

Draft Guidance on Cefixime

This draft guidance, once finalized, will represent the Food and Drug Administration's (FDA's) current thinking on this topic.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: Cefixime

Form/Route: Suspension/Oral

Recommended studies: 2 studies

1. Type of study: Fasting
Design: Single-dose, two way crossover *in-vivo*
Strength: 200 mg/5 mL
Subjects: Normal healthy males and females, general population
Additional comments: Females should not be pregnant or lactating, and if applicable, should practice abstention or contraception during the study.

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2. Type of Study: Fed
Design: Single-dose, two way crossover *in-vivo*
Strength: 200 mg/5 mL
Subjects: Normal healthy males and females, general population
Additional Comments: Please see comment above.
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Analytes to measure (in appropriate biological fluid): Cefixime in plasma

Bioequivalence based on (90% CI): Cefixime

Waiver request of in-vivo testing: 100 mg/5 mL based on (i) acceptable bioequivalence studies on the 200 mg/5 mL strength, (ii) proportional similarity of the formulations across all strengths, and (iii) acceptable in vitro dissolution testing of all strengths.

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.fda.gov/cder/ogd/index.htm>. 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. A dosage unit for a suspension is the labeled strength (5 mL). A total of 12 units from 12 different bottles should be used. Specification will be determined upon review of the application.