

Draft Guidance on Ertugliflozin; Metformin Hydrochloride

October 2024

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Active Ingredients:	Ertugliflozin; Metformin hydrochloride
Dosage Form:	Tablet
Route:	Oral
Strengths:	2.5 mg; 500 mg, 2.5 mg; 1 gm, 7.5 mg; 500 mg, 7.5 mg; 1 gm
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: 7.5 mg; 1 gm
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.

Analytes to measure: Ertugliflozin and metformin in plasma

Bioequivalence based on (90% CI): Ertugliflozin and metformin

Waiver request of in vivo testing: 2.5 mg; 500 mg, 2.5 mg; 1 gm, 7.5 mg; 500 mg strengths based on (i) acceptable bioequivalence study on the 7.5 mg; 1 gm 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_209806

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*.