



**Your strategic API source
for niche molecules**



Your Partner in API's
Manufacturing



Lebsa is an innovative European laboratory dedicated to the development, manufacture and distribution of **Active Pharmaceutical Ingredients (API)** with **68 years of experience.**

Based in Barcelona, Spain, Lebsa is a trusted partner. **We are flexible, customer-centred, committed to successful long-term business relationship and continuously** seeking out new challenges, focusing on niche molecules, R&D development and up to date flow-chemistry processes.

Lebsa has always had high standards: in innovation, engineering, supply-chain management, regulatory support, quality control and customer service.



The challenging API industry

Lebsa provides solutions
for these challenges.

Following the **highest standards**, Lebsa offers a product range of **niche, small-molecule APIs** while actively researching and innovating in order to bring new products to the market.

Besides, Lebsa facilities and expertise allow us to accept **API CDMO (Contract Development and Manufacturing Organization)** requests, in addition to our existing product catalogue.

Pharmaceutical companies are facing new and more complex challenges every day. From stronger regulatory boundaries to the urgent need for new developments.

We are on the brink of a revolution in human healthcare, and **the opportunities will be immense**. API manufacturers must rise to meet the challenge.

European quality API manufacturing

Lebsa also provides the **API regulatory support you need** to get your products to market on time **overcoming barriers** and **speed up time-to-market** processes.

Lebsa has outstanding expertise in **registering APIs in Europe, Canada, Taiwan, China, Russia, Brazil and Japan, among others**.

Facilities

Lebsa's European facilities feature cutting-edge technology, equipment and automated processes, as well as state-of-the-art validated IT systems, including SAP®, Opentext® and SCADA Intouch®.

Lebsa doesn't outsource production. Everything is done within their facilities, with **complete control** over operations and production management. Lebsa's own secure IT systems for supply-chain and quality control management assures **total data integrity**.

Our **highly trained technical team** is ready to work closely with you at every stage, offering total confidentiality, rapid responses and full regulatory support.

Technical details

Reactors:

All fitted with distillation units

Glass lined:	1,6 m ³	4
With rectification column		

Stainless steel:	2,5 m ³	1
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Centrifuges:

Stainless steel AISI 904	850 mm	4
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Dryers:

Vacuum Tray Dryer	2,5 m ³	1
Vacuum Bicone Dryer	1,5 m ³	1

Reaction conditions:

Temperature	-20 °C to 160	°C
Pressure	Up to 5	bar
Vacuum	Up to 1	mbar

Utility services:

Water steam boiler	2.000 kg/h	1
Chiller	500 kW	2
Scrubber	1.500 m ³ /h	2
Vacuum pump	300 m ³ /h	3



cGMP Pilot Plant

Lebsa's cGMP pilot plant is fully automated and develops API in a scalable, secure and environmentally friendly way. Open to third parties, it also offers analytical services and full regulatory support.

Lebsa's **experience and flexibility** allows us to manufacture from a laboratory scale of just a few grams to some kilograms in our GMP pilot plant (10, 63 and 250 litres reactors) or to industrial quantities (1600 or 2500 litres reactors). Compliance with the highest cGMP standards is always assured.

Technical features



Kilolab

Triple Wall glass reactor and distillation unit	10 litres
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Vacuum oven	2 trays
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Operation Conditions

Temperature	-85 °C to 200 °C
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Pressure	Up to 0,5 bar
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Vacuum	Up to 1mbar
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Reactors

All fitted with distillation units

Glass lined	250 litres
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With rectification column

Hastelloy C-22	63 litres
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Filter Dryers

Hastelloy C-22	0,1m ² / 60 litres
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Operation Conditions

Temperature	-20°C to 160 °C
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Pressure	Up to 3nbar
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Vacuum	Up to 1mbar
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API Research & Development with purpose

Our active R&D department is constantly focusing on **improving our molecular-synthesis processes**, shortening times and improving the final quality and stability of the molecules.

At the same time, we are incorporating **new green manufacturing technologies**, such as **flow chemistry**.



LABORATORY: R&D

Reactor

Microwave Synthesizer

Operation Conditions

Temperature 40-300 °C

Pressure 0-30 bar

Analytical equipment

HPLC/UHPLC with mass detector

LABORATORY: ANALYTICAL EQUIPMENT

High pressure liquid chromatography

Gas chromatography instrument with Head Space

UV/Vis Spectrophotometer 160

Infra Red Spectrophotometer FT-IR 8400S

Titration (Potentiometer) DL 25

Karl Fischer Apparatus DL 31

Melting Point Apparatus FP 61

Muffle furnaces

Vacuum furnace V0400

Polarimeter Carl Zeiss

Refractometer

Quality and compliance

Lebsa's concept of quality goes beyond certification. It extends to everything we do – and how we do it.

Our dedicated Quality Assurance (QA) and Regulatory Affairs department ensures complete compliance with regulations and manages the documentation for pharmaceutical registration records.



Our own QMS (Quality Management System) is based on the GMP ICH Q7A guide and the ISO 9001:2015 standard. Its objective is continuous improvement in all areas of the company.

All standard Lebsa products have EU DMF and/or CEP, which guarantee a robust manufacturing process, data integrity and traceability.

Transparency is total: Lebsa is regularly audited and inspected by clients and healthcare regulatory agencies. We work only with approved and trusted providers to assure full supply chain control.



We adhere to the highest standards of **health and safety in the pharmaceutical industry**. We work tirelessly to prevent risks, ensuring that industrial safety is paramount and fully incorporated into all operations.

Sustainability

Our respect for the environment drives us to follow best practices and achieve full certification in managing solid and liquid waste.



Standard APIs



Main Therapeutic Application	Product Name	CAS-RN	Specification	Documentation
ANTIPSYCHOTICS	Amisulpride	71675-85-9	Ph. Eur.	EDMF / CEP
	Sulpiride	15676-16-1	Ph. Eur.	EDMF
	Tiapride hydrochloride	51012-33-0	Ph. Eur.	EDMF / CEP
ANTIVERTIGO	Betahistine dihydrochloride	5579-84-0	Ph. Eur. / USP	EDMF / CEP
	Betahistine dimesilate	54856-23-4	Ph. Eur.	EDMF
ANTIEMETIC	Bromopride	4093-35-0	Ph. Brazil	EDMF
ANTISEPTICS	Dequalinium chloride	522-51-0	Ph. Eur.	EDMF / JDMF / CEP* <small>*submitted in 2019</small>
	Dibrompropamide isetionate	614-87-9	Ph. Eur.	EDMF
	Picloxydine dihydrochloride	19803-62-4	In-house	EDMF
	Propamide isetionate	140-63-6	In-house	EDMF
DIAGNOSTIC AGENT	Histamine dihydrochloride	56-92-8	Ph. Eur.	EDMF / CEP
	Histamine diphosphate monohydrate	51-74-1	In-house	EDMF
ANTIHYPERTENSIVE	Lacidipine	103890-78-4	B.P.	EDMF

APIs Pipeline

ANTIDEPRESSANT	Mianserin	24219-97-4	Ph. Eur	Technical package
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Are you looking for
other molecules?

For more information about our R&D
pipeline and CDMO services

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