

Contains Nonbinding Recommendations
Draft – Not for Implementation
Draft Guidance on Ulipristal Acetate
October 2024

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Active Ingredient:	Ulipristal acetate
Dosage Form:	Tablet
Route:	Oral
Strength:	30 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: 30 mg
Subjects: Healthy non-pregnant, non-lactating females
Additional comments: Ensure an adequate washout period between treatments in the crossover study due to the long elimination half-life of ulipristal. Alternatively, a parallel study design may be considered.

Analyte to measure: Ulipristal acetate in plasma

Bioequivalence based on (90% CI): Ulipristal acetate

Waiver request of in vivo testing: Not applicable

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 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*.