

marino Solving the unsolvable with radical innovation













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Marinomed at a glance

Solving the unsolvable with radical innovation





...with a lean set-up...

generation

In-house Out-sourced Innovation Production • Product Marketing . development Distribution . Marketed products and clinical pipeline technology Patents and IP

Asset light business model

Out-licensing of

...and a strong management team



Reasons to be invested



Transforming treatments with radical innovation

- Carragelose[®] is a radical innovation for the treatment respiratory viral infections
- Carragelose[®] is in the frontline in combating the current pandemic and those to come
- Marinosolv[®] platform solubilization of otherwise insoluble compounds
- Transforming the entire markets with Marinosolv[®] allergic rhinitis market as first target
- Record revenues, solid financials and strong outlook for 2020 and beyond

Carragelose[®] – broad band virus blocker

Carragelose® blocks viral attachment to cells via an unspecific physical mechanism

More than 200 respiratory viruses infect the nasal mucosa

Carragelose[®] directly binds to respiratory viruses and creates a protective layer

Physical mode of action

•

Inhibition of attachment of viruses to mucosa cells → Infection prohibited

 Similar to wool blocking a burdock to hook itself onto textiles, Carragelose[®] binds to the virus and in this

High molecular weight hinders Carragelose[®] from

way blocks the virus from attaching to cells

crossing the nasal mucosa

Carragelose[®] creates a protective physical barrier on the nasal and oral mucosa thus inhibiting the attachment of viruses to cells and the viral replication





Clinical data on Coronaviruses and others



Carragelose[®] is successfully treating virus related symptoms since 2008

Carragelose[®] binds Corona viruses (no data for Sars-CoV-2) Post hoc clinical data from 2 double blind placebo controlled trials

Group	Rhino	Corona ¹	Influenza A
Carragelose®	8.8 ± 0.6	9.0 ± 0.7	8.7 ± 1.0
Placebo	10.7 ± 0.7	12.9 ± 1.0	12.0 ± 1.2
Reduction of duration	-1.9 days*	-3.9 days**	-3.3 days*
			*p<0.05 ** p<0.01

Clinically proven reduction of duration of flu like symptoms – by almost **4 days for Corona** viruses

I. Carragelose[®] neutralizes SARS-CoV-2



New Data published on July 16th, 2020



The neutralizing effect of Carragelose is comparable to serum from a patient who recently recovered from COVID-19.

II. Carragelose^{\mathbb{R}} significantly inhibit SARS-CoV-2 replication in tissue culture

New Data published on August 21st from independent group in Argentina/USA

SARS-CoV-2 virus replication inhibition (Log₁₀ TD₅₀)



SARS-CoV-2 viral titer after treatment with iota carrageenan and P2 (3 replicates per treatment)

Composition

1.2 mg/mL iota-carrageenan, 5 mg/mL sodium chloride, pH 6-7. Vero E6 were pre-treated with dilutions of Iota-Carrageenan with sample P2 (placebo without iota-carrageenan) to get 600 µg/mL, $60 \mu g/mL$, $6 \mu g/mL$, and $0.6 \mu g/mL$ final iotacarrageenan concentration for 2 h. After a 2 h pretreatment, cells were infected with SARS-CoV-2 and incubated for 48h in the presence of the same dilutions of iota-carrageenan. Supernatants were harvested and virus vield determined by an end point dilution assay (TCID50). Controls consisted of untreated infected cells or infected cells treated with P2 (no iota-carrageenan). Results were determined using the Reed and Muench formula and expressed as log TCID₅₀/mL. Dotted line shows the limit of detection (LOD). Testing of samples was performed in triplicate.

Carragelose[®] reduces SARS-CoV-2 virus replication by more than 99,99% or 4,75logs even when the product is diluted by a factor of 1:200

Study of the efficacy and safety of topical Ivermectine and Iota-carrageenan



Positive data on the prophylaxis of COVID-19 for health care personnel

Group	Study participants	Incidence of SARS-CoV-2 (PCR)	Intervention Iota + Ivermectin group
Iota-carrageenan + Ivermectin	131	0	topical medication, iota- carrageenan and Ivermectin, 5 times a day. PPEs used as suggested by OMS.
Standard of care	98	11	Control group PPEs used as suggested by OMS
		p < 0.0001	Clinical trial registration and identifier number NCT04425850

...valuable tool, to **optimize the protection of the people most exposed to contracting the disease,** such as health care personnel

Carragelose[®]: safe and available innovation



Carragelose[®] - a new frontrunner in pandemic countermeasures

- Carragelose[®] products nasal spray, throat spray and lozenges bear a huge potential in the fight against the Coronavirus
- Today, Carragelose[®] products are marketed in more than 40 countries through distribution partners of Marinomed
- Marinomed continues to expand regional coverage through ongoing discussions with existing and new potential partners
- Carragelose[®] products may be one of very few established options when the next pandemic strikes



Marinomed updated pipeline





¹ Carragelose® Xylometazoline combination; ² Dissolved Corticosteroid; ³ Dissolved Tacrolimus

Dynamic addressable Allergic Rhinitis (AR) market

Marinosolv[®] disrupts a multibillion dollar market with strong continued growth prospects



20 Sales 5 sbn 10 5 2019 2020 2021 2022 2023 2024 2025 2026 2027 2028 Global allergic rhinitis market Intranasal corticosteroids market

Attractive market with >5% CAGR²









Intranasal corticosteroids are recommended as first-line therapy for moderate to severe AR

Corticosteroids are projected to overtake antihistamines as the market leading AR category in 2028 with existing marketed products; Budesolv or Flutisolv may even boost the growth to higher %

Onset of action for marketed products takes over 5 days with slow symptom relief



No significant improvement in corticosteroid treatment options in ${\sim}20~{\rm years}$

Marinosolv[®] enabled corticosteroid is Clinically Proven

First Marinosolv®-enabled product achieves superior results in clinical phase III trial

64µg / spray 10µg / spray

Budesolv* – First aqueous steroid solution

Secondary endpoint met

- Goal: Demonstrate a significant reduction of TNSS compared to placebo and the originator on **Day 1**
- Budesolv shows significant reduction of TNSS compared to both other treatments within 3 hours
- Allows for meaningful differentiation and marketing claim

Primary endpoint met

- Goal: Result to demonstrate a reduction of total nasal symptom score (TNSS) score noninferior / equivalent to the originator on **Day 8**
- Result: Both active treatments show equal significant superiority compared to placebo on Day 8
- Allows hybrid application in EU (make use of originator's claims and add additional benefits)

USPs of Budesolv - dissolved corticosteroid

- ~85% reduced dose compared to originator
- First corticosteroid to show activity after a few hours
- Lower cost of production compared to suspensions
- Preservative free
- Patent protected

Marinosolv[®]-enabled corticosteroids change the potential of the compound class





Marinosolv[®] with advanced pipeline



Advanced pipeline of Marinosolv® enabled compounds



Upcoming milestones and value inflection points...



Preliminary Timetable



Double digit growth

Growth path of Carragelose® maintained in 2020







Margin

	H1 2020	H1 2019
Sale of goods	2.0	1.5
Cost of goods sold	(1.5)	(1.0)
Gross result	0.5	0.4
Gross margin	26.4%	28.7%

Comments

- Double digit growth in revenue was achieved from the sale of products in the Carragelose[®] segment
- First revenues from services provided through application of the Marinosolv[®] platform were generated
- Return of certain regions (extra ordinary revenue in Q4 2019) gives opportunity to address underserved markets
- COVID-19 pandemic leads to strong order book
 challenging the supply chain
- Various measures in progress to increase capacities with suppliers and improve speed of the supply chain



Significant investments into R&D



Planned continued losses through high R&D investments are sufficiently funded

R&D Expenses



Comments

- Significant investments in R&D as planned, primarily relating to R&D personnel expenses and external services for clinical trials
- COVID-19 pandemic resulting in short-term delays
 - Allergy challenge trials are currently not possible
 - Planned start for the Tacrosolv phase II not before 2021
- Significant cash position available to advance pipeline products until commercialization
- Carragelose[®] segment margin contributing to financing
- Financial flexibility available through undrawn tranches from EIB loan (€11 million) and expected refinancing of real estate
- Expected to be funded until break-even

Supply chain improvements

Strong focus on eliminating bottlenecks in supply

Two main drivers for supply chain improvements:

1. Close collaboration with manufacturing organizations

- Improvement of long-term planning to reserve capacity
- Become preferred partner to receive production slots that become available
- · Get multiple suppliers approved where possible

2. Eliminate key bottlenecks in production

- Build up stock of primary packaging with long lead times
- Source primary packaging on own account (see inventory development) and with a long-term view
- Increase flexibility through management of stocks between primary packaging supplier and manufacturing organizations



Inventory



Positive Outlook 2020



Investments in Marinosolv® and commercialisation of Carragelose®

Further strong demand for Carragelose® products expected

- Lab data and clinical data for Sars-CoV-2 are growth drivers
- Full focus on conversion of data into marketing & sales
- Expansion of capacities continues
- Ongoing discussions with renowned new partners

Marinosolv[®] products drive value further

- First deal for Budesolv expected in H2/2020
- New applications for nose, lung, eyes and gastrointestinal have huge potential for further growth



Stay Healthy!

...and further reduce the risk by following these rules







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