



Hospital Products Division  
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March 22, 2004

Division of Dockets Management  
Food and Drug Administration  
Department of Health and Human Services  
Room 1061, HFA-305  
5630 Fishers Lane  
Rockville, MD 20852

**RE:           Docket 2003P-0519**  
  
**Citizen's Petition for ANDA Suitability**  
  
**Ondansetron Hydrochloride Injection Premixed**

In accordance with 21 CFR 10.30(g), Abbott Laboratories hereby request the Commissioner of Food and Drugs to withdraw without prejudice to resubmission at a later date, our Citizen's Petition for ANDA Suitability of Ondansetron Hydrochloride Injection Premixed (8, 12, 16, 20 and 24 mg/50 mL). This petition was filed on October 31, 2003 and assigned docket number 2003P-0519.

If you require further information, please feel free to contact me.

Sincerely,

ABBOTT LABORATORIES

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