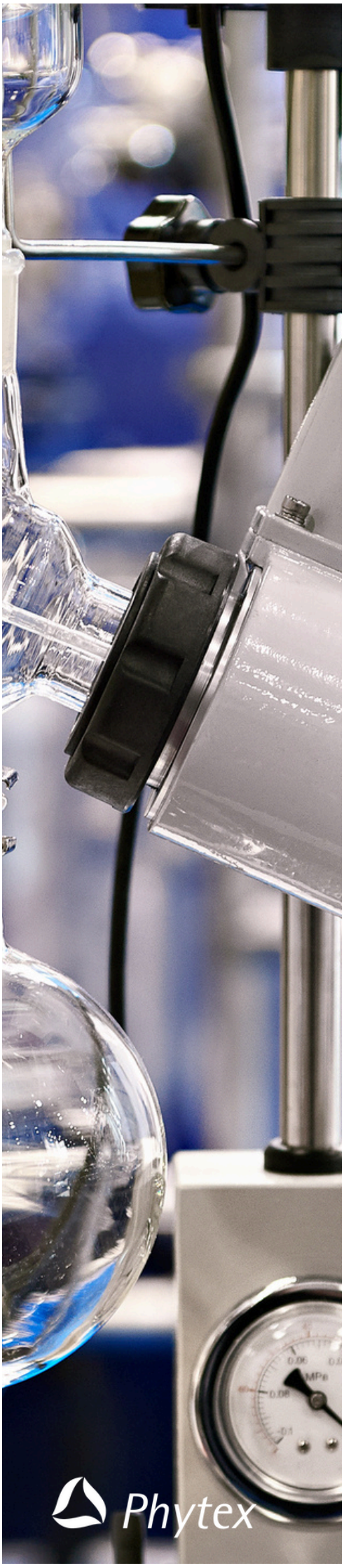

Our API products

USDMF 7415

Scopolamine (Hyoscine) Hydrobromide
USP & EP specification
Active (1988) Type II USDMF & CEP (2023)
Active Type I MF (Health Canada)

USDMF

Atropine Base
USP and EP specification
Type II USDMF*

USDMF 7408

Scopolamine Base
EP specification
Active (1988) Type II USDMF

USDMF

Atropine Sulfate
USP specification
Type II USDMF*

USDMF 10880

Castanospermine
USP specification
Active (1988) Type II USDMF

USDMF 7410

Methscopolamine Bromide
USP specification
Active (1988) Type II USDMF

*DMF in development

Our intermediates & KSM

Available now

Scopine HCl
Specification upon request
Tiotropium Br Intermediate

Available now

Scopolamine HBr
CP & USP specification
Intermediate grade

Our API pipeline

Evaluation ongoing

Cochicine Base
USP, EP & CP specification

Evaluation ongoing

Solasodine Base
Specification upon request

Exceptional track record at global standards

With **40 years of experience** in Good Manufacturing Practice, Phytex maintains an exceptional track record of compliance recognised by all international regulators of the markets we work in. These include the Australian Therapeutic Goods Administration (**TGA**), United States Food and Drug Administration (**USFDA**), Health Canada (**UC**), European Directorate Quality for the Quality of Medicines (**EDQM**) and Medicines and Healthcare products Regulatory Association (**MHRA**).

Our expertise lies in alkaloids and glycosides used in pharmaceutical marketed prescription medicines, medical research and product development; all manufactured to United States (USP) and European Pharmacopeia (EP) specifications.

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