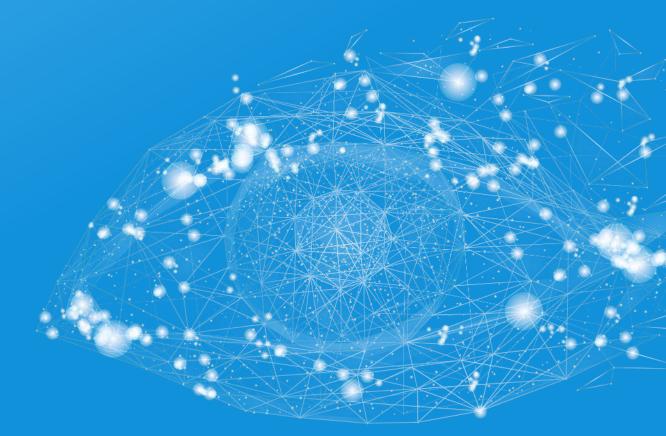
Marinomed programs at a glance

Business Development
October 2022





Marinomed at a glance

Lean, public company with track record of successfully inventing, partnering and marketing products creating sustainable value for patients and stakeholders

Company

Founded in 2006

Marketed products since 2008

IPO in 2019

Virology Immunology

Unmet medical need

Immunology:

Rare disease: herpetic stromal keratitis

Virology:

No broad-spectrum antiviral treatment for viral pneumonia

Patented technologies

Immunology:

Marinosolv® - solubilisation technology for hydrophobic small molecules/peptides

Virology:

Virus blocking Carragelose®

Successful partnering

Virology:

6 products with >25 M units sold in 40+ countries

Immunology:

First licence deal

Solv4U: technology partnerships





Marinomed has built a proven track record

Successes in the therapeutic areas

Immunology (Marinosolv®)

• 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



Virology (Carragelose®)

- **Discovery** of the efficacy of Carragelose® against respiratory viruses
- Performance of a series of clinical studies to support the efficacy
- Generating sustainable revenues and contribution margins
- Patent protected
- Successful partnering in more than 40 countries across the globe; examples:



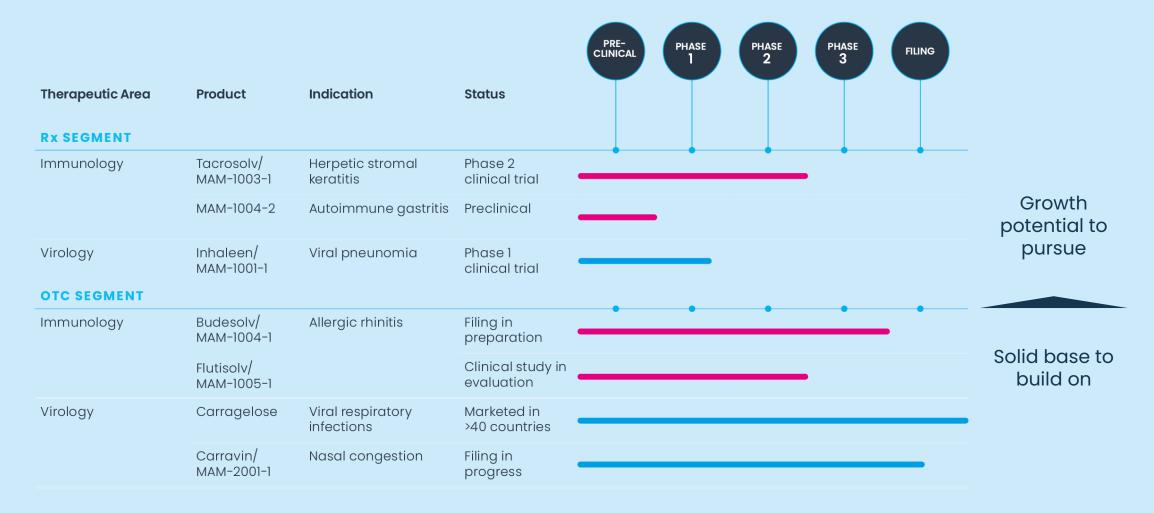








Strong pipeline for future growth



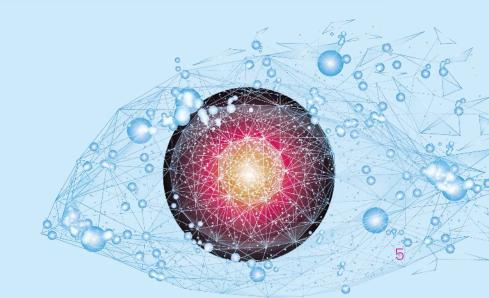


Tacrosolv (MAM-1003-1):

Highlights & strategy

- Tacrolimus is a potent immunosuppressant with a very low solubility in aqueous formulations
- The goal is to make it available for a multitude of ophthalmic inflammatory diseases
- Marinomed has developed a formulation with substantially increased solubility and bioavailability enabled by Marinosolv® technology
- Increased bioavailability enables a reduction of dose as preclinically demonstrated
- Anti-inflammatory activity clinically shown in model indication (allergic conjunctivitis)
- Significant improvement of HSK (herpetic stromal keratitis) indicators was shown in tacrolimus add-on therapy in a reference study

- Looking for an experienced partner in ophthalmology for further clinical development in HSK or other relevant diseases
- Available for in-licensing
- IP protection (Marinosolv®) until 2036
- Classical license deal favored





Autoimmune Gastritis (MAM-1004-2):

Program highlights & strategy

- Autoimmune gastritis is an **inflammatory** disease of the stomach's mucosa leading to the destruction of parietal cells
- The goal is to develop the first medication for this unmet medical need
- Loss of parietal cells leads to a pH increase in the stomach
- Increased risk to develop pernicious anemia and gastric tumors
- "Female disorder" women are more affected than men
- Difficult to diagnose due to diffuse symptoms (biopsies are required)
- Establishment of biomarker pattern to identify patients early in the disease to allow a precision therapy restore the mucosa and prevent further worsening of the condition

- Looking for partner in the field of biomarker diagnostic
- Apply for external funding, national and EU
- IP protection (Marinosolv®) until 2036



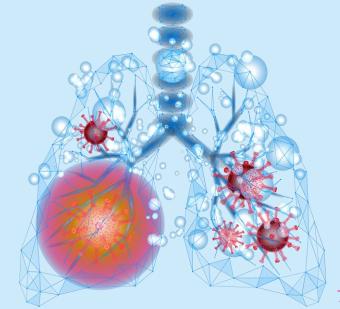


Inhaleen (MAM-1000-1):

Highlights & strategy

- Inhalable formulation based on iota-carrageenan
- The goal is the prevention and therapy of viral infections of the lung
- It is expected that a combination of iota-carrageenan with an antiviral drug will result in an increased efficacy
- Already demonstrated with the combination of iotacarrageenan and zanamivir in an influenza infection model
- Currently the inhalation of iota-carrageenan is tested in hospitalized COVID-19 patients in several clinics in Austria
- Treatment of viral respiratory infections caused by major pathogenic viral families
- Substantial data package already available for iotacarrageenan

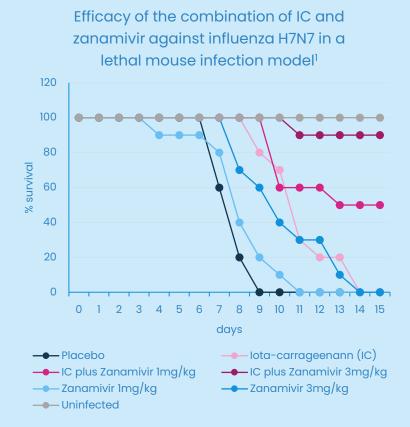
- Looking for an experienced partner to develop a broad-spectrum inhalable drug-device combination
- Available for in-licensing
- Classical license deal favored





Inhaleen (MAM-1000-1):

Broad-spectrum inhalable antiviral product combining at least two different mode of actions for optimal treatment of viral pneumonia



Rationale for the use of iotacarrageenan (IC)

- Clinically proven to significantly reduce viral load¹
- Clinical effectiveness shown for human Rhinoviruses, Coronaviruses and Influenzaviruses²
- Synergistic efficacy in a mouse model shown for combination with the anti- influenza drug Zanamivir*,3

Next steps

- Evaluate & identify antiviral combination with IC
- Carry out additional toxicology studies to enable early clinical study

Virus type	Virus-blocking activity demonstrated
SARS-CoV-2	X
SARS-CoV-1	X
Human coronavirus, endemic	X
Influenza A (human)	X
Influenza A (avian)	X
Influenza B	X
Parainfluenza virus	X
Respiratory syncytial virus	X
Human metapneumovirus	X
Human rhinovirus A and B	X
Human adenovirus	X

Antiviral inhibition by IC as shown in in vitro, in vivo or clinical studies³



²Könighofer et al., Multidisciplinary Respiratory Medicine (2014); 9: 57

³Morokutti-Kurz et al., PLOS ONE (2015) | DOI:10.1371/journal.pone.0128794. *Zanamivir is licensed for the treatment of uncomplicated influenza under the tradename Relenza

Budesolv (MAM-1004-1):

Highlights & strategy

- Budesolv is a Marinosolv®-based nasal spray of budesonide to treat allergic rhinitis
- Substantially improved **solubility** of API and thereby strongly increased bioavailability
- Phase 3 clinical study successfully completed
- Dose reduced by >85% compared to the marketed product
- Early onset of action shown
- Licensed to a first development and marketing partner in
 Greater China
- Further addressable indication identified

- Available for in-licensing outside China
- IP protection (Marinosolv®) until 2036
- Classical license deal favored





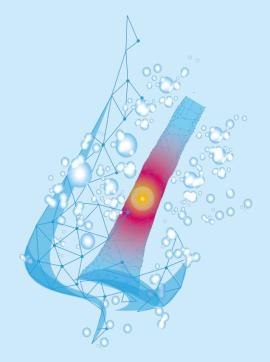


Flutisolv (MAM-1005-1):

Highlights & strategy

- Flutisoly is a Marinosoly®-based formulation of fluticasone proprionate
- Substance well established in various indications characterized by an inflammation including allergic rhinitis, asthma, COPD, or atopic dermatitis
- Substantially improved **solubility** of API
- Strongly increased **bioavailability** shown in ex-vivo models
- Stability data indicate storage at room temperature is possible
- Phase 2/3 clinical study in the indication allergic rhinitis in evaluation

- Available for partnering outside ChinaIP protection (Marinosolv®) until 2036
- Classical license deal favored



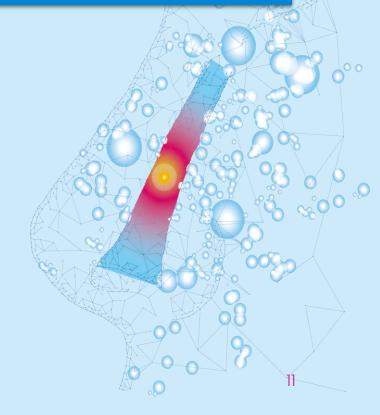


Carravin (MAM-2001-1):

Product highlights & strategy

- A nasal spray combining the virus-blocking properties of iota-carrageenan and the decongestant properties of xylometazoline
- Currently in registration process in several European countries
- Product should provide **immediate relief** of the symptom blocked nose
- Data demonstrate that the two components are **not** interfering

- Available for in-licensing
- Classical license deal favored





SOLV4U Technology partnerships

Marinosolv® solubilization technology to improve drug delivery and bioavailability

About Marinosoly®

- Patent protected drug delivery platform
- Based on **micelles** formed by excipients
- Clinically validated in phase 2 and 3 studies

Suitable for

- Hydrophobic small molecules and peptides
- All stages of NCE/API development
- Re-formulation and re-purposing of established APIs
- Life cycle management of APIs facing patent expiry
- Differentiation via USFDA 505(b)(2) pathway
- Locally applied, locally acting drugs for sensitive tissues
- Systemic administration routes

We offer

- Customized formulation development
- Feasibility studies
- Production of samples for early stage tox/PK studies
- Formulation optimization
- Upscale and tech transfer of manufacturing process

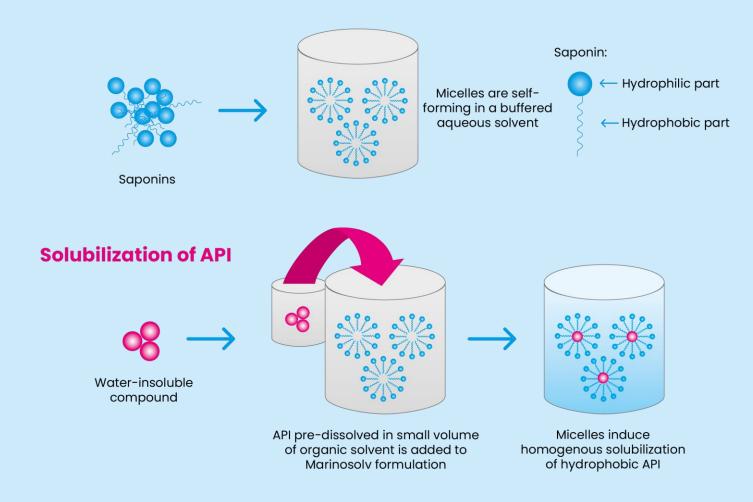




The technology: Marinosolv® works in two steps

Solubilization of water-insoluble compounds in a micelle forming aqueous system

Marinosoly formulation





Solv4U: Formulation development

Feasibility study and development of scalable manufacturing process

PHASE I FEASIBILITY

- Establishment of analytical method
- Design, prepare and evaluate of 30 different combinations of Marinosolv® formulation with target compound
- 6-day stability data
- Report
- Solubilized Marinosolv®-API formulation samples for early stage tox/PK studies

PHASE II

FORMULATION DEVELOPMENT

- Optimize formulation and evaluate stability according to ICH guidelines
- Provide non-GMP material for preclinical studies

PHASE III

SUPPORT CLINICAL & SAFETY STUDIES

- Develop an up-scalable production process
- Provide support for CDMOs
- Selection of potential primary packaging and other product requirements

PHASE IV TECH TRANSFER

- Support tech transfer to CMO selected by client
- Establish a production schedule

 LICENSE MODEL based on milestones and royalties from the Clinical Development phase and beyond



Carragelose

Expertise in Virology – brought to market

Mode of action







Carragelose forms a barrier on the nasal mucosa and unspecifically entraps viruses

Prophylactic and therapeutic efficacy clinically validated

Broadly active across several respiratory virus families and excellent safety profile

Partner Logos (sample)























Product pictures (sample)





East Asia







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