</td>
<td>• Age 6-11 years: 20 mg/day</td>
<td>• Not approved for children and adolescents</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Age ≥ 12 years: 30 mg/day</td>
<td>• Age ≥ 12 years: 30 mg/day</td>
<td>• Approved for treatment of MDD in adolescents (age 12-17 years); 20 mg/day</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fluoxetine*</td>
<td>Prozac®</td>
<td>• Age 6-11 years: 5-10 mg/day</td>
<td>• Age ≥ 6 years: 60 mg/day</td>
<td>• Approved for treatment of MDD (age 8-18 years): 20 mg/day</td>
<td>Once daily</td>
<td></td>
<td></td>
<td>Increased risk compared to placebo of suicidal thinking and behavior (suicidality) in children, adolescents, and young adults in short-term studies of major depressive disorder (MDD) and other psychiatric disorders</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Age ≥ 12 years: 10 mg/day</td>
<td></td>
<td>Approved for treatment of OCD (age 7-17 years): 60 mg/day</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Paroxetine*</td>
<td>Paxil®</td>
<td>• Children: Not recommended</td>
<td>• Children: Not recommended</td>
<td>Not approved for children and adolescents</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Age ≥ 12 years: 10 mg</td>
<td>• Age ≥ 12 years: 40 mg</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Paxil®CR</td>
<td>• Children: Not recommended</td>
<td>• Children: Not recommended</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Age ≥ 12 years: 25 mg</td>
<td>• Age ≥ 12 years: 50 mg</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fluvoxamine*</td>
<td>Luvox®</td>
<td>Age ≥ 8 years: 25 mg/day</td>
<td>• Age 8-11 years: 200 mg/day</td>
<td>Approved for treatment of OCD (age 8-17 years): 200 mg/day</td>
<td>Daily doses &gt;50 mg should be divided</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Luvax®CR</td>
<td>Age ≥ 8 years: 100 mg/day</td>
<td>• Age 12-17 years: 300 mg/day</td>
<td>Approved for treatment of OCD (age 8-17 years): 300 mg/day</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sertraline*</td>
<td>Zoloft®</td>
<td>• Age 6-12 years: 12.5-25 mg/day</td>
<td>• Age ≥ 6 years: 200 mg/day</td>
<td>Approved for treatment of OCD (age 6-17 years): 200 mg/day</td>
<td>Once daily</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Age 13-17 years: 25-50 mg/day</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Vilazodone</td>
<td>Viibryd®</td>
<td>Insufficient Evidence</td>
<td>Insufficient Evidence</td>
<td>Not approved for children and adolescents</td>
<td></td>
<td></td>
<td>Insufficient Evidence</td>
<td></td>
</tr>
</tbody>
</table>

* Generic available

+ CR, controlled-release

** From Boxed Warning in FDA approved labeling for Antidepressants (SSRIs, SNRIs and Other Mechanisms): Patients of all ages who are started on antidepressant therapy should be monitored appropriately and observed closely for clinical worsening, suicidality, or unusual changes in behavior. Families and caregivers should be advised of the need for close observation and communication with the prescriber. Both patients and families should be encouraged to contact the clinician if depression worsens, the patient demonstrates suicidal behavior or verbalizations, or if medication side effects occur. The appropriate utilization of non-physician clinical personnel who are knowledgeable of the patient population can aid in increasing the frequency of contact between the clinic and the patient/parent.
### Antidepressants, SNRIs

<table>
<thead>
<tr>
<th>Drug (generic)</th>
<th>Drug (brand)</th>
<th>Initial Dosage</th>
<th>Literature Based Maximum Dosage</th>
<th>FDA Approved Maximum Dosage for Children and Adolescents</th>
<th>Schedule</th>
<th>Patient Monitoring Parameters</th>
<th>Black Box Warning</th>
<th>Warnings and Precautions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Venlafaxine*</td>
<td>Effexor®</td>
<td>Age 7-17 years: 37.5 mg/day</td>
<td>• Age 7-11 years: 150 mg/day</td>
<td>Not approved for children and adolescents</td>
<td>IR: Two to three times daily</td>
<td>• Pregnancy test – as clinically indicated</td>
<td>Increased risk compared to placebo of suicidal thinking and behavior (suicidality) in children, adolescents, and young adults in short-term studies of major depressive disorder (MDD) and other psychiatric disorders</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Effexor®XR</td>
<td></td>
<td>• Age 12-17 years: 375 mg/day</td>
<td></td>
<td>XR: Once daily</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Duloxetine</td>
<td>Cymbalta®</td>
<td>Age 7-17 years: 30 mg/day</td>
<td>Age 7-17 years: 120 mg/day</td>
<td>Approved for treatment of Generalized Anxiety Disorder Age 7-17 years: 120 mg/day</td>
<td>Once or twice daily</td>
<td>• Monitor for emergence of suicidal ideation or behavior</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Desvenlafaxine</td>
<td>Pristiq®</td>
<td>• Children: Insufficient Evidence</td>
<td>• Age 12-17 years: 50 mg/day</td>
<td>Not approved for children and adolescents</td>
<td>Once daily</td>
<td>• Blood pressure during dosage titration and as clinically indicated</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Levomilnacipram</td>
<td>Fetzima®</td>
<td>Insufficient Evidence</td>
<td>Age 10-17 years: 3 mg/kg/day or 200 mg/day, whichever is less</td>
<td>Approved for treatment of OCD: Age 10-17 years: 3 mg/kg/day or 200 mg/day, whichever is less</td>
<td>Once daily</td>
<td>• CBC and EKG at baseline and as clinically indicated for Clomipramine</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Clomipramine*</td>
<td>Anafranil®</td>
<td>Age 10-17 years: 25 mg/day</td>
<td>Age 10-17 years: 7.5 mg/day</td>
<td>Not approved for children and adolescents</td>
<td></td>
<td>• Monitor weight and height</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Antidepressants, Other Mechanisms

<table>
<thead>
<tr>
<th>Drug (generic)</th>
<th>Drug (brand)</th>
<th>Initial Dosage</th>
<th>Literature Based Maximum Dosage</th>
<th>FDA Approved Maximum Dosage for Children and Adolescents</th>
<th>Schedule</th>
<th>Patient Monitoring Parameters</th>
<th>Black Box Warning</th>
<th>Warnings and Precautions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mirtazapine*</td>
<td>Remeron®</td>
<td>Age ≥ 3 years: 7.5 mg/day</td>
<td>Age ≥ 3 years: 45 mg/day</td>
<td>Not approved for children and adolescents</td>
<td>Once daily</td>
<td>• Pregnancy test – as clinically indicated</td>
<td>Increased the risk compared to placebo of suicidal thinking and behavior (suicidality) in children, adolescents, and young adults in short-term studies of major depressive disorder (MDD) and other psychiatric disorders</td>
<td></td>
</tr>
<tr>
<td>Vortioxetine</td>
<td>Trintellix®</td>
<td>Insufficient Evidence</td>
<td>Insufficient Evidence</td>
<td>Not approved for children and adolescents</td>
<td>Insufficient Evidence</td>
<td>• Serum cholesterol levels</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

* Generic Available
+ XR, extended-release
# Psychotropic Medication Utilization Parameters

## Antipsychotics: Second Generation (Atypical)

<table>
<thead>
<tr>
<th>Drug (generic)</th>
<th>Drug (brand)+</th>
<th>Initial Dosage</th>
<th>Literature Based Maximum Dosage</th>
<th>FDA Approved Maximum Dosage for Children and Adolescents</th>
<th>Schedule</th>
<th>Patient Monitoring Parameters</th>
<th>Black Box Warning</th>
<th>Warnings and Precautions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abilify®</td>
<td>Abilify® (oral solution)</td>
<td>Age ≥ 4 years: 2 mg/day</td>
<td>• Age 4-11 years: 15 mg/day</td>
<td>• Approved for treatment of Bipolar Mania or Mixed Episodes (age 10-17 years) and Schizophrenia (13-17 years): 30 mg/day</td>
<td>Once daily</td>
<td>• Fasting plasma glucose level or hemoglobin A1C – at baseline, at 3 months, then every 6 months.</td>
<td>Increased the risk of suicidal thoughts and behavior in short-term studies in children, adolescents, and young adults with major depressive disorder and other psychiatric disorders.</td>
<td></td>
</tr>
<tr>
<td>discerpin®</td>
<td>discerpin® (oral disintegrating tab)</td>
<td>Age ≥ 10 years: 2.5 mg/day</td>
<td>• Age 10-17 years: 50 mg/day</td>
<td>• Approved for treatment of Intractability Associated with Autistic Disorder (age 6-17 years): 15 mg/day</td>
<td>IR: One to three times daily</td>
<td>CBC as clinically indicated.</td>
<td>None related to youth</td>
<td>Extrapyramidal side effects</td>
</tr>
<tr>
<td>loraprazole®</td>
<td>loraprazole® (brand only)</td>
<td>Age ≥ 6 years: 20 mg/day</td>
<td></td>
<td>• Approved for treatment of Schizophrenia (13-17 years): 800 mg/day</td>
<td>XR: Once daily</td>
<td>Pregnancy test – as clinically indicated.</td>
<td>None related to youth</td>
<td>Neuroleptic Malignant Syndrome</td>
</tr>
<tr>
<td>Zyprexa®</td>
<td>Zyprexa® (brand only)</td>
<td>Age ≥ 4 years: 1.25 mg/day</td>
<td>• Age 4-5 years: 12.5 mg/day</td>
<td>Approved for treatment of Bipolar Mania or Mixed Episodes and Schizophrenia (age 13-17 years): 20 mg/day</td>
<td>Once daily</td>
<td>Blood pressure, pulse rate, height, weight and BMI measurement – at every visit</td>
<td>None related to youth</td>
<td>Tardive Dyskinesia</td>
</tr>
<tr>
<td>Zydis®</td>
<td>Zydis® (brand only)</td>
<td>Age ≥ 13 years: 20 mg/day</td>
<td></td>
<td>Approved for treatment of Schizophrenia (age 13-17 years): 80 mg/day</td>
<td>IR: One to two times daily</td>
<td>• Sexual function – inquire for evidence of galactorrhea/amenorrhea, menstrual disturbance, libido disturbance or erectile/ejaculatory disturbances in males (Parson has been reported with paliperidone, risperidone and ziprasidone).</td>
<td>None related to youth</td>
<td>Hyperglycemia and Diabetes Mellitus</td>
</tr>
<tr>
<td>Clozaril®</td>
<td>Clozaril® (oral solution)</td>
<td>Age ≥ 16 years: 6.25 mg/day</td>
<td>• Age 16-17 years: 12.5 mg/day</td>
<td>Approved for treatment of Schizophrenia (age 13-17 years) and Bipolar Mania or Mixed Episodes (age 10-17 years): 600 mg/day</td>
<td>IR: One to two times daily</td>
<td>None related to youth</td>
<td>None related to youth</td>
<td>Prolactinemia and gynecomastia (most common with risperidone and paliperidone)</td>
</tr>
<tr>
<td>FazaClob® (oral disintegrating tablet)</td>
<td></td>
<td>Age ≥ 12 years: 6.25 mg/day</td>
<td></td>
<td>Approved for treatment of irritability associated with autistic disorder (age 5-16 years): 3 mg/day</td>
<td>IR: One to two times daily</td>
<td>None related to youth</td>
<td></td>
<td>Orthostatic Hypotension</td>
</tr>
<tr>
<td>Veralox® (oral suspension)</td>
<td></td>
<td>Age ≥ 10 years: 5 mg/day</td>
<td></td>
<td>Approved for treatment of Bipolar Mania or Mixed Episodes and Schizophrenia (age 13-17 years): 20 mg/day</td>
<td>IR: One to two times daily</td>
<td>None related to youth</td>
<td></td>
<td>Leukopenia, neutropenia, and agranulocytosis</td>
</tr>
<tr>
<td>Amoxapine®</td>
<td>Amoxapine® (brand only)</td>
<td>Age ≥ 10 years: 2.5 mg twice daily</td>
<td></td>
<td>Approved for acute treatment of Bipolar Mania and Mixed Episodes (age 10-17 years): 10 mg twice daily</td>
<td>Twice daily</td>
<td>Avoid eating or drinking for 10 minutes after sublingual administration</td>
<td>None related to youth</td>
<td>Lowers seizure threshold</td>
</tr>
<tr>
<td>iloperidone**</td>
<td>Fanapt® (brand only)</td>
<td>Insufficient Evidence</td>
<td>Insufficient Evidence</td>
<td>Not approved for children and adolescents</td>
<td>Insufficient Evidence</td>
<td>Vision questionnaire – ask whether the patient has experienced a change in vision and should specifically ask about distance vision and blurry vision-yearly.</td>
<td>None related to youth</td>
<td>Cognitive and motor impairment potential</td>
</tr>
<tr>
<td>Paliperidone®</td>
<td>Invegta® (brand only)</td>
<td>Children: Insufficient Evidence</td>
<td>Adolescents: Insufficient Evidence</td>
<td>Approved for treatment of Schizophrenia (age 12-17 years):</td>
<td>Insufficient Evidence</td>
<td>None related to youth</td>
<td>None related to youth</td>
<td>Hyperthermia</td>
</tr>
<tr>
<td>Ziprasidone®</td>
<td>Geodon® (brand only)</td>
<td>Bipolar Disorder (age 10-17 years): 20 mg/day</td>
<td></td>
<td>• Weight ≤ 56 kg: 40 mg/day</td>
<td>Once daily</td>
<td>None related to youth</td>
<td>None related to youth</td>
<td>Extrapyramidal side effects</td>
</tr>
<tr>
<td>Lurasidone</td>
<td>Latuda® (brand only)</td>
<td>Insufficient Evidence</td>
<td>Insufficient Evidence</td>
<td>Approved for treatment of Bipolar Mania (age 10-17 years): 30 mg/day</td>
<td>Insufficient Evidence</td>
<td>None related to youth</td>
<td>None related to youth</td>
<td>Olanzapine can cause a rare but serious skin reaction known as DRESS (Drug Reaction with Eosinophilia and Systemic Symptoms). Presence of a fever with a rash and swollen lymph glands, or swelling to the face requires immediate medical attention.</td>
</tr>
<tr>
<td>Brexpiprazole</td>
<td>Rexulti® (brand only)</td>
<td>Insufficient Evidence</td>
<td>Insufficient Evidence</td>
<td>Not approved for children and adolescents</td>
<td>Insufficient Evidence</td>
<td>None related to youth</td>
<td>None related to youth</td>
<td>Antidepressants increased the risk of suicidal thoughts and behavior in children, adolescents, and young adults in short-term studies.</td>
</tr>
</tbody>
</table>

* Generic available
+ XR, extended-release

** While iloperidone alone can cause QTc prolongation, concomitant administration with a CYP2D6 inhibitor (e.g., paroxetine) or a CYP3A4 inhibitor (e.g., ketoconazole) can double QTc prolongation compared with administering iloperidone alone.

No long-acting injectable antipsychotic formulations are FDA-approved for use in children and adolescents.

March 2016
(Tables Updated July 2016)
## Antipsychotics: First Generation (Typical)

<table>
<thead>
<tr>
<th>Drug (generic)</th>
<th>Drug (brand)</th>
<th>Initial Dosage</th>
<th>Literature Based Maximum Dosage</th>
<th>FDA Approved Maximum Dosage for Children and Adolescents</th>
<th>Schedule</th>
<th>Patient Monitoring Parameters</th>
<th>Black Box Warning</th>
<th>Warnings and Precautions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chlorpromazine*</td>
<td>Thorazine®</td>
<td>• Age &gt; 6 months: 0.25 mg/lb every 4-6 hours, as needed • Adolescents: 10-25 mg/dose every 4-6 hours</td>
<td>• Age &lt; 5 years: 40 mg/day • Age 5-12 years: 75 mg/day • Age &gt; 12 years: 800 mg/day</td>
<td>Approved for treatment of severe behavioral problems (age 6 months-12 years) • Outpatient Children: 0.55 mg/kg every 4-6 hours, as needed • Inpatient Children: 500 mg/day</td>
<td>One to six times daily</td>
<td></td>
<td>None related to youth</td>
<td>• Tardive Dyskinesia • Neuroleptic Malignant Syndrome • Leukopenia, neutropenia, and agranulocytosis • Drowsiness</td>
</tr>
<tr>
<td>Haloperidol*</td>
<td>Haldol®</td>
<td>• Age 3-12 years weighing ○ 15-40 kg: 0.025-0.05 mg/kg/day ○ ≥ 40 kg: 1 mg/day • Age ≥ 12: 1 mg/day</td>
<td>• Age 3-12 years: 0.15 mg/kg/day or 6 mg/day, whichever is less • Age &gt;12 years ○ Acute agitation: 10 mg/dose ○ Psychosis: 15 mg/day ○ Tourette’s Disorder: 15 mg/day</td>
<td>Approved for treatment of Psychotic Disorders, Tourette’s Disorder, and severe behavioral problems (age ≥3 years): • Psychosis: 0.15 mg/kg/day • Tourette’s Disorder and severe behavioral problems: 0.075 mg/kg/day • Severeely disturbed children: 6 mg/day</td>
<td>One to three times daily</td>
<td>Same as Second Generation Antipsychotics</td>
<td>None related to youth</td>
<td>• Orthostatic hypotension • EKG changes • Extrapyramidal symptoms • Ocular changes • Hyperprolactinemia • Anticholinergic effects (constipation, dry mouth, blurred vision, urinary retention)</td>
</tr>
<tr>
<td>Perphenazine*</td>
<td>Trilafon®</td>
<td>• Age 6-12 years: Insufficient Evidence • Age &gt; 12 years: 4-16 mg two to four times daily</td>
<td>• Age 6-12 years: Insufficient Evidence • Age &gt; 12 years: 64 mg/day</td>
<td>Approved for treatment of Psychotic Disorders (age ≥12 years): • Outpatient: 24 mg/day • Inpatient: 64 mg/day</td>
<td>Two to four times daily</td>
<td></td>
<td>None related to youth</td>
<td>• Risk of prolonged QTc interval and torsades de pointes (particularly with pimozide)</td>
</tr>
</tbody>
</table>
| Pimozide | Orap® | Age ≥7 years: 0.05 mg/kg | • Age 7-12 years: 6 mg/day or 0.2 mg/kg/day, whichever is less • Age ≥ 12 years: 10 mg/day or 0.2 mg/kg/day, whichever is less | Approved for treatment of Tourette’s Disorder (age ≥12 years): 10 mg/day or 0.2 mg/kg/day, whichever is less | Once or twice daily | | None | |#### *Generic available
<table>
<thead>
<tr>
<th>Drug (generic)</th>
<th>Drug (brand)+</th>
<th>Initial Dosage</th>
<th>Target Dosage Range</th>
<th>Literature Based Maximum Dosage</th>
<th>FDA Approved Maximum Dosage for Children and Adolescents</th>
<th>Schedule</th>
<th>Patient Monitoring Parameters</th>
<th>Black Box Warning</th>
<th>Warnings and Precautions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Epitol® (tab)</td>
<td>Tegretol® (tab, oral suspension, chewable)</td>
<td>• Age 4-5: 10-20 mg/kg/day</td>
<td>10-20 mg/kg/day</td>
<td>• Age 4-5: 35 mg/kg/day</td>
<td>Approved for treatment of Seizure Disorders in all ages</td>
<td>Two to four times daily</td>
<td>• CBC with differential — baseline and 1 to 2 weeks after each dose increase, annually, and as clinically indicated</td>
<td>Stevens-Johnson Syndrome</td>
<td>Aplastic anemia</td>
</tr>
<tr>
<td>Tegretol® (cr)</td>
<td>• Age 6-12: 10 mg/kg/day or 200 mg/day</td>
<td>• Ages 6-12: 400-800 mg/day</td>
<td>• Ages 6-12: 800 mg/day</td>
<td>• Age 13-15: 1000 mg/day</td>
<td>• Age 6-15: 1000 mg/day</td>
<td>Twice daily</td>
<td>• Electrolytes — baseline and 1 to 2 weeks after each dose increase, annually, and as clinically indicated</td>
<td>Suicidality</td>
<td>Teratogenicity</td>
</tr>
<tr>
<td>Carbatrol® (extended release capsule)</td>
<td>• Age ≥ 13: 400 mg/day</td>
<td>• Age ≥ 13: 1200 mg/day</td>
<td>• Age 15: 1200 mg/day</td>
<td>• Age &gt;15: 1200 mg/day</td>
<td>• Carbamazepine levels — obtain 1 week after initiation and 3-4 weeks after dose adjustment, then as clinically indicated</td>
<td>Two to three times daily</td>
<td>• Pregnancy Test — baseline as appropriate, and as clinically indicated</td>
<td>Neutropenia and granulocytosis</td>
<td></td>
</tr>
<tr>
<td>Equetro® (extended release capsule)</td>
<td>• Age ≥ 13: 400 mg/day</td>
<td>• Age ≥ 13: 1200 mg/day</td>
<td>• Age 15: 1200 mg/day</td>
<td>• Age &gt;15: 1200 mg/day</td>
<td>• Carbamazepine levels — obtain 1 week after initiation and 3-4 weeks after dose adjustment, then as clinically indicated</td>
<td>Two to three times daily</td>
<td>• For patients with Asian descent; genetic test for HLA-B*1502 at baseline (prior to the initiation of carbamazepine). May use results of previously completed testing. Patients testing positive for the allele should not use carbamazepine unless benefit outweighs the risk</td>
<td>Hyponatremia</td>
<td></td>
</tr>
<tr>
<td>Divalproex Sodium</td>
<td></td>
<td>Age ≥6: 10-15 mg/kg/day</td>
<td>Age ≥6: Serum level: 125 µg/mL, or 60 mg/kg/day</td>
<td>Maximum dose based upon serum level: 50-100 µg/mL, or 60 mg/kg/day</td>
<td>Approved for treatment of Seizure Disorders (age ≥ 10 years)</td>
<td>One to three times daily</td>
<td>• Lithium levels</td>
<td>Stevens-Johnson Syndrome</td>
<td></td>
</tr>
<tr>
<td>Depakote® delayed-release tablets</td>
<td>• Age ≥6: Lesser of 15-20 mg/kg/day or 150mg twice per day</td>
<td>• Age ≥6: Serum level: 1.2 mg/dL, or 1800 mg</td>
<td>Dose adjustment based upon serum level</td>
<td>One to four times daily</td>
<td>• Lithium Levels — one week (i.e., 5-7 days) after initiation or dosage change, 3 months after initiation, and as clinically indicated; for maintenance treatment every 8 months, and as clinically indicated</td>
<td>• For patients with a history of severe debilitation, or sodium depletion</td>
<td>Aplastic anemia/ granulocytosis</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Equetro®</td>
<td>• Age 6-11: 10 mg/kg/day or 200 mg/day</td>
<td>• Age 6-11: 40-80 mg/day</td>
<td>• Age 6-11: 80 mg/day</td>
<td>• Age 6-15: 1000 mg/day</td>
<td>• Carbamazepine levels — obtain 1 week after initiation and 3-4 weeks after dose adjustment, then as clinically indicated</td>
<td>Two to four times daily</td>
<td>• For patients with Asian descent; genetic test for HLA-A*3101 genetic testing at baseline for those to be considered at high risk (most common in Asian, Native American, European, and Latin American descents)</td>
<td>Neutropenia and granulocytosis</td>
<td></td>
</tr>
<tr>
<td>Lithium*</td>
<td>• Age ≥ 12: Lesser of 15-20 mg/kg/day or 300 mg twice per day</td>
<td>12 hour post dose serum level: 0.6-1.2 mg/dL</td>
<td>• Age ≥12: Serum level: 1.2 mg/dL, or 1800 mg</td>
<td>Maximum dose based upon 12 hour post dose serum level: 1.2 mg/dL, or 1800 mg</td>
<td>Approved for treatment of manic episodes and maintenance of Bipolar Disorder (age ≥ 12 years)</td>
<td>One to four times daily</td>
<td>• Lithium levels — one week (i.e., 5-7 days) after initiation or dosage change, 3 months after initiation, and as clinically indicated; for maintenance treatment every 8 months, and as clinically indicated</td>
<td>Aplastic anemia/ granulocytosis</td>
<td></td>
</tr>
<tr>
<td>Lithobid®(ER)</td>
<td>• Age 6-11: 10 mg/kg/day or 200 mg/day</td>
<td>• Age 6-11: 40-80 mg/day</td>
<td>• Age 6-11: 80 mg/day</td>
<td>• Age 6-15: 1000 mg/day</td>
<td>• Carbamazepine levels — obtain 1 week after initiation and 3-4 weeks after dose adjustment, then as clinically indicated</td>
<td>Two to four times daily</td>
<td>• For patients with Asian descent; genetic test for HLA-B*1502 at baseline (prior to the initiation of carbamazepine). May use results of previously completed testing. Patients testing positive for the allele should not use carbamazepine unless benefit outweighs the risk</td>
<td>Neutropenia and granulocytosis</td>
<td></td>
</tr>
</tbody>
</table>

(Continued on Page 16)
## Mood Stabilizers (continued)

<table>
<thead>
<tr>
<th>Drug (generic)</th>
<th>Drug (brand)*</th>
<th>Initial Dosage</th>
<th>Target Dosage Range</th>
<th>Literature Based Maximum Dosage for Children and Adolescents</th>
<th>FDA Approved Maximum Dosage for Children and Adolescents</th>
<th>Schedule</th>
<th>Patient Monitoring Parameters</th>
<th>Black Box Warning</th>
<th>Warnings and Precautions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lamotrigine*</td>
<td>Lamictal®</td>
<td>Age 6-11 years: 25-5 mg/day</td>
<td>Age 6-11 years:</td>
<td>• Monotherapy: 4.5-7.5 mg/kg/day</td>
<td>Approved for adjunctive therapy for Seizure Disorders:</td>
<td>Once or twice daily</td>
<td>Safety and effectiveness for treatment of Bipolar Disorder in patients younger than 18 years had not been established</td>
<td>None</td>
<td>• Dermatological reactions</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Age ≥12 years: 25 mg/day (increase by 25 mg every 2 weeks)</td>
<td>6-11 years: 1-3 mg/kg/day</td>
<td>With Valproate: 1-3 mg/kg/day</td>
<td>Age 2-12: 400 mg/day</td>
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<tr>
<td></td>
<td></td>
<td>Age ≥12 years:</td>
<td>With Valproate and EIAEDs: 1-5 mg/kg/day</td>
<td></td>
<td>Age 2-12: 500 mg/day</td>
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<tr>
<td></td>
<td></td>
<td>Age ≥12 years:</td>
<td>With EIAED’s: 5-15 mg/kg/day</td>
<td></td>
<td>Use &gt; 200 mg/day in adults for bipolar depression has not conferred additional efficacy</td>
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<td></td>
<td></td>
<td>Age ≥12 years:</td>
<td>With Valproate: 100-200 mg/day</td>
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<tr>
<td></td>
<td></td>
<td>Age ≥12 years:</td>
<td>With Valproate and EIAEDs: 100-400 mg/day</td>
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<td></td>
<td></td>
<td>Age ≥12 years:</td>
<td>With EIAEDs: 300-500 mg/day</td>
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<td></td>
<td></td>
<td>Age ≥6 years:</td>
<td>With Valproate: 100-200 mg/day</td>
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<td></td>
<td></td>
<td>Age ≥6 years:</td>
<td>With Valproate and EIAEDs: 100-400 mg/day</td>
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<tr>
<td></td>
<td></td>
<td>Age ≥6 years:</td>
<td>With Valproate: 100-400 mg/day</td>
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<tr>
<td></td>
<td></td>
<td>Age ≥6 years:</td>
<td>With Valproate and EIAEDs: 100-400 mg/day</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>** Lamotrigine clearance</td>
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<td></td>
<td></td>
<td>Age ≥6 years:</td>
<td>With Valproate: 100-400 mg/day</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>** Withdrawal seizures</td>
</tr>
<tr>
<td>Oxcarbazepine*</td>
<td>Trileptal®</td>
<td>6-10 mg/kg/day</td>
<td>Monotherapy (based on weight):</td>
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<tr>
<td></td>
<td></td>
<td>Age 7-12 years: 60 mg/kg/day or 1500 mg/day</td>
<td>• Monotherapy: 20-34.9 kg: 600-900 mg/day</td>
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<tr>
<td></td>
<td></td>
<td>Age 13-17 years: 60 mg/kg/day or 2100 mg/day</td>
<td>25-34.9 kg: 900-1200 mg/day</td>
<td>Approved for treatment of Seizure Disorders as mono-therapy (age ≥ 4 years), or as adjunctive therapy (age ≥ 2 years): 60 mg/kg/day or 1800 mg/day</td>
<td>Approved for treatment of Seizure Disorders as monotherapy (age ≥ 4 years), or as adjunctive therapy (age ≥ 2 years):</td>
<td>Twice daily</td>
<td>Safety and effectiveness for treatment of Bipolar Disorder in patients younger than 18 years had not been established</td>
<td>None</td>
<td>** Drug-drug interaction potential</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>35-44.9 kg: 1000-1500 mg/day</td>
<td>60 mg/kg/day or 1800 mg/day</td>
<td>Safety and effectiveness for treatment of Bipolar Disorder in patients younger than 18 years had not been established</td>
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<td></td>
<td>45-49 kg: 1200-1500 mg/day</td>
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<td></td>
<td>50-59.9 kg: 1200-1800 mg/day</td>
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<td></td>
<td>60-69.9 kg: 1200-2100 mg/day</td>
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<td></td>
<td>≥70 kg: 1500-2100 mg/day</td>
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<td></td>
<td></td>
<td>** Multi-organ hypersensitivity</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>** Hematologic events</td>
</tr>
</tbody>
</table>

* Generic Available
** EIAED's - Enzyme Inducing Anti-Epileptic Drugs (e.g. Carbamazepine, Phenytoin, Phenytoin, Primidone)
+EIAED’s - Enzyme Inducing Anti-Epileptic Drugs (e.g. Carbamazepine, Phenytoin, Phenytoin, Primidone)
# Sedatives/Hypnotics

<table>
<thead>
<tr>
<th>Drug (generic)</th>
<th>Drug (brand)</th>
<th>Initial Dosage</th>
<th>Literature Based Maximum Dosage(^*)</th>
<th>FDA Approved Maximum Dosage for Children and Adolescents</th>
<th>Schedule</th>
<th>Black Box Warning(^*)</th>
<th>Warnings and Precautions</th>
</tr>
</thead>
</table>
| Diphenhydramine* | Benadryl® | • Age 3-5 years: 6.25-12.5 mg (1mg/kg max)  
  • Age 5-11 years: 12.5-25 mg  
  • Age ≥12 years: 25-50 mg | • 25-37 lbs: 12.5 mg  
  • 38-49 lbs: 19 mg  
  • 50-99 lbs: 25 mg  
  • ≥100 lbs: 50 mg | Evidence suggests that tolerance develops to the hypnotic effects of diphenhydramine within 5-7 nights of continuous use. | Approved for treatment of insomnia (age ≥12 years): 50 mg at bedtime | Once at bedtime | None |
| Trazodone* | Desyrel® | • Children: Insufficient Evidence  
  • Adolescents: 25 mg | • Children Insufficient Evidence  
  • Adolescents: 100 mg/day | Not approved for children or adolescents as a hypnotic | Once at bedtime | Increased the risk compared to placebo of suicidal thinking and behavior (Suicidality) in children, adolescents, and young adults in short-term studies of major depressive disorder (MDD) and other psychiatric disorders | • Serotonin Syndrome  
• Contraindicated for use within 14 days of an MAOI  
• Suicidal ideation  
• Activation of mania/hypomania  
• Discontinuation syndrome  
• Abnormal bleeding  
• QT prolongation and risk of sudden death  
• Orthostatic hypotension and syncope  
• Abnormal bleeding  
• Priapism  
• Hypomania  
• Cognitive and motor impairment |
| Melatonin | No brand name | • Age 3-5 years: 0.5mg  
  • Age ≥6 years: 1mg | Regulated by FDA as a dietary supplement and not as a medication (no FDA approved indications) | | Once at bedtime or alternatively, give 5-6 hrs before Dim Light Melatonin Onset (DLMO) | None |
| Ramelteon | Rozerem® | • Age 3-5 years: 1mg | Not approved for children or adolescents | | | |
| Hydroxyzine* | Vistaril® | • Age 3-5 years: 25 mg  
  • Age ≥6 years: 50mg | Approved for treatment of anxiety and tension:  
• Age <6 years: 50 mg/day in divided doses  
• Age ≥6 years: 50-100 mg/day in divided doses | Approved as a sedative when used as a premedication and following general anesthesia: 0.6 mg/kg | Once at bedtime | | |

\(^*\) Generic Available  
\(^*\) Maximum doses for the sedative/hypnotics are based upon night time doses to induce sleep in a child with severe insomnia.
Glossary

ANC = ABSOLUTE NEUTROPHIL COUNT

BMI = Body Mass Index. A measure of body fat based upon height and weight.

CBC = Complete blood count. Lab test used to monitor for abnormalities in blood cells, e.g., for anemia.

Cp = Plasma concentration

Serum creatinine = A lab test used to calculate an estimate of kidney function.

EKG = Electrocardiogram

EEG = Electroencephalogram

EPS = Extrapyramidal side effects. These are adverse effects upon movement, including stiffness, tremor, and severe muscle spasm

FDA = U.S. Food and Drug Administration

Hemoglobin A1c = A laboratory measurement of the amount of glucose in the hemoglobin of the red blood cells. Provides a measure of average glucose over the previous 3 months.

LFTs = Liver function tests

MAOIs = Monoamine Oxidase Inhibitors

MRI = Magnetic resonance imaging

PRN = as needed

Prolactin = A hormone produced by the pituitary gland

TFTs = Thyroid Function Tests
**References**


Biederman et al. A prospective open-label trial of lamotrigine monotherapy in children and adolescents with bipolar disorder. CNS Neurosci Ther Apr 2010;16(2):91-102


References (continued)


References (continued)


Web Link References


When to seek referral or consultation with a child or adolescent psychiatrist. American Academy of Child and Adolescent Psychiatry, 2003. http://www.aacap.org/AACAP/Member_Resources/Practice_Information/When_to_Seek_Referral_or_Consultation_with_a_CAP.aspx
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Dr. Blader has received funding as a consultant/researcher from Supernus Pharmaceuticals and research funding through his employer institution from Supernus.
Dr. Crismon has served on a one day advisory committee regarding biosimilars with funding through his employer institution from Amgen.
Dr. Kratochvil has received research funding through his employer institution from AstraZeneca, Abbott, Forest, Lilly, Neurex, Novartis, Pfizer, Otsuka, Seaside, Shire, and Somaset. He has received funding as an advisor through his employer institution from Abbott, AstraZeneca, Forest, Lilly, Pfizer, Seaside, and Shire Pharmaceuticals and, through his employer institution has received support for serving on the Data Safety Monitoring Boards for Neurex, Otsuka, Pfizer and Seaside Pharmaceuticals.
Dr. Lopez holds stock in Lilly, Merck, Proctor & Gamble, and Pfizer Pharmaceuticals.
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The other members of the working group do not have any financial relationships to disclose.
Disclaimer

The authors of this document have worked to ensure that all information in the parameters is accurate at the time of publication and consistent with general psychiatric and medical standards and consistent with FDA labeling and information in the biomedical literature.

However, as medical research and practice continue to advance, therapeutic standards may change, and the clinician is encouraged to keep up with the current literature in psychiatry and clinical psychopharmacology. In addition, not all potential adverse drug reactions or complications are listed in the tables, and the clinician should consult the official FDA labeling and other authoritative reference sources for complete information.

These parameters are not a substitute for clinical judgement, and specific situations may require a specific therapeutic intervention not included in these parameters.

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Web Address for the March 2016 Psychotropic Medication Utilization Parameters for Children and Youth in Foster Care

http://www.dfps.state.tx.us/Child_Protection/Medical_Services/guide-psychotropic.asp