orexo

A profitable specialty pharmaceutical company – with the aim of becoming a leader within treatment of addiction











Nordic-American Life Science Conference, Dec. 5, 2019

Robert DeLuca, President Orexo US Inc.

Nasdaq Stockholm: ORX US OTC Market: ORXOY (ADR)

Company Overview









Orexo is a Swedish specialty pharmaceutical company

- Focuses on **treatment of addiction** in all phases, from prevention to treatment
- Broad pipeline with multiple assets based on innovative drug delivery technologies and digital therapeutics
- Developed four commercial products with worldwide approval
- Direct presence in the **US opioid dependence market**, one of the biggest health crises in US history, with lead asset **Zubsolv**®
- Strategic focus on **portfolio expansion** through development and licensing/M&A
- Growing profitability and strong financial position to support future growth
- Top two largest shareholders¹: **Novo Holdings** (27.2%) and **HealthCap** (10.2%)



Corporate Headquarters (Uppsala, Sweden) Corporate functions and Development



US Headquarters (Morristown, New Jersey) Commercial subsidiary incl. fully owned field force

Net Revenues EBITDA LTM, USD, Q319 LTM, USD,

87_m

Cash Position,

USD, Sep. 30, 2019

¹ As of September 30, 2019 Note: USD based on average exchange rate from Q2, 2019 SEK/USD 9.5



Products & Pipeline



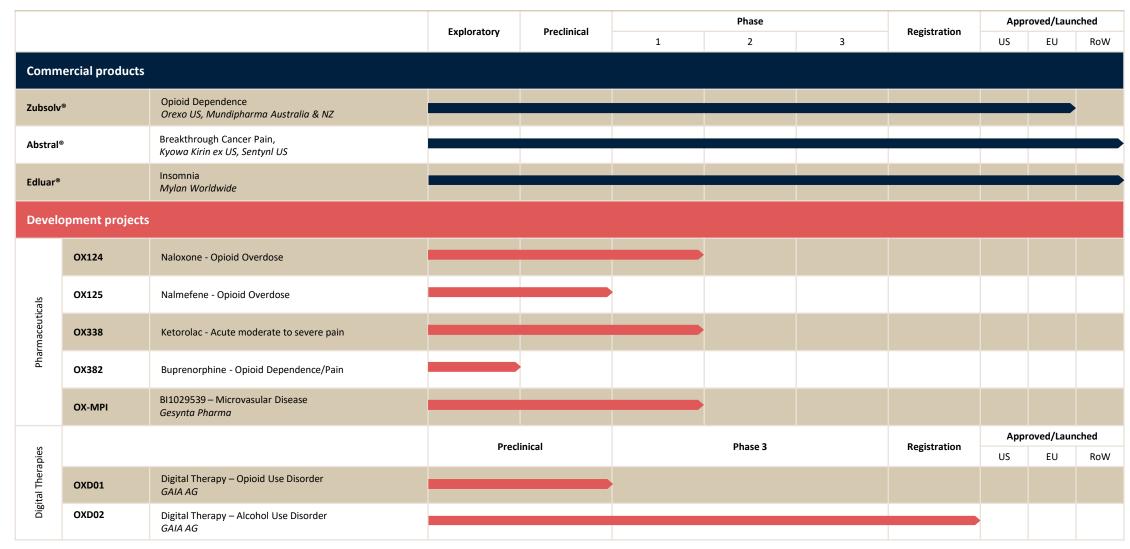






Pipeline of multiple potential assets

The partnership with GAIA has extended Orexo's technology platform to also include Digital Therapies





OX124 and OX125 – the projects at a glance

Unmet medical need

- 70,200 people died from a drug overdose in the US in 2017
- Synthetic opioids, such as fentanyl, are now the leading cause of death, which are more potent and stay in the body for a longer time than heroin
- Narcan® Nasal Spray, the leading naloxone rescue drug, is effective, but has shortcomings:
 - ~34% of overdose patients require more than one dose of Narcan
 - Half life of 2 hours (vs. 8-10 for fentanyl) bears risk of 2nd overdoses

OX124 and OX125 concept

- Unique and improved nasal formulations of naloxone (OX124) and nalmefene (OX125) to specifically address the challenges arising from the fentanyl crisis
 - Ability to reverse effect of most powerful synthetic opioids
 - Longer duration than currently approved formulations to reduce need for 2nd doses



Committed to find new solutions to improve treatment of addiction

Orexo has recently closed two deals with GAIA AG, a worldleader within digital therapeutics (DTx)

GAIA develops scientifically proven DTx

- GAIA has demonstrated digital therapies can have impressive improvement in treatment outcomes from supporting behavioral change and adherence to treatment
- GAIA's platform has proven effectiveness in numerous RCTs with over 10,000 patients
- GAIA has 12+ years R&D experience, 20+ CE & FDA compliant products
- GAIA's products use artificial intelligence and offer unique individualization of the treatment



OXD01 (opioid use disorder, OUD)



- Complementary product
- Significant synergies across the entire value chain
- Expected US launch in 2022
- Holds the global rights

OXD02 vorvida® (alcohol use disorder, AUD)



- US license deal
- Synergistic asset
- FDA discussion required whether an additional trial will be needed
- Launch in 2020 or 2021



Key Market & Sales







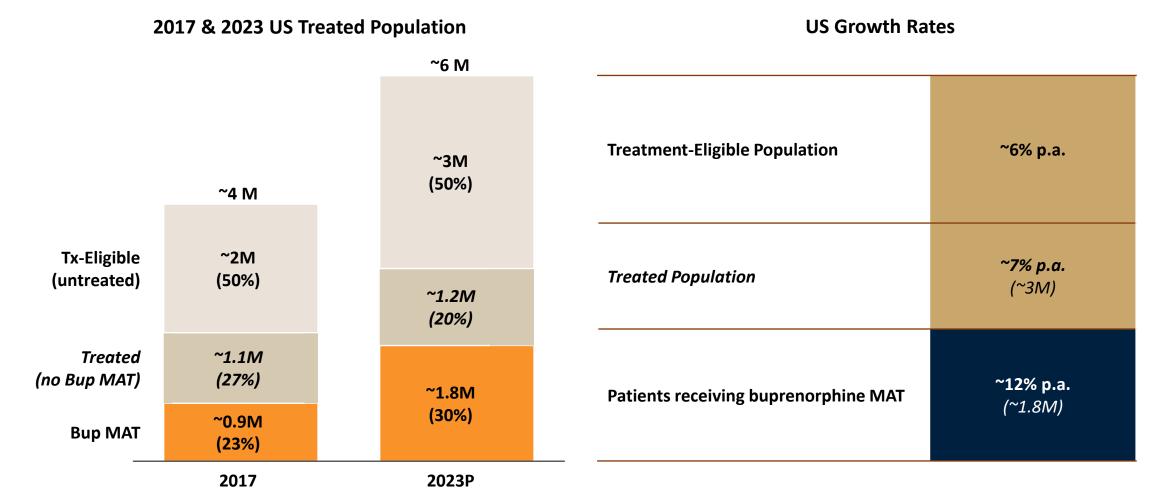


Orexo focuses on the number one health crisis in the US





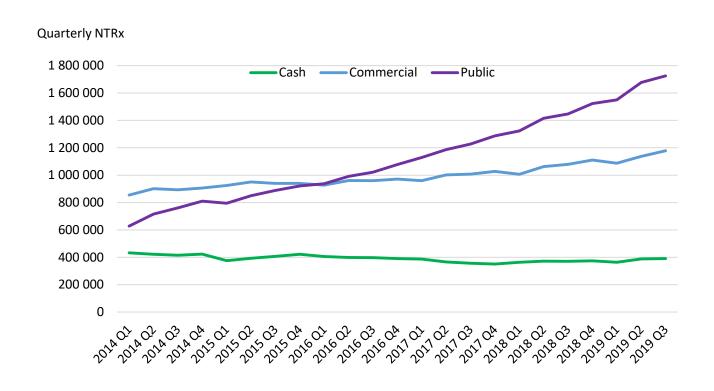
US buprenorphine MAT population likely to double by 2023





YTD Year-over-year market growth at 14 percent

New market highs in Public and Commercial NTRx in Q3 and Zubsolv grows 16 percent in open business



Q3 YoY Growth		
Market	Zubsolv [®] total	Zubsolv®
+19%	-5%	+17%
+11%	+1%	+16%
+4%	-33%	

Note: Quarterly NTRx levels =Total prescriptions adjusted to 30 tablet/film scripts

Note: Historical quarters restated due to IMS recategorization

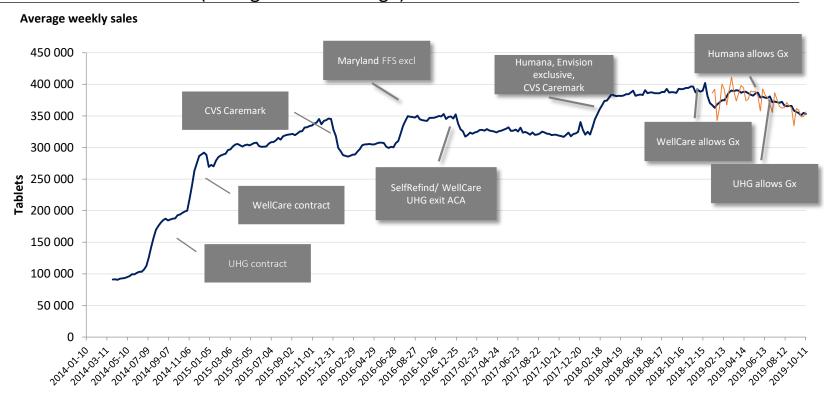
*Open Business segment includes all payers, excluding current & recently former exclusive payers

Source: Orexo analysis, IMS data



In Q3 2019 Zubsolv® sales grew in SEK by 10 percent despite declined volume due to formulary changes

Zubsolv Tablet Volume (rolling 4 week average)



Q3 2019 Zubsolv volume declined due to loss of exclusive contracts

- Volume declined 4% since Q318 and 5% from Q219
- WellCare, Humana and UHG change from exclusivity and declining cash segment have had a negative effect of ~18% on overall volume growth vs Q318 and 9% since Q2 19
- Decline from loss of three exclusive contracts continues but at a slowing pace
- "Open business*" grew 16 percent from Q318 and no change since Q219

^{*}Open Business segment includes all payers, excluding current & recently former exclusive payers

Note: Weekly prescription data is based on extrapolation and is associated with uncertainties and may differ between sources

Source: Orexo analysis, IMS NPA weekly data



The entry of generic film caused the loss of exclusive contracts, but recent market dynamic will offer opportunities

WellCare removes exclusivity on Zubsolv®

Humana removes exclusivity on Zubsolv

United removes exclusivity on Zubsolv

NJ enacted law for Medicaid to reimburse Zubsolv

Ohio, Texas, Florida DC, BCBS NC reimburse Zubsolv

Delaware removes Zubsolv from formulary (small)

Zubsolv reimbursement stable entering 2020*

2019

Brixadi launch blocked until Nov. 2020

2020

CVS Caremark first major PBM to block branded Suboxone® Film

Indivior to discontinue authorized generic ending Sandoz partnership

^{*} As per October 24th 2019



Financials









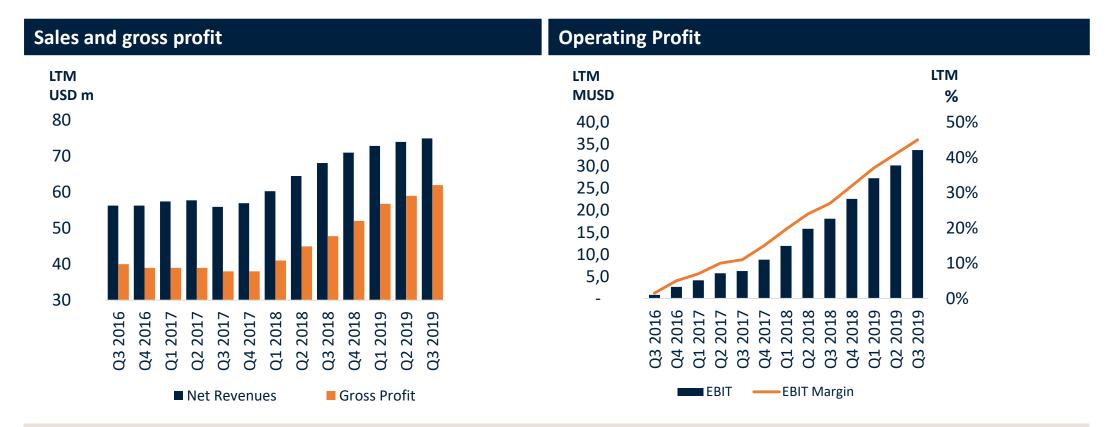
Continued strong performance in Q3 2019

Group Net Revenues LTM ¹	Growth
USD 87 m	12%
83.4% from Zubsolv®	(Zubsolv 20%)
Group EBITDA LTM ¹	Growth
USD 24 m	110%
US EBIT LTM ¹	Growth
USD 33 m	101%
Cash position Q3 2019	Net Cash position Q3 2019
USD 85 m	USD 55 m

 $^{^{1}}$ LTM, Last Twelve Months Note: USD based on average exchange rate from Q3, 2019 SEK/USD 9.6



Orexo US Operating Margin (LTM) grew to 45.2%



- Net Sales grew to USD 74.6 m from USD 68.1 m in Q318
- Strong growth in EBIT contribution to USD 33.7 m increasing from USD 18.1 m in Q318 driven by organizational efficiencies and by lower COGS
- US EBIT margin of 45.2% LTM in Q319 increasing from 26.6% in Q318, EBIT margin in Q3 reached 51.1%

Note: COGS converted from SEK to USD using monthly average exchange rates for the period.



Outlook









Strong value drivers for long-term growth

Future value drivers

1. Growing key market

14 percent growth YTD in 2019 in a market addressing one of the largest health crises ever in America and a growing global concern

2. Strong financial position and profitability

Fueled by the US commercial organization which is an important cash and profitability contributor on a Group level

3. Strong track record of developing products

Orexo has developed four products with worldwide approval

4. M&A and business development

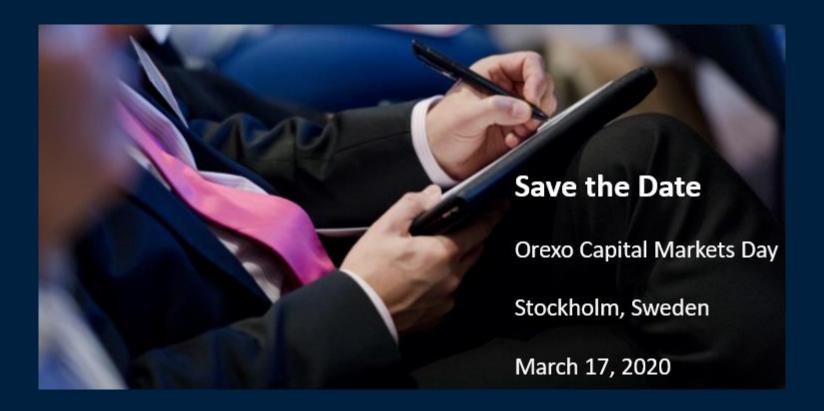
Add commercial stage products in the US to leverage the commercial infrastructure and expand sales

5. Expanding pipeline

Growing pipeline including several interesting projects to embrace all aspects of opioid addiction



Thank You **Q&A**



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