Nicardipine Hydrochloride Injection

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In accordance with the Rules and Procedures of the Council of Experts, the Small Molecules 2 Expert Committee has revised the Nicardipine Hydrochloride Injection monograph. The purpose of this revision is to widen the *Nicardipine pyridine analog* impurity limit from NMT 0.9% to NMT 2.5% in the Organic Impurities test to accommodate FDA-approved drug products with different limits.

The Nicardipine Hydrochloride Injection Revision Bulletin supersedes the currently official monograph.

Should you have any questions, please contact Robyn Fales, Scientist IV (240-221-2047 or rnf@usp.org).