Investor Presentation

November 2022





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Marinomed investment highlights

Solid existing business, broad late-stage pipeline

- Highly experienced management team
- Solid growth perspective with existing product portfolio
- Lean and efficient business model with cost efficient approach along entire value chain
- Focus on of early clinical data to facilitate partnering with **milestones and royalties**
- Long term growth perspective with Marinosolv products and SOLV4U business unit



Marinomed management and ownership



Total number of shares 1,506,162

Acropora Beteiligungs GmbH

Dr. Hermann Unger

DI Dr. Andreas Grassauer (CEO)

Mag. Dr. Eva Prieschl-Grassauer (CSO)

Free Float

Prime Market of the Vienna Stock exchange MARI:AV; ATMARINOMED6; MARI.VI - part of ATPX, AP8, AXGP, NAP8, TAP8, WBI

Team with long industry expertise and background

Left to right: Pascal Schmidt, CFO; Eva Prieschl-Grassauer, CSO and co-founder; Andreas Grassauer, CEO and co-founder





Marinomed has built a proven track record

Successes in the therapeutic areas

Virology

- Carragelose effective against respiratory viruses
- Series of **clinical studies** supports efficacy
- Generating sustainable revenues and contribution margins
- Patent protected
- Marketed via partners in >40 countries





Perrigo



South East Asia

Immunology

• First **aqueous steroid solution**, all other major competitors are suspensions



- Dose reduced by >85% compared to originator
- Primary and secondary endpoint met in **successful** clinical Phase III
- Patent protected
- Partnered





The US market Cough, Cold & Allergy (CCA)

With \$7.9bn the North American Cough, Cold & Allergy market is almost double the size of Western Europe (\$4.3bn)



CHC market by category 2021

Leading CCA brands 2021 (global sales)

	Brand	Owner	\$mn	21/20
1	Vicks	P&G	1418	+12%
2	Halls	Mondelez	724	+2%
3	Mucinex	Reckitt	650	+13%
4	Claritin	Bayer	615	+7%
5	Zyrtex	J&J, GSK)	584	+7%

With P&G we target a \$7.9bn market - and P&G leads with the Vicks brand

Marinomed Strategy

Building on solid existing business, entering new markets

	2022+	2023/24	2025+
	MaximizingGeographical expansion	Expand in OTCGeographical expansion	 Expand in Rx Progressing technology partnerships
Future revenue			Tacrosolv Inflammatory eye diseases
i uture revenue			Inhaleen Viral pneumonia
Milestones	Carravin	Budesolv Allergic rhinitis	
	Viral respiratory infections		S
Recurring revenue	Carragelose Viral respiratory infections		Technology partnerships
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Tacrosolv in inflammatory eye diseases

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



Tacrolimus, (FK506)

- Tacrolimus is a macrolide calcineurin inhibitor
- Practically insoluble in water
- ~100 times more active than cyclosporine

Tacrosolv

- Fully solubilized Tacrolimus
- Better bioavailability than suspensions
- Clinical proof of concept established in phase II trial

Tacrosolv – best in class calcineurin inhibitor – fully solubilized for ophthalmic indications

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Tacrosolv validated in clinical phase II trial¹

Anti-inflammatory activity in model indication allergic conjunctivitis shown¹



SIGNIFICANT REDUCTION³

- of ocular symptoms on day 8 of treatment compared to day 1
- of ocular symptoms after one week of treatment at 3.5 hours after challenge begin
- of nasal symptoms on day 8 of treatment



*TOSS: total ocular symptom score: itchy eyes, watery eyes, redness of eyes, gritty feeling
1) Data on file from phase II clinical trial sponsored by Marinomed
2) Baseline corrected
3) In higher dose group

Herpetic stromal keratitis

Potentially blinding condition characterized by recurrent infections of the cornea

HERPETIC STROMAL KERATITIS (HSK)

- Pathology caused by
 - a local viral cytopathic effect induced by herpetic viruses (mainly Herpes simplex) AND
 - host immunological and reparative response in the stroma of the cornea¹
- Much of morbidity due to CD4+ T-cell destruction in inflammatory response to the virus²
- Shows highest impact on vision loss compared to other keratitis subtypes³







Neovascularization due to HSK Most common infectious cause of unilateral blindness and vision impairment in industrialized world¹



Sibley D, Larkin DFP; Update on Herpes simplex keratitis management; Eye (2020) 34:2219-2226
 Farooq A et al; Herpes simplex epithelial and stromal keratitis: an epidemiological update; Surv Ophthalmol (2021), 57(5):448-462
 Koganti et al., Pathobiology and treatment of viral keratitis; Exp Eye Res (2021); 205:108483

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Reference Study¹: add-on therapy with tacrolimus in HSK

Primary and key secondary endpoints were met

Treatment plan		CORNEAL HAZINESS	CORNEALEDEMA
<u>All patients</u>		2.00- SE 2.00-	3.00- 2.50- 9 2.00- 9 2.00-
Oral antiviral treatment 400 mg Acylovir tablets • twice daily for interstitial keratitis • five times daily if accompanied by iridocyclitis		Correct haziness	o Correct defenses -00.1 e -00.1 e
<u>Control group</u>	<u>Tacrolimus group</u>	00- BEPORE 30 1W 2W 3W 4W TIME	00- BEFORE 3D 1W 2W 3W 4W TIME
	 1% prednisolone eye drops every two hours with a two- 		eal haziness and edema scores at baseline, tervention in the case and the control group:
1% prednisolone eye drops every two hours with a two-	hour dose reduction every week	Significant improvements in (Primary endpoints)	case group vs control: - Standar
hour dose reduction every week	• 0.5 mg/ml Prograf additionally 4 x daily (1 drop)	Corneal Haziness P =	0.001 after 1 week 5tandar 0.001 after 1 week 0.05 % to
	(diluted with balanced salt solution)	Other significant improveme (Secondary endpoints)	nts in case group vs control:

(Secondary endpoints) Visual Acuity P < 0.001 after 2 weeks Corneal Vascularization P < 0.01 after 2 weeks

Standard treatment

Standard treatment +

0.05 % topical tacrolimus

Tacrosolv Development Plan 2022/23

Estimated Timelines

2022/23

Apply for Orphan Drug Designation

- Start preparing the application process at EMA with a full service CRO
- Will allow access to benefits, e.g., scientific advice and protocol assistance regarding the overall development plan, fee & tax reductions

2023

Orphan Drug Designation

 Start application process for an EMA protocol assistance with a full service CRO once Orphan Drug Status is granted

2022/23

Pharmaceutical development

- Preparation for clinical study
- Manufacturing of study medication

2023

Preclinical development

• 14-days local tolerance

Initiate Phase II study in HSK patients

Solv4U: Solution for drugs with low solubility

Development for external customers- potential upside w/o own risk



Initial revenues followed by a licence model with milestones and royalties



Strong pipeline – still more to come



Continuous double digit growth

Strong Carragelose business and upfront from first Marinosolv-deal

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



Historical quarterly performance (m€)



Comments Key growth drivers

- Revenues rose from €3.2m to €4.9m (+52%)
- Solv4U generated other revenues

New business

- Intense regulatory work with new partners (Korea and USA)
- Solv4U unit moving into second stage after successful feasibility studies with compounds from external partners

Positive outlook

- "Old" viruses in combination with new variants of SARS-CoV-2 keep demand high and challenge the supply chain
- Various measures in progress to compensate COVID-19 and war related bottlenecks and increased lead times for raw material

Comfortable cash reach

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Low cash drain due to profitable revenues and inflows from convertible bond program



- Ended December 2021 with €5.8m in cash
- Raised +€7.3m in net cash through EIB (€6.0m), NÖBEG (€0.2m) and CNFP¹ (€1.2m)
- Received **+€0.7m** in milestones
- Earned **+€1.9m** net cash through profitable sale of goods
- Spent **-€4.7m** in operations, mainly R&D and personnel
- Results in **€11.0m** cash position
 - → Low cash burn in the first half year 2022

This keeps us busy

Various projects require significant attention, but are rewarding

Co-development and definition of regulatory path	Execution with Luoxin for the Chinese territory	 Delays through strict lock-downs in Shanghai Very constructive collaboration
Regulatory work to achieve market authorization for Carragelose products	with the following partners: • P&G for USA • Hanmi for South Korea • M8 (former Moksha8) for Brazil and Mexico	 Good progress with all new partners Heavy support by Marinomed with documentation and data First launches in 2023/2024 season
Transition from MDD to MDR	With legal manufacturer and CMOs	 Intense work to compile significantly increased documentation requirements including new studies
Remain capable to fulfill customer demand	 Own supply chain activities and suppliers 	 Packaging material and stock keeping optimization
Ongoing R&D efforts	Currently mostly internal R&D teams	 Read-out for completed studies and preparation for next phase and clinical trials
Solv ⁴⁰	Various customers	 Moving into follow-on projects

Outlook and timeline 2022/2023

News flow / milestones

News flow to date	 February – Draw down of €6m EIB financing tranche April – New agreement with Hanmi for marketing of Carragelose in South Korea April – Publication of strong 2021 financials May – Procter & Gamble (P&G) for Carragelose products in the US June – New Supervisory Board Members Elisabeth Lackner and Ulrich Kinzel Aug – H1/22 Record sales +52% revenue growth 		
Milestones for 2022 and beyond	 Short term (up to 9 months) Carragelose – regulatory progress Additional Solv⁴⁰ technology collaborations 	 Mid term (6 – 12 months) Second Budesolv deal Carragelose launches in new territories New technology-based product candidates 	 Longer term (9 – 18+ months) Orphan drug designation EMA + FDA New technology partnerships Accelerated revenue growth

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Aim to build second pillar growth with entering new market segments



www.marinomed.com





Sustainability

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Marinomed contributes to sustainability goals and adheres to ESG



Autoimmune gastritis (AIG) Chronic inflammatory disorder of stomach lining without specific antiinflammatory treatment available

What is AIG?	Standard of care	Prevalence	
 Inflammation of gastric mucosa due to an autoimmune reaction 	 No effective anti-inflammatory treatment available 	 Highly variable due to different use of diagnostic tools 	
 Destruction of parietal cells and subsequently increased pH 	 oral / systemic iron and vitamin B12 supplementation¹ 	 Between 2–5% of the total population² 	
 Iron and vitamin B12 deficiency lead to anaemia and pernicious anemia 			
 Increased risk for gastric tumors¹ 			
	Introduction of precision medicine in AIG	•	
	Unspecific symptoms require identification of relevant biomarkers to define patient population that would		
	 benefit most from treatment, and 		
	 increase probability of success of clinical development 		



Budesolv

A new fast acting, low dose nasal therapy for allergic rhinits

• First aqueous steroid solution, all other major competitors are suspensions



- Dose reduced by >85% compared to originator
- Primary and secondary endpoint met in successful clinical Phase III
- Patent protected
- 1st partnership in place with



With Budesolv we target the multibillion \$ allergic rhinitis market – next step: regulatory advancement and partnering for Europe and US



Cough and cold is back

Strong season 2022/23 expected – and has already started



