

Marinomed Biotech AG Corporate Presentation

October 2024



Disclaimer

This presentation (the “Presentation”) was prepared by Marinomed Biotech AG.

The information contained in this Presentation has not been independently verified and no representation or warranty expressed or implied is made as to, and no reliance should be placed on, the fairness, accuracy, completeness or correctness of this information or opinions contained herein. Neither Marinomed Biotech AG nor any officer or employee of Marinomed Biotech AG or any person connected with them accepts any liability whatsoever for any direct, indirect or consequential damages or losses arising from any use of this Presentation or its contents or otherwise arising in connection therewith. Marinomed Biotech AG undertakes no obligation to update or correct any information contained herein or to otherwise advise as to any future changes to it.

Certain statements contained in this document may be statements of future expectations and other forward looking statements that are based on management’s current view and assumptions and involve known and unknown risks and uncertainties that could cause actual results, performance or events to differ materially from those expressed or implied in such statements.

Certain figures in this presentation have been rounded in accordance with commercial principles and practice. Such figures that have been rounded in various tables may not necessarily add up to the exact total given in the respective table.

This Presentation does not constitute or contain any investment advice. It is not and shall not be construed as an offer, invitation, recommendation or solicitation to sell, issue, purchase or subscribe for any securities in any jurisdiction or to enter into any transaction.

By accessing this Presentation, you represent, warrant and undertake that you have read and agree to comply with and to be bound by the contents of this disclaimer.

Logos and images of medical products in this presentation are for illustration purposes only and do not constitute advertising. Note: Information on effects and possible undesirable side effects is provided by the directions for use, doctor or pharmacist.

Company

Equity Story, Overview, Business Model



Equity Story

Solid existing business, growth perspective through late-stage pipeline assets



Experienced management team



Recurring revenue from marketed products



Late-stage assets in partnering



Lean and efficient business model



Restructuring ongoing – **long-term growth perspective**



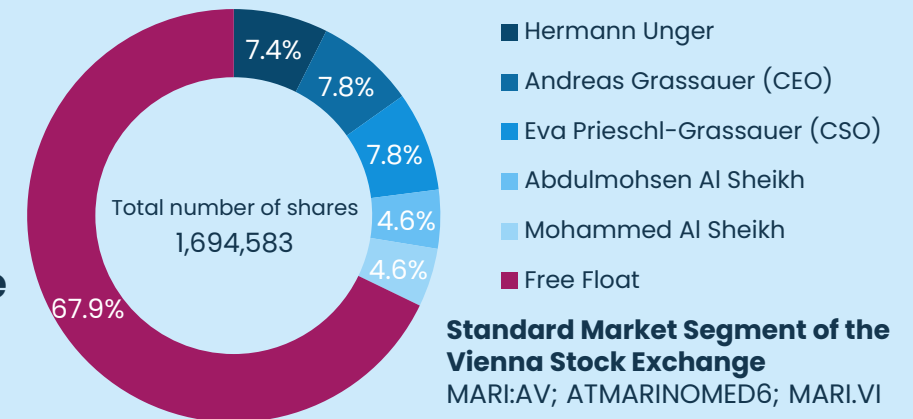
Andreas Grassauer
Chief Executive Officer



Eva Prieschl-Grassauer
Chief Scientific Officer



Pascal Schmidt
Chief Financial Officer

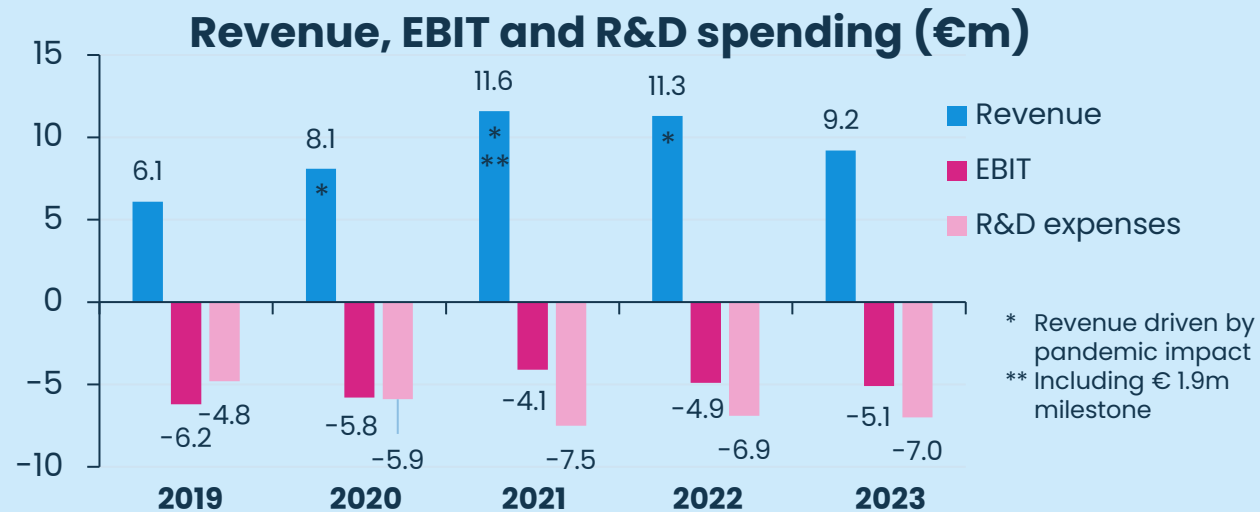


Note: Rounding differences possible



Marinomed at a glance

Publicly listed biopharmaceutical company located in Korneuburg, Austria



VIROLOGY
Revenue-generating
OTC portfolio

IMMUNOLOGY
High-value products in
late-stage development

SOLV4U
Solubilization technology
partnerships

Business Model

	Development phase	OTC (over-the-counter)	Rx (prescription)
 Generation of intellectual property	Idea & preclinical research		
	Early clinical development		
	Late clinical development		 
	Market authorization		
 Commercialization through partners	Manufacturing		
	Distribution & marketing		
	Vigilance		

 Marinomed

 Partner

**Revenue through license deals
(upfront, milestones & royalties)**
Sale of goods
Sale of assets

Technologies & Pipeline



Technologies & Therapeutic Areas

Carragelose®

Universal **blocking of viruses and allergens** as well as moistening of mucosal tissues

Cough & cold portfolio*

Viral respiratory infections



Allergy nasal spray*

Mild allergic rhinitis



Eye drops

Dry, irritated eyes



Marinosolv®

Solubilization of poorly water-soluble compounds and improving **local onset of action**

Budesolv

Allergic rhinitis



Tacrosolv

Inflammatory eye diseases



Solv4U Technology partnerships

Solv4U

VIROLOGY



IMMUNOLOGY



SOLV4U



Pipeline

Including late-stage projects with low risk and meaningful upside potential

Pharmaceutical Products

Therapeutic Area	Product Indication	Status	Pre-clinical	Phase I	Phase II	Phase III	Filing	
IMMUNOLOGY	MAM-1004-1/Budesolv Treatment of severe allergic rhinitis	Filing in preparation						
	MAM-1003-1/Tacrosolv Severe inflammatory eye diseases	Phase II clinical study						
VIROLOGY	MAM-2001-1/Carravin Nasal congestion	Partnering in progress						

OTC Medical Devices

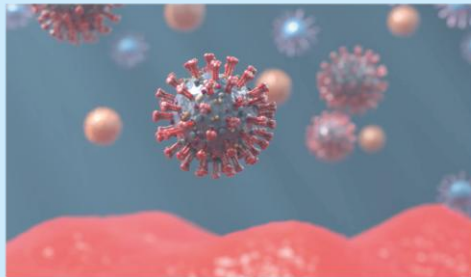
Therapeutic Area	Product Indication	Status	Pre-clinical	Clinical studies	Certification
IMMUNOLOGY	MAM-1001-4 nasal spray Prophylaxis of mild allergic rhinitis	First launch			
	MAM-1001-3 eye drops Dry, irritated eyes	Pre-launch			
VIROLOGY	MAM-1001-1/Inhaleen Viral pneumonia	Clinical studies			



Carragelose®

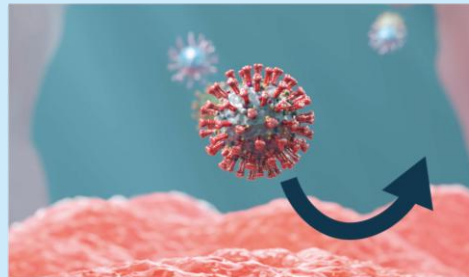
Universal blocking of viruses and allergens as well as moistening of mucosal tissues

Carragelose®



Without Carragelose:
Virus / allergens interact with mucosal cell

Carragelose®



With Carragelose:
Physical barrier prevents interactions of viruses and allergens with mucosa

- Polymer extracted from **red seaweed**
- Forms non-specific layer that **protects mucosa** from viruses & allergens → multi-purpose
- Favorable **safety profile**
- **Clinically validated¹ & patent protected**
- **Marketed** product portfolio

Assets

- **Virus-blocking OTC product portfolio**
 - Partnered in >40 countries
- **MAM-1001-4 nasal spray**
 - First launch in Austria (April 2024)
- **MAM-1001-3 eye drops**
 - Launch planned in 2024
- **MAM-2001-1/Carravin**
 - Ready for partnering
- **MAM-1001-1/Inhaleen**
 - Initial clinical studies performed



Carragelose® OTC portfolio

Revenue-generating product portfolio targeting cough/cold, hayfever, and dry eyes

Extensive clinical data¹

- Forms non-specific, moisturizing layer that protects mucosa from viruses & allergens and provides relief for dry eyes → targeting several indications
- **Broadly-active virus-blocking effectiveness** of Carragelose demonstrated in several clinical trials, including **>1,000 participants**
- Protective layer may not only prevent an infection but also inhibits viral spread → **prophylactic** and **therapeutic** effect
- Favorable safety profile

Partnering

- Nasal sprays, throat spray, lozenges & eye drops
- Partnered in >40 countries, including major players such as P&G in the U.S. or Reckitt in UK
- Global patent protection

Revenues (€m)



Universal blocking of viruses and allergens as well as moistening of mucosal tissues

Recent developments

- Received **MDR certificates** for first Carragelose products
- Completed **clinical study** for moisturizing eye drops & published first results
- Expanded partnership with existing partner and leading consumer healthcare player for **Europe** and other **selected countries**
- New distribution partnerships for **Gulf region** and **Eastern Europe**
- **Launch** of virus-blocking nasal spray with partner M8 in Mexico
- **Launch** of allergen-blocking nasal spray in Austria
- Evaluation of **strategic options** for whole Carragelose business

Next steps

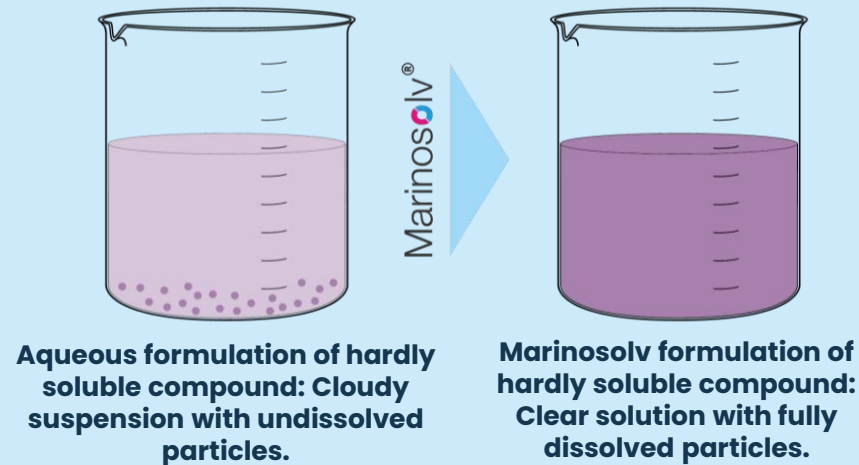
- **P&G:** supporting partner in FDA-process; preparation of possible launch
- Progress new partnership with **leading consumer healthcare player** in Europe and support market access **in Southeast Asia** with Favorex/DKSH or **Eastern Europe** with VitaPlus
- Evaluation of clinical study results and **launch** of moisturizing Carragelose **eye drops** in Austria
- **Drive BD processes** for moisturizing eye drops and allergen-blocking nasal spray
- **Carragelose business:** Start of contract negotiations for the **sale of the whole Carragelose portfolio** in form of an asset deal



Marinosolv

Marinosolv®

Solubilization of poorly water-soluble compounds - improving local onset of action



- Significantly increases **bioavailability** in target tissue
- Allows **dose reduction & faster onset of action**
- **Clinically validated^{1,2} & patent protected**
- Suitable for **sensitive tissues** such as eyes and nose
- Reduces pharmaceutical compounds in wastewater
- Applicable to a wide range of compounds

Assets

- **MAM-1004-1/Budesolv:**
 - Solubilized budesonide for treating allergic rhinitis
 - Post-phase-III
- **MAM-1003-1/Tacrosolv:**
 - Solubilized tacrolimus for treating inflammatory eye diseases
 - Post-phase-II
- **Solv4U**
 - Marinosolv solubilization technology partnerships for external customers
 - Several successful projects & three long-term partnerships established

Budesolv: anti-allergic product post phase III

The first low-dose, preservative-free nasal steroid with strong symptom relief after first dose

Budesolv → fully solubilized Budesonide (corticosteroid) targeting allergic rhinitis

- ✓ **Onset of action in hours:** First-in-class fast acting corticosteroid nasal spray
- ✓ **High bioavailability** in nasal mucosa with low systemic concentration
- ✓ **Lower dose:** Less side effects, less wastewater contamination
- ✓ **Preservative-free:** Improved local tolerability, higher patient acceptance
- ✓ **Clinical validation:** successful phase III study¹
- ✓ **Patent-protected until 2043:** Innovation in allergic rhinitis

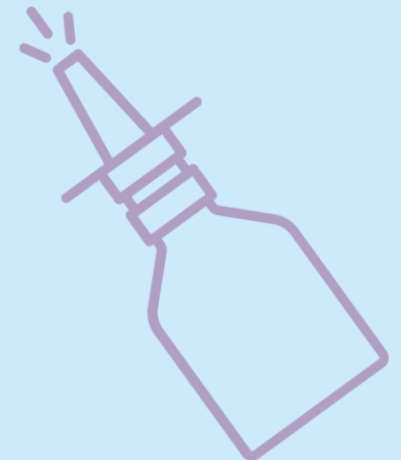
Recent developments:

- Innovation in formulation & primary packaging → **improved stability**
- **Partnership with Luoxin:** progressing towards next milestone

Next steps:

- **Add further partnerships in 2024**
- **Complete development towards marketing authorization with partners**
- **Support partner Luoxin for China**

Allergic rhinitis market
expected to
grow from 16.1 to 22.4 bn US\$
by 2031
(CAGR 4.8%)²



¹ Zieglmayer P, Schmutz R, Lemell P, et al. Fast effectiveness of a solubilized low-dose budesonide nasal spray in allergic rhinitis. Clin Exp Allergy. 2020; 50: 1065–1077. <https://doi.org/10.1111/cea.13691>

² Coherent Market Insights: Allergic Rhinitis Treatment Market Analysis, 06/2024

Tacrosolv: targeting ocular inflammation

Novel low-dose, preservative-free eye drops post-phase II

Tacrosolv → fully solubilized Tacrolimus (immunosuppressor) targeting anterior inflammatory eye diseases and beyond

- ✓ **Increased bioavailability** in cornea and conjunctiva
- ✓ **Lower dose** compared to marketed product in Korea and Japan
- ✓ **Preservative-free**
- ✓ **Clinical proof of concept** established in phase II trial: Significant reduction² of ocular symptoms compared to placebo in model indication allergic conjunctivitis

Recent developments:

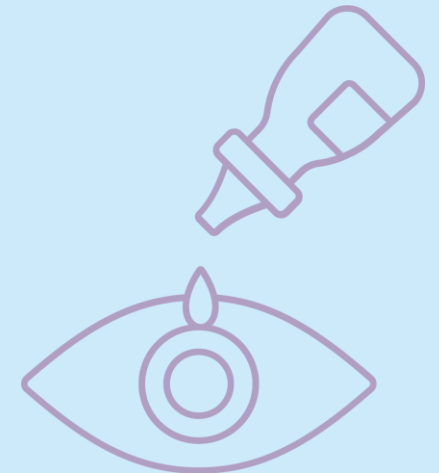
- **Improved formulation & primary packaging**
- Published **results from clinical phase II** study in peer-reviewed journal *Clinical Ophthalmology*¹

Next steps:

- **Late-stage clinical development** together with partner – enable near-term partnership

Attractive ophthalmic market

- Dry eye disease: US\$ 13bn (2032) (CAGR: 7.1%)³
- Allergic conjunctivitis: US\$ 5bn (2032) (CAGR: 5.6%)⁴



¹ Sladek, S. et al., *Clin Ophthalmol.* 2024;18:2797–2811; doi: <https://doi.org/10.2147/OPTH.S476163> <https://www.dovepress.com/alleviation-of-allergic-rhinoconjunctivitis-symptoms-in-participants-t-peer-reviewed-fulltext-article-OPTH>

² In higher dose group

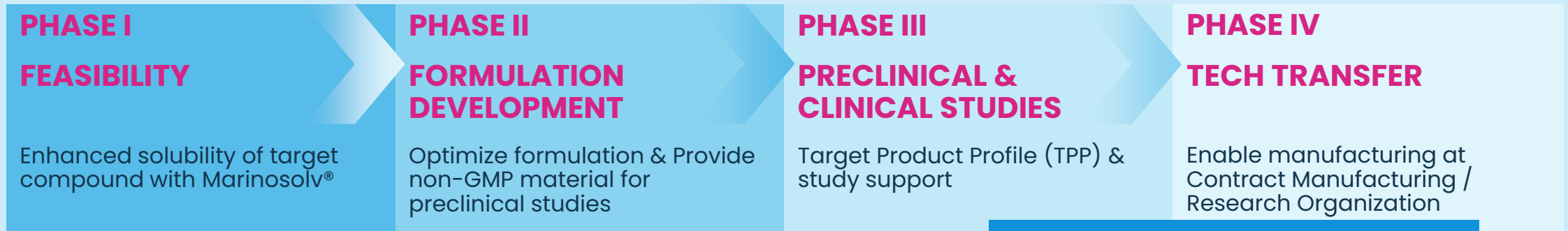
³ fortunebusinessinsights.com: Dry eye syndrome market size, share & industry analysis, 06/2024

⁴ fortunebusinessinsights.com: Allergic conjunctivitis market size, share & industry analysis, 06/2024

Solv4U overview



Solubilization technology partnerships based on Marinosolv®



LICENSE MODEL

Milestones and royalties through clinical development phase and beyond

Recent developments

- **Long-term partnerships** with **SPH Sine** for China, **Aché Laboratórios** for Latin America and **Unither Pharmaceuticals** for France
- Similar deals on the horizon

Next steps

- Drive **long-term partnerships**
- Add additional partnerships



Financials

*H1 2024**

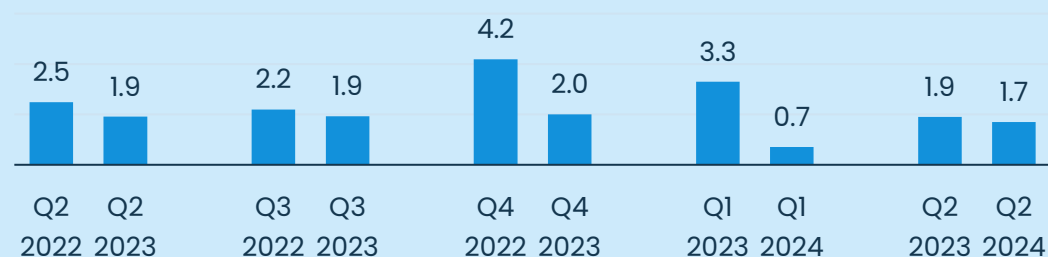
**Selected, preliminary H1 figures – Publication of the half-year financial report 2024 planned after the three-month restructuring phase*



H1 2024: Preliminary figures*

All amounts in € thousand	H1 2024*	H1 2023
Revenues	2,463.9	5,228.5
Operating result (EBIT)	-3,225.3	-2,906.4
R&D expenses	-3,137.1	-3,712.4
Cash flow utilized by operating activities	-1,890.0	-2,275.4
	30.06.2024	31.12.2023
Cash and cash equivalents	911.5	2,588.8
Balance sheet total	11,323.0	14,611.7

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



Revenues

- Revenues for Q2 2024 at € 1.7m (Q1 2024: € 0.7m)
- Revenues include milestone from new European contract
- Order intake currently on pre-pandemic levels, but reviving
- Outlook
 - Robust pharmacy sales reduce customers' stocks and increase likelihood of new orders for the next season
 - Carragelose-based allergy spray launched, eye drops following shortly – business development ongoing
 - Ongoing registration processes with P&G, DKSH, VitaPlus and GAIA with revenue potential in the 2024/25 season
 - Business development processes for Carragelose franchise, Budesolv and Tacrosolv ongoing

Cash

- Ended Q2 2024 with €0.9m in cash
- Received +€0.5m milestone from new partnership with leading consumer healthcare player for partnership in Europe
- Raised +€0.8m through capital increase on September 18th, 2024

Cash drain stabilized, inflow from milestone and capital increase – cash sufficient to fund restructuring proceedings

Outlook



Restructuring proceedings

On August 14, 2024, restructuring proceedings without self-administration were opened for Marinomed Biotech AG

→ Inability to raise funds required at short notice to secure the Company's liquidity

→ Revenue expectations for 2024 financial year could not be realized as anticipated

Successful first step:

- First court hearing (September 26th, 2024) with decision to continue the Company

Next steps:

- Development and implementation of a restructuring plan to sustainably secure the Company's financial stability
- The Company's operations are to be continued as part of the restructuring proceedings
- Marinomed shares continue to be listed in standard market segment of the Vienna Stock Exchange, normal trading possible
- Publication of the half-year financial report 2024 planned after the three-month restructuring phase
- Continued cost saving measures: Concentrate on UGB reporting, suspend publication of voluntary quarterly reports
- More information on the restructuring proceedings of Marinomed Biotech AG can be found here:
<https://www.marinomed.com/en/investors-esg/restructuring-proceedings>



Summary project status (October 2024)

	Project	Status/next steps
Immunology	MAM-1001-4 nasal spray	Launched in Austria; drive BD-process for further partnerships
	MAM-1001-3 eye drops	Completed clinical study in Spain and launch planned in 2024; drive BD-process
	MAM-1004-1/Budesolv	Additional partnerships in 2024; work on registration with Luoxin for Greater China
	MAM-1003-1/Tacrosolv	Late-stage clinical development – enable first partnership
Virology	Carragelose OTC portfolio	Progressing partnership with leading consumer healthcare player for Europe and other selected countries; P&G: support in FDA process & preparations for launch Close gaps with new partners (e.g., Eastern Europe)
	MAM-2001-1/Carravin	Partnering in progress
	MAM-1001-1/Inhaleen	Prepare for certification as medical device
Solv4U	Solv4U	Progress the three long-term partnerships Add more partnerships



Preparing for long-term perspective

Turning innovation into revenue as basis for the continuation of the Company

**Agreement with
creditors &
completion of
restructuring
proceedings**

**Sale of
Carragelose
portfolio**

**Conclude license
agreements for
Budesolv &
Tacrosolv**

**Expand Solv4U
technology
partnerships**

Focus on generating near-term cash flows & successful continuation of Company



Financial calendar & IR contact

Financial Calendar 2024

TBD*	Publication of the Results H1 2024
Canceled**	Publication of the Results Q1-Q3 2024

Upcoming Events

November 25-27, 2024	Deutsches Eigenkapitalforum Frankfurt
----------------------	---------------------------------------



Lucia Ziegler

Investor & Public Relations
phone: +43 2262 90300 158
e-mail: ir@marinomed.com

www.marinomed.com

