Active ingredient: Doxazosin Mesylate

Form/Route: Extended Release Tablets/Oral

Recommended studies: 2 studies

1. Type of study: Fasting
   Design: single-dose, two-way crossover in-vivo
   Strength: 8 mg
   Subjects: Normal healthy males and females, general population. Females should not be pregnant, and if applicable, should practice abstention or contraception during the study.
   Additional Comments:

2. Type of study: Fed
   Design: single-dose, two-way crossover in-vivo
   Strength: 8 mg
   Subjects: Normal healthy males and females, general population. Females should not be pregnant, and if applicable, should practice abstention or contraception during the study.
   Additional comments:

Analyzer to measure (in appropriate biological fluid): Doxazosin in plasma.

Bioequivalence based on (90% CI): Doxazosin

Waiver request of in-vivo testing: 4 mg based on (i) acceptable bioequivalence studies on the 8 mg strength, (ii) proportionally similar across both strengths, and (iii) acceptable in vitro dissolution testing of both strengths.

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.fda.gov/cder/ogd/index.htm. 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.

In addition to the method above, for modified release products, dissolution profiles on 12 dosage units each of test and reference products generated using USP Apparatus I at 100 rpm and/or Apparatus II at 50 rpm in at least three dissolution media (pH 1.2, 4.5 and 6.8 buffer) should be submitted in the application. Agitation speeds may have to be increased if appropriate. It is acceptable to add a small amount of surfactant, if necessary. Please include early sampling times of 1, 2, and 4 hours and continue every 2 hours until at least 80% of the drug is released, to provide assurance against premature release of drug (dose dumping) from the formulation. Specifications will be determined upon review of the data submitted in the application.

Finalized May 2008