



# Q1FY22 Earnings Report

May 10, 2022 | Nasdaq: COLL

## Forward-Looking Statements

This presentation contains forward-looking statements within the meaning of The Private Securities Litigation Reform Act of 1995. We may, in some cases, use terms such as "predicts," "forecasts," "believes," "potential," "proposed," "continue," "estimates," "anticipates," "expects," "plans," "intends," "may," "could," "might," "should" or other words that convey uncertainty of future events or outcomes to identify these forward-looking statements. Examples of forward-looking statements contained in this presentation include, among others, statements related to our full-year 2022 financial guidance, including total projected product revenue, adjusted operating expenses and adjusted EBITDA, current and future market opportunities for our products and our assumptions related thereto, expectations (financial or otherwise) and intentions, and other statements that are not historical facts. Such statements are subject to numerous important factors, risks and uncertainties that may cause actual events or results, performance, or achievements to differ materially from the company's current expectations. Actual results may differ materially from management's expectations and such forward-looking statements in this presentation could be affected as a result of various important factors, including risks relating to, among others: risks related to the ability to realize the anticipated benefits of our acquisition of BDSI, including the possibility that the expected benefits from the BDSI acquisition will not be realized or will not be realized within the expected time period; the risk that BDSI's business will not be integrated successfully; negative effects of the consummation of the BDSI acquisition on the market price of our common stock and/or operating results; unknown liabilities; risks related to future opportunities and plans for the products acquired with BDSI, including uncertainty of the expected financial performance of such products; the impact of the COVID-19 pandemic on our ability to conduct our business, reach our customers, and supply the market with our products; our ability to commercialize and grow sales of our products; our ability to manage our relationships with licensors; the success of competing products that are or become available; our ability to obtain and maintain regulatory approval of our products and any product candidates, and any related restrictions, limitations, and/or warnings in the label of an approved product; the size of the markets for our products and product candidates, and our ability to service those markets; our ability to obtain reimbursement and third-party payor contracts for our products; the rate and degree of market acceptance of our products and product candidates; the costs of commercialization activities, including marketing, sales and distribution; changing market conditions for our products; the outcome of any patent infringement, opioid-related or other litigation that may be brought by or against us, including litigation with Purdue Pharma, L.P.; the outcome of any governmental investigation related to our business; our ability to secure adequate supplies of active pharmaceutical ingredient for each of our products and manufacture adequate supplies of commercially saleable inventory; our ability to obtain funding for our operations and business development; regulatory developments in the U.S.; our expectations regarding our ability to obtain and maintain sufficient intellectual property protection for our products; our ability to comply with stringent U.S. and foreign government regulation in the manufacture of pharmaceutical products, including U.S. Drug Enforcement Agency, or DEA, compliance; our customer concentration; and the accuracy of our estimates regarding expenses, revenue, capital requirements and need for additional financing. These and other risks are described under the heading "Risk Factors" in our Annual Reports on Form 10-K and Quarterly Reports on Form 10-Q and other filings with the SEC. Any forward-looking statements that we make in this presentation speak only as of the date of this presentation. We assume no obligation to update our forward-looking statements whether as a result of new information, future events or otherwise, after the date of this presentation.

## Non-GAAP Financial Measures

To supplement our financial results presented on a GAAP basis, we have included information about certain non-GAAP financial measures such as adjusted EBITDA and adjusted operating expenses. We use these non-GAAP financial measures to understand, manage and evaluate our business as we believe they provide additional information on the performance of our business. We believe that the presentation of these non-GAAP financial measures, taken in conjunction with our results under GAAP, provide analysts, investors, lenders and other third parties insight into our view and assessment of our ongoing operating performance. In addition, we believe that the presentation of these non-GAAP financial measures, when viewed with our results under GAAP and the accompanying reconciliations, provide supplementary information that may be useful to analysts, investors, lenders, and other third parties in assessing our performance and results from period to period. We report these non-GAAP financial measures to portray the results of our operations prior to considering certain income statement elements. These non-GAAP financial measures should be considered in addition to, and not as a substitute for, or superior to, net income or other financial measures calculated in accordance with GAAP.

Adjusted EBITDA is a non-GAAP financial measure that represents GAAP net income adjusted to exclude interest expense, interest income, the benefit from or provision for income taxes, depreciation, amortization, stock-based compensation, and other adjustments to reflect changes that occur in our business but do not represent ongoing operations. Adjusted EBITDA, as used by us, may be calculated differently from, and therefore may not be comparable to, similarly titled measures used by other companies.

There are several limitations related to the use of adjusted EBITDA rather than net income, which is the nearest GAAP equivalent, such as:

- adjusted EBITDA excludes depreciation and amortization, and, although these are non-cash expenses, the assets being depreciated or amortized may have to be replaced in the future, the cash requirements for which are not reflected in adjusted EBITDA;
- we exclude stock-based compensation expense from adjusted EBITDA although (a) it has been, and will continue to be for the foreseeable future, a significant recurring expense for our business and an important part of our compensation strategy and (b) if we did not pay out a portion of our compensation in the form of stock-based compensation, the cash salary expense included in operating expenses would be higher, which would affect our cash position;
- adjusted EBITDA does not reflect changes in, or cash requirements for, working capital needs;
- adjusted EBITDA does not reflect the benefit from or provision for income taxes or the cash requirements to pay taxes;
- adjusted EBITDA does not reflect historical cash expenditures or future requirements for capital expenditures or contractual commitments;
- we exclude restructuring expenses from adjusted EBITDA. Restructuring expenses primarily include employee severance and contract termination costs that are not related to acquisitions. The amount and/or frequency of these restructuring expenses are not part of our underlying business;
- we exclude litigation settlements from adjusted EBITDA, as well as any applicable income items or credit adjustments due to subsequent changes in estimates. This does not include our legal fees to defend claims, which are expensed as incurred;
- we exclude acquisition related expenses as the amount and/or frequency of these expenses are not part of our underlying business. Acquisition related expenses include transaction costs, which primarily consisted of financial advisory, banking, legal, and regulatory fees, and other consulting fees, incurred to complete the acquisition, employee-related expenses (severance cost and benefits) for terminated employees after the acquisition, and miscellaneous other acquisition expenses incurred; and
- we exclude recognition of the step-up basis in inventory from acquisitions as the amount and/or frequency of these expenses are not part of our underlying business.

Adjusted operating expenses is a non-GAAP financial measure that represents GAAP operating expenses adjusted to exclude stock-based compensation expense, and other adjustments to reflect changes that occur in our business but do not represent ongoing operations.

The Company has not provided a reconciliation of its full-year 2022 guidance for adjusted EBITDA or adjusted operating expenses to the most directly comparable forward-looking GAAP measures because it is unable to predict, without unreasonable efforts, the timing and amount of items that would be included in such a reconciliation, including, but not limited to, stock-based compensation expense. These items are uncertain and depend on various factors that could have a material impact on GAAP net income and operating expenses for the guidance period.

## Mission Driven

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Building a leading, diversified specialty pharmaceutical company committed to improving the lives of people living with serious medical conditions

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## GUIDED BY OUR CORE VALUES



# Strong Culture and a Commitment to Our Communities

## EXTERNAL RECOGNITION



## COMMUNITY PARTNERSHIPS



# Q1 2022 Key Business Highlights



Closed strategically and financially transformative BDSI acquisition



On track to exceed targeted run rate synergies of at least \$75M



Seamlessly integrated BDSI core operations and achieved day-one field force readiness



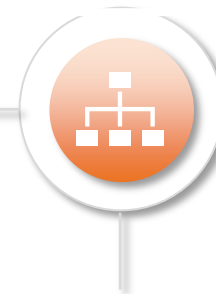
Grew Belbuca<sup>®</sup> TRx 4% Y/Y and Xtampza<sup>®</sup> ER TRx 3% Y/Y



Received approval from FDA for the prior approval supplement for a new Nucynta<sup>®</sup> ER manufacturing site



Resolved all 27 pending opioid-related lawsuits



Strengthened Executive team and Board of Directors

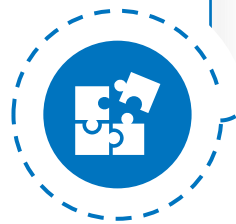
# Collegium 3-Phase Action Agenda

## PHASE 1

TODAY – 6/30/22

### SEAMLESS INTEGRATION

1. Executed with no disruptions to core operations
2. Achieved day one field force readiness
3. Realized majority of targeted run rate synergies



## PHASE 2

7/1/22 – 12/31/22

### GENERATE MOMENTUM

1. Grow Belbuca and Xtampza ER TRxs
2. Complete Xtampza ER contract renegotiations
3. Achieve remainder of target cost synergies
4. Synthesize Elyxyb™ launch learnings



## PHASE 3

2023

### ACCELERATE

1. Propelled by Xtampza ER gross-to-net of <65% in January 2023
2. Driven by Belbuca and Xtampza ER TRx growth
3. Bolstered by fully synergized cost structure



# 2022 Priorities



## GROW TOP LINE

**Grow** Belbuca and Xtampza ER

**Maximize** Nucynta Franchise and Symproic®

**Launch** Elyxyb

**Renegotiate** Xtampza ER contracts



## ACCELERATE BOTTOM LINE

**Exceed** targeted run rate synergies of at least \$75 million

**Maintain** financial discipline



## DEPLOY CAPITAL

**Business development** focused on commercial-stage neurology assets

**Rapidly** pay down debt

**Opportunistically** return capital to shareholders

# Capital Allocation Priorities

1

## FOCUSED BUSINESS DEVELOPMENT

- Commercial-stage neurology assets with \$150 million peak sales potential

2

## RAPIDLY PAYDOWN DEBT

- New \$650M Pharmakon loan issued on 3/22/22<sup>2</sup>
- \$100M to be repaid in first 12 months<sup>1</sup>
- >\$450M to be repaid in first 36 months<sup>1</sup>

3

## OPPORTUNISTICALLY RETURN CAPITAL TO SHAREHOLDERS

- >\$50M remaining on \$100M share repurchase program<sup>2</sup>



# 2022 is Off to a Strong Start



Well-positioned to achieve record full-year revenue and adjusted EBITDA



Completed strategically and financially transformative acquisition of BDSI



On track to exceed targeted run rate synergies of at least \$75M



Strong balance sheet; expect accelerating cash flow



Strategically investing in the growth of our business



Resolved all 27 pending opioid-related lawsuits

Growth and Value Creation

# Financial Highlights

Colleen Tupper, Executive Vice President & Chief Financial Officer

# Financial Highlights: Leveraging Cost Structure and Generating Cash

## Adjusted Opex<sup>1</sup>

Q1FY22 ADJUSTED OPEX,  
30.1% OF REVENUE

↓ **130 BASIS POINTS**

OVER Q1FY21

## Cash Balance<sup>2</sup>

Q1FY22

**\$106.7 MILLION**

FOLLOWING CLOSE OF BDSI  
ACQUISITION

## Operating Cash Flow

EXPECT OPERATING CASH FLOW TO

**ACCELERATE**

FOLLOWING CLOSE OF BDSI  
ACQUISITION

# Capital Allocation Priorities

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# Collegium is in a Strong Financial Position and Executing to Plan<sup>1,2</sup>

## REVENUE GROWTH & SCALE

Est. 2022 revenue of **\$450-465M**  
~+65% y/y at mid-point

## ROBUST CASH FLOWS<sup>4</sup>

Est. 2022 Adj. EBITDA of **\$235-250M**  
~+105% y/y at mid-point

## SIGNIFICANT COST LEVERAGE<sup>3</sup>

Est. 2022 Adj. Op Ex of **\$130-140M**  
~+33% y/y at mid-point

Expect to grow revenue approximately 2x the  
rate of operating expenses

## RAPID DELEVERAGING OF BALANCE SHEET<sup>5</sup>

First year deleveraging of term loan of **\$100M**, full  
paydown over 4 years

Est. 2022 YE Debt/EBITDA ratio **<3.0x**

1. This financial data was provided by Collegium in its press release filed with the SEC on April 5, 2022.
2. Percent change year-over-year is calculated based on financial data provided by Collegium on form 10-K filed with the SEC on February 24, 2022, compared to the mid-point of the guidance ranges provided by Collegium in its press release filed with the SEC on April 5, 2022.
3. Adjusted operating expenses is a non-GAAP financial measure. See Non-GAAP Financial Measures on Slide 2.
4. Adjusted EBITDA is a non-GAAP financial measure. See Non-GAAP Financial Measures on Slide 2.
5. Details regarding the Pharmakon term-loan debt amortization schedule provided by Collegium on form SC TO-C filed with the SEC on February 14, 2022.

# Commercial Update

Scott Dreyer, Executive Vice President & Chief Commercial Officer

# The Leader in Responsible Pain Management

## PORTFOLIO SPANS THE CONTINUUM OF CARE

ACUTE

CHRONIC

 **NUCYNTA**  
(tapentadol) TABLETS 

 **BELBUCA**   
(buprenorphine) Buccal Film

 **NUCYNTA ER**  
(tapentadol) EXTENDED-RELEASE  
TABLETS 

 **Xtampza ER**  
(oxycodone) EXTENDED-RELEASE  
CAPSULES 

~50% market share of  
the branded ER market<sup>1</sup>

**Highly differentiated  
products** that are viewed  
favorably by physicians,  
with a high future intent  
to prescribe<sup>2</sup>

**Pain portfolio distinctly  
positioned** and sources  
differently

# Belbuca Expanded Market Position over Q1 2021



<b>Total Prescriptions<sup>1</sup></b>	
<b>116,900</b>	<b>+4%</b> Q1'22 vs. Q1'21
<b>Branded Extended-Release Market Share<sup>1,2</sup></b>	
<b>17.4%</b>	<b>+2%</b> Q1'22 vs. Q1'21
<b>Unique Prescribers<sup>3</sup></b>	
<b>9,000</b>	<b>+8%</b> Q1'22 vs. Q1'21



# Xtampza ER Expanded Market Position over Q1 2021



<b>Total Prescriptions<sup>1</sup></b>	
<b>166,400</b>	<b>+3%</b> Q1'22 vs. Q1'21
<b>Oxycodone Extended-Release Market Share<sup>1,2</sup></b>	
<b>34.2%</b>	<b>+3.6%</b> Q1'22 vs. Q1'21
<b>Unique Prescribers<sup>3</sup></b>	
<b>18,700</b>	<b>+4%</b> Q1'22 vs. Q1'21

# Nucynta Franchise and Symproic are Contributors



**Branded Extended-Release Market Share<sup>1,2</sup>**

**5.6%**

Stable  
Q1'22 vs. Q1'21

**Total Prescriptions<sup>1</sup>**

**17,800**

**+13.7%**  
Q1'22 vs. Q1'21

# A Strategic Foothold in Neurology



## NEUROLOGY IS A TARGET ADJACENCY

- Business development priority
- Aligned to organizational capabilities
- Complementary to pain



## ELYXYB LAUNCH OPPORTUNITY

- Focused and Phased Approach
  - 25 territories, 3,500 targets
  - ~15% of acute migraine market
- Where we choose to play, we will play to win
- Success-gated expansion

# 2022 Commercial Priorities



## Grow

Belbuca and  
Xtampza ER



## Maximize

potential of  
Nucynta  
Franchise and  
Symproic



## Launch

Elyxyb in focused  
and phased  
approach



## Achieve

gross-to-net of  
<65% for  
Xtampza ER  
beginning in  
January 2023

Q&A

# Building a Leading, Diversified Specialty Pharmaceutical Company



# Non-GAAP Reconciliations

**Collegium Pharmaceutical, Inc.**  
**Reconciliation of GAAP Net Income to Adjusted EBITDA**  
(in thousands)  
(unaudited)

	<b>Three months ended March 31,</b>	
	<b>2022</b>	<b>2021</b>
GAAP net (loss) income	\$ (13,069)	\$ 15,662
Adjustments:		
Interest expense	5,831	5,721
Interest income	(4)	(3)
(Benefit from) provision for income taxes	(2,773)	(188)
Depreciation	715	439
Amortization	18,923	16,795
Stock-based compensation expense	6,135	6,879
Acquisition related expenses	27,167	-
Recognition of step-up basis in inventory	603	-
Total adjustments	\$ 56,597	\$ 29,643
Adjusted EBITDA	\$ 43,528	\$ 45,305



**Collegium Pharmaceutical, Inc.**  
**Reconciliation of GAAP Operating Expenses to Adjusted Operating Expenses**  
(in thousands)  
(unaudited)

	<b>Three months ended March 31,</b>	
	<b>2022</b>	<b>2021</b>
GAAP Operating expenses	\$ 58,511	\$ 34,406
Adjustments:		
Stock-based compensation	6,135	6,879
Acquisition related expenses	27,167	-
Total adjustments	33,302	6,879
Adjusted operating expenses	\$ 25,209	\$ 27,527