CREATING VALUE FOR INVESTORS THROUGH NEXT PHASE OF GROWTH



DISCLAIMER

- * This Presentation has been prepared by Laboratorios Farmacéuticos ROVI, S.A. (the "Company") and comprises the slides for a presentation concerning the Company and its subsidiaries (the "Group"). For the purposes of this disclaimer, "Presentation" means this document, its contents or any part of it, any oral presentation, any question or answer session and any written or oral material discussed or distributed during the Presentation meeting or otherwise in connection with it.
- * This Presentation does not constitute or form part of, and should not be construed as, any offer to sell or issue or invitation to purchase or subscribe for, or any solicitation of any offer to purchase or subscribe for, any securities of the Company, nor shall it or any part of it nor the fact of its distribution form the basis of, or be relied on in connection with, any contract or investment decision.
- + The information contained in this Presentation does not purport to be comprehensive. None of the Company, its respective subsidiaries or affiliates, or its or their respective directors, officers, employees, advisers or agents accepts any responsibility or liability whatsoever for, or makes any representation or warranty, express or implied, as to the truth, fullness, accuracy or completeness of the information in this Presentation (or whether any information has been omitted from the Presentation) or any other information relating to the Group, whether written, oral or in a visual or electronic form, and howsoever transmitted or made available or for any loss howsoever arising from any use of this Presentation or its contents or otherwise arising in connection therewith. Each of such persons accordingly disclaims all and any liability whatsoever, whether arising in tort, contract or otherwise in respect of this Presentation or any such information.
- * The information in this Presentation may include forward-looking statements, which are based on current expectations, projections and assumptions about future events. These forward-looking statements as well as those included in any other information discussed in the Presentation are subject to known or unknown risks, uncertainties and assumptions about the Group and its investments, including, among other things, the development of its business, its growth plan, trends in its operating industry, its future capital expenditures and acquisitions. In light of these risks, uncertainties and assumptions, the events in the forward-looking statements may not occur and actual results, performance or achievements may materially differ from any future results, performance or achievements that may be expressed or implied in this Presentation. No representation or warranty is made that any forward-looking statement will come to pass. Forward-looking statements speak as of the date of this Presentation and no one undertakes to publicly update or revise any such forward-looking statement, whether as a result of new information, future events or otherwise. Accordingly, undue reliance should not be placed on any forward-looking statement contained in this Presentation.
- * To the extent available, the industry, market and competitive position data contained in this Presentation come from official or third party sources. Third party industry publications, studies and surveys generally state that the data contained therein have been obtained from sources believed to be reliable, but that there is no guarantee of the accuracy or completeness of such data. While the Company reasonably believes that each of these publications, studies and surveys has been prepared by a reputable source, the Company has not independently verified the data contained therein. In addition, certain of the industry, market and competitive position data contained in this Presentation come from the Company's own internal research and estimates based on the knowledge and experience of the Company's management in the markets in which the Group operates. While the Company reasonably believes that such research and estimates are reasonable and reliable, they, and their underlying methodology and assumptions, have not been verified by any independent source for accuracy or completeness and are subject to change. Accordingly, undue reliance should not be placed on any of the industry, market or competitive position data contained in this Presentation.

 This Presentation also includes certain alternative performance measures ("APMs") that have not been prepared under IFRS-EU and have not been reviewed or audited by the Company's auditors nor by any independent expert. Moreover, the way the Group defines and calculates these measures may differ to the way similar measures are calculated by other companies. Accordingly, they may not be comparable.
- * Certain financial and statistical information contained in this Presentation is subject to rounding adjustments. Accordingly, any discrepancies between the totals and the sums of the amounts listed are due to rounding. Certain financial information and operating data relating to the Company contained in this Presentation has not been audited and in some cases is based on management information and estimates, and is subject to change.
- * No reliance may or should be placed by any person for any purposes whatsoever on this Presentation, or on its completeness, accuracy or fairness. The information in this Presentation is in summary draft form for discussion purposes only. The information and opinions contained in this Presentation are provided as at the date of the Presentation and are subject to verification, correction, completion and change without notice. In giving this Presentation, none of the Company, its subsidiaries or affiliates, or its or their respective directors, officers, employees, advisers or agents, undertakes any obligation to amend, correct or update this Presentation or to provide the recipient with access to any additional information that may arise in connection with it.



OVERVIEW



1,419 Employees as of Dec 2020

c.250Specialty
Pharma Sales
Force

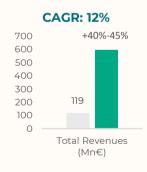
6 Fully Invested Manufacturing Facilities

Мкт Сар (€м)1



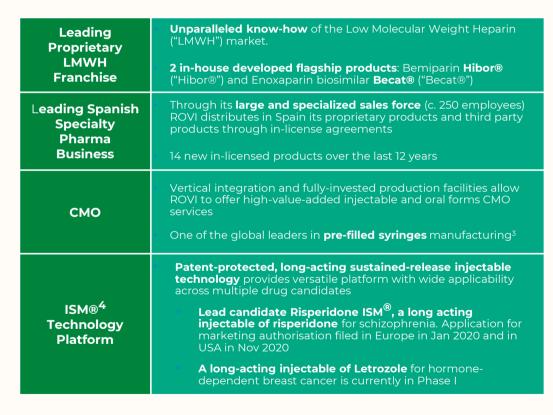
■Dec 2007-IPO ■Current

TOTAL REVENUES (€M)²

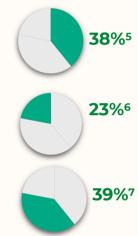


■ Dec 2007-IPO ■2021E

Solid specialty pharma growth story coupled with strong potential from the ISM® Platform



Revenue 9M 2021²



In regulatory process in Europe and United Estates

Currently in Clinical Development Stage

1. CapIQ as of Nov 3, 2021.



- 2. Total revenues include sales from products and services, royalties and government grants.
- 3. In terms of annual number of units manufactured. Offers filling and finishing; does not manufacture the syringe itself.
- 4. ISM® stands for "In-Situ Microimplants" technology.

- 5. Includes revenues from Hibor®and Becat®.
- 6. Includes sales of goods excluding Hibor® and Becat®.
- 7. Includes sales of services.

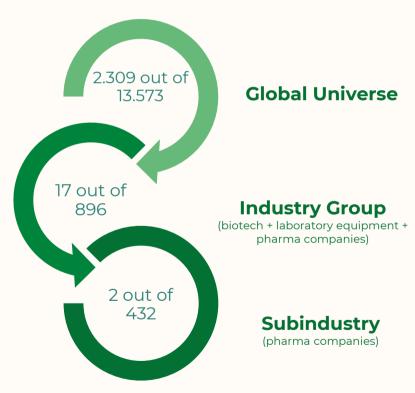
ROVI improves its ESG Rating in 2021

Rovi has obtained an ESG Rating 2021 of 18,4
(low risk between 10 and 20)



2nd position out of 432 companies

(in the sub-industry "pharmaceuticals")





Key Company Highlights



Well-balanced pan-European specialty pharma business with diversified growth drivers

Unparalleled proprietary heparin franchise with strong European footprint

Leading Spanish specialty pharma franchise

High-value-added global CMO business with differentiated capabilities

Proprietary ISM® Platform opens up new avenues of growth

Ownership of technology and vertical integration enhance competitive position

Potential wide applicability of ISM® technology to new chronic therapeutic areas

Sound financial policy supported by strong track record



Well-Balanced European Specialty Pharma Company with Three Diversified Growth Drivers

Leading Proprietary LMWH Franchise

- Developed and successfully launched proprietary LMWH bemiparin, the 2^{nd 1} leading LMWH in Spain
- Developed enoxaparin biosimilar, one of the first to reach the market
- Vertically integrated, well positioned to benefit from significant economies of scale

Leading Spanish Specialty Pharma Franchise

- Strong market leadership in Spain
- Partner of choice for in-licensing for leading global players
- Highly skilled c.250 person sales force

High-Value-Added CMO Services

- One of the global leaders in pre-filled injectables manufacturing
- Fully-invested production facilities
- Help absorb fixed costs and overheads, providing for highly cost-competitive manufacturing position
- Particularly strengthens the LMWH franchise which relies on ROVI's production capabilities

GROWTH DRIVERS

- Continue gaining branded LMWH market share through bemiparin and enoxaparin biosimilar in Spain and abroad
- Launch enoxaparin biosimilar across more European countries and other international markets
- Roll-out of pan-European commercial network
- Second active principle heparin plant in Granada which will allow ROVI to double its capacity

- Leverage on leadership position in Spain
- Maintain strong sales performance and operational excellence
- New in-licensing opportunities with global players in specialty therapeutic areas

- Drive volume growth from existing customers
- Additional CMO customers given strong economies of scale
- Agreement with Moderna



Bemiparin Hibor® is ROVI's First Internally-Developed Flagship Heparin Product

Unparalleled Know-How of the Heparin Market

- ROVI has been engaged in the development of heparin-based drugs for over 70 years
- Bemiparin Hibor® is a **Low Molecular Weight Heparin** (LMWH)

#2 market position in Spain with a c.32% ¹ market share and marketed in 59 countries in total

Only 2nd generation LMWH; clinically differentiated from other competitor (such as Sanofi's Clexane / Lovenox)

• Vertically integrated structure with its own LMWH manufacturing plant

Bemiparin Hibor® Global Sales



Marketed in **59** countries

Pending registration in **13** countries

Bemiparin HIBOR® is the LMWH with the highest anti Xa/IIa ratio, which may lead to a higher antithrombotic activity without increasing the bleeding risk

More convenient treatment: 1 daily injection needed in comparison to Sanofi's (Clexane / Lovenox) treatment (which needs 2) ²

Established international network supported by long-term contracts with leading local pharma distributors

In-house legal team with regulatory knowhow has achieved marketing authorisations worldwide

International Bemiparin sales in ROVI are expected to decrease due to our focus on the enoxaparin biosimilar outside Spain



Enoxaparin €2.6bn Global Market Represents an Untapped Opportunity to Be Explored with ROVI's biosimilar Becat®

European Market Represents an Attractive Opportunity

- Enoxaparin (such as Clexane / Lovenox) is the world leading LMWH
- Europe is the largest Enoxaparin market worldwide (>50%)1



European Uncrowded Competitive Landscape

CTEXANE COLEMAN

Enoxaparin

biosimilar

Becat®

- Originator product developed by Sanofi Aventis
- Patent expired in 2011 (high entry barriers: first biosimilar entered the market 6 years after patent expiry)
- ROVI markets its internally-developed enoxaparin biosimilar
- Launched in Sep'17 with total sales of €101.4Mn in 2020 and €89.9Mn in 9M 2021

In the long term, biosimilars tend to reach a 50%-70% share of the reference product market²

Well-Positioned for Long-Term Leadership in LMWH

- ROVI aims to become one of **Europe's top players in a €1.3bn** market
- ROVI's **competitive advantages** within the LMWH market:





Strong growth potential of Enoxaparin Biosimilar Becat®

Well-Established Network to Minimize Timeto-Market

Direct Marketed in Germany, UK, Italy, Spain, Portugal and **Poland**

Approved in 26 countries in Europe and 24 in the Rest of the World

28 countries

Enoxaparin Biosimilar Becat® Sales Ramp-up



Stage I of Commercial Strategy



...the largest

enoxaparin market²

with **€1.3bn** sales

ROVI will directly market enoxaparin biosimilar Becat® in 7 European countries...











...which account for $c.75\%^{3}$ of the European market

In the long-term. biosimilars tend to reach



...of the reference product market

Launches in 9M 2021

ROVI launched enoxaparin in Canada, Malasya, Albania, North Macedonia, Guatemala, El Salvador, Honduras, Georgia and Bahamas.



















Spanish Market Leadership Positions ROVI as the Partner of Choice for Global Pharma Players in Spain

Our Strong Market Leadership in Spain... ...allows us to be the Partner of Choice for Global Pharma Players in Spain

Presence in the Spanish market since 1946

Well-known proprietary portfolio driving strong leadership position

Franchise focused business: 20 proprietary and 28 in-licensed products

Multiple Strategic Alliances





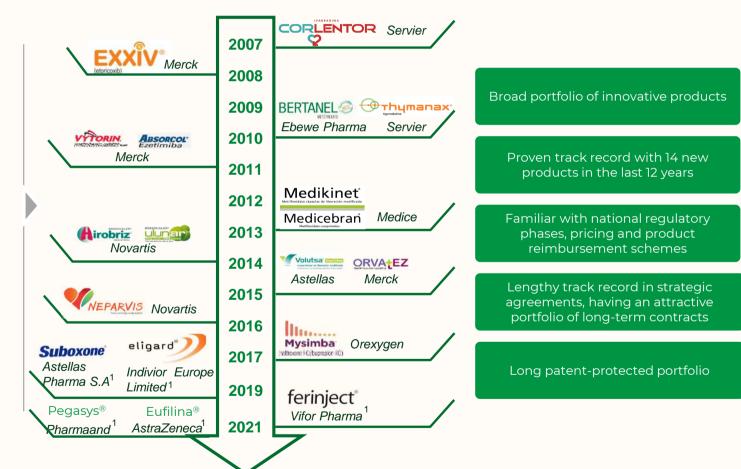






One of the largest specialty pharma sales forces in Spain with c.250 employees

Strong knowledge of the Spanish regulatory framework





High-Value-Added Global CMO Services

Customer-oriented business model

High-value-added service with pre-filled syringes toll manufacturing

Differentiated capabilities drive significant barriers to entry

Revenue visibility on the back of long-term agreements

International sales represent c.97% of CMO business

Clean regulatory track record at manufacturing plants with multiple GMP / FDA approvals

CMO Targets

Toll manufacturing sales increased **39%** in **2020** and **187%** in **9M 2021**

ROVI expects the CMO business to increase by between 2 and 2.5 times, including production of the COVID-19 vaccine.

ROVI aims to become a longterm manufacturing partner for Moderna



ROVI strengthens its collaboration with Moderna (1/2)





Fill-Finish manufacturing

- Investment in 2 new lines for compounding, filling, automatic visual inspection and labeling at ROVI's San Sebastián de los Reyes (Madrid) facility
- These lines will **more than double** the number of vials for which there is fill-finish capacity at this facility
- Supply to markets outside the United States

DARA 2

Expected to come into operation in Q4 2021

DARA 3

• Expected to come into operation between Q2 and Q3 2022



ROVI strengthens its collaboration with Moderna (2/2)



Manufacture of the active substance

- Installation of a new line in Granada
- Production capacity equivalent to more than 100 million doses per year
- Greater vertical integration of the vaccine production process
- Supply to markets outside the United States



ISM® Platform opens up new avenues of growth for ROVI

Overview

- Internally-developed and patented innovative drug-release technology, ISM®, which allows for the sustained release of compounds administered by injection
- Based on two separate syringes respectively containing (a) the drug and polymer (solid state) and (b) the solvent (liquid state)
- Potential wide applicability of ISM® technology to new chronic therapeutic areas, including psychiatry and oncology
- 505(b)(2) path of approval for candidates leveraging ISM® technology

Product	Potential Indication	Current Situation	Key Milestones
Risperidone ISM® Risperidone, monthly	Schizophrenia	Phase III	In approval process in Europe and in USA
Letrozole ISM® Long acting Letrozole	Breast Cancer	Phase I	Phase I started in November 2017
Risperidone, quarterly	Schizophrenia	Non-Clinical	
Concentrated on	improving posology fo	or already approved con	anounds which

Concentrated on improving posology for already approved compounds, which benefits risk / reward profile

Multiple FDA / GMP approved facilities to support the platform

Key Company Highlights of ISM® Platform

1 Predictability	Pop PK ² model & simulations already validated for Risperidone ISM® in Clinical Program	Expected high success rate in Phase III in new developments
2 Usability	Improved stability	No cold chain needed
3 Flexibility	Selecting the most convenient posology depending on clinical needs	From 1 to 6-month administration
Improved Clinical Management	Long-acting injection (1-6 months) plasma therapeutic levels from day 1	Rapid onset & sustained clinical effect
5 Vertical Integration	Technological barriers (e.g. power filling) Strong IP Manufacturing capabilities	Protected technology Fully integrated manufacturing plants



^{1.} ISM® stands for In Situ Microimplants®

DK stands for pharmacokinetic

Risperidone ISM®: Attractive Schizophrenia Market with Strong Growth Prospects

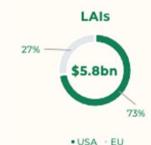
Attractive Schizophrenia Market

- Chronic and progressive disease
- Affects 21m people worldwide with a relatively high lifetime prevalence¹
- Strict compliance needed to avoid relapses
- LAIs² are becoming the gold standard for treatment given improved adherence and effectiveness

MAT Q3-19 Schizophrenia Market Value US & EU³



- Largest schizophrenia LAIs market
- MAT Q3 2015 MAT Q3 2019 CAGR of 20.0%
- Higher prices than other markets
- LAIs penetration: 5.8% (in monthly treatments)4





- Second largest schizophrenia LAIs market
- MAT Q3 2015 MAT Q3 2019 CAGR of **8.5%**
- Relatively low competition due to fewer drug options
- LAIs penetration: 8.4% (in monthly treatments)4

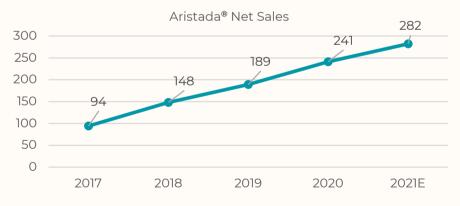
Solid Grounds for Success for a Risperidone LAI

LAI schizophrenia market presents key features for a successful launch High treatment switching rate

Focused group of psychiatrists to target

Increasing penetration of LAIs across treatment paradigm

Ample Market with Room for New Entrants: Alkermes Success Story⁵



Due to current low penetration, schizophrenia LAIs sales are expected to drive future market growth



LAIs stands for Long Acting Injectables. Iqvia Midas MAT Q3 2019.

Risperidone ISM®: Fast Onset Long-Acting Injectable of Risperidone

Superior Value Proposition When Compared to Alternatives

Fully supervised monthly injection

Clinical Convenience of Risperidone

Therapeutic plasma levels from 2 hours post dose aimed at PANSS reduction at day 8

- Ongoing monitoring of nonadherence through regular interactions between patient and medical staff
- Reduce the risk of accidental or deliberate overdose
- Proven efficacy and safety of Risperidone¹
- Well-known drug among psychiatrists for the treatment of schizophrenia
- Fast onset of action to achieve therapeutic plasma levels from the beginning
- Achieving significant PANSS³ reduction in unstable schizophrenia patients at day 8
- No need to supplement with oral medication or loading dose

Strong Product Expected to Offer Superior Characteristics

	RISPERDAL CONSTA® (Risperidone)	INVEGA SUSTENNA®/ XEPLION® (Paliperidone)	INVEGA TRINZA® / TREVICTA® (Paliperidone)	ABILIFY MAINTENA® (Aripiprazole)	ARISTADA® (Aripiprazole Lauroxil)	PERSERIS® (Risperidone Atrigel®)	Risperidone ISM® (Risperidone)
Once Monthly Administration ⁴	×	✓	Quarterly	✓	✓	✓	√ 12-14
No Oral Supplementation / Loading dose ⁴	×	×	After ≥4 months Inv. Sustenna/ Xeplion	×	×	✓	√ 12-14
Therapeutic Levels ² within First 2 Hours ⁴	×	×	×	×	×	✓	√ 12,13
Currently Marketed in Europe ^{5, 6}	✓	✓	✓	✓	×	×	Targeted
Stability at Room Temperature ⁴	×	✓	✓	✓	✓	×	✓
PANSS Reduction from Day 8 ¹⁵	x 7	× 8	× 4	× 9	x 10	x 11	√ 14

7.Kane et al. Am J Psychiatry 2003 8.Pandina et al. J Clin Psycopharmacol 2010. 9.Kane J et al. J Clin Psychiatry 2014. 10.Meltzer H et al. J Clin Psychiatry 2015. 11.Nasser A et al. J Clin Psycopharmacol 2016. 12.Llaudó J et al. Int Clin Psychopharmacol 2016



^{1.} Leucht et al. Am J Psychiatry 201

^{2.}The therapeutic concentration range of risperidone is quite wide and can vary from 10 ng/mL to 80 ng/mL or even higher (Remington et al. Am J Psychiatry 2006).

^{3.}PANSS: positive and negative syndrome scale. Scale used to evaluate the symptoms of patients with schizophrenia. 4.Drugs@FDA:FDA Approved Drug Product. Available at: https://www.accessdata.fda.gov/scripts/cder/daf/index.cfm

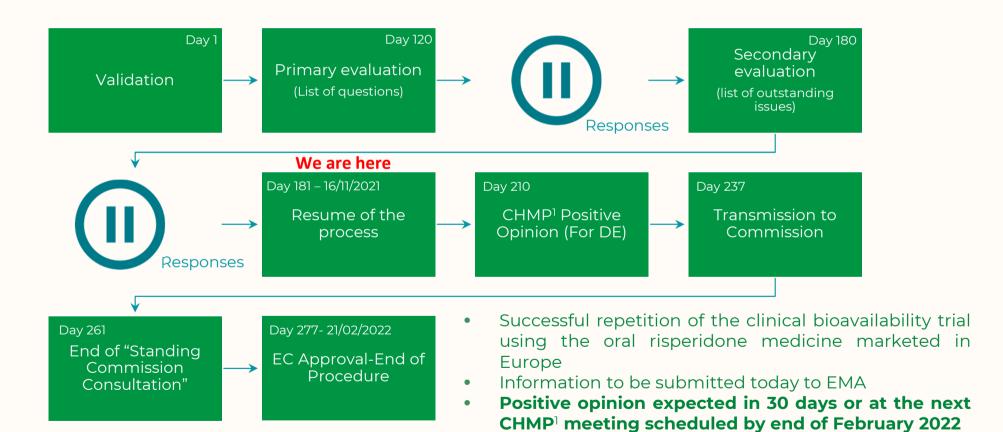
^{5.}Only applies to Risperidal Consta: Heads of Medicines Agencies. MRI Product Index. Available at: http://mri.cts-mrp.eu/Human/

^{6.}European Medicines Agency. European Public Assessment Reports. Available at https://www.ema.europa.eu/en/medicines

European regulatory process - Where are we?



Resume of regulatory process



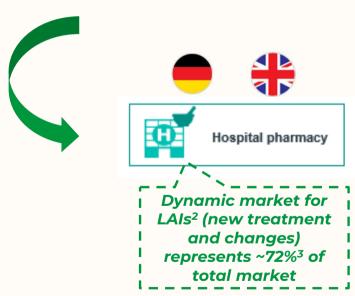
Therapeutic Indication – Covering the Unmet Medical Need

Requested Therapeutic Indication to EMA



Treatment of Schizophrenia in adults

Risperidone ISM® would be the **ONLY¹ Long-Acting Injectable Antipsychotic** that can be administered to **unstable patients** with severe or moderate psychotic symptoms suffering a relapse

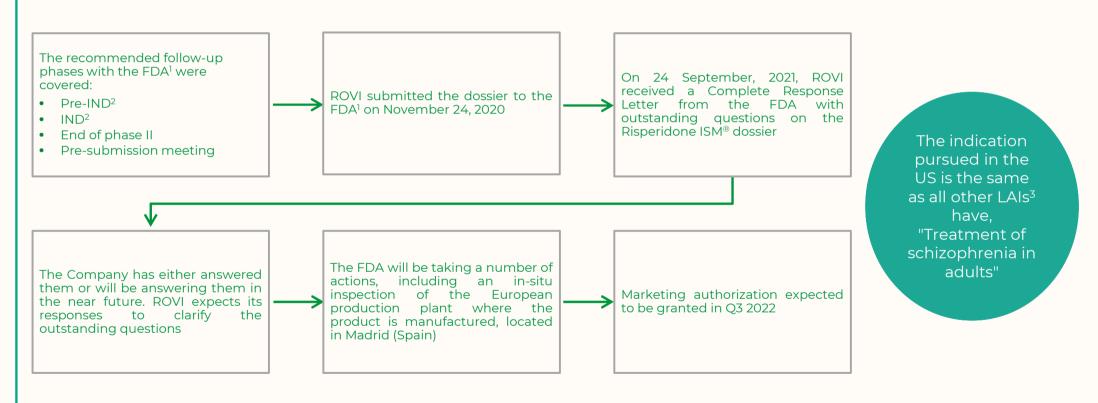


- Dynamic market originating in hospitals
- In Germany and UK, around 72% of prescriptions in hospitals correspond to patients who start treatment with a LAI²
- Competitive advantage because of the potential unique indication of Risperidone ISM®

7 Source: 101/1A

USA regulatory process – Where are we?





Letrozole ISM®: Second ISM® Candidate in Phase I Trial

Overview

Hormone receptor-targeting drugs offer a unique opportunity to leverage ISM® technology. Aromatase Inhibitors (AI) Letrozole and Anastrozole are used in HR+ breast cancer as they block the production of estrogen in post-menopausal women.

- Oral Letrozole is the gold standard treatment for HR+ breast cancer
- Current posology of Als is daily oral potential for Letrozole ISM® targeting a long-acting injection to meaningfully disrupt the market and improve patient outcomes
- Currently, there is no LAI approved for Letrozole in the market

Upcoming Key Catalyst:

- The company has initiated discussions with the FDA to review Letrozole ISM® Phase I results as well as the next steps for continuing its clinical development
- 505(b)(2) path of approval for candidates leveraging ISM® technology

Expected Value Proposition:

Rapid & Sustained Estrogen Suppression Targeting a LONG ACTING Injection

Superior Efficacy

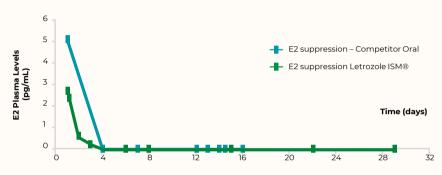
- Preliminary results suggest that sustained long-term hormone suppression therapy (HT) may obtain a superior clinical outcome in breast cancer compared to an oral daily dosage treatment
- Early discontinuation and non-adherence to HT are common and associated with increased mortality – improved treatment compliance with Letrozole ISM® has potential to enhance treatment

Improved Safety Profile

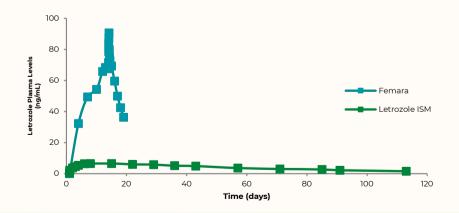
- Sustained lower effective doses (compared to oral treatment) could reduce adverse side effects (bone mass loss, bone/joint/muscle pain, dyslipidemia) due to lower exposure to drug
- Better safety profile has potential to positively impact treatment duration adherence

Preliminary Phase I Results

Rapid and sustained estrogen suppression with lower doses

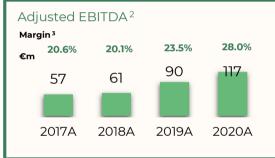


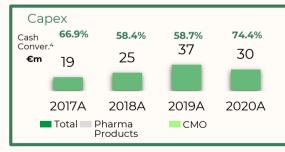
Rapid and sustained Letrozole plasma levels



Sound Financial Policy Supported by Strong Track Record







ISM® Platform

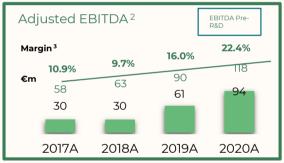






ROVI Group







Proven track record of Specialty Pharma business



^{6.} Calculated as R&D revenues minus R&D expenses, which include Specialty Pharma <u>R&D expenses of enoxaparin</u>

9M 2021 results (1/2)

 REVENUE
 EBITDA
 EBIT

 463.5 (+53%)
 139.5 (+100%)
 123.3 (+123%)

 Net profit
 Capex
 Net cash

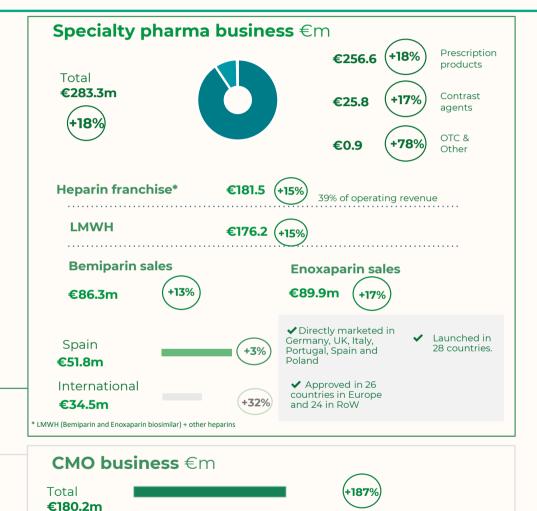
 98.9 (+111%)
 22.0 (30%)
 74.4

2022 operating revenue guidance: ROVI expects a mid-single-digit growth rate for the operating revenue

For 2021, ROVI is upgrading again its 2021 operating revenue guidance from the range between 35% and 40% to the 40% and 45% range.

Operating revenue €m







9M 2021 results (2/2)

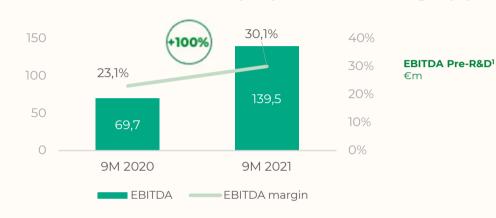
Gross profit (€m) and gross margin (%)

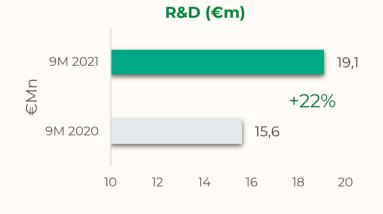


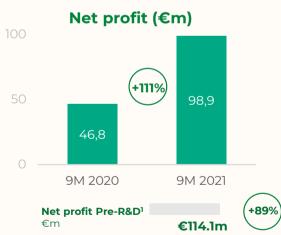
SG&A (€m)



EBITDA (€m) and EBITDA margin (%)







+86%

€158.6m

Outlook 2022



2022 operating revenue growth rate

Mid-single-digit

The key Growth levers in 2022

Specialty Pharma

Bemiparir

Biosimilar of Enoxaparin

Launches such as Neparvis® and Volutsa®

Existing portfolio of specialty pharmaceuticals

СМО

Spare capacity in the manufacturing plants

New customers to be acquired

Agreement with Moderna

Given the uncertainties associated to the development of the Covid-19 pandemic, it is not yet possible to make a precise assessment of the impact that the pandemic will have on 2022.



2023 long term guidance expected to be achieved in 2021

	2018	2021	LT indicative guidance 2023	
Operating revenues	€303.2Mn	€588Mn - €609Mn	X2 €606.4Mn	Expected to be achieved in 2021
EBITDA pre-R&D	€63.0Mn	€158.6Mn in 9M 2021 EBITDA pre-R&D	X2.5 €157.5Mn	Already achieved in 9M 2021

Next phase of growth achieved through several key levers (Enoxaparin biosimilar, Risperidone ISM®, Letrozole-ISM®, M&A activities, Moderna agreement)...

... underpinned by solid specialty pharma and toll manufacturing businesses



For further information, please contact:

Juan López-Belmonte Chairman and Chief Executive Officer +34 91 3756235 www.rovi.es

Javier López-Belmonte Vicepresident and Chief Financial Officer +34 91 3756266 www.rovi.es

Marta Campos Head of Investor Relations +34 91 2444422 mcampos@rovi.es www.rovi.es

Antonio Marquina Investor Relations +34 674 315 715 amarquina@rovi.es www.rovi.es

