In accordance with the Rules and Procedures of the Council of Experts, the Small Molecules 5 Expert Committee has revised the Budesonide monograph. The purpose of this revision is to widen the limit of “Budesonide glyoxal (epimers)” in the Organic Impurities test from NMT 0.07% to NMT 0.10% to be consistent with the FDA-approved specification. Existing references to reagents have been updated for consistency with the reagent entry.

The Budesonide Revision Bulletin supersedes the currently official monograph.

Should you have any questions, please contact Pavani Jagu, Associate Scientific Liaison (+91 40 44488968 or pavani.jagu@usp.org).