

**Track Changes  
from Chapter 3 Section N v1.19.1  
to Chapter 3 Section N v1.20.1**

<b>Chapter</b>	<b>Section</b>	<b>Page(s) in version 1.20.1</b>	<b>Change</b>
3	—	—	Hyperlinks in this section have been revised to reflect up-to-date locations.
3	N0415	N-9	<ul style="list-style-type: none"> <li>• Facilities may wish to identify a resource that their staff consistently use to identify pharmacological classification as assessors should be able to identify the source(s) used to support coding the MDS 3.0.</li> <li>• Assessors should consult the manufacturer's package insert, which may contain the medication's pharmacological classification. They can also work with the resident's pharmacist to confirm the medication classification(s) for a resident's medication(s).</li> </ul>
3	N0415	N-9	<ul style="list-style-type: none"> <li>• Anticoagulants such as Target Specific Oral Anticoagulants (TSOACs), which may or may not require laboratory monitoring, should be coded in N0415E, Anticoagulant.</li> <li>• Do not code flushes to keep an IV access patent in N0415E, Anticoagulant.</li> <li>• Code a medication even if it was given only once during the look-back period.</li> </ul>
3	N0415	N-9	<ul style="list-style-type: none"> <li>• Over-the-counter sleeping medications are not coded as hypnotics, as they are not categorized as hypnotic medications.</li> <li>• <del>In circumstances where reference materials vary in identifying a medication's therapeutic category and/or pharmacological classification, consult the resources/links cited in this section or consult the medication package insert, which is available through the facility's pharmacy or the manufacturer's website. If necessary, request input from the consulting pharmacist.</del></li> </ul>

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3	N0450	N-15	<p><b>Coding Tips and Special Populations (N0450B and N0450C)</b></p> <ul style="list-style-type: none"><li>• Compliance with the requirement to perform a GDR may be met if, for example, <del>W</del>ithin the first year in which a resident is admitted on an antipsychotic medication or after the facility has initiated an antipsychotic medication, the facility <del>must attempts</del> a GDR in two separate quarters (with at least one month between the attempts), unless physician documentation is present in the medical record indicating that a GDR is clinically contraindicated. <del>After the first year, a GDR must be attempted at least annually, unless clinically contraindicated.</del> Information on GDR and tapering of medications can be found in the <b>State Operations Manual, Appendix PP, Guidance to Surveyors for Long Term Care Facilities</b> (see F758) in accordance with 42 CFR 483.45. The <b>State Operations Manual</b> can be found at <a href="https://www.cms.gov/regulations-and-guidance/guidance/manuals/internet-only-manuals-ioms-items/cms1201984">https://www.cms.gov/regulations-and-guidance/guidance/manuals/internet-only-manuals-ioms-items/cms1201984</a>.</li></ul>