



Q3FY22 Earnings Report

November 3, 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; 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. 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.

In our quarterly and annual reports, earnings press releases and conference calls, we may discuss the following financial measures that are not calculated in accordance with GAAP, to supplement our consolidated financial statements presented on a GAAP basis.

Adjusted EBITDA

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 (i.e., the adjustment to record inventory from historic cost to fair value at acquisition) as the adjustment does not reflect the ongoing expense associated with sale of our products as part of our underlying business.

Adjusted Operating Expenses

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.

Adjusted Net Income and Adjusted Earnings Per Share

Adjusted net income is a non-GAAP financial measure that represents GAAP net income adjusted to exclude significant income and expense items that are non-cash or not indicative of ongoing operations, including consideration of the tax effect of the adjustments. Adjusted earnings per share is a non-GAAP financial measure that represents adjusted net income per share. Adjusted weighted-average shares - diluted is calculated in accordance with the treasury stock, if-converted, or contingently issuable accounting methods, depending on the nature of the security.

Reconciliations of adjusted EBITDA, adjusted operating expenses, adjusted net income, and adjusted earnings per share to the most directly comparable GAAP financial measures are included in this presentation.

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.

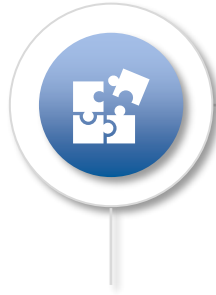
Mission Driven

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

GUIDED BY OUR CORE VALUES



First Nine Months 2022 Key Business Highlights



Completed BDSI acquisition



Grew Belbuca® and Xtampza® ER market share



Resolved all 27 pending opioid industry-related lawsuits



Increased run rate synergies target to \$85M



Renegotiated Xtampza ER contracts

Collegium 3-Phase Action Agenda

PHASE 1

✓ COMPLETED

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
2. Complete Xtampza ER contract renegotiations
3. Achieve remainder of target 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



Top Capital Allocation Priority: Business Development

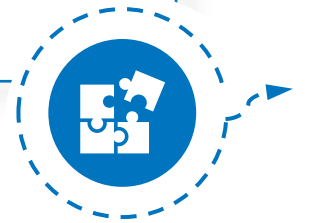
STRONG TRACK RECORD

- ✓ Nucynta Franchise (February 2020)
- ✓ BDISI (March 2022)



BUSINESS DEVELOPMENT FOCUS

- Differentiated commercial-stage assets
 - Peak sales potential >\$150M
 - Exclusivity into 2030s



Financial Highlights

Colleen Tupper, Executive Vice President & Chief Financial Officer

Q3FY22 Financial Highlights

Record Revenue¹

Q3FY22
TOTAL PRODUCT REVENUE

\$127.0 MILLION

+61% OVER Q3FY21

Record Adjusted EBITDA^{1,2}

Q3FY22
ADJUSTED EBITDA

\$74.9 MILLION

+100% OVER Q3FY21

Rapid Deleveraging^{2,3}

ESTIMATED
NET DEBT/ADJUSTED EBITDA RATIO

<3.0x

BY 2022 YEAR-END

Updated 2022 Financial Guidance¹

	Prior	Updated
Total Product Revenues	\$450.0 to \$465.0 million	\$455.0 to \$465.0 million
Total Adjusted Operating Expenses² (Excluding Stock-Based Compensation)	\$125.0 to \$135.0 million	\$125.0 to \$130.0 million
Total Adjusted EBITDA³ (Excluding Stock-Based Compensation and Acquisition Related Expenses)	\$245.0 to \$255.0 million	\$250.0 to \$255.0 million

Capital Allocation Priorities

1

FOCUSED BUSINESS DEVELOPMENT

- Commercial-stage assets:
 - With >\$150 million peak sales potential
 - Differentiated and durable with exclusivity into 2030s

2

RAPIDLY PAYDOWN DEBT

- \$650M Pharmakon loan issued on 3/22/22¹
- \$100M to be repaid in first 12 months²
- >\$450M to be repaid in first 36 months²

3

OPPORTUNISTICALLY RETURN CAPITAL TO SHAREHOLDERS

- Returned \$10M to shareholder in Q3FY22 and October 2022
- >\$42M remaining on \$100M share repurchase program³

Commercial Update

Scott Dreyer, Executive Vice President & Chief Commercial Officer

Successfully Completed Xtampza ER Contract Renegotiations

Plan	% Total Prescriptions	OER Formulary Status	
		2022	2023
Plan 1	27%	Exclusive	Exclusive
Plan 2	15%	Exclusive	Exclusive
Plan 3	2%	Exclusive	Exclusive
Plan 4	4%	Parity	Parity
Plan 5	6%	Exclusive	Non-formulary ¹

Xtampza ER gross-to-net <65% effective January 1, 2023

The Leader in Responsible Pain Management

Strong and Growing Market Position

50%


Branded ER
Market Share^{1,2}

+3% Over Q3FY21


Large Prescriber Bases

Unique prescribers in Q3FY22³


BELBUCA™ (buprenorphine) Buccal Film  **~9,100**


Xtampza ER (oxycodone) EXTENDED-RELEASE CAPSULES  **~19,300**


NUCYNTA™ (tapentadol) TABLETS 

NUCYNTA ER (tapentadol) EXTENDED-RELEASE TABLETS  **~13,100**

Collegium 3-Phase Action Agenda: *Q3FY22 Accomplishments and Looking Ahead*

7/1/22 – 12/31/22

PHASE 2 **GENERATE MOMENTUM**

- 1. Grow Belbuca and Xtampza ER**
 - ✓ Fully trained pain salesforce with only active promotion in pain
 - ✓ Laser focused on execution of plan to drive prescription growth
- 2. Complete Xtampza ER contract renegotiations**
3. Achieve remainder of target synergies
4. Synthesize Elyxyb launch learnings



2023

PHASE 3 **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



Building a Leading, Diversified Specialty Pharmaceutical Company



DIVERSE AND DURABLE PORTFOLIO

- Durable growth drivers
- Leader in responsible pain market



STRONG FINANCIAL POSITION

- FY22 revenue expected to grow ~66% Y/Y¹
- Significant cost leverage: FY22 revenue expected to grow >2x rate of OPEX¹
- Est. 2022 Year-End Net Debt/Adjusted EBITDA ratio <3.0x^{1,2,3}



LONG-TERM VALUE CREATION

- Focused business development
- Rapid debt pay-down
- Return capital to shareholders

1. Percent change year-over-year, growth rates and financial ratios are calculated based on financial data provided by Collegium on Form 10-Q filed with the SEC on November 3, 2022, compared to the mid-point of the guidance ranges provided by Collegium in its press release filed with the SEC on November 3, 2022.
2. Adjusted EBITDA is a non-GAAP financial measure. See Non-GAAP Financial Measures on Slide 3. The net debt/adjusted EBITDA ratio is calculated based on financial data provided by Collegium on Form 10-Q filed with the SEC on November 3, 2022 compared to the mid-point of the guidance ranges provided by Collegium in its press release filed with the SEC on November 3, 2022.
3. 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.

Q&A

Quarterly Non-GAAP Reconciliations

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

	Three Months Ended	
	September 30,	
	2022	2021
GAAP Net income (loss)	\$ 457	\$ 8,046
Adjustments:		
Interest expense	19,046	5,115
Interest income	(11)	(3)
Provision for (benefit from) income taxes	975	991
Depreciation	488	448
Amortization	37,552	16,796
Stock-based compensation expense	5,377	5,948
Acquisition related expenses	463	—
Recognition of step-up basis in inventory	10,519	—
Total adjustments	<u>\$ 74,409</u>	<u>\$ 29,295</u>
Adjusted EBITDA	<u>\$ 74,866</u>	<u>\$ 37,341</u>

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

	Three Months Ended	
	September 30,	
	2022	2021
GAAP Operating expenses	\$ 38,372	\$ 31,964
Adjustments:		
Stock-based compensation	5,377	5,948
Acquisition related expenses	463	—
Total adjustments	\$ 5,840	\$ 5,948
Adjusted operating expenses	\$ 32,532	\$ 26,016

Collegium Pharmaceutical, Inc.
Reconciliation of GAAP Net Income to Adjusted Net Income and Adjusted Earnings Per Share
(in thousands, except share and per share amounts)
(unaudited)

	Three Months Ended	
	September 30,	
	2022	2021
GAAP Net income (loss)	\$ 457	\$ 8,046
Adjustments:		
Non-cash interest expense	2,467	833
Amortization	37,552	16,796
Stock-based compensation expense	5,377	5,948
Acquisition related expenses	463	—
Recognition of step-up basis in inventory	10,519	—
Income tax effect of above adjustments (1)	(14,290)	(5,899)
Total adjustments	\$ 42,088	\$ 17,678
Non-GAAP adjusted net income	\$ 42,545	\$ 25,724
GAAP Weighted-average shares — diluted (2)	39,495,453	41,186,308
Adjusted diluted earnings per share	\$ 1.10	\$ 0.65

1. The income tax effect of the adjustments was calculated by applying our blended federal and state statutory rate of 26% to the items that have a tax effect. As such, the non-GAAP effective tax rates for the three months ended September 30, 2022 and 2021 were 25.3% and 25.0%, respectively, and the non-GAAP effective tax rates for the nine months ended September 30, 2022 and 2021 were 25.4% and 16.8%, respectively.
2. Adjusted weighted-average shares - diluted were calculated using the "if-converted" method for the Convertible Senior Notes in accordance with ASC 260, Earnings per Share. As such, for the three and nine months ended September 30, 2022 and 2021 adjusted earnings per share includes 4,925,134 shares related to the assumed conversion of the Convertible Senior Notes and the associated cash interest expense added-back to non-GAAP adjusted net income. In addition, for the nine months ended September 30, 2022, adjusted earnings per share also includes other potentially dilutive securities to the extent that they are not antidilutive given that non-GAAP adjusted net income was in an income position.