



NDA 17-673/S-065  
NDA 17-789/S-051

Abbott Laboratories  
Hospital Products Division  
200 Abbott Park Road, D-389, J45-2  
Abbott Park, IL 60064-6157

Attention: Jean Kirkleit Davis  
Manager, Regulatory Affairs

Dear Ms. Davis:

Please refer to your supplemental new drug applications dated August 21, 1998, received August 25, 1998, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Aminosyn 5%, 7%, 8.5%, 8.5% w/Electrolytes & 10% and Aminosyn 3.5%, 3.5% M and 7% w/ Electrolytes.

We acknowledge receipt of your submissions dated September 6, 2002, and March 27, 2003.

Your submission of March 27, 2003, constituted a complete response to our November 20, 2001, action letter.

These supplemental new drug applications provide for a revised **PRECAUTIONS** section. A **Geriatric Use** subsection is added in accordance with 21CFR 201.57(f)(10). We note that you have incorporated all the changes that were approved on October 25, 2002, for supplements NDA 17-673/S-066 and NDA 17-789/S-052 (Package Insert, Overwrap Labels, and Immediate Container Labels).

We have completed our review of these supplemental new drug applications, as amended, and they are approved, effective on the date of this letter,

If you issue a letter communicating important information about this drug product (i.e., a "Dear Health Care Professional" letter), we request that you submit a copy of the letter to this NDA and a copy to the following address:

MEDWATCH, HFD-410  
FDA  
5600 Fishers Lane  
Rockville, MD 20857

We remind you that you must comply with the requirements for an approved NDA set forth under 21 CFR 314.80 and 314.81.

NDA 17-673/S-065

NDA 17-789/S-051

Page 2

If you have any questions, call Victoria Kao, Regulatory Project Manager, at (301) 827-7410.

Sincerely,

*{See appended electronic signature page}*

Bob Rappaport, M.D.  
Acting Director  
Division of Anesthetic, Critical Care, and  
Addiction Drug Products  
Office of Drug Evaluation II  
Center for Drug Evaluation and Research

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/s/

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Bob Rappaport  
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