
**UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, DC 20549**

FORM 8-K

CURRENT REPORT

Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): November 4, 2021

Baudax Bio, Inc.

(Exact name of registrant as specified in its charter)

Pennsylvania
(State or other jurisdiction of
incorporation)

001-39101
(Commission
File Number)

47-4639500
(I.R.S. Employer
Identification No.)

490 Lapp Road, Malvern, Pennsylvania

(Address of principal executive offices)

19355

(Zip Code)

Registrant's telephone number, including area code: (484) 395-2470

Not Applicable

(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions (see General Instruction A.2. below):

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
- Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Securities registered pursuant to Section 12(b) of the Act:

<u>Title of Each Class</u>	<u>Trading Symbol</u>	<u>Name of Exchange on Which Registered</u>
Common Stock, par value \$0.01	BXRX	Nasdaq Capital Market

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

Item 2.02 Results of Operations and Financial Condition.

On November 4, 2021, Baudax Bio, Inc. (the “Company”) issued a press release announcing its financial results for the third quarter ended September 30, 2021. A copy of the press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K. The Company has scheduled a conference call and webcast for 8:00 a.m. Eastern time on November 4, 2021 to discuss these financial results and business updates.

The information disclosed under Item 2.02, including Exhibit 99.1, is being furnished and shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”), or otherwise subject to the liabilities of that section, and shall not be deemed to be incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by specific reference in such filing.

Item 8.01 Other Events.

On November 4, 2021, the Company updated information reflected in a slide presentation, which is attached as Exhibit 99.2 to this Current Report on Form 8-K and is incorporated herein by reference. Representatives of the Company will use the updated presentation in various meetings with investors from time to time.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

The following exhibits are being filed herewith:

Exhibit No.	Document
99.1	Press Release of Baudax Bio, Inc., dated November 4, 2021.
99.2	Investor Presentation of Baudax Bio, Inc., dated November 4, 2021.
104	Cover Page Interactive Data file (embedded within the Inline XBRL document).

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

Baudax Bio, Inc

By: /s/ Gerri A. Henwood
Name: *Gerri A. Henwood*
Title: *President and Chief Executive Officer*

Date: November 4, 2021



Baudax Bio Reports Third Quarter 2021 Financial Results and Business Highlights

ANJESO® Revenue Up 40% in Q3 2021 over Q2 2021 and Up Over 300% in Q3 2021 over Q3 2020

Solid Growth in End User Demand Units, September 2021 Strongest Month Launch-to-Date, Indicating Growing Adoption of ANJESO

Neuromuscular Blocking Agent BX1000 Completes Clinical Phase of Dose Escalation Study

Management to Host Investor Conference Call and Webcast Today at 8:00 a.m. ET

MALVERN, Pa., November 4, 2021 -- Baudax Bio, Inc. (NASDAQ:BRRX) (the “Company”), a pharmaceutical company focused on commercializing and developing innovative products for acute care settings, today reported financial results for the three and nine months ended September 30, 2021, and provided key metrics around the ongoing commercial rollout of ANJESO® (meloxicam) injection and updated status of neuromuscular blocking agent (NMBA) BX1000 dose escalation study.

“We are extremely pleased with the continued solid growth we saw during the third quarter in both sales and end-user demand for ANJESO, the significant uptick in sales to new accounts, which grew an estimated 66% quarter-over-quarter and September results that represented our best month to date,” said Gerri Henwood, President and CEO of Baudax Bio. “In addition, we are working to advance our pipeline, which includes novel neuromuscular blocking agents and a reversal agent that have the potential to meaningfully reduce time of “onset” of neuromuscular blockade, and “offset” for procedure recovery time, resulting in potentially valuable cost savings to surgical centers. The clinical dosing portion of the dose escalation study has been completed and we look forward to sharing the results in the near future.”

Third Quarter 2021 and Recent Business Highlights

ANJESO

🕒 **ANJESO U.S. Commercialization.** ANJESO is indicated for the management of moderate to severe pain, alone or in combination with other non-NSAID analgesics. During the third quarter of 2021, demand for ANJESO demonstrated solid growth and deepening usage patterns, with revenue increasing 40% for the third quarter of 2021 compared to the second quarter of 2021 and increasing over 300% for the third quarter of 2021 compared to the third quarter of 2020.

Despite extensive outbreaks of COVID-19 delta in much of the south and selected other regions, and its impact on elective surgeries in July and August, there was growth in estimated demand units sold to all customers in the third quarter of 2021 compared to the second quarter of 2021 of approximately 16% and vials sold to end-user hospitals and ambulatory surgery centers (ASCs) combined was up 17% in the same period. Vials sold to existing hospitals increased 11% in the third quarter of 2021 compared to the second quarter of 2021 and vials sold to existing ASCs increased 66% in the same period. In addition, the month of September 2021 was our single largest month of ANJESO end-user demand units sold launch-to-date for the product.

🕒 **Summer COVID-19 Breakthrough.** COVID-19 related impacts continue to periodically and regionally affect the number of elective surgeries performed, as well as access for field activities. While there were early signs of elective surgeries gradually returning to pre-COVID levels late in the second quarter of 2021, the COVID-19 delta variant had a significant impact on elective surgeries in July and August because of its effects on how hospitals allocated bed space and resources specifically for COVID-19 patients, especially in the Southern U.S. (e.g., Texas, Florida and Alabama, etc.), which comprises approximately 40% of Baudax Bio’s target hospitals.

Pipeline

⌚ **BX1000 clinical update and outlook.** Baudax Bio holds exclusive global rights to two NMBAs and a proprietary chemical reversal agent specific to both of these NMBAs. The Company believes that these agents allow for a very rapid induction of neuromuscular blockade for surgical settings, which especially when used with the proprietary reversal agent, is followed by a rapid reversal of the neuromuscular blockade. BX1000, the Company's intermediate duration NMBA drug candidate, has undergone an earlier Phase 1 trial prior to the Company licensing of the drug. The second study, a dose escalation trial, recently completed its clinical dosing in the trial and is in the data analysis stage, which Baudax Bio believes will be completed in the near future.

Corporate and Financial

⌚ **Amended Articles of Incorporation.** In July 2021, Baudax Bio held a Special Meeting of Shareholders where it secured the required number of shareholder votes to adopt an amendment to its Amended and Restated Articles of Incorporation to increase the number of authorized shares of common stock from 100 million shares to 190 million shares.

Third Quarter 2021 Financial Results

As of September 30, 2021, Baudax Bio had cash, cash equivalents and short-term investments of \$24.9 million.

Net product revenue, recognized according to U.S. GAAP, for the three months ended September 30, 2021 was \$0.3 million, related to sales of ANJESO in the U.S., compared to \$0.1 million for the three months ended September 30, 2020. While utilizing the title model of distribution, product revenue is recognized as shipments are made to the Company's 3PL provider. The increase of \$0.2 million was attributable to securing additional formulary approvals and generating trial and adoption of ANJESO, as well as increased end-user demand leading to increased purchasing by direct customers.

Cost of sales for both the three months ended September 30, 2021 and 2020 was \$0.5 million, and consisted of product costs, royalty expense and certain fixed costs associated with the manufacturing of ANJESO, including supply chain and quality costs. Certain product costs of ANJESO units recognized as revenue during the three months ended September 30, 2021 and 2020 were expensed prior to the FDA approval of ANJESO in February 2020, and therefore are not included in cost of sales during the related periods. Baudax Bio expects that over time, product costs in cost of sales will increase as sales increase and inventory associated with the units manufactured prior to FDA approval are sold.

Research and development expenses for the three months ended September 30, 2021 were \$0.7 million, compared to \$1.5 million for the three months ended September 30, 2020, a decrease of \$0.8 million. This decrease was primarily a result of a decrease in personnel costs of \$0.5 million as well as a decrease in clinical costs of \$0.5 million.

Selling, general and administrative expenses for the three months ended September 30, 2021 were \$11.1 million, compared to \$13.8 million for the same prior year period, a decrease of \$2.7 million. This decrease was primarily a result of a decrease in personnel costs of \$3.1 million.

Baudax Bio reported a net loss, including a non-cash charge of \$5.7 million, of \$17.0 million, or \$(0.20) per diluted share, for the three months ended September 30, 2021. Adjusted net loss* for the three months ended September 30, 2021 was \$11.3 million.

Nine Months Ended September 30, 2021 Financial Results

Net product revenue, recognized according to U.S. GAAP, for the nine months ended September 30, 2021 was \$0.7 million, related to sales of ANJESO in the U.S, compared to \$0.4 million for the nine months ended September 30, 2020. While utilizing the title model of distribution, product revenue is recognized as shipments are made to the Company's 3PL provider. The increase of \$0.3 million was attributable to securing additional formulary approvals and generating trial and adoption of ANJESO, as well as increased end-user demand leading to increased purchasing by direct customers.

Cost of sales for the nine months ended September 30, 2021 was \$1.9 million, compared to \$1.2 million for the nine months ended September 30, 2020 and consisted of product costs, royalty expense and certain fixed costs associated with the manufacturing of ANJESO including supply chain and quality costs. Certain product costs of ANJESO units recognized as revenue during the nine months ended September 30, 2021 and 2020 were expensed prior to the FDA approval of ANJESO in February 2020, and therefore are not included in cost of sales during the related periods. Baudax Bio expects that over time, product costs in cost of sales will increase as sales increase and inventory associated with the units manufactured prior to FDA approval are sold.

Research and development expenses for the nine months ended September 30, 2021 were \$2.6 million, compared to \$5.9 million for the nine months ended September 30, 2020, a decrease of \$3.3 million. Baudax Bio's research and development expenses decreased primarily from a decrease in personnel costs of \$2.6 million and a decrease of \$0.8 million in pre-commercialization manufacturing costs for ANJESO.

Selling, general and administrative expenses for the nine months ended September 30, 2021 were \$33.8 million, compared to \$33.0 million for the same prior year period, an increase of \$0.8 million. The increase was primarily a result of the prior period of 2020 included \$1.5 million in reimbursed general and administrative expenses related to the Transition Services Agreement with Recro Pharma, which ended on December 31, 2020. This increase as well as an increase in public company costs of \$1.2 million were partially offset by a decrease in personnel expenses of \$1.9 million.

Baudax reported a net loss, including a non-cash charge of \$14.9 million, of \$49.2 million, or \$(0.66) per share, for the nine months ended September 30, 2021. Adjusted net loss* was \$34.3 million.

*Adjusted net loss is a non-GAAP financial measure (see reconciliation of non-GAAP financial measures in this release).

Non-GAAP Financial Measures

To supplement the Company's financial results determined by U.S. generally accepted accounting principles ("GAAP"), the Company is reporting certain non-GAAP information for its business, including adjusted net loss. Adjusted net loss is net loss as determined under GAAP, excluding the changes in fair values of contingent consideration and warrant valuations, gain on extinguishment of debt, interest, depreciation, amortization, and stock-based compensation. The Company believes this non-GAAP financial measure is helpful in understanding its business as it is useful to investors in allowing for greater transparency of supplemental information used by management. This measure is used by investors, as well as management in assessing the Company's performance. Non-GAAP financial measures should be considered in addition to, but not as a substitute for, reported GAAP results. Further, Non-GAAP financial measures, even if similarly titled, may not be calculated in the same manner by all companies, and therefore should not be compared. Please see the section of this press release titled "Reconciliation of GAAP to Non-GAAP Financial Measures" for a reconciliation of non-GAAP adjusted net loss to its most directly comparable GAAP measure.

Conference Call Information

Baudax Bio will host a conference call today, Thursday, November 4, 2021, at 8:00 a.m. Eastern Time, to discuss the third quarter 2021 financial results and recent corporate achievements. To access the conference call, please dial (866) 220-5595 (local) or (615) 622-8062 (international) at least 10 minutes prior to the start time and refer to conference ID 9972669. A live audio webcast of the call will be available under "Events" in the News & Investors section of the Company's website, <https://www.baudaxbio.com/news-and-investors/events>. An archived webcast will be available on the Company's website approximately two hours after the event.

About ANJESO®

ANJESO (meloxicam) injection is a proprietary, long-acting, preferential COX-2 inhibitor that possesses analgesic, anti-inflammatory and antipyretic activities, which are believed to be related to the inhibition of cyclooxygenase type 2 pathway (COX-2) and subsequent reduction in prostaglandin biosynthesis. ANJESO is indicated for the management of moderate to severe pain, alone or in combination with other non-NSAID analgesics. Because of the delayed onset of analgesia, ANJESO alone is not recommended for use when rapid onset of analgesia is required. ANJESO is supported by two pivotal Phase III clinical efficacy trials, a large double-blind, placebo-controlled Phase III safety trial and four Phase II clinical efficacy trials, as well as other safety studies. As a non-opioid, Baudax Bio believes ANJESO has the potential to overcome many of the issues associated with commonly prescribed opioid therapeutics, including respiratory depression, constipation, excessive nausea and vomiting, as well as having no addictive potential, while maintaining meaningful analgesic effects for relief of pain. ANJESO was designed using the NanoCrystal® platform, a technology that enables enhanced bioavailability of poorly water-soluble drug compounds. NanoCrystal® is a registered trademark of Alkermes Pharma Ireland Limited (APIL).

About Baudax Bio

Baudax Bio is a pharmaceutical company focused on commercializing and developing innovative products for acute care settings. ANJESO is the first and only 24-hour, intravenous (IV) COX-2 preferential non-steroidal anti-inflammatory (NSAID) for the management of moderate to severe pain. In addition to ANJESO, Baudax Bio has a pipeline of other innovative pharmaceutical assets including two novel neuromuscular blocking agents (NMBAs) and a proprietary chemical reversal agent specific to these NMBAs. For more information, please visit www.baudaxbio.com.

Forward Looking Statements

This press release contains forward-looking statements that involve risks and uncertainties. Such forward-looking statements reflect Baudax Bio's expectations about its future performance and opportunities that involve substantial risks and uncertainties. When used herein, the words "anticipate," "believe," "estimate," "may," "upcoming," "plan," "target," "goal," "intend," and "expect," and similar expressions, as they relate to Baudax Bio or its management, are intended to identify such forward-looking statements. These forward-looking statements are based on information available to Baudax Bio as of the date of publication on this internet site and are subject to a number of risks, uncertainties, and other factors that could cause Baudax Bio's performance to differ materially from those expressed in, or implied by, these forward-looking statements. These risks and uncertainties include, among other things, risks related to the ongoing economic and social consequences of the COVID-19 pandemic, including any adverse impact on the commercial launch of ANJESO® or disruption in supply chain, Baudax Bio's ability to maintain regulatory approval for ANJESO, Baudax Bio's ability to successfully commercialize ANJESO; the acceptance of ANJESO by the medical community, including physicians, patients, health care providers and hospital formularies; Baudax Bio's ability and that of Baudax Bio's third party manufacturers to successfully scale-up the commercial manufacturing process for ANJESO, Baudax Bio's ability to produce commercial supply in quantities and quality sufficient to satisfy market demand for ANJESO, Baudax Bio's ability to raise future financing for continued product development, payment of milestones and ANJESO commercialization, Baudax Bio's ability to pay its debt and satisfy conditions necessary to access future tranches of debt, Baudax Bio's ability to comply with the financial and other covenants under its credit facility, Baudax Bio's ability to manage costs and execute on its operational and budget plans, the accuracy of Baudax Bio's estimates of the potential market for ANJESO, Baudax Bio's ability to achieve its financial goals; and Baudax Bio's ability to obtain, maintain and successfully enforce adequate patent and other intellectual property protection. These forward-looking statements should be considered together with the risks and uncertainties that may affect Baudax Bio's business and future results included in Baudax Bio's filings with the Securities and Exchange Commission at www.sec.gov. These forward-looking statements are based on information currently available to Baudax Bio, and Baudax Bio assumes no obligation to update any forward-looking statements except as required by applicable law.

CONTACTS:

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BAUDAX BIO, INC.
Consolidated Balance Sheets
(Unaudited)

(amounts in thousands, except share and per share data)	September 30, 2021	December 31, 2020
Assets		
Current assets:		
Cash and cash equivalents	\$ 14,772	\$ 30,342
Short-term investments	10,150	—
Accounts receivable, net	422	51
Inventory	4,310	2,978
Prepaid expenses and other current assets	1,484	3,346
Total current assets	31,138	36,717
Property, plant and equipment, net	4,933	5,052
Intangible assets, net	22,322	24,254
Goodwill	2,127	2,127
Other long-term assets	979	583
Total assets	\$ 61,499	\$ 68,733
Liabilities and Shareholders' Deficit		
Current liabilities:		
Accounts payable	\$ 1,819	\$ 3,653
Accrued expenses and other current liabilities	4,568	5,326
Current portion of long-term debt, net	1,389	683
Current portion of contingent consideration	6,666	8,467
Total current liabilities	14,442	18,129
Long-term debt, net	6,913	8,469
Long-term portion of contingent consideration	60,059	56,576
Other long-term liabilities	813	358
Total liabilities	82,227	83,532
Commitments and contingencies		
Shareholders' deficit:		
Preferred stock, \$0.01 par value. Authorized, 10,000,000 shares; none issued and outstanding	—	—
Common stock, \$0.01 par value. Authorized, 190,000,000 shares; issued and outstanding, 84,423,342 shares at September 30, 2021 and 48,688,480 shares at December 31, 2020	844	487
Additional paid-in capital	139,951	97,034
Accumulated deficit	(161,523)	(112,320)
Total shareholders' deficit	(20,728)	(14,799)
Total liabilities and shareholders' deficit	\$ 61,499	\$ 68,733

BAUDAX BIO, INC.
Consolidated Statements of Operations
(Unaudited)

(amounts in thousands, except share and per share data)	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2021	2020	2021	2020
Revenue, net	\$ 281	\$ 68	\$ 680	\$ 417
Operating expenses:				
Cost of sales	462	540	1,869	1,190
Research and development	658	1,469	2,623	5,889
Selling, general and administrative	11,074	13,763	33,770	33,026
Amortization of intangible assets	644	643	1,932	1,502
Change in warrant valuation	(6)	(11,182)	(47)	2,863
Change in contingent consideration valuation	3,829	(17,427)	9,551	14,252
Total operating expenses	16,661	(12,194)	49,698	58,722
Operating (loss) income	(16,380)	12,262	(49,018)	(58,305)
Other expense:				
Other expense, net	(582)	(577)	(185)	(753)
Net (loss) income	<u>\$ (16,962)</u>	<u>\$ 11,685</u>	<u>\$ (49,203)</u>	<u>\$ (59,058)</u>
Per share information:				
Net (loss) income per share of common stock, basic	<u>\$ (0.20)</u>	<u>\$ 0.64</u>	<u>\$ (0.66)</u>	<u>\$ (3.84)</u>
Net (loss) income per share of common stock, diluted	<u>\$ (0.20)</u>	<u>\$ 0.62</u>	<u>\$ (0.66)</u>	<u>\$ (3.84)</u>
Weighted average common shares outstanding, basic	<u>84,400,156</u>	<u>18,374,604</u>	<u>74,008,574</u>	<u>15,366,861</u>
Weighted average common shares outstanding, diluted	<u>84,400,156</u>	<u>18,768,376</u>	<u>74,008,574</u>	<u>15,366,861</u>

BAUDAX BIO, INC.
Reconciliation of GAAP to Non-GAAP Measures
(Unaudited)

To supplement the Company's financial results determined by U.S. generally accepted accounting principles ("GAAP"), the Company has disclosed in the tables below the following non-GAAP information about adjusted net loss.

Adjusted net loss is net loss as determined under GAAP, excluding the changes in fair values of contingent consideration and warrant valuations, gain on extinguishment of debt, interest, depreciation, amortization, and stock-based compensation.

The Company believes that non-GAAP financial measures are helpful in understanding its business as it is useful to investors in allowing for greater transparency of supplemental information used by management. Adjusted net loss is used by investors, as well as management in assessing the Company's performance. Non-GAAP financial measures should be considered in addition to, but not as a substitute for, reported GAAP results. Further, Non-GAAP financial measures, even if similarly titled, may not be calculated in the same manner by all companies, and therefore should not be compared.

(amounts in thousands)	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2021	2020	2021	2020
Net loss (GAAP)	\$ (16,962)	\$ 11,685	\$ (49,203)	\$ (59,058)
Stock-based compensation	925	2,489	4,132	7,431
Non-cash interest expense	215	230	673	306
Gain on extinguishment of debt	—	—	(1,553)	—
Depreciation expense	46	104	195	315
Amortization expense	644	643	1,932	1,502
Change in warrant valuation	(6)	(11,182)	(47)	2,863
Change in contingent consideration valuation	3,829	(17,427)	9,551	14,252
Adjusted net loss (non-GAAP)	<u>\$ (11,309)</u>	<u>\$ (13,458)</u>	<u>\$ (34,320)</u>	<u>\$ (32,389)</u>

Q'3 2021 Conference Call Update Slides

November 4, 2021

Forward Looking Statements

This presentation contains forward-looking statements that involve risks and uncertainties. Such forward-looking statements reflect Baudax Bio's expectations about its future performance, ability and opportunities that involve substantial risks and uncertainties. When used herein, the words "anticipate," "believe," "estimate," "may," "upcoming," "plan," "target," "goal," "intend," and "expect," and similar expressions, as they relate to Baudax Bio or its management, are intended to identify such forward-looking statements. These forward-looking statements are based on information available to Baudax Bio as of the date hereof and are subject to a number of risks, uncertainties, and other factors that could cause Baudax Bio's performance to differ materially from those expressed in, or implied by, these forward-looking statements. These risks and uncertainties include, among other things, risks related to the ongoing economic and social consequences of the COVID-19 pandemic, including any adverse impact on the commercial launch of ANJESO® or disruption in supply chain, Baudax Bio's ability to achieve its strategic objectives, including achievement of sales forecasts and financial projections; Baudax Bio's ability to maintain regulatory approval for ANJESO, Baudax Bio's ability to successfully commercialize ANJESO; the acceptance of ANJESO by the medical community, including physicians, patients, health care providers and hospital formularies; Baudax Bio's ability and that of Baudax Bio's third party manufacturers to successfully scale-up the commercial manufacturing process for ANJESO, the accuracy of Baudax Bio's sales estimates, Baudax Bio's ability to produce commercial supply in quantities and quality sufficient to satisfy market demand for ANJESO, Baudax Bio's ability to raise future financing for continued product development, payment of milestones and ANJESO commercialization, Baudax Bio's ability to pay its debt and satisfy conditions necessary to access future tranches of debt, Baudax Bio's ability to comply with the financial and other covenants under its credit facility, Baudax Bio's ability to manage costs and execute on its operational and budget plans, the accuracy of Baudax Bio's estimates of the potential market for ANJESO, Baudax Bio's ability to achieve its financial goals; and Baudax Bio's ability to obtain, maintain and successfully enforce adequate patent and other intellectual property protection. These forward-looking statements should be considered together with the risks and uncertainties that may affect Baudax Bio's business and future results included in Baudax Bio's filings with the Securities and Exchange Commission at www.sec.gov. These forward-looking statements are based on information currently available to Baudax Bio, and Baudax Bio assumes no obligation to update any forward-looking statements except as required by applicable law.

Executive Summary

Tracking 2021 exit rate for more robust 2022 growth

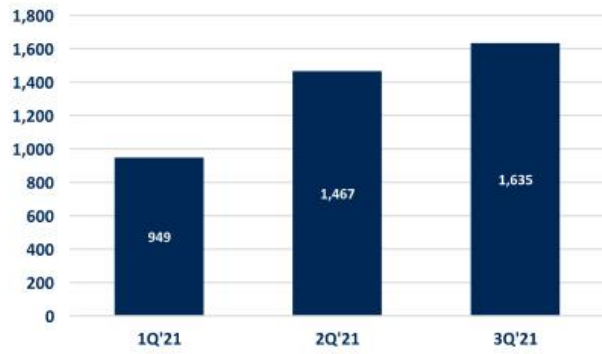
ANJESO growth accelerating in September after summer delta variant peak

Peer-to-Peer programs and National Accounts are helping ANJESO gain momentum and setting the stage for Q4 and 2022

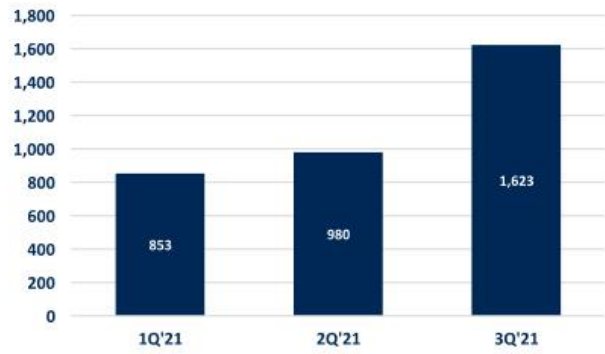
Deepening Usage In Existing Hospital And ASC Customers Contributed To Strong Q'3 2021 Sales

11% growth in existing Hospital sales; 66% growth in sales to existing ASC customers

Quarterly Hospital Sales (Existing)

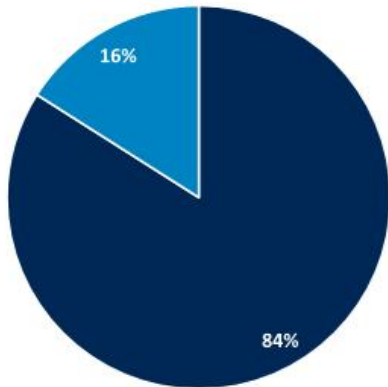


Quarterly ASC Sales (Existing)



150+ Formulary Wins by end Q'3; P&T Success Rate 84%

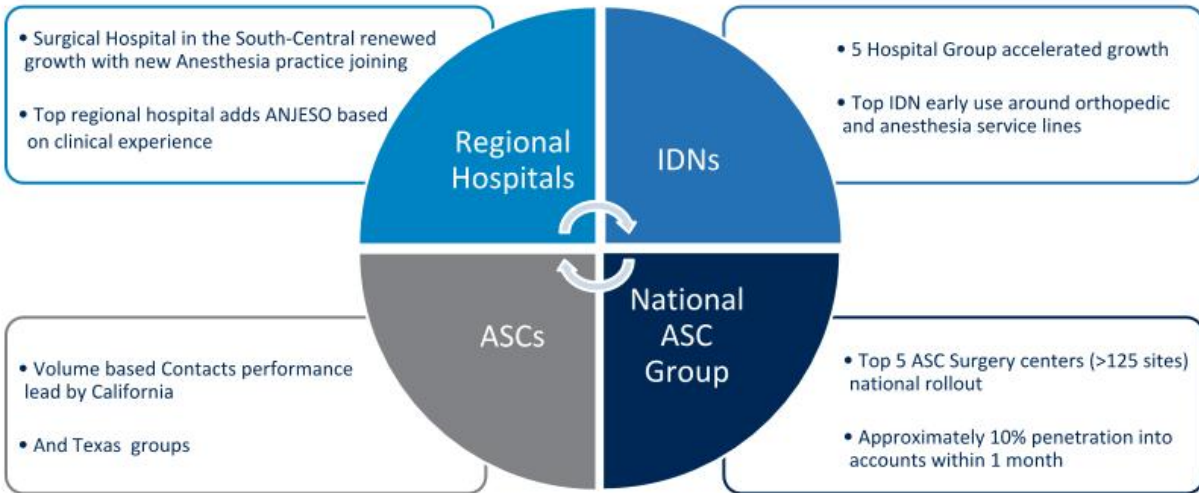
LTD P&T Review Results



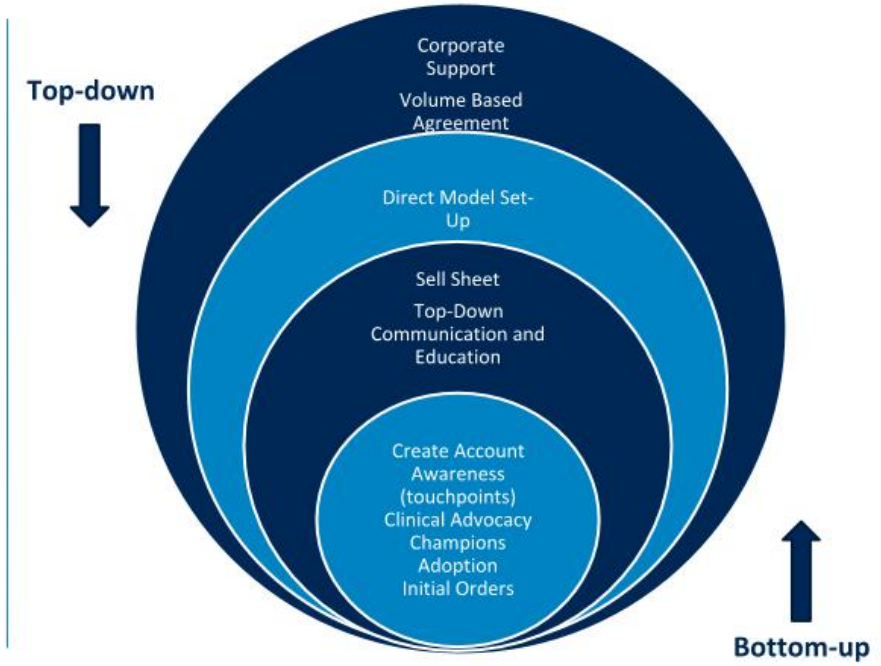
■ Approved ■ Denied

- 70% of upcoming P&T Reviews are Hospital
- Top 5 ASC and services provider with >125 ASCs nationwide approved addition of ANJESO to formulary:
 - Only 8 of these ASCs are included in our formulary wins to date
 - Additional ASCs progressing for approval and onboarding of ANJESO
 - Significant opportunity for broader adoption as we add more active formulary ASCs

Q3 Highlights Demonstrated Success Across Arenas



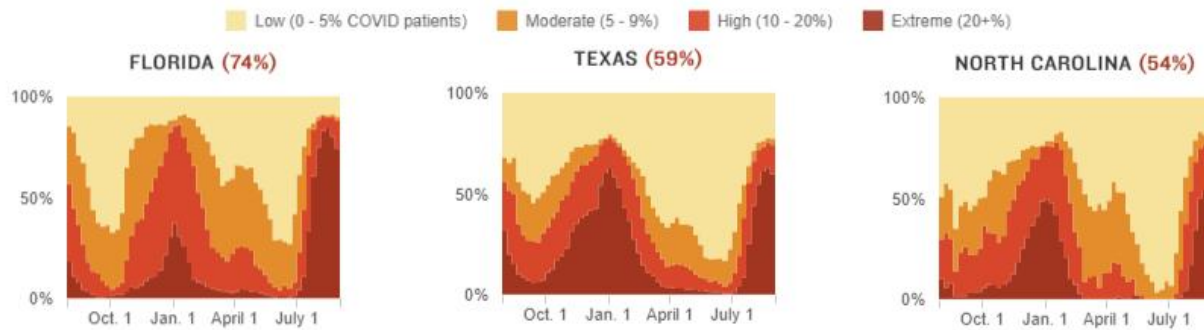
Our Strategic Approach To Large Accounts



Impact of the Pandemic In Our Most Valuable Areas

Florida, Texas and North Carolina Are Our Top 3 States With Over 40% Of ANJESO Sales

- These states have been among the highest % of hospitals suffering from “extreme” stress based on COVID hospitalizations vs. available beds & ICU capacity



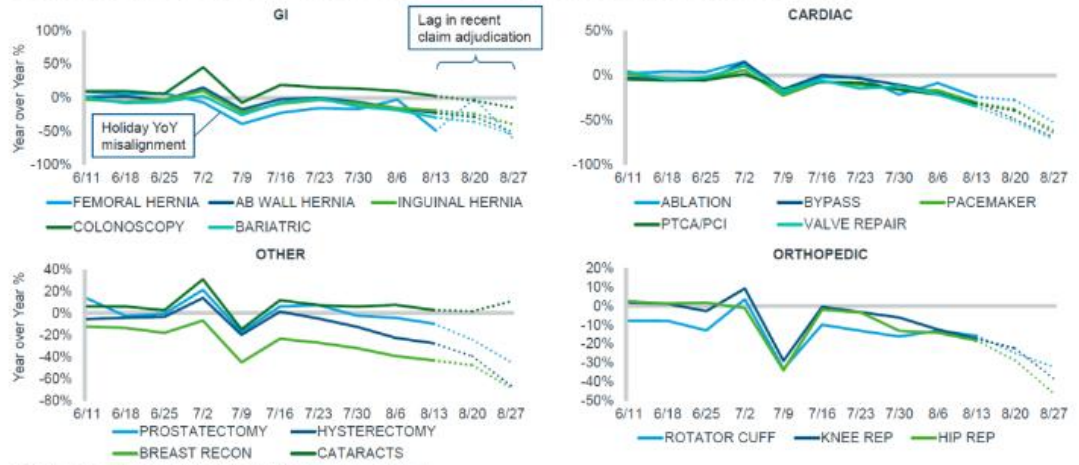
CA, AZ, AR, AL and MO are also heavily impacted and account for another 30% of sales

Orthopedic & Other Procedures Declined Significantly During Summer Covid Flare

ANJESO Utilization by Top Procedures:

- 46% Ortho
- 22% Pain/RFA
- 16% Uro/Gyn

Elective procedures show consistency across procedure type



*See Appendix for Elective Procedure Actual vs. Calibrated 2020 volumes as reference
 Source: IQVIA: Medical Claims Data Analysis, 2021, Week 35 2019 - 8/8/2019; Week 35 2020 - 9/4/2020; Week 34 2021 - 8/27/2021. Elective procedures based on IQVIA custom analysis; Data includes claims from inpatient & outpatient care sites; Latest two weeks are estimates using a recency data factor that is adjusted out as all claims are received; See Appendix for further details

COVID-19 Market Impact - w/e August 27, 2021



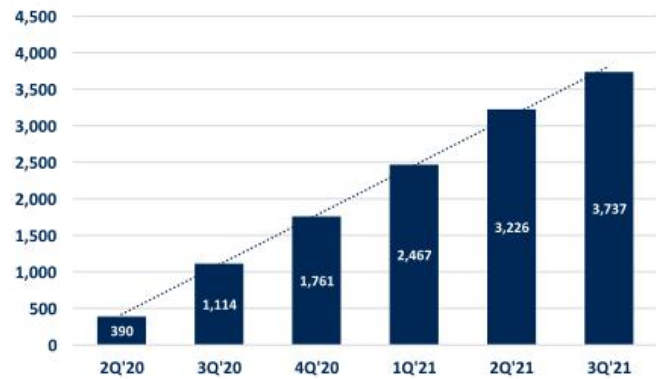
And Sales Continue To Show Quarter Over Quarter Growth in Q'3 2021

September rebound and overall growth in spite of July-August CV-19 Delta Impact

Monthly ANJESO® Units*
(Actuals thru 9/30/21)



Units Sold by Quarter



Average Weekly Sales (2021):

Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep
124	235	206	203	237	293	260	277	318

Income Statement – Q3'21 versus Q3'20

(amounts in thousands)	For the Three Months Ended September 30,		For the Nine Months Ended September 30,	
	2021	2020	2021	2020
Revenue, net	\$ 281	\$ 68	\$ 680	\$ 417
Operating expenses:				
Cost of sales	462	540	1,869	1,190
Research and development	658	1,469	2,623	5,889
Selling, general and administrative	11,074	13,763	33,770	33,026
Amortization of intangible assets	644	643	1,932	1,502
Change in warrant valuation	(6)	(11,182)	(47)	2,863
Change in contingent consideration valuation	3,829	(17,427)	9,551	14,252
Operating (loss) income	16,661	(12,194)	49,698	58,722
Operating loss	(16,380)	12,262	(49,018)	(58,305)
Other expense:				
Other expense, net	(582)	(577)	(185)	(753)
Net (loss) income	\$ (16,962)	\$ 11,685	\$ (49,203)	\$ (59,058)

The logo for Baudax Bio, featuring the company name in a dark blue, sans-serif font. The 'x' in 'Baudax' has a small blue cross-like symbol above it. The background of the slide is split diagonally from the top-left to the bottom-right, with a dark blue upper-left section and a lighter blue lower-right section.

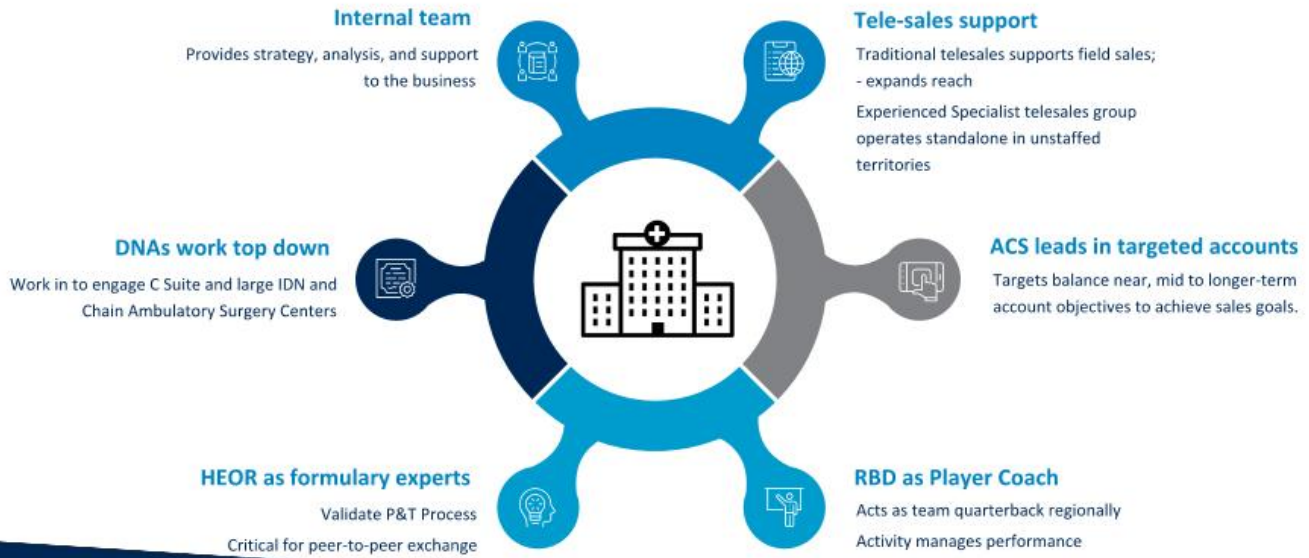
Baudax BIO™

Closing Q'4 And Preparing For 2022

Commercial Roles Work Closely Together

Field Role Integration is Critical

Field Leadership Meeting 9/27

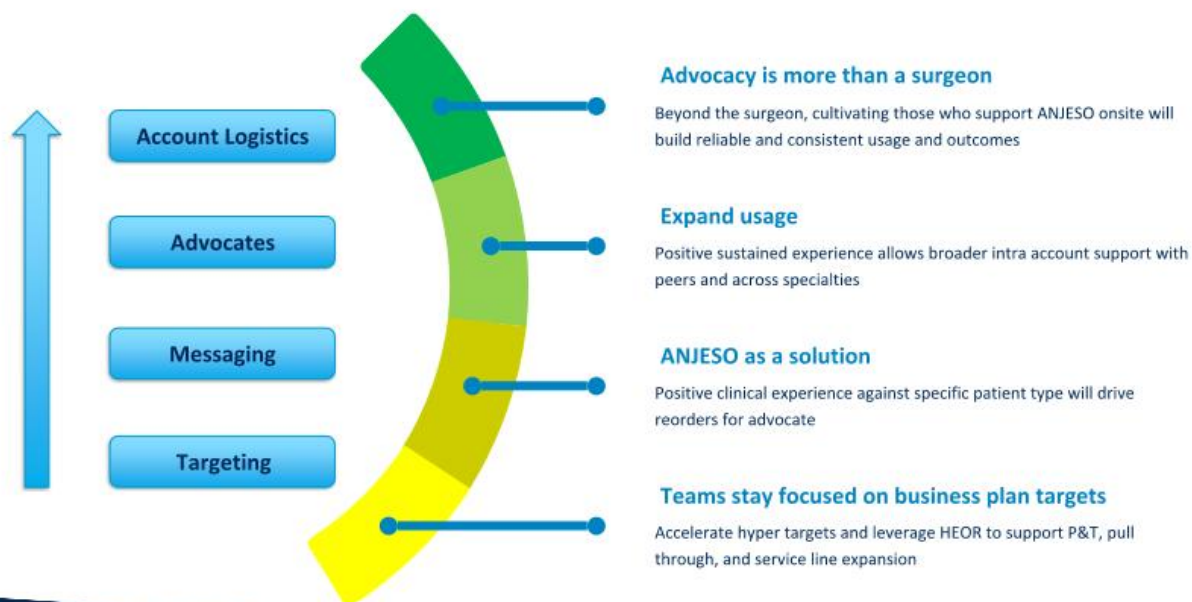


Tailored Communication By Specialty Is Key To Increasing Likelihood To Trial

Evolving the Sophistication of our Messages



Early Trial Accounts Are Opportunity To Expand Usage



Stage Is Set For 2022

