A pharmaceutical & digital therapeutics company

November, 2022

Orexo supports the UN's Agenda 2030 with a focus on:



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Orexo – our business areas



HQ & Pipeline

Development of improved drugs based on well-known substances combined with innovative proprietary Drug Delivery technologies, such as amorphOX[®].



US Pharma

Commercial US Pharma platform since 2013, incl. market access team and sales representatives who on a daily basis visit physicians, medical clinics and minor hospitals.



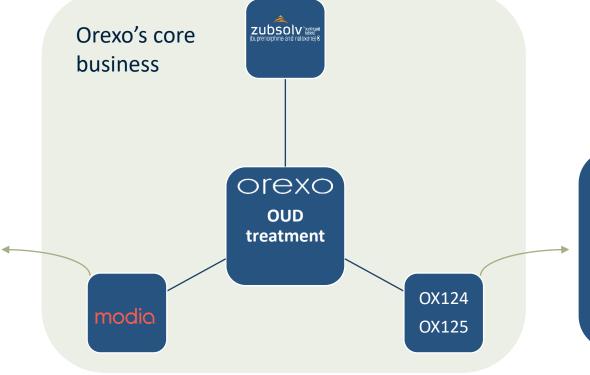
Digital Therapeutics

Evidence-based digital therapies grounded in cognitive behavioral therapy techniques, offer better treatment access for patients and improve their outcomes.

A strong focus on opioid misuse

DTx

- Scalable reimbursement platform
- DTx in mental health with co-morbidities to OUD
 VORV!DA*



amorphOX®

- OX640
- Technology outlicense and partnering in development

Pioneer in future growth area with proprietary reimbursement platform and products Profitable and stable US Pharma business with long presence in significantly underserved market and significant near-term milestones from MODIA[®] study and OX124 approval enabling a comprehensive treatment solution to OUD

World leading technology enabling new profit generating partnerships

Commercial products and development pipeline

Pharmaceuticals										
Product/ Project		Exploratory	Preclinical	Clinical development phases		- Registration	Approve US	ed and/or La	aunched RoW	
ZUBSOLV [®] Opioid Use Disorder/ sublingual platform	ACCOrd Make it better					-				
Abstral [®] Breaktrough Cancer Pain/ sublingual platform	KYOWA KIRIN									
Edluar® Insomnia/ sublingual platform	📶 Mylan									
OX124 Naloxone, Opioid Overdose/ amorphOX®							Q4 2022			
OX125 Nalmefene, Opioid Overdose/ amorphOX®										
OX640 Adrenaline, allergic reactions/ amorphOX [®]					•		2025			
OX-MPI Endometriosis										
Digital Therapeutics										
		Explor	ratory		Developme	ent	Registration			
modia	GAIA									
VORV!DA°	GAIA									
deprexís	GAIA									

US market: Commercial products targeting a large unmet need



Opioid depender	Alcohol misuse	Depression		
zubsolv modia OX124 (expected 2024)		VORV!DA	deprexís®	
Overdose deaths surpassed 107 000 in 2021, fueled by use of synthetic opioids such as fentanyl	10m are misusing opioids	17m are heavy alcohol users	20m are suffering from at least one major depression annually	
120000 80000 40000 2017 2018 2019 2020 2021 Total overdose deaths All opioid overdoses Synthetic opioids eg fentanyl	1.4m in treatment	1.5m in treatment	10m in treatment	

ZUBSOLV[®] an important foundation to Orexo's commercial operations

Net revenues since launch 2013 (SEK m)



m



¹ Last Twelve Months

Market dynamic

Sales volume stabilizing during 2022

- New reimbursement agreements in NY and Kentucky main growth drivers
- Decline in largest payers UHG and Humana after market leader lost exclusivity in 2019
- All market access maintained 2023 Market growth double digit pre-covid, but slowed down to 5% due to Covid-19 and Fentanyl crisis
- Significant increase in funding expected in 2023 and beyond





562 MSEK revenues LTM (Q3 2022)

Orexo is at the leading edge of digitalization in the pursuit to take DTx from its infancy to become a natural part of healthcare

"In less than a decade, DTx companies have completely disrupted the healthcare scene for the better."

The Future of Digital Therapeutics and The Impact On Care, The Linus Group, May 2021

Establishing Reimbursement Orexo working in tight collaboration with world leaders in digital health to make DTx accessible to all patients. However, universal reimbursement processes still to be established

✓ Disruptive technology

Through pilot programs, trials and real world evidence collection, Orexo is working with payers and leading healthcare organizations to build confidence in the value of our DTx to healthcare.

Clinically proven DTx

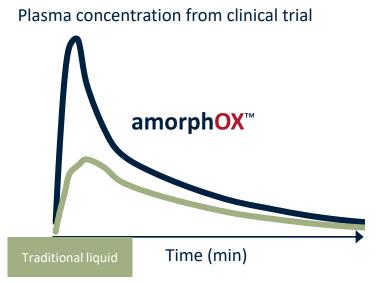
Rooted in cognitive behavioral therapy techniques and based on AI technology offering a highly individualized intervention.

	modia	VORV!DA	deprexis
Instructions for use	Opioid dependence	Alcohol misuse	Depression
Clinical evidence	Ongoing randomized clinical trial incl. 437 patients	Evaluated in 1 randomized clinical trial, > 600 patients	Evaluated in 13 randomized clinical trials, > 2.800 patients
Length of treatment	6 months	6 months	3 months
Treatment method	Along with current standard of care including medication	Standalone or as a complement to current standard of care	Standalone or as a complement to current standard of care
FDA clearance	Will apply for a 510 k clearance, meanwhile launched under FDA's Public Health Emergency Use Authorization (EUA)	FDA cleared under the EUA	FDA cleared under the enforcement discretion

Orexo's development platform building on the proprietary amorphOX® technology

Validated in humans

 Superior pharmacokinetic properties with more rapid onset, higher peak and overall exposure, lower variability

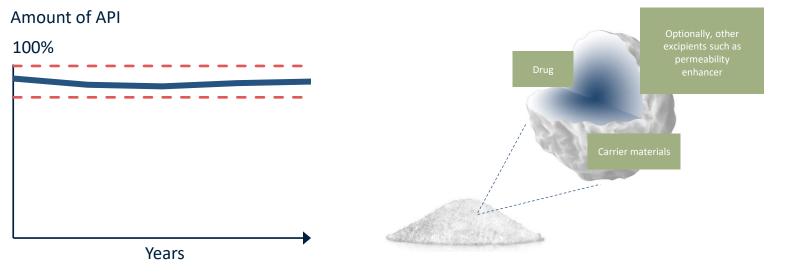


Excellent stability

 Excellent stability even under accelerated conditions and proven to work on a broad scope of API's

Wide applicability

 Powder technology that works with a broad scope of small and large molecules, such as peptides and proteins



OX124 & OX125 – overdose rescue medications

- Significant health issue in the US with >107.000 deaths from overdose in 2021
- Based on amorphOX[®] and designed to treat overdoses caused by synthetic opioids, such as fentanyl
- OX124 clinically differentiated to market leader and GX of market leader
- OX124 on track to be filed with FDA in Q4 2022





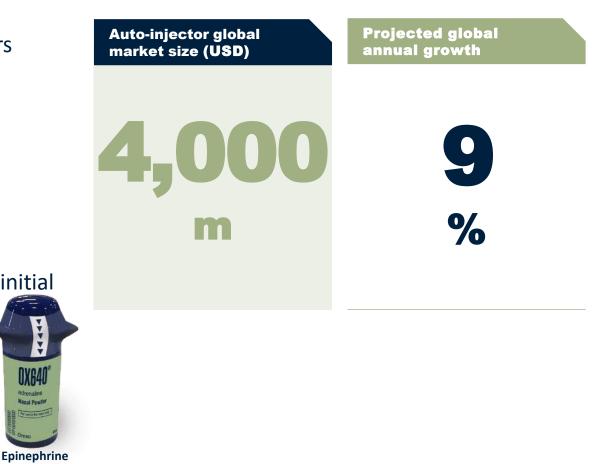
OX640 – emergency treatment of allergic reactions

- First line treatment today: intramuscular auto-injectors
- OX640 offers clear differentiation
 - Needle free alternative based on amorphOX[®]
 - Chemically stable and more robust formulation
 - Optimized manufacturing
 - Free of antioxidants or preservatives
- No meaningful innovation for decades
- OX640 could be ready for FDA filing in 2025 based on initial FDA feedback on clinical evidence required

OX64

Adrenaline

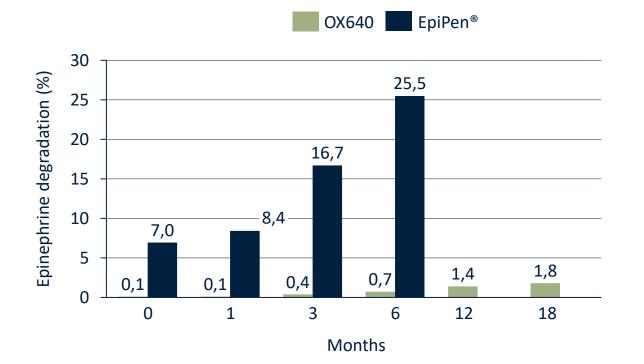
Positive data showed in the clinical phase 1 study



OX640: The power to transform emergency treatment of allergic reactions



OX640: Nasal epinephrine enabled by amorphOX® - remains stable in any life situation





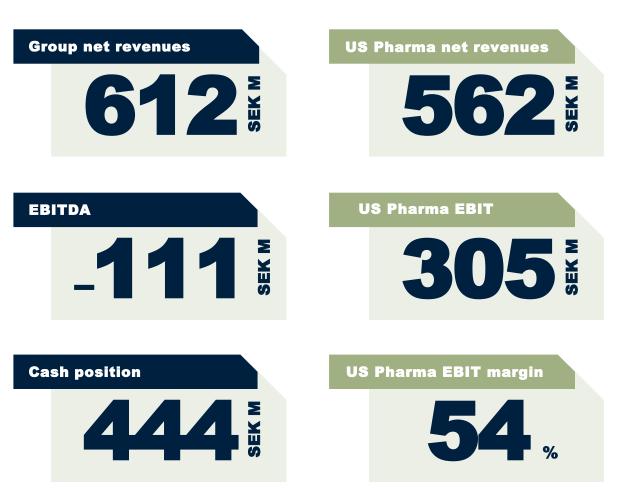




OX640 will work indpendent on temperature, from a hot car in Texas to a winterday in Colorado

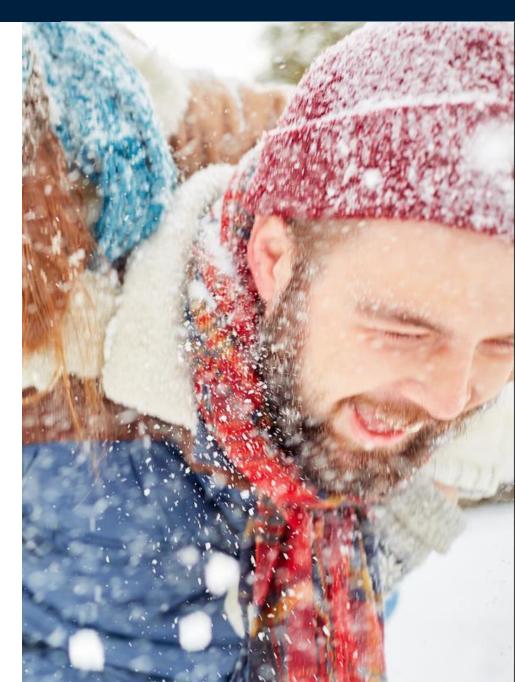
Q3 2022 LTM¹ – A transformative period building for future growth

- Significant investments in establishing the DTx business, conducting clinical trials and to protect ZUBSOLV[®] IP rights
- Recurring business is financed from ZUBSOLV® profit and royalty contributions



Why Orexo?

- Corporate profitability in sight
 - Main external cost drivers will diminish during H1 2023
 - No new activity driving external expenses to be initiated without certainty of associated revenues
 - Significant focus on cost efficiency
- ZUBSOLV[®] sales stabilized and opportunities to grow
 - Settlements providing USD 54 billions announced
- DTx revenues expected to build from Q4 and beyond
 - Reimbursement model confirmed
- **R&D pipeline** is expected to result in revenue generating partnerships 2023
 - OX124 filing with the FDA in Q4 2022 (to be commercialized by Orexo)
 - OX640 and amorphOX[®] partnering discussions on-going



For more information please visit <u>www.orexo.com</u>. You can also follow Orexo at Twitter @orexoabpubl, LinkedIn and YouTube.

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