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Food and Drug Administration  
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**Re: Docket 2004P-0006  
Petition for Stay of Action**

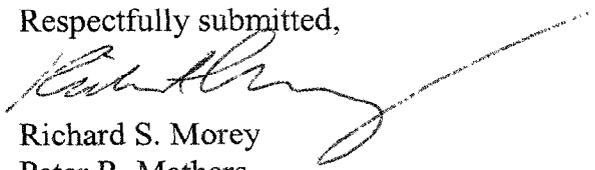
The undersigned submit this supplement to the above referenced Petition for Stay of Action filed on January 6, 2004. This supplement encloses the labeling supplements referenced in the Petition for Stay of Action and subsequently filed by Purdue Pharma L.P. to its approved New Drug Application 20-553 for OxyContin® (oxycodone HCl controlled-release) Tablets, 10 mg, 20 mg, 40 mg, 80 mg, and 160 mg ("OxyContin").

Specifically, the following documents are enclosed:

- Exhibit A December 19, 2003 Changes being Effected Supplement to NDA 20-553
- Exhibit B January 7, 2003 Changes being Effected Supplement to NDA 20-553
- Exhibit C January 13, 2003 Expedited Supplement to NDA 20-553

The copy of the December 19, 2003 CBE Supplement attached as Exhibit A to the Petition for Stay inadvertently omitted the "clean" copy of the revised package insert and included only the version that highlighted the changes being made. Both versions were included in the original Supplement and, therefore, an additional copy of the entire December 19, 2003 Supplement is enclosed with this submission as Exhibit A.

Respectfully submitted,



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2004P-0006

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