

DEPARTMENT OF HEALTH & HUMAN SERVICES
Centers for Medicare & Medicaid Services
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Baltimore, Maryland 21244-1850



CENTER FOR DRUG AND HEALTH PLAN CHOICE

TO: All Part D Plan Sponsors

FROM: Cynthia G. Tudor, Ph.D., Director, Medicare Drug Benefit and C & D Data Group

SUBJECT: CY 2010 Formulary Enhancement Window and Marketing Information

DATE: September 24, 2009

CMS is providing Part D sponsors an opportunity to enhance their CY 2010 Part D formularies prior to the Annual Coordinated Election Period. In addition to making formulary enhancements, Part D sponsors must also utilize this submission window to modify any applicable prior authorization (PA) and/or step therapy (ST) criteria that CMS could not approve as submitted (a separate CMS communication will be sent at a later date).

Formularies and PA and ST text files must be submitted between 12:00 AM EDT on October 5, 2009 and 5:00 PM EDT on October 7, 2009. Any technical difficulties encountered upon upload or during validation of your formulary should be brought to the attention of the HPMS help desk (800-220-2028, hpms@cms.hhs.gov) prior to the window closing. No consideration will be given for late submissions due to technical difficulties unless HPMS assistance was sought in ample time to troubleshoot the problems before the deadline.

Formulary changes must be limited to enhancements, as described in this memo, as well as modifications to PA and/or ST criteria as directed by CMS in a forthcoming email notification. Any formulary files that contain non-allowable changes, including the addition of a limited access indicator, or changes not otherwise described within this memo will be denied. Finally, failure to correct applicable PA and/or ST criteria review concerns with this submission will result in suppression in the Medicare Prescription Drug Plan Finder.

Please refer to the following formulary and marketing questions and answers regarding this formulary window. If you have any questions, please email the Part D Formularies mailbox at PartDformularies@cms.hhs.gov.

Q1: Will CMS provide an update to the Formulary Reference File (FRF) for this enhancement window?

A1: Yes. The October update to the FRF will be available in the HPMS CY 2010 Formulary Submission Module on September 28, 2009.

Q2: What types of formulary changes can be made during this submission window?

A2: This submission window is for formulary enhancements only. The types of formulary changes that CMS considers enhancements are outlined in Attachment 1. If you are adding drugs to your formulary, you must ensure that:

- each category and class has at least two Part D drugs (if more than one drug exists);
- you have Part D claims data to support the addition of any Part D drugs to a specialty tier;
- all quantity limits allow for FDA-approved dosing regimens; and
- any PA/ST applied to drugs within the protected classes are consistent with current CMS policy.

Q3: What changes can be made to the PA and ST formulary attachments during the October window?

A3: If you receive a subsequent notification regarding PA and/or ST criteria that are determined to be unacceptable by CMS reviewers, you must modify the specific criteria as necessary. The changes must be limited to those elements identified by CMS reviewers. PA and ST criteria changes that were not part of the CMS communication are not permitted (except for removal of PA or ST requirements).

Q4: How does this submission window relate to the Medicare Prescription Drug Plan Finder (MPDPF)?

A4: CY 2010 formulary approvals and denials will be entered into HPMS on October 20, 2009. Pricing data submissions due during the Monday, October 26, 2009 – Tuesday, October 27, 2009 submission window should reflect the most recently approved HPMS formulary data as of October 21, 2009. If your October submission is conditionally approved, the updated formulary information will be available for display in MPDPF on November 9, 2009.

Q5: If a Part D sponsor wishes to make further enhancements beyond the October submission window, can these enhancements be included in plan marketing materials and websites?

A5: Yes. If an organization has decided to enhance their formulary outside of an HPMS formulary submission window, these enhancements can be included on marketed formulary information. However, the enhancements must be limited to those listed in Attachment 1. Further, an HPMS formulary file reflective of these enhancements must be submitted during the next available submission window.

Q6: Should the supplemental formulary files (free first fill, partial gap, and home infusion) be updated with this submission?

A6: When submitting this formulary update, you are given a choice either to use the last supplemental file(s) submitted or upload new file(s). New supplemental files should only be submitted if you are making changes to your HPMS formulary file that would necessitate a new file upload.

Attachment 1.

Formulary Enhancements
1. Addition of Part D drugs, with or without utilization management
2. Moving drugs to a more favorable beneficiary cost-sharing tier
3. Removal of prior authorization requirements
4. Changing PA Type from 1 (PA applies) to 2 (PA applies to new starts only) or 3 (Part B versus Part D PA only, if a Part B versus Part D PA is appropriate)
5. Removal of quantity limit restrictions
6. Making existing quantity limits less restrictive (e.g. increasing the allowable quantity limit amount without changing the quantity limit days)
7. Step therapy (ST) enhancements: <ul style="list-style-type: none">• Removal of entire ST protocol (e.g. removal of step therapy requirements for the stepped drug(s) and the corresponding removal of step edits from all prerequisite drugs)• Removal of ST requirements for a drug(s) within the highest step level of a protocol (e.g. removal of step requirements for one step 2 drug within a step therapy protocol containing two step levels and more than one step 2 drug)• Addition of prerequisite step 1 drugs to existing ST protocols (i.e. the new step 1 drug <i>or</i> the existing step 1 drugs would qualify the member for the step 2 drug)• Changing ST Type from 1 (ST applies) to 2 (ST applies to new starts only)