### **CENTER FOR DRUG EVALUATION AND RESEARCH**

### **APPROVAL PACKAGE FOR:**

### **APPLICATION NUMBER**

### 21-372

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**DEPARTMENT OF HEALTH & HUMAN SERVICES** 

**Public Health Service** 

Food and Drug Administration Rockville, MD 20857

NDA 21-372

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Helsinn Healthcare SA c/o August Consulting Attention: Craig Lehmann, Pharm.D. 515 Capital of Texas Highway, Suite 150 Austin, TX 78746

Dear Dr. Lehmann:

We have received your new drug application (NDA) submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for the following:

Name of Drug Product: Palonosetron Hydrochloride Intravenous Injection, 0.25 mg

Review Priority Classification: (S) Standard

Date of Application: September 26, 2002

Date of Receipt: September 27, 2002

Our Reference Number: NDA 21-372

Unless we notify you within 60 days of the receipt date that the application is not sufficiently complete to permit a substantive review, we will file the application on November 26, 2002 in accordance with 21 CFR 314.101(a). If we file the application, the user fee goal date will be July 27, 2003.

Under 21 CFR 314.102(c), you may request an informal conference with this Division (to be held approximately 90 days from the above receipt date) for a brief report on the status of the review but not on the ultimate approvability of the application. Alternatively, you may choose to receive a report by telephone.

Please cite the NDA number listed above at the top of the first page of any communications concerning this application. Address all communications concerning this NDA as follows:

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NDA 21-372 Page 2

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<u>U.S. Postal Service/Courier/Overnight Mail:</u> Center for Drug Evaluation and Research Division of Gastrointestinal and Coagulation Drug Products, HFD-180 Attention: Division Document Room, 6B-24 5600 Fishers Lane Rockville, Maryland 20857

If you have any questions, call me at (301) 827-7310.

Sincerely,

{See appended electronic signature page}

Brian Strongin, R.Ph., M.B.A. Regulatory Health Project Manager Division of Gastrointestinal and Coagulation Drug Products Office of Drug Evaluation III Center for Drug Evaluation and Research

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This is a representation of an electronic record that was signed electronically and this page is the manifestation of the electronic signature.

/s/ Brian Strongin 11/7/02 01:16:40 PM

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