



Candesartan Cilexetil and Hydrochlorothiazide Tablets

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Expert Committee	Small Molecules 2

In accordance with the Rules and Procedures of the Council of Experts, the Small Molecules 2 Expert Committee has revised the Candesartan Cilexetil and Hydrochlorothiazide Tablets monograph. The purpose of this revision is to add *Dissolution Test 2* to accommodate FDA-approved drug products with different dissolution conditions and/or tolerances than the existing dissolution test(s). *Labeling* information has been incorporated to support the inclusion of *Dissolution Test 2*.

- *Dissolution Test 2* was validated using the Kromasil CN brand of column with L10 packing. The typical retention times for hydrochlorothiazide and candesartan cilexetil are about 4.8 and 13.5 min, respectively.

The Candesartan Cilexetil and Hydrochlorothiazide Tablets Revision Bulletin supersedes the currently official monograph.

Should you have any questions, please contact Jasmine Lawrence, Scientist IV (301-230-6363 or jasmine.lawrence@usp.org).