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June 4, 2007

**(Filed Electronically)**

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**Re: Docket No. 2004P-0006**

**PETITION FOR STAY OF ACTION**

The undersigned submit this Petition for Stay pursuant to 21 C.F.R. § 10.35, 21 C.F.R. Part 314, and 21 U.S.C. § 355 on behalf of Purdue Pharma L.P. ("Purdue"), holder of approved New Drug Application 20-553 for OxyContin<sup>®</sup> (oxycodone HCl controlled-release) Tablets, 10 mg, 15 mg, 20 mg, 30 mg, 40 mg, 60 mg, 80 mg, and 160 mg ("OxyContin"). This Petition for Stay requests that the Commissioner of Food and Drugs stay the Agency's implementation of its March 23, 2004 Decision on Purdue's January 6, 2004 Petition for Stay (PSA1), Docket No. 2004P-0006.

**I. Decision Involved**

The March 23, 2004 Decision on Purdue's January 6, 2004 Petition for Stay (PSA1), Docket No. 2004P-0006, attached hereto as Exhibit 1.

**II. Action Requested**

By this Petition for Stay, the undersigned hereby request that FDA stay implementation of the Agency's March 23, 2004 Decision on Purdue's January 6, 2004 Petition for Stay (PSA1), Docket No. 2004P-0006, until the Agency has properly considered the issues raised in the Citizen Petition filed to this Docket on June 4, 2007 and has acted on that Petition.

2007P-0232

PSA1