

Acorda Therapeutics 2021 Q1 Earnings Call

May 6, 2021



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SCIENCE.**
ACORDA
THERAPEUTICS

Forward Looking Statement

This press release includes forward-looking statements. All statements, other than statements of historical facts, regarding management's expectations, beliefs, goals, plans or prospects should be considered forward-looking. These statements are subject to risks and uncertainties that could cause actual results to differ materially, including: we may not be able to successfully market AMPYRA, INBRIJA or any other products under development; the COVID-19 pandemic, including related quarantines and travel restrictions, and the potential for the illness to affect our employees or consultants or those that work for other companies we rely upon, could have a material adverse effect on our business operations or product sales; our ability to raise additional funds to finance our operations, repay outstanding indebtedness or satisfy other obligations, and our ability to control our costs or reduce planned expenditures; risks associated with the trading of our common stock and our reverse stock split; risks related to our workforce, including our ability to realize the expected benefits of our corporate restructuring; risks associated with complex, regulated manufacturing processes for pharmaceuticals, which could affect whether we have sufficient commercial supply of INBRIJA to meet market demand; our reliance on third-party manufacturers for the production of commercial supplies of AMPYRA and INBRIJA; third party payers (including governmental agencies) may not reimburse for the use of INBRIJA or our other products at acceptable rates or at all and may impose restrictive prior authorization requirements that limit or block prescriptions; competition for INBRIJA, AMPYRA and other products we may develop and market in the future, including increasing competition and accompanying loss of revenues in the U.S. from generic versions of AMPYRA (dalfampridine) following our loss of patent exclusivity; the ability to realize the benefits anticipated from acquisitions, among other reasons because acquired development programs are generally subject to all the risks inherent in the drug development process and our knowledge of the risks specifically relevant to acquired programs generally improves over time; the risk of unfavorable results from future studies of INBRIJA (levodopa inhalation powder) or from our other research and development programs, or any other acquired or in-licensed programs; the occurrence of adverse safety events with our products; the outcome (by judgment or settlement) and costs of legal, administrative or regulatory proceedings, investigations or inspections, including, without limitation, collective, representative or class action litigation; failure to protect our intellectual property, to defend against the intellectual property claims of others or to obtain third party intellectual property licenses needed for the commercialization of our products; and failure to comply with regulatory requirements could result in adverse action by regulatory agencies.

These and other risks are described in greater detail in our filings with the Securities and Exchange Commission. We may not actually achieve the goals or plans described in our forward-looking statements, and investors should not place undue reliance on these statements. Forward-looking statements made in this press release are made only as of the date hereof, and we disclaim any intent or obligation to update any forward-looking statements as a result of developments occurring after the date of this presentation.

INBRIJA Net Sales – Q1 2021



Inbrija®
(levodopa inhalation powder)

\$5M Q1 2021 net revenue

13% ↑ over Q1 2020

AMPYRA Net Sales – Q1 2021



\$20.3M Q1 2021
net revenue
Flat to Q1 2020



Inbrija  TM
(levodopa inhalation powder)
42 mg capsules

INBRIJA Growth Q1 2020 – Q1 2021

2020



\$4.4M
net sales



2957
TRx



6955
cartons



2021

\$5M
net sales

↑ 13%

3106
TRx

↑ 5%

8679
cartons

↑ 25%

Organic growth of **25%**

Regular Users are Key Contributors to Growth



~44%

of patients are
regular users
(taking INBRIJA
≥1x per day)



~70%

Regular users
still on at one
year



Average
consumption of
regular users is
2.8x per day
Up from 2.5x
per day April '20

INBRIJA 2021 Growth Catalysts

Prescription Conversions

- Increased from ~50% to ~67% in 2020
- Increased access and coverage
- Streamlined SP network

Patient Experience

- Improved training materials
- Nurse Educators
 - Refills increased 20%
- Patient Ambassadors

Physician Reengagement

- > 550 new HCPs prescribed in 2020
- Field team returned to in-person visits



Think MS
Think Walking
Think AMPYRA

ampyra[®]
(dalfampridine) ER tablets

Selected Important Safety Information

AMPYRA is contraindicated in patients with history of seizures, moderate or severe renal impairment (CrCl ≤ 50 mL/min), or history of hypersensitivity to AMPYRA or 4-aminocoumarin.

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Real Patients. Real Results.
Examples of Timed 25-Foot Walk videos now playing in the interactive panels of this booth.

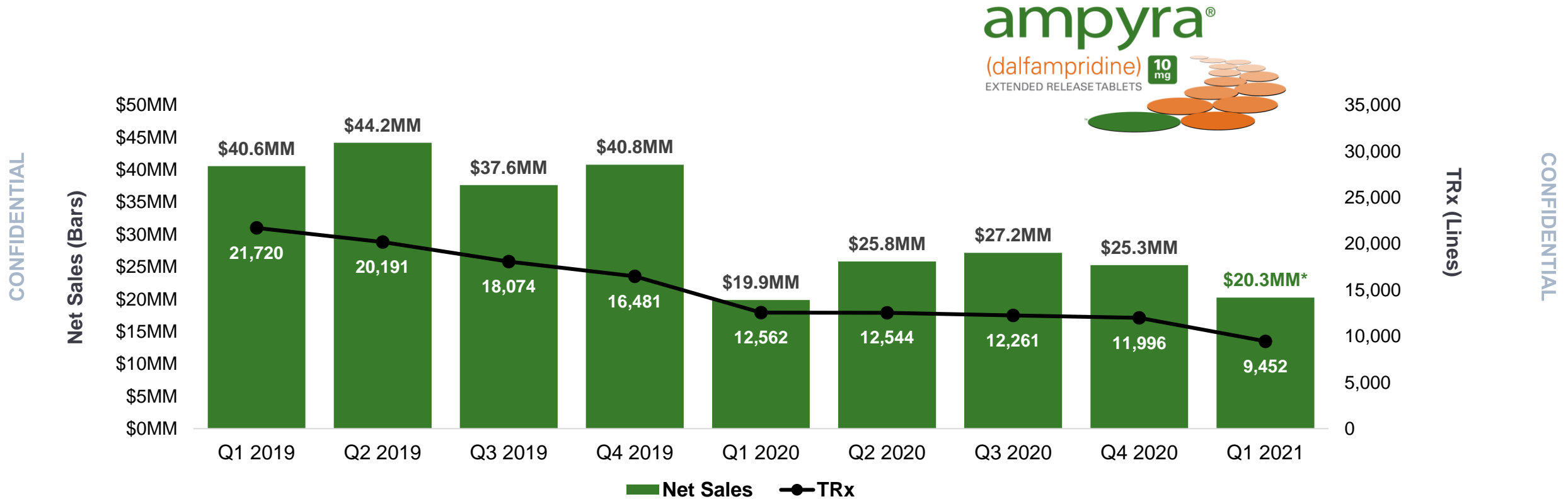
Selected Important Safety Information:
AMPYRA is contraindicated in patients with history of seizures or with epileptiform activity on an EEG, as these patients could be at increased risk of seizures.

For more information, see Important Safety Information and Full Prescribing Information.

Please see additional Important Safety Information and Full Prescribing Information within this booth.

Ampyra[®]

AMPYRA Franchise Strength



Q1 2021 Delivered Stable Ampyra Net Sales For the First Time Since LOE



2021 Financials and Goals

Goal: Cash flow neutral by YE 2022



AMPYRA
durable



INBRIJA
poised
for growth



FAMPYRA
royalties



Ardsley
facility



Fiscal
discipline
on OpEx

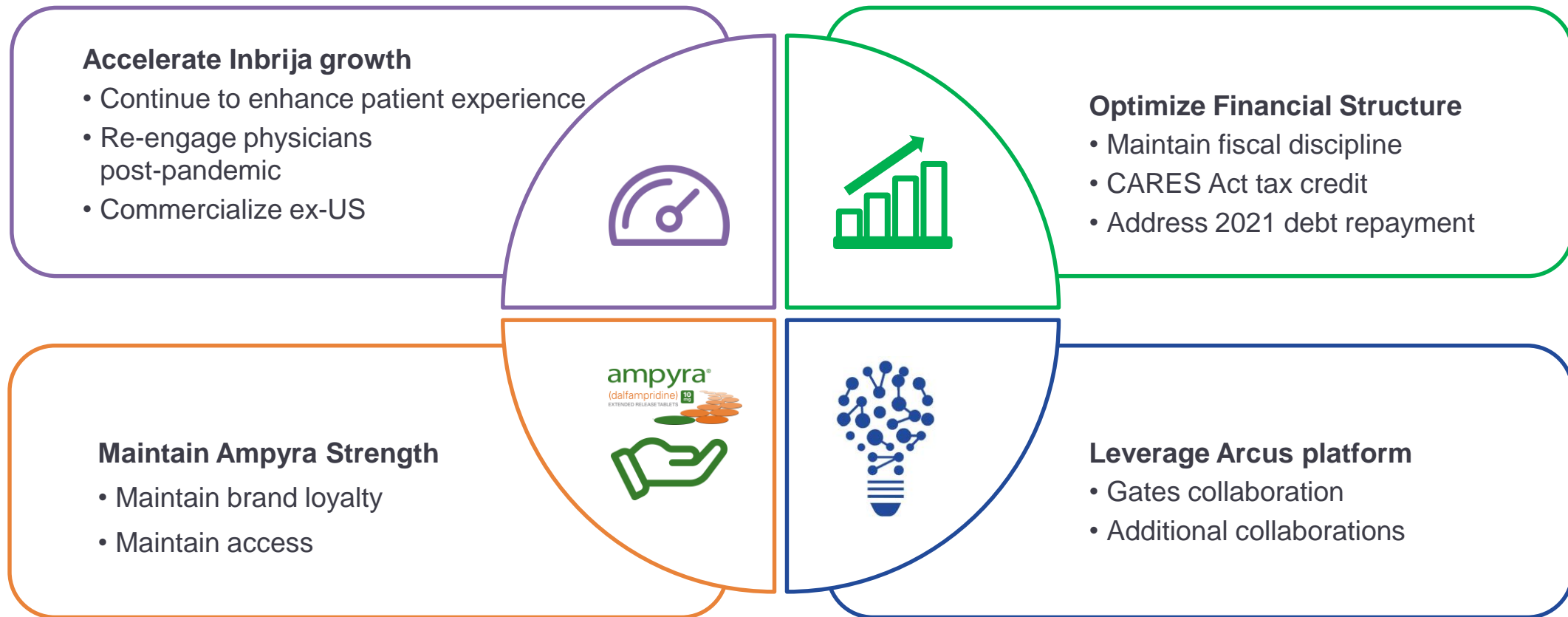
Q1 2021 Financial Summary

(\$ in millions)	1Q'21	1Q'20	Δ Q/Q
Net Inbrija Revenue	5.0	4.4	13.6%
Net Ampyra Revenue	20.3	20.1	1.0%
R&D	4.7	7.7	(39.0%)
SG&A	34.0	41.1	(17.3%)
GAAP Net Loss	(33.5)	(6.5)	415.4%
Non-GAAP Net Loss	(23.3)	(24.4)	(4.5%)
Cash, Cash Equivalents, Short-Term Investments and Restricted Cash*	148.4	126.3	17.5%

This slide contains GAAP and non-GAAP financial measures. Non-GAAP net (loss) excludes certain items. Information regarding our use of non-GAAP measures, a description of excluded items, and a reconciliation of those measures to GAAP is available in our financial results press release dated May 6, 2021, which is available in the investor relations section of our website at www.acorda.com.

*Includes marketable securities.

Building Long Term Value



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