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Draft Guidance on Midodrine Hydrochloride

October 2024

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Active Ingredient: Midodrine hydrochloride

Dosage Form: Tablet

Route: Oral

Strengths: 2.5 mg, 5 mg, 10 mg

Recommended Study: One in vivo bioequivalence study with pharmacokinetic endpoints

1. Type of study: Fasting
   Design: Single-dose, two-treatment, two-period crossover in vivo
   Strength: 5 mg
   Subjects: Healthy males and healthy females not of reproductive potential
   Additional comments: Due to safety reasons, the 5 mg dose is more appropriate for a single dose in vivo bioequivalence study in subjects.

Analyte to measure: Midodrine in plasma

Bioequivalence based on (90% CI): Midodrine

Waiver request of in vivo testing: 2.5 mg and 10 mg strengths based on (i) acceptable bioequivalence study on the 5 mg strength (ii) acceptable in vitro dissolution testing across 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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[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.*