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The following materials were distributed by Enzon Pharmaceuticals, Inc. ("Enzon") and NPS Pharmaceuticals, Inc. ("NPS") to attendees of the Sun Trust Robinson Humphrey 32nd Annual Institutional Investor Conference held on April 8, 2003, in Atlanta, Georgia to discuss the proposed business combination between Enzon and NPS.







Safe Harbor

Cautionary Statement For The Purpose Of The "Safe Harbor" Provisions Of The Private Securities Litigation Reform Act Of 1995

This presentation contains forward-looking statements within the meaning of the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995. These statements are based on management's current expectations and beliefs and are subject to a number of factors and uncertainties that could cause actual results to differ materially from those described in the forward-looking statements. The forward-looking statements contained in this presentation include statements about future financial and operating results and the proposed NPS/Enzon merger. These statements are not guarantees of future performance, involve certain risks, uncertainties and assumptions that are difficult to predict, and are based upon assumptions as to future events that may not prove accurate. Therefore, actual outcomes and results may differ materially from what is expressed herein. For example, if either of the companies do not receive required stockholder or governmental approvals or fail to satisfy other conditions to closing, the transaction will not be consummated. In any forward-looking statement in which NPS or Enzon expresses an expectation or belief as to future results, such expectation or belief is expressed in good faith and believed to have a reasonable basis, but there can be no assurance that the statement or expectation or belief will result or be achieved or accomplished. The following factors, among others, could cause actual results to differ materially from those described in the forward-looking statements: the risk that the NPS and Enzon businesses will not be integrated successfully; costs related to the proposed merger, failure of the NPS or Enzon stockholders to approve the proposed merger; and other economic, business, competitive and/or regulatory factors affecting NPS' and Enzon's businesses generally as set forth in NPS's and Enzon's filings with the SEC, including their Annual Reports on Form 10-K for their respective most recent fiscal years, especially in the Management's Discussion and Analysis section, their most recent Quarterly Reports on Form 10-Q and their Current Reports on Form 8-K. NPS and Enzon are under no obligation to (and expressly disclaim any such obligation to) update or alter their forward-looking statements whether as a result of new information, future events or otherwise.





Safe Harbor continued

Additional Information And Where To Find It

In connection with the proposed NPS-Enzon merger, NPS, Enzon and Momentum Merger Corporation have caused to be filed a joint proxy statement/prospectus with the SEC in connection with the transaction described herein. INVESTORS AND SECURITY HOLDERS ARE URGED TO READ THE JOINT PROXY STATEMENT/PROSPECTUS BECAUSE IT CONTAINS IMPORTANT INFORMATION ABOUT THE TRANSACTION DESCRIBED HEREIN. Investors and security holders may obtain a free copy of the joint proxy statement/prospectus and other documents filed by NPS and Enzon with the SEC at the SEC's web site at http://www.sec.gov or by contacting NPS at 801-583-4939 and through NPS' website at http://www.npsp.com, or by contacting Enzon at 908-541-8678 and through Enzon's website at http://www.enzon.com.

NPS and its directors and executive officers may be deemed to be participants in the solicitation of proxies from the stockholders of NPS and Enzon in connection with the transaction described herein. Information regarding the special interests of these directors and executive officers in the transaction described is included in the joint proxy statement/prospectus described above. Additional information regarding these directors and executive officers is also included in NPS' 2002 Annual Report on Form 10-K, which was filed with the SEC on March 21, 2003. This document is available free of charge at the SEC's web site at http://www.sec.gov or by contacting NPS at 801-583-4939 and through NPS' website at http://www.npsp.com.

Enzon and its directors and executive officers also may be deemed to be participants in the solicitation of proxies from the stockholders of Enzon and NPS in connection with the transaction described herein. Information regarding the special interests of these directors and executive officers in the transaction described herein is included in the joint proxy statement/prospectus described above. Additional information regarding these directors and executive officers is also included in Enzon's proxy statement for its 2002 Annual Meeting of Stockholders, which was filed with the SEC on or about October 28, 2002. This document is available free of charge at the SEC's web site at http://www.sec.gov or by contacting Enzon at 908-541-8678 and through Enzon's website at http://www.enzon.com.





Our Mutual Goal





To Build a Sustainable Top-Tier Biotech Based On:

- A deep, diversified and sustainable pipeline
- A clearly defined pathway to profitability
- A fully integrated infrastructure and stable financial position





A Fully Integrated Biotechnology Leader







Synergies Expand and Accelerate Value Creation

- Expand pipeline development
 - e.g., CNS product opportunities
- · Accelerate program development
 - e.g., ALX-0600 for multiple indications
- · Leverage financial strength and commercial capacity
 - e.g., in-license products and technologies, and optimize PREOS partnership





The Merger Creates a Top-Tier Biotech

- Management with a proven record of building businesses
- Drug discovery and development expertise
- Manufacturing capacity and experience
- Commercial infrastructure
- Strong, dependable revenues





Post Merger Strengths

- Fully integrated:
 - From drug discovery through manufacturing and commercialization
- Innovative and robust pipeline:
 - ~\$150M R&D budget *
 - Phase III: 2 programs
 - Phase II: 3 programs
 - >10 early stage programs
 - Multiple platform technologies

*Based upon 2003 pro-forma financials





Post Merger Metrics

Comparable Company Analysis^(a)

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		2003 ESI.		Products		
Company	Market Value	Revenue	R&D	Marketed	Phase III	
Millennium	\$2,152.2	\$402.5	\$496.5	2		
Celgene	\$1,942.9	\$171.0	\$ 89.8	1	1	
Amylin	\$1,334.2	\$ 49.4	\$111.8	0	1	
Neurocrine	\$1,297.1	\$ 92.0	\$110.5	0	1	
ıcos	\$1,055.7	\$105.5	\$157.5	1	1	
Enzon/NPS		~\$200.0	>\$150.0	5	2	

(a) Projected Financial Information as per Wall Street equity research and calendarized to reflect 12/31 year end





Pipeline, Products & Technologies





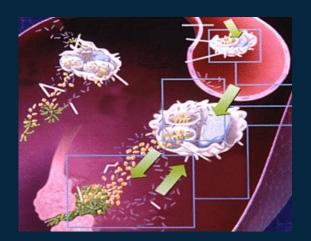
Product Pipeline & Marketed Products Preclinical/ **Product** Research Phase I Phase II Phase III Marketed **Partner PEG-INTRON** Schering-Plough **ABELCET** Proprietary **ADAGEN Proprietary ONCASPAR Proprietary DEPOCYT Proprietary** Cinacalcet HCl Amgen/Kirin **PREOS Proprietary PROTHECAN Proprietary** ALX-0600 Proprietary Cinacalcet HCl Amgen/Kirin NPS 1776 **Proprietary** NPS 1506 **Proprietary** Calcilytics **GSK PEG-Cytotoxics Proprietary** Gly-T Inhibitors Janssen mGluRs AstraZeneca SCA's Micromet Inhaled Leuprolide Nektar = NPS = Enzon

NPS

ENZON

5 Marketed Products

- PEG-INTRON®
- ABELCET®
- ONCASPAR®
- DEPOCYT®
- ADAGEN®







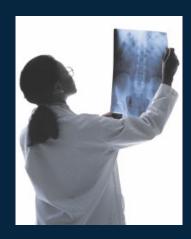
PEG-INTRON: A Strong Revenue Base

- U.S. Pegylated alpha-interferon scripts increasing
- Roche share of new U.S. scripts holding at approximately 20%
- Japan represents significant upside beginning in 2005
- Solid intellectual property position
- Potential for price increases and expanded indications





2 Phase III Clinical Products



PREOS™



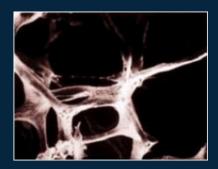
Cinacalcet HCI



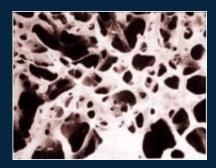


PREOS™ (Intact Human Parathyroid Hormone)

- Stimulates natural bone growth, with potential for:
 - Stronger, healthier bones
 - A lower risk of fracture



 Recent PaTH data support Phase II results and possible PREOS and bisphosphonate combination







PREOS™

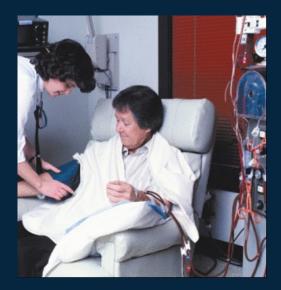
- Expect to compete in a large and growing market
- Pivotal Phase III study to be completed in September 2003
- FDA submission targeted for mid-2004
- Launch anticipated by late 2005
- Combined company more able to aggressively execute development program and negotiate an optimal partnership





Cinacalcet HCI

- Novel treatment for hyperparathyroidism (HPT)
- Market opportunity (U.S.)
 - Primary HPT500,000 patients
 - Secondary HPT280,000 dialysis patients
 - Secondary HPT800,000 predialysis patients







Cinacalcet HCl

- Amgen's Phase III program on-going and on-track
- Amgen confirms 2H 03 NDA filing
- First-in-class molecule in growing market
- Potential for significant royalty stream





Operational Overview





Combined Management Team Includes:

Hunter Jackson, Ph.D. (NPS)

Executive Chairman of the Board

Arthur J. Higgins (Enzon)

Chief Executive Officer

Ulrich Grau, Ph.D. (Enzon)

Executive Vice President, Research and Development

Kenneth J. Zuerblis (Enzon)

Executive Vice President, Finance, Chief Financial Officer & Secretary

Thomas B. Marriott, Ph.D. (NPS)

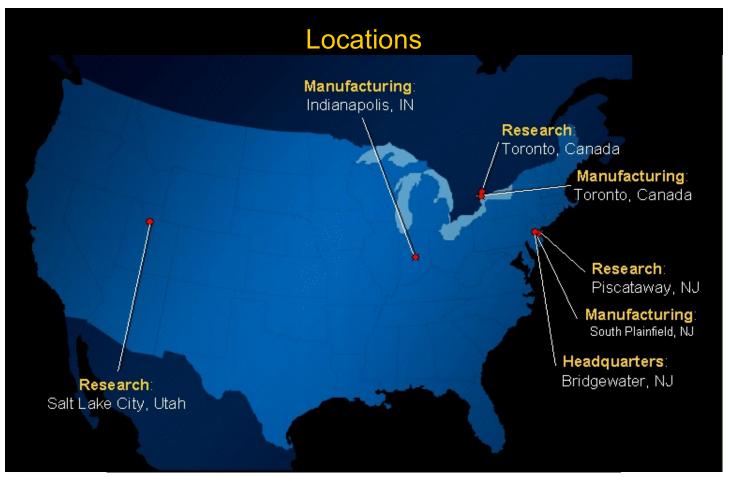
Executive Vice President, Development

David L. Clark (NPS)

Executive Vice President, Corporate Communications and Investor Relations











Selected Pro-Forma Financials

<i>(US\$ in millions)</i> Pro Forma 2002		
\$	31.5	
\$	<u>82.6</u>	
\$	114.1 *	
\$	48.0	
\$	102.0	
\$	400.0	
\$	380.0	
	Pro For \$ \$ \$ \$	





Expected Milestones and News Flow Type 2003 2004 **Clinical Data** Phase III Cinacalcet HCl data in secondary HPT File Cinacalcet HCI NDA Regulatory PREOS 2-year rat toxicology study data **Clinical Data** PREOS TOP Study completed **Clinical Data** Phase IIa results from PROTHECAN Clinical Data **Clinical Data** Initiate additional PII/III ALX-0600 (SBS) Market Data HCV maintenance studies File PREOS NDA Regulatory

= Enzon

Prothecan Phase III program
Cinacalcet HCl approval



Regulatory

Clinical Data

= NPS



2007... Where We're Going

- Revenue > \$500M
- Strong, balanced clinical pipeline
- R&D budget > \$180M
- EBITDA > \$100M
- Industry leading growth rate
- Cash > \$500M









