

## Draft Guidance on Acetaminophen; Oxycodone Hydrochloride

This draft guidance, once finalized, will represent the Food and Drug Administration's (FDA's) current thinking on this topic.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

**Form/Route:** Tablets/Oral

### Recommended studies:

Acetaminophen; Oxycodone Hydrochloride Tablets are a DESI<sup>1</sup> effective drug for which there are no known or suspected bioequivalence problems, and as such is rated “AA” in the FDA/CDER’s Approved Drug Products with Therapeutic Equivalence Evaluations (“Orange Book”).

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**Analytes to measure (in appropriate biological fluid):** Not Applicable

**Bioequivalence based on (90% CI):** Not Applicable

**Waiver request of *in-vivo* testing:** 325 mg/2.5 mg, 325 mg/5 mg, 325 mg/7.5 mg, 325 mg/10 mg, 500 mg/2.5 mg, 500 mg/7.5 mg, and 650 mg/10 mg pursuant to 21 CFR 320.22 (c) provided that the *in-vitro* dissolution profiles of the proposed product are comparable to those of the reference product.

### 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.accessdata.fda.gov/scripts/cder/dissolution/>. 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.

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<sup>1</sup> Drug Efficacy Study Implementation