

Draft Guidance on Atropine Sulfate; Diphenoxylate Hydrochloride

This draft guidance, when finalized, will represent the current thinking of the Food and Drug Administration (FDA, or the Agency) on this topic. It does not establish any rights for any person and is not binding on FDA or the public. You can use an alternative approach if it satisfies the requirements of the applicable statutes and regulations. To discuss an alternative approach, contact the Office of Generic Drugs.

Active Ingredient: Atropine sulfate; Diphenoxylate hydrochloride

Dosage Form; Route: Tablet; oral

Recommended Studies: Atropine Sulfate and Diphenoxylate Hydrochloride Tablet is a DESI¹ effective drug for which there are no known or suspected bioequivalence problems, and as such is rated “AA” in FDA/CDER’s Approved Drug Products with Therapeutic Equivalence Evaluations (“Orange Book”).

Analytes to measure (in appropriate biological fluid): Not applicable

Bioequivalence based on (90% CI): Not applicable

Waiver request of in vivo testing: 0.025 mg/2.5 mg pursuant to 21 CFR 320.22 (c) provided the in-vitro dissolution profiles of your product are comparable to those of the reference product.

Dissolution test method and sampling times: The dissolution information for this drug product can be found on the FDA-Recommended Dissolution Methods web site, available to the public at the following location: <http://www.accessdata.fda.gov/scripts/cder/dissolution/>. Conduct comparative dissolution testing on 12 dosage units each of all strengths of the test and reference products. Specifications will be determined upon review of the abbreviated new drug application (ANDA).

¹ Drug Efficacy Study Implementation