

#### **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 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
  - RediTrex methotrexate product line
- Five Phase II products in development with upcoming study milestones
- Proven record of successful product development and product acquisition
- Strong financial position and positive net cash flows from operations



#### **Mission & Strategy**

Mission: Advance Patient Care

through delivery of high quality medicines

Strategy: Build a portfolio of

Specialized Biopharmaceutical Brands



#### **Product Portfolio**

Product Development:







**Product Acquisition:** 











## **IVACETADOTE**°

- 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





#### **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





Pre-empt post-op pain Have you given CALDOLOR yet? MAKE THE DECISION BEFORE INCISION attached full Prescribing Information

#### **CALDŌLOR®**

- New, easy-to-administer formulation
- Designed to help address National Opioid Crisis
- First and only FDA-approved pre-mixed bag of ibuprofen, ready for administration without further dilution
- Completed a successful soft-launch, gaining early acceptance and growing demand
- National launch underway, using sales organization of 50 professionals
- 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
- Expanding Managed Care coverage





# **Commercial Portfolio Expansion Strategy**







ACQUIRE
Under-Promoted,
Approved Brands



DEVELOP

Early-Stage Candidates

### PRODUCT PORTFOLIO



### **Acquisition Initiative**



#### GOAL TO ADD ONE NEW PRODUCT PER YEAR

through business development initiative or internal product development



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.





<sup>\*</sup> 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 seeing significant growth
- Recently FDA-approved and now preparing to launch





#### **Ifetroban 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
- Cumberland is collaborating with Vanderbilt, Harvard, Scripps and other academic centers
- Cumberland successfully manufactures both IV and oral formulations



## **Ifetroban Development Pipeline**

Portaban<sup>™</sup> (portal hypertension)

Phase I Phase II Phase III Boxaban® (aspirin-exacerbated respiratory disease) Vasculan<sup>™</sup> (systemic sclerosis) Dyscorban<sup>™</sup> (Duchenne muscular dystrophy) Hepatoren® (hepatorenal syndrome)



NDA

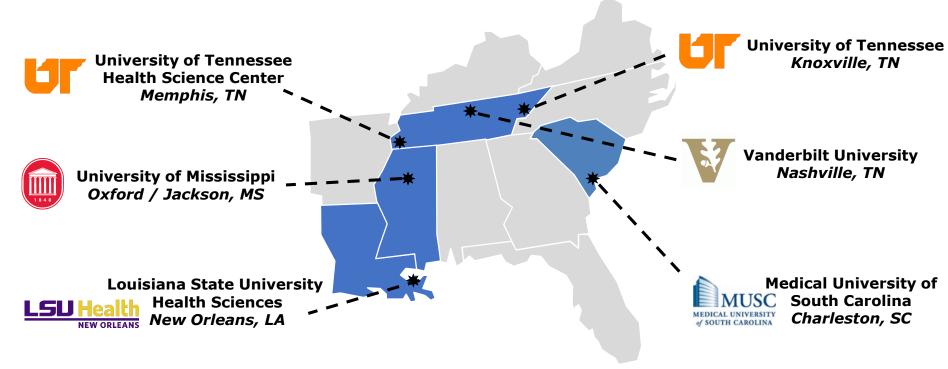
# **Duchenne Muscular Dystrophy (DMD)**

- A rare, fatal, genetic neuromuscular disease characterized by the progressive loss of muscle which results in deterioration of the skeletal, heart and lung muscles
- 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 underway for the treatment of DMD cardiomyopathy



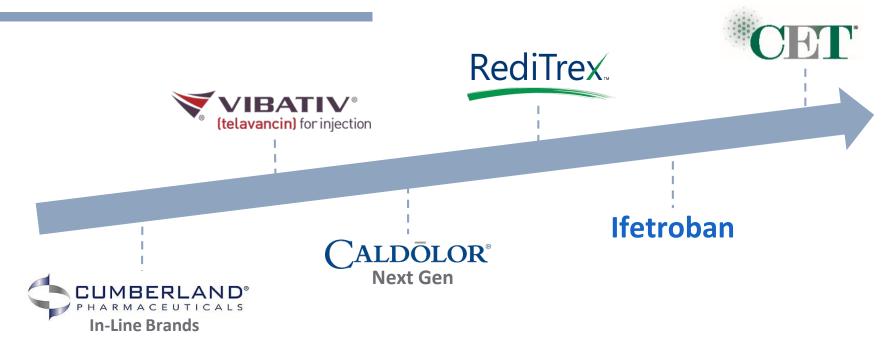








#### **Expanding Our Product Portfolio**



**Deploying a Multifaceted Strategy to Create Value** 



## Financial Overview

(\$ in millions, except per diluted share)	2019
Net Revenues	\$47.5
Cost of Products Sold	8.8
Gross Profit	\$38.7
Selling & Marketing	\$21.4
Research & Development	6.5
General Administrative	10.3
Amortization	4.1
Operating Income (Loss)	(\$3.6)
Adjusted Earnings*	5.0
Adjusted Earnings* per diluted share	\$0.32

<sup>\*</sup>Represents a non-GAAP financial measure.



#### **Summary Balance Sheet**

(\$ IN MILLIONS)	As of Dec 31, 2019
CASH & SECURITIES	\$28.2
TOTAL ASSETS	104.5
TOTAL LIABILITIES	53.5
RETAINED EARNINGS	1.2
TOTAL EQUITY	51.1

<sup>\*\$20</sup> million available on revolving line of credit



<sup>\*</sup>Tax carry forward credits of \$44 million available

<sup>\*</sup>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



Strong financial position with positive net cash flows from operations



Valuation gap given assets, cash, sales, and pipeline



