

Draft Guidance on Cefpodoxime Proxetil

This draft guidance, once finalized, will represent the Food and Drug Administration's (FDA's) current thinking on this topic. It does not create or confer any rights for or on any person and does not operate to bind FDA or the public. You can use an alternative approach if the approach satisfies the requirements of the applicable statutes and regulations. If you want to discuss an alternative approach, contact the Office of Generic Drugs.

Active ingredient: Cefpodoxime Proxetil

Form/Route: Tablet; Oral

Recommended studies: 2 studies

1. Type of study: Fasting
Design: Single-dose, two-way crossover in vivo
Strength: EQ 200mg base
Subjects: Healthy males and non-pregnant females, general population
Additional Comments: Subjects should be screened for hypersensitivity reactions to Cefpodoxime, other cephalosporins and penicillins prior to the BE study.

2. Type of study: Fed
Design: Single-dose, two-way crossover in vivo
Strength: EQ 200mg base
Subjects: Healthy males and non-pregnant females, general population
Additional Comments: Please refer to the Amantadine Hydrochloride Tablet Draft Guidance for additional information regarding fed studies

Analytes to measure (in appropriate biological fluid): Cefpodoxime in plasma

Bioequivalence based on (90% CI): Cefpodoxime

Waiver request of in-vivo testing: EQ 100mg base based on (i) acceptable bioequivalence studies on the EQ 200mg base strength, (ii) acceptable in vitro dissolution testing of all strengths, and (iii) proportional similarity of the formulations across all strengths. Please refer to the Mirtazapine Tablet Draft Guidance for additional information regarding waivers of in vivo testing.

Dissolution test method and sampling times:

Please note that a **Dissolution Methods Database** is available to the public at the OGD website at <http://www.accessdata.fda.gov/scripts/cder/dissolution/>. Please find the dissolution information for this product at this website. Please 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 application.