Investor Presentation Full Year 2021

Andreas Grassauer, CEO, Pascal Schmidt, CFO Korneuburg, 13th April 2022





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Agenda

Highlights 2021

 Financials 2021
 Marinomed delivered and presents augmented strategy

 Product portfolio
 Investment highlights & outlook

• Financial calendar, contact



Highlights 2021

Delivering on our promises

€11.6m

Record revenues

+43% versus 2020

Budesolv

1st deal

2m USD upfront

Tacrosolv

Clinical data demonstrate fast onset of action Carragelose

Proven neutralization activity against SARS-CoV-2 and VoCs



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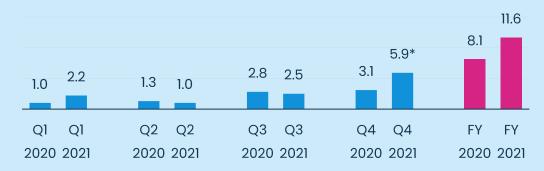
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Continuous double digit growth

Strong Carragelose business with upfront from first Marinosolv-deal

Y-o-Y comparison of quarterly revenues (m€)



Historical performance (revenues and EBIT)



Comments

Key growth drivers

- Sale of goods rose from €7.5m to €9.0m (+20%)
- Other revenues increased from €0.6m to €2.6m (+342%); therein €1.9m milestone from Budesolv licensing in Q4

New business

- New partners in Brazil, Mexico, Finland, Norway, Sweden and France
- Newly established Solv^{4U} unit performs feasibility studies for external partners

Positive outlook

- COVID-19 pandemic continues to fuel order book, challenging the supply chain
- Various measures in progress to compensate COVID-19 related bottlenecks and increased lead times for raw material



Statement of profit or loss (IFRS)

Higher revenues – better margins – more R&D invest – improved EBIT

€m		FY 2021	FY 2020
Revenues	1	11.6	8.1
Other income	2	1.6	1.2
Materials expenses		-6.4	-5.4
Services expenses	3	-3.8	-3.4
Personnel expenses	3	-4.5	-4.1
Depreciation and amortisation	3	-0.6	-0.4
Other expenses		-2.1	-1.8
Operating result		-4.1	-5.8
Financial result	4	-1.5	-0.2
Profit/loss before taxes		-5.7	-6.0
Taxes on income		-0.2	-0.0
Profit/loss for the period		-5.9	-6.0

	FY 2021	FY 2020
Sale of goods	9.0	7.5
Milestones	2.1	0.0
License revenues	0.4	0.4
Other revenues	0.1	0.2
Total revenue	11.6	8.1
Cost of goods sold	-6.1	-5.2
Margin	32.1%	30.3%

2 Primarily consisting of research premium and grant income

3		FY 2021	FY 2020
	Personnel expenses	-2.0	-1.8
	Services expenses	-3.0	-2.7
	Materials expenses	-0.4	-0.2
	Other expenses*	-2.2	-1.2
	Total R&D expenses	-7.5	-5.9

4 Thereof €0.4m interest paid



Statement of financial position (IFRS)

Assets

€m	FY 2021	FY 2020
Assets		
Intangible assets	2.0	2.1
Property, plant and equipment ①	6.4	6.0
Deposits and other non-current receivables	0.0	0.0
Total non-current assets	8.5	8.1
Inventories 2	1.0	0.9
Trade and other receivables 3	6.0	5.3
Cash and cash equivalents 4	5.8	9.2
Total current assets	12.9	15.4
Total assets	21.3	23.5

Includes fully recognized headquarter, incl. land and building (€5.7m) therein directly R&D related* (€0.5m)

2	Inventories€m	FY 2021	FY 2020
	Goods for sale	0.1	0.1
	Raw materials	0.9	0.8
	Total inventories	1.0	0.9

- (3) High balance of trade receivables (€3.4m) due to high number of deliveries in December, also includes part of the upfront payment from Luoxin, received in January as well as receivables from research premium (€0.9m)
- 4 Last tranche from EIB loan (€6m) received in February (after closing date)



Statement of financial position (IFRS)

Equity and liabilities

€m		FY 2021	FY 2020
Equity and liabilities			
Share capital		1.5	1.5
Capital reserves (D	42.1	41.4
Accumulated deficit		-43.4	-37.5
Total capital and reserves		0.2	5.4
Borrowings	2	15.0	12.5
Other non-current liabilities		0.1	0.1
Total non-current liabilities		15.1	12.5
Borrowings	3	0.8	0.4
Trade payables		2.0	2.0
Current contract liabilities and other current liabilities	4)	3.3	2.5
Provisions	5	-	0.8
Total current liabilities		6.0	5.6
Total equity and liabilities		21.3	23.5

- Capital increases through exercise of employee stock options and convertible note funding program
- Primarily related to first and second tranche of EIB loan (€9.0m) and ERP/aws and NÖBEG real estate refinancing (€4.8m)
 - Third tranche of EIB loan (€6.0m) was drawn down in February; additional real estate refinancing (€0.2m) subject to final accounting of investment
- 3 Short term borrowings primarily related to interest on EIB loan, convertible note, which was converted in January, and outstanding loan to WAW
- 4 Current liabilities consist mostly of deferred income from subsidised COVID-19 trial and contract liabilities for clinical studies and employee related provisions
- 5 The use/reversal of the warranty provision is related to the waiver of commercialization rights by a European licensing partner



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Marinomed achievements

Towards higher values

Corporate

2006:

Founding and first financing

2017:

Convertible bond (€7m) at €20m valuation

2019:

IPO and EIB financing (>€37m) at **€75m valuation**

2021:

At year end: €131m valuation

Carragelose virus blocker

2008:

Clinically validated

2010 - 2012:

Further clinical studies, incl. in children (1 year and older)

2020:

Carragelose active against **SARS-CoV-2**

2021:

Portfolio management: adding top tier partners

Marinosolv solubilization technology

2016:

Platform established

2020:

Clinically validated

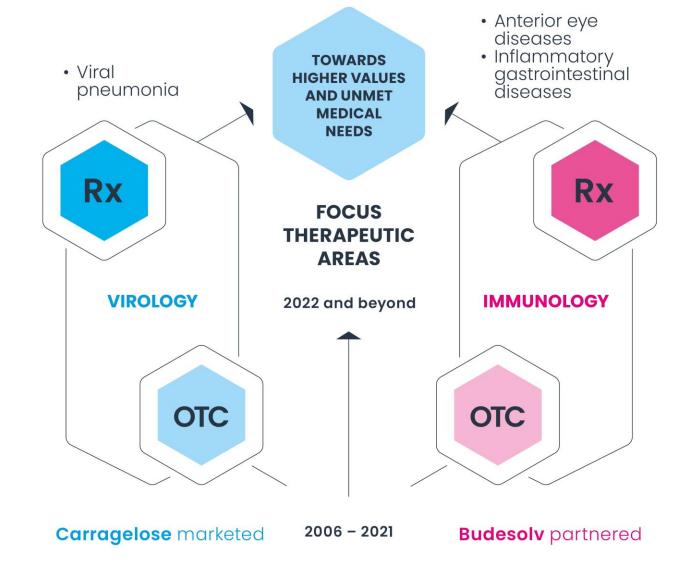
2021:

- Solv⁴⁰ launch with technology partnership projects
- First **licensing deal** for Budesolv

ACHIEVEMENTS 2006 - 2021



Strategy 2025





Strategy 2025

Key initiatives 2022 - more to come

Therapeutic area: Virology

Viral infectious diseases

Besides viral pneumonia, Marinomed has started an additional program that targets a potentially life-threatening disease caused by viruses. Details are currently not disclosed.

Key initiatives

- Establish GMP process for pharma-grade polymers
- Prepare clinical trials
- Secure IP
- Engage in partnerships including co-development

Therapeutic area: Immunology

Autoreactive immune disorders

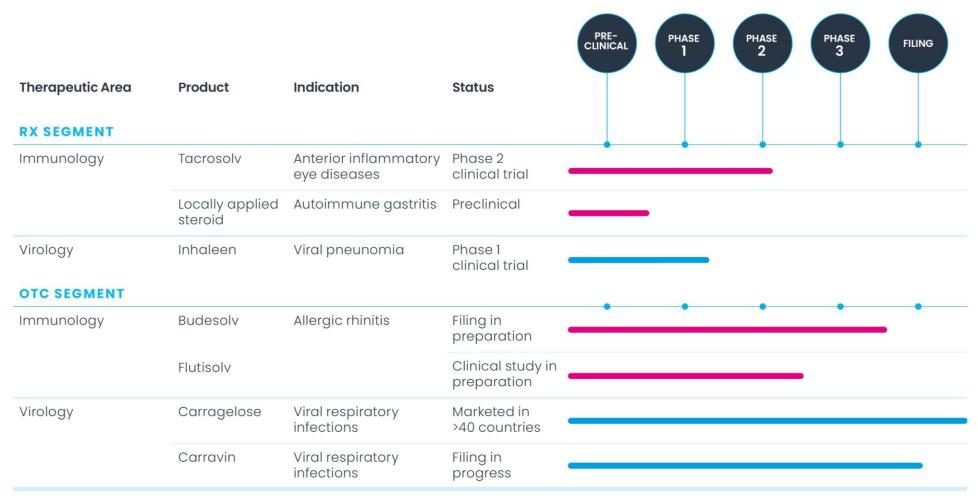
Autoreactive immune disorders are characterized by an exaggerated adaptive or innate immune system. Marinomed actively works on a project targeting an autoreactive immune disorder with a high unmet medical need. Details are currently not disclosed.

Key initiatives

- Engage in partnership for eye delivery and co-development
- Define clinical development plan and prepare initiation of clinical study
- Secure IP
- Board with international KOLs established



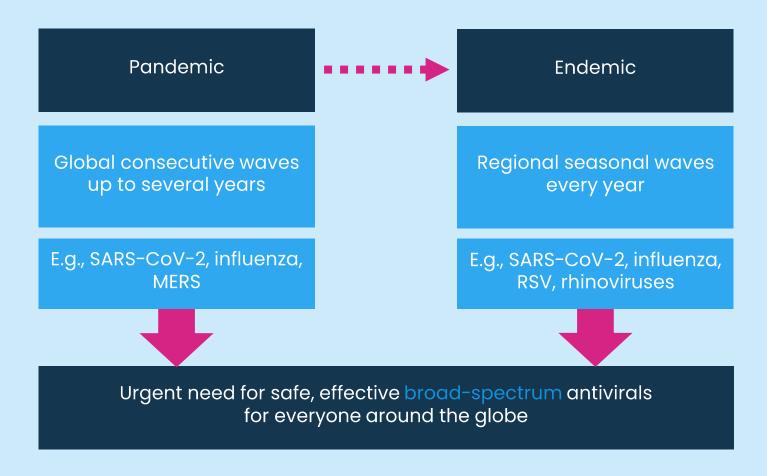
Pipeline overview





INHALeeN – inhalable broad-spectrum antiviral

Addressing the global need for improved treatment options for respiratory infections across virus families





INHALeeN

A potential first-in-class pharmaceutical for prophylaxis and therapy at the onset of outbreaks

Scientific evidence

- Inhibition of replication of SARS-CoV-2 and its VoCs as well as rhinoviruses, influenza, RSV, hMPV, PIV and adenoviruses demonstrated in-vitro
- Clinical study with inhalable formulation in SARS-CoV-2 positive patients ongoing

Strategic impact

- Suitable for strategic stockpiling to allow roll out right-away at outbreak
- Usable in pre/post-exposure prophylaxis and treatment, e.g., in healthcare workers and at-risk population

Focus 2022

- Conclude selection process for API manufacturer
- Engage in partnership for co-development



Budesolv

First deal for Chinese territory – more business development to do

First deal generated first milestone payment

Activities at Chinese partner

- Register ingredients in territory
- Establish production
- Apply for regulatory approval planned for 2024
- Launch planned for 2025

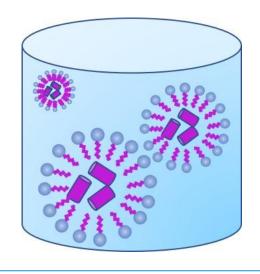
Focus 2022 at Marinomed

 Additional regulatory preparations and partnering activities ongoing for Europe and other territories



Tacrosolv

Best in class macrolide immunosuppressant – fully solubilized with Marinosolv® technology



Solubilized
Tacrolimus
(Liquid formulation)
Micelle enabled
solubilization in
aqueous Marinosolv

formulation

- Tacrolimus is a macrolide calcineurin inhibitor
- Practically insoluble in water
- ~100 times more active than cyclosporine
- No water-soluble formulation available today
- Key ophthalmic indications currently not addressable

- Fully solubilized Tacrolimus in aqueous formulation
- Better bioavailability than suspensions
- Clinical proof of concept established in phase II trial with Tacrosolv
- Patent protection



Tacrosolv – primary indication

Target population & market sizes – offering the potential for the treatment of several inflammatory anterior eye diseases

Tacrosolv – assessed target diseases

Dry eye disease

- Many patients, but only < 1% treated with high price products
- Several treatments available
- Market US-centered

Allergic conjunctivitis

- Disease affects > 15% of population
- Several cheap treatments available

Herpetic stromal keratitis

- · Rare disease
- If untreated may cause blindness
- High recurrency rate
- Support by KOLs

Cornea transplantation

- Rare disease
- Prolonged clinical trials
- May serve as a secondary indication

Recurrent stromal herpes simplex keratitis (HSK) is the main cause of visual loss from HSK

- Current standard of care includes 10 weeks treatment with topical steroids that may lead to AEs like increased intraocular pressure
- Failure in treatment occurs within 6 weeks of completing steroid treatment
- Addition of Tacrolimus on top of SoC indicates significant benefits in visual acuity, corneal vascularization, corneal edema and haziness



Investment Highlights

Target therapeutic areas

Virology

Achievements

- lota-carrageenan is effective against SARS-CoV-2 and all prevalent variants of concern
- Proven track record safe, top tier partners, partnered in more than 40 countries, generating margins

Next steps

- Target more severe indications (i.e., COVID-19) and higher medical need
- Establish polymer as active pharmaceutical ingredient – establish Rx products
- Foster upside potential through additional indication(s)

Immunology

Achievements

- Two important completed clinical trials (Budesolv, Tacrosolv) demonstrating safety and supporting
 - Better bioavailability
 - Significant drug dose reduction
 - Increased efficacy
- Licensing deal for commercial validation of platform

Next steps

- Address autoreactive immune disorders with high unmet medical need
- Partner with pharma companies to create value
- Engage in further partnerships for Budesolv



Strong outlook for 2022 and beyond

Both therapeutic areas offer significant upside potential



R&D

- INHALeeN as first Rx product in viral infectious diseases supported by KOLs
- Tacrosolv confirmed as attractive by KOLs, primary indication defined



Business development

- Carragelose allows for additional partnerships/ product launches and is expected to continue its revenue growth
- Tacrosolv to be partnered for next development steps



Organization

- Moving to Rx as key value driver
- R&D headcount to grow



Financials

- Revenues from sale of products expected to show growth again; revenues from milestones targeted, but subject to licensing
- R&D spending in 2022 slower as clinical studies are supposed to start in 2023
- Further project candidates available but subject to financing evaluating options incl. capital increase



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Financial calendar & IR contact

Financial Calendar

April 13, 2022	Publication of the Annual Report 2021
May 23, 2022	Publication of the Results Q1 2022
June 5, 2022	Record date AGM
June 15, 2022	AGM
August 25, 2022	Publication of the Results H1 2022
November 21, 2022	Publication of the Results Q1-Q3 2022

Stephanie Kniep

Investor Relations phone: +43 2262 90300 226 e-mail: IR@marinomed.com



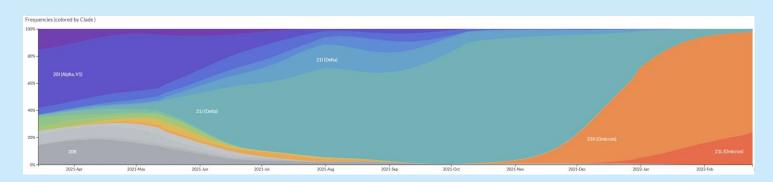


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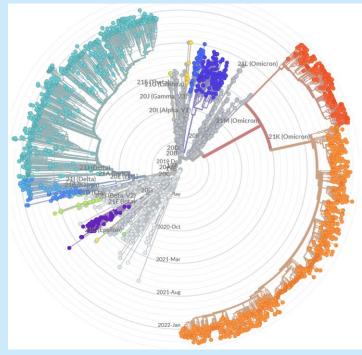
Carragelose

SARS-CoV-2 will evolve and evade immunity



Expected future developments

- 1. Ongoing manifestations of severe disease with high levels of infection
- 2. Transition to an epidemic seasonal disease like influenza
- 3. Transition into an endemic disease similar to other coronavirus infections



CARRAGELOSE will continue to be an option in the fight against COVID-19



Legend: Alpha (201) Delta (211) Delta (213) Omicron (21K) Omicron (21L)