PHARMACEUTICALS

Corporate Presentation

Nasdaq CPIX

Safe Harbor Statement

This presentation contains forward-looking statements concerning approved products and our product development, our technology, our competitors, our intellectual property, our financial condition and our plans for research and development programs that involve risks, uncertainties and assumptions. These statements are based on the current estimates and assumptions of the management of Cumberland Pharmaceuticals as of the date of this presentation and are subject to uncertainty and changes in circumstances. Given these uncertainties, you should not place undue reliance upon these forwardlooking statements. Such forward-looking statements are subject to risks, uncertainties, assumptions and other factors that may cause the actual results of Cumberland Pharmaceuticals to be materially different from those reflected in such forward-looking statements.

Important factors that could cause actual results to differ materially from those indicated by such forward-looking statements include, among others, those set forth under the headings "Risk factors" and "Management's discussion and analysis of financial condition and results of operations" in our Form 10-K and Form 10-Q Reports on file with the SEC. The Company does not undertake any obligation to release publicly any revisions to these statements to reflect events forward-looking or circumstances after the date hereof or to reflect the occurrence of unanticipated events. All statements contained in this presentation are made only as of the date of this presentation. For more information on our brands, including full prescribing and safety information, please see the links to the product websites which can be found at www.cumberlandpharma.com.



Company Overview

- Specialty pharmaceutical company
 - Portfolio of seven FDA approved products
 - Promoted by two national sales forces
- Several near-term catalysts for new growth opportunities
 - Vibativ post-acquisition integration and market expansion
 - Next Generation Caldolor product launch
 - RediTrex methotrexate product line launch
- Phase II candidates in development with upcoming study milestones
- Proven record of successful product development and product acquisition



Mission & Strategy

Mission: Advance Patient Care through delivery of high quality medicines

Strategy: Build a portfolio of Specialized biopharmaceutical brands



Product Portfolio

Product Development:







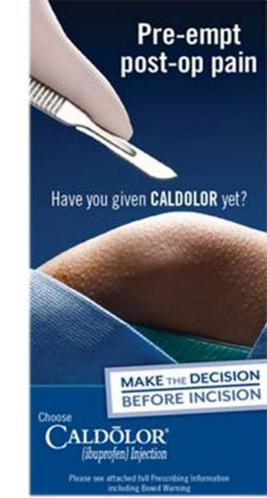
- IV treatment for America's leading cause of poisoning
- Treats liver toxicity associated with acetaminophen overdose
- Developed and registered by Cumberland
- Acetadote now standard of care
- Cumberland developed unique EDTA free formulation
- Maintaining significant market share



*National Poison Data System, American Association of Poison Centers

CALDŌLOR®

- Injectable delivery of ibuprofen
- Developed and registered by Cumberland
- Antipyretic, analgesic & anti-inflammatory properties
- Evaluated in published studies with ~ 2,000 patients
- Over 2.3 million doses administered
- Pediatric labeling approved by FDA
- Study in newborns recently completed

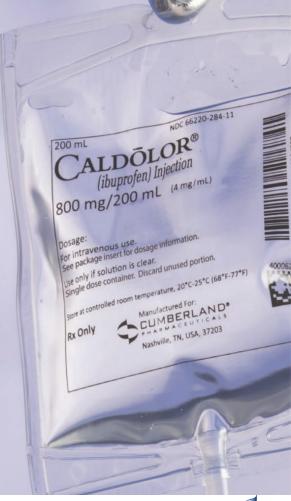




*Symphony Source Health

CALDŌLOR®

- New, ready to administer without further dilution
- Designed to help address National Opioid Crisis
- First and only FDA-approved pre-mixed bag of ibuprofen
- Completed a soft-launch
- National launch underway, gaining early acceptance and growing demand
- Aim to significantly grow Caldolor's sales volume over time with the advantages of this ready-to-use product





- Unique crystalline formulation of lactulose
- Prescription strength laxative
- Clinically proven increases in patient satisfaction
- Repositioned to reflect branded status
- New pricing allowed co-pay support and improved Managed Care coverage





Commercial Portfolio Expansion Strategy





EXPAND Existing Products



ACQUIRE Under-Promoted, Approved Brands



DEVELOP *Early-Stage Candidates*

PRODUCT PORTFOLIO

Acquisition Initiative



Active, ongoing initiative to identify, evaluate and acquire/license new products into the portfolio

Source opportunities through direct efforts and intermediaries

Seek commercial and late stage development assets that fit our strategy and focus

- Branded, Rx products in hospital acute care or gastroenterology
- Sales of **\$5-25 million or larger** with attractive margins





- Acquired following development by Astellas & Theravance
- Injectable antibiotic that treat serious, life-threatening infections
- Hospital product that aligns well with our current infrastructure
- New data demonstrates superiority over vancomycin in select patients with bacterial pneumonia.

* Published in *Infectious Disease and Therapy*







- New injectable delivery of methotrexate
- Designed for the treatment of arthritis and psoriasis
- Widely used throughout Europe with a strong brand presence
- The U.S. methotrexate market is demonstrating **significant growth**
- **Recently FDA-approved** and now preparing to launch





lfetroban Overview

- Cumberland's first new chemical entity (NCE)
- A potent, selective antagonist of thromboxane receptor (TPr)
- Initially developed by Bristol-Myers Squibb as an anti-platelet agent
- Safety is well-established in 26 clinical studies with over 1,300 subjects
- Collaborating with Vanderbilt, Harvard, Scripps and other academic centers
- Cumberland successfully manufactures both IV and oral formulations



Ifetroban Development Pipeline

Preclinical

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Phase I

ILESE III

NDA

16961

Boxaban[®] (aspirin-exacerbated respiratory disease)

Vasculan[™] (systemic sclerosis)

Dyscorban[™] (Duchenne muscular dystrophy)

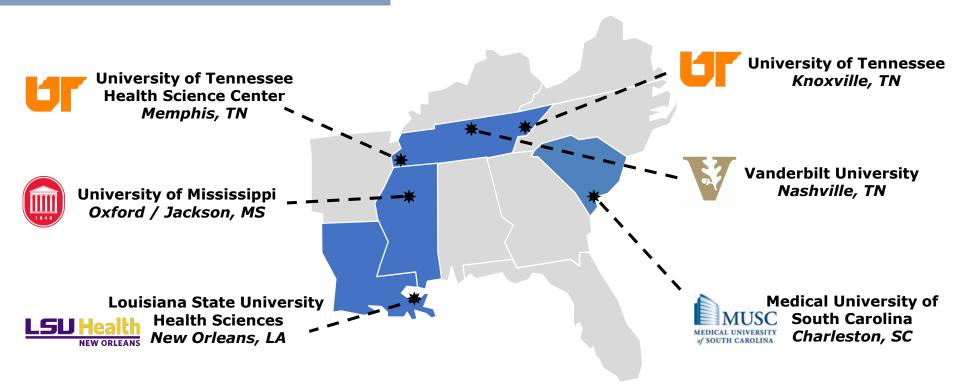
Duchenne Muscular Dystrophy (DMD)

- A rare, fatal, genetic neuromuscular disease characterized by the progressive loss of muscle which results in deterioration of the skeleton, heart and lungs
- Cumberland is investigating ifetroban for the treatment of cardiomyopathy associated with DMD
- New data demonstrates ifetroban could prevent cardiac fibrosis and improve cardiac function – published Journal American Heart Association
- The FDA awarded just over \$1 million in Orphan Drug Grant funding for this unmet medical need
- IND cleared and Phase II study is now underway



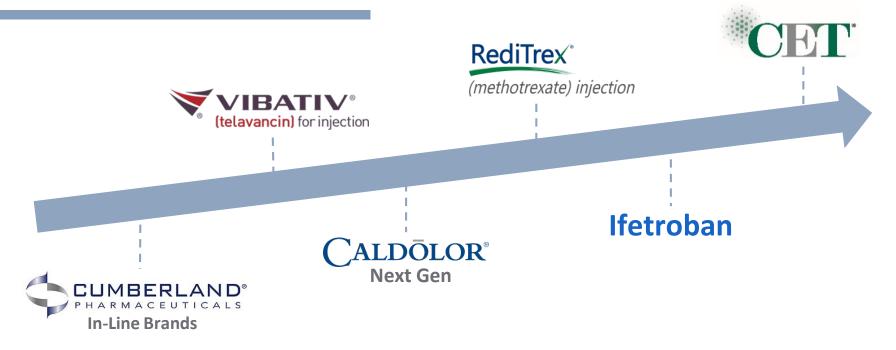








Expanding Our Product Portfolio



Deploying a Multifaceted Strategy to Create Value

Financial Overview

(\$ in millions, except per diluted share)	2020
Net Revenues	\$8.3
Cost of Products Sold	1.6
Gross Profit	\$6.7
Selling & Marketing	\$3.7
Research & Development	1.7
General Administrative	2.0
Amortization	1.1
Discontinued Operations	0.8
Adjusted Earnings*	0.2
Adjusted Earnings [*] per diluted share	\$0.01



*Represents a non-GAAP financial measure.

Summary Balance Sheet

(\$ IN MILLIONS)	As of Mar 31, 2020
CASH & SECURITIES	\$27.0
TOTAL ASSETS	97.3
TOTAL LIABILITIES	47.5
TOTAL EQUITY	49.8

*\$20 million available on revolving line of credit *Tax carry forward credits of \$44 million available *Continued Share Repurchase Program

Cumberland Moving Forward



Diverse product portfolio with 7 FDA approved brands



Proven development and commercialization capabilities



Various initiatives in place to support near-term growth



Phase II products in development with upcoming study milestones



Valuation gap given assets, cash, sales, and pipeline



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