## Guidance on Lisinopril; Hydrochlorothiazide

This guidance represents 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:** Lisinopril; Hydrochlorothiazide

Form/Route: Tablet/Oral

**Recommended studies:** 2 studies

1. Type of study: Fasting

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

Strength: 25 mg/20 mg

Subjects: Normal healthy males and females, general population

Additional Comments: Female subjects enrolled in the BE studies should not be

pregnant, and if applicable, should practice abstention or contraception during the study.

2. Type of study: Fed

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

Strength: 25 mg/20 mg

Subjects: Normal healthy males and females, general population

Additional comments: Please see comment above.

Analytes to measure: Lisinopril and Hydrochlorothiazide in plasma.

Bioequivalence based on (90% CI): Lisinopril and Hydrochlorothiazide

Waiver request of in-vivo testing: 12.5/10 mg and 12.5 mg/20 mg based on (i) acceptable bioequivalence studies on the 25 mg/20 mg strength, (ii) proportional similarity of the formulations across all strengths, and (iii) acceptable in vitro dissolution testing of all strengths.

## Dissolution test method and sampling times:

Please note that a **Dissolution Methods Database** is available to the public at the OGD website at <a href="http://www.fda.gov/cder/ogd/index.htm">http://www.fda.gov/cder/ogd/index.htm</a>. 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.