

Draft Guidance on Glimepiride

October 2024

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Active Ingredient: Glimepiride

Dosage Form: Tablet

Route: Oral

Strengths: 1 mg, 2 mg, 4 mg

Recommended Study: One in vivo bioequivalence study with pharmacokinetic endpoints

1. Type of study: Fed

Design: Single-dose, two-treatment, two-period crossover in vivo

Strength: 1 mg

Subjects: Healthy males and non-pregnant, non-lactating females

Additional comments: The drug product should be administered with 240 mL of a 20% glucose solution in water, followed by 60 mL of the glucose solution administered every 15 minutes for up to 4 hours after dosing. Monitor blood glucose concentrations and signs and symptoms of hypoglycemia during the study. Implement appropriate hypoglycemia management protocol.

Analyte to measure: Glimepiride in plasma

Bioequivalence based on (90% CI): Glimepiride

Waiver request of in vivo testing: 2 mg and 4 mg strengths based on (i) an acceptable bioequivalence study on the 1 mg strength, (ii) acceptable in vitro dissolution testing of all strengths, and (iii) proportional similarity of the formulations across all strengths

Dissolution test method and sampling times: The dissolution information for this drug product can be found in the FDA's Dissolution Methods database, <http://www.accessdata.fda.gov/scripts/cder/dissolution/>. Conduct comparative dissolution testing on 12 dosage units for each of all strengths of the test product and reference listed drug (RLD).¹ Specifications will be determined upon review of the abbreviated new drug application.

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Unique Agency Identifier: PSG_020496

¹ If the RLD is not available, refer to the most recent version of the FDA guidance for industry on *Referencing Approved Drug Products in ANDA Submissions*.