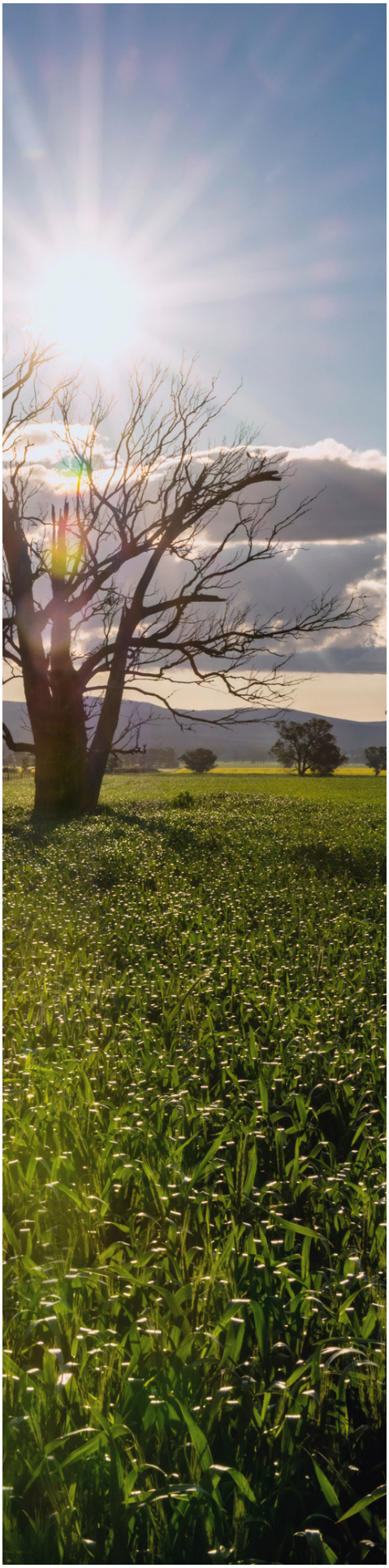
Phytex Australia

Leading global phytochemist manufacturers



Phytex is an Australian owned and operated company specialising in the extraction, isolation and purification of active pharmaceutical ingredients (APIs) derived from Australian native flora. Founded in 1979 and headquartered in Sydney Australia, Phytex is a leading supplier to major global pharmaceutical manufacturers and research organisations.

Zero product recalls since 1979

Experts in extraction, isolation and purification from natural sources

Australian owned, operated and sourced

Guaranteed consistent quality, supply and delivery

Phytex reliability and project technical solutions enable our customers to focus on critical and technical challenges in the product formulation, application, clinical trials and expansion processes.

Security of supply is underpinned by **strong supplier partnerships**. Phytex has maintained relationships with key raw material suppliers for more than 25 years, for the shared benefit of customers, manufacturers and Australian farmers.

With a highly flexible operational capacity, Phytex specialises in supporting **early-stage drug product development** requirements with the ability to **scale production to commercial quantities** (kilograms to tonnes) and meet the expectations of global market operators.

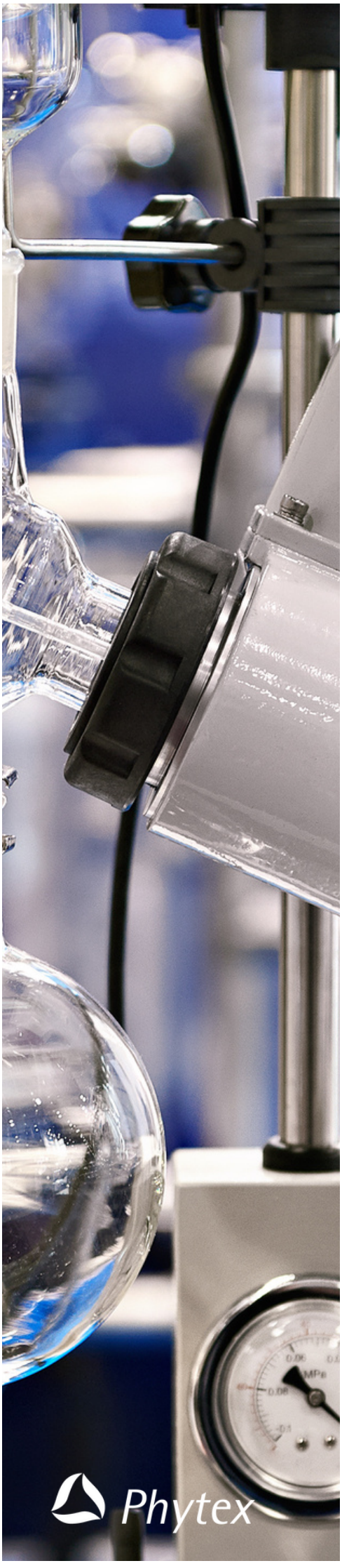
An experienced and trusted partner

Phytex has a long history of partnering and supplying APIs to global pharmaceutical clients to support their research, development and formulation for both clinical trials and commercial needs.

Phytex also has the onsite expertise and capacity for:

- **contract manufacturing** for the pharmaceutical, food and research industries
- **outsourcing stages of development**, such as the extraction and isolation manufacturing processes
- **research and development, regulatory navigation and commercial scaling consultation** for the entire drug development journey

Exploration	Supply	Manufacture	Support
Pre-development	Quality	Finished APIs	Technical
NDA/ ANDA Support	Stability	Contract manufacturing	Commercial
	Security		



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

Evaluation ongoing
Solasodine Base
Specification upon request

Available 2024
Tropine HCl
Specification upon request

Our API pipeline

Available 2024
Scopolamine N-Butyl Bromide
USP & EP specification

Available 2024
Anisodamine Hydrobromide
CP specification

Available 2025
Cochicine Base
USP, EP & CP specification

Available 2025
Tiotropium Bromide
USP & EP specification

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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