Sciv4U

Solv4U technology partnerships

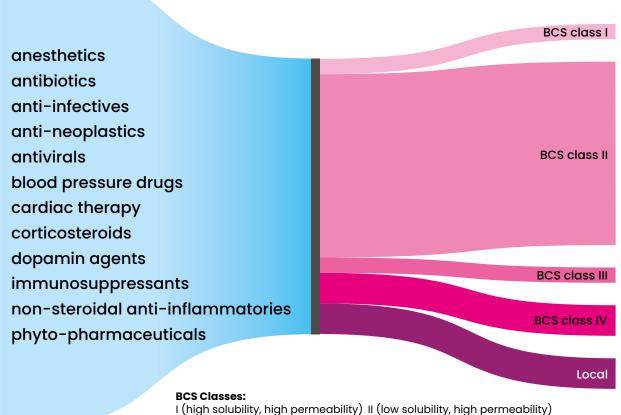
SOLVE THE UN(DIS)SOLVABLE



Solve the un(dis)solvable www.solv4u.com

Solubility enhancement

Marinosolv® solubilization APIs (biopharmaceutics enables the of from mainly BCS classification system) classes and validated the technology 11 IV. Marinomed has development through clinically validated its own and well tolerated product projects, improving established APIs. Furthermore, several compounds have been solubilized successfully in different client projects.



I (high solubility, high permeability) II (low solubility, high permeability) III (high solubility, low permeability) IV (low solubility, low permeability) Local (topical used only)

Using Marinosolv[®] as formulation technology offers several new applications for highly insoluble substances in eyes, lungs, as well as several other sensitive target tissues, resulting in high local activity and low systemic side effects. Marinosolv[®] may be used for re-formulation and/or re-purposing of established API's enabling life cycle extension of locally applied, locally acting drugs but also for systemic administration routes.





Local Systemic (e.g. intranasal/ocular) (e.g. oral/intravenous)





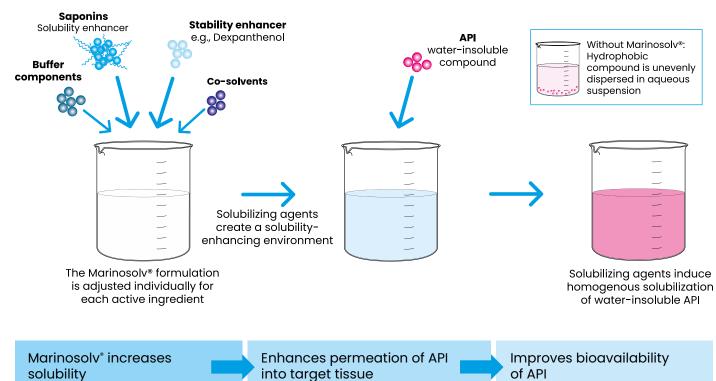


Liquid solution

Mode of action

Saponins (e.g., Escin, Glycyrrhizin), stability enhancers (e.g., Dexpanthenol) and selected solvents are used as solubility-enhancing agents in aqueous formulations. The different constituents of the Marinosolv® matrix are individually composed in buffered solutions for an optimized formulation of each drug substance.

Marinosolv® formulation & solubilization of water-insoluble API



Patients will benefit from	Manufacturers will benefit from
Faster onset of action	Easily scalable process
Dose reduction	Increased sustainability of manufacturing
Lower possible side effects	Option for preservative-free formulations
Well-tolerable local and systemic administration	Aseptic filtering of formulation

Suspension in water/buffer





Solution enabled by Marinosolv®



Support from early stage to product manufacturing

Solv4U partnerships address formulation challenges in small molecules and peptide drug development and rely on a clinically proven technology. Formulations are customized for individual compounds to achieve an optimal result. Formulation services include a feasibility study with clearly defined scope and timelines. All formulations are custom-made and designed in close cooperation with our clients, taking into consideration the physicochemical uniqueness of each compound as well as the requirements for later development stages and manufacturing.

PHASEI FEASIBILITY

- **PHASE II FORMULATION** DEVELOPMENT
- Optimize formulation & stability studies
- Provide non-GMP material for

Design & prepare Marinosolv® formulations with target compound

analytical method

Proof of enhanced

compound with **Marinosolv®**

• Establishment of

solubility of target

• Evaluate approx. 30 different variations

- perfom indicative
- preclinical studies

PHASE III MANUFACTURING **PROCESS**

- Joint definition of TTP & specifications
- Support of preclinical & clinical studies

PHASE IV TECH TRANSFER

 Support tech transfer to CMO/ CRO based on selection of the client

LICENCE MODEL

Based on milestones and royalties from the clinical development phase and beyond

About Marinomed Biotech AG

Marinomed Biotech AG is an Austrian, science-based biotech company with globally marketed therapeutics and a growing development pipeline. The Company focuses on the development of innovative products based on two patent-protected technology platforms, Marinosolv® and Carragelose®. Marinomed is listed on the Vienna Stock Exchange (VSE: MARI).



For more information, please contact our Business Development team: E-Mail: bd@marinomed.com Tel: +43 2262 90300 www.solv4u.com

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