



**ADRAGOS**  
PHARMA

**YOUR END-TO-END CDMO**

# Customer Centric Global CDMO

Focus on Drug Products

**CDMO**  
Full service

**6**  
Countries<sup>1</sup>

**1,400+**  
FTEs

LIVRON  
France

HALDEN  
Norway

LEIPZIG  
Germany

KAWAGOE  
Japan

ATHENS  
Greece

JURA  
Switzerland

**\$5bn+**  
Product value<sup>2</sup>

**100**  
Customers

**80+**  
Countries supplied

# Our Value Proposition



## Tailor Precision at Scale

We combine multinational operational strength with **personalized** service. Our **6 GMP** facilities within the EU, US, and Japan offer **local insights**, as we know that **one size doesn't fit all**.



## Legacy Meets Innovation

With **3 centuries of collective experience**, our team from leading pharma, biotech, and strategy sectors deliver innovative solutions grounded in established practices.



## R&D as a Profit Center

Our **proactive and value-driven** R&D focuses on transforming science into regulatory-ready, **scalable and market relevant products**, specifically value-added generics.



## Co-Ownership of Success

We **invest in your products success** beyond standard timelines. From tech transfers to lifecycle management, we offer more than just a Gantt chart—we provide **full attention and dedication**.



## Masters of Market Adaptability

Our established presence in **Japan**, the world's most demanding market, demonstrates our adaptability, commitment to quality, and long-term vision.

# Adragos History



# Our Expertise

End-to-End: from Development to Manufacturing

		Athens, Greece	Jura, Switzerland	Kawagoe, Japan	Leipzig, Germany	Livron, France	Halden, Norway
Drug Development	Custom API Synthesis	✓*					
	Drug Development Services	✓		✓			
	HPAPI Handling	✓					
	Clinical Trial Materials	✓*	✓ ✓		✓		✓
Manufacturing Capabilities	Lyophilization		✓ ✓				
	Sterile Liquids		✓ ✓	✓		✓	✓
	Non-Sterile Liquids				✓		✓
	Creams/Ointments				✓		✓
	Tablets			✓			
	Suppositories					✓	
Other Services	Visual Inspection		✓ ✓	✓ ✓		✓	✓
	Packaging Solutions		✓ ✓	✓ ✓	✓	✓	✓
	Controlled Drug Handling	✓	✓	✓		✓	✓
	Out-licensing	✓				✓	

# ATHENS, GREECE

With over 25 years of experience, our team specialises in the **development of Value Added Medicines and NCEs for global markets** and has been recently upgraded with HPAPI capabilities.



## OUR EXPERTISE

- 150+ submissions of eCTD Dossiers
- Product approvals in 20+ countries
- Expertise in NCEs, VAM and Generics
- Handling HPAPIs and controlled substances
- Out-licensing dossiers
- 70+ professionals engaged in pharmaceutical R&D
- Center of Excellence of galenic development
- Development of all dosage forms



## CERTIFICATIONS

- EU-GMP
- Compliance with FDA regulations



Aggelos  
Karatzas

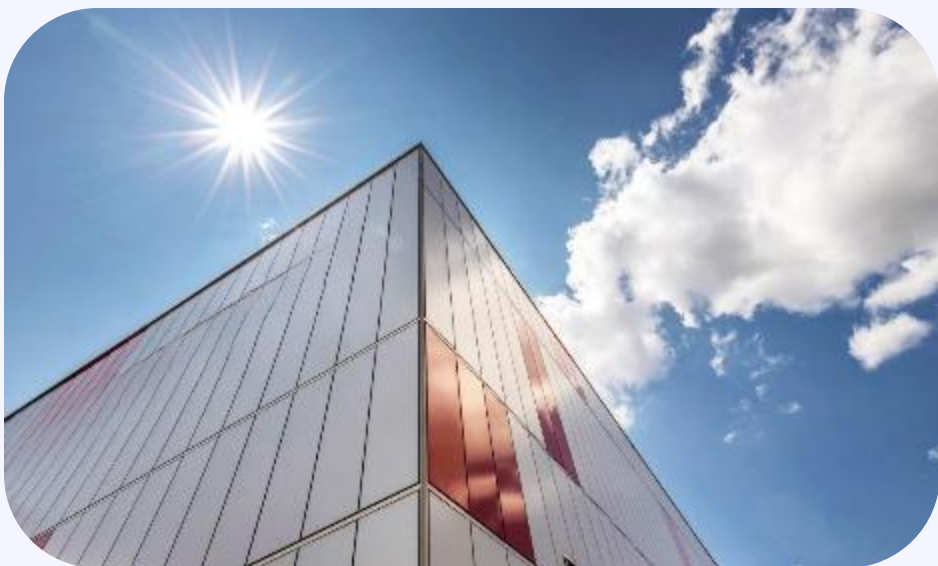
Site Head



Stella  
Chronaiou

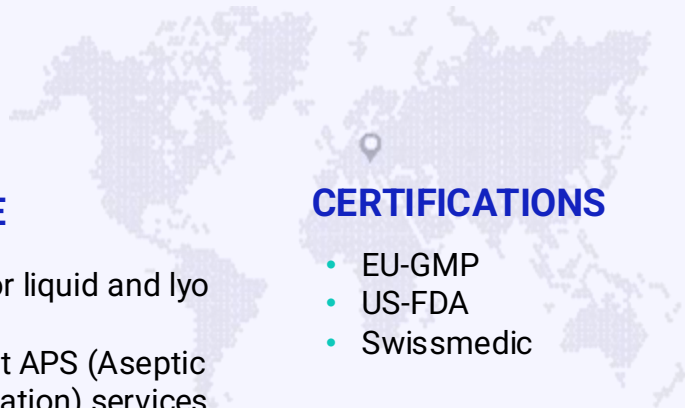
Customer Service





# JURA, SWITZERLAND

With over 25 years of specialization in **aseptic fill-and-finish** for both liquid and lyophilized vials, Jura offers expertise in **Clinical Trial Materials** (CTM) and small-to-medium scale commercial manufacturing.



Mike  
Stradling

Site Head



Marie-Sophie  
Quittet

Customer Service

## OUR EXPERTISE

- Aseptic F&F for liquid and lyo vials
- Fully compliant APS (Aseptic Process Simulation) services
- Biologics and Small Molecules
- 3-month response from order to batch
- Handling limited quantities of high valued materials
- No batch size restrictions: up to 74,000 vials
- Controlled drugs handling

## CERTIFICATIONS

- EU-GMP
- US-FDA
- Swissmedic



# KAWAGOE, JAPAN

With over 50 years of experience, our cutting-edge facility features one of Japan's largest cold storage warehouses, enabling top-tier visual inspection and packaging services for successful entry into the Japanese market.



## OUR EXPERTISE

- Visual Inspection and packaging of Small Molecules and Biological products
- Strong knowledge of local market, regulations, and supply chain
- Fluent English-speaking team
- Proven track record with Big Pharma
- Controlled drugs handling
- Manufacturing of OSD and ampoules

## CERTIFICATIONS

- Japan-PMDA
- Certified under EU-GMP standards
- Compliance with FDA regulations



Masanori  
Kurogome

Site Head



Munetaka  
Hirosaki

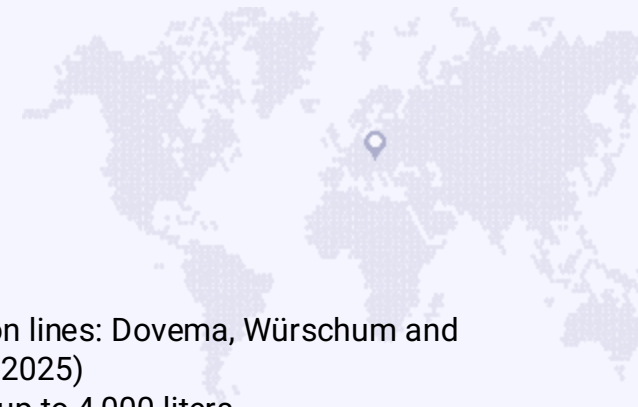
Customer Service





# LEIPZIG, GERMANY

With 100 years of experience, this facility specialises in the production of semi-solid and non-sterile liquid pharmaceuticals.



Florian  
Prell

Site Head



Franziska  
Flauder

Customer  
Service

## OUR CAPABILITIES

### Non-Sterile Liquids

- 3 production lines: Dovema, Würschum and Groninger (2025)
- Batch size up to 4,000 liters
- Plastic and Glass Bottles from 20ml to 700ml

### Semi-Solids

- 2 production semi-solid lines: Marchesini and Norden
- Batch sizes from 15 to 1250 liters
- Laminate tubes and plastic tubes (25g-200g) and aluminum tubes (2g-150g)

## OUR EXPERTISE

- Controlled drugs handling
- High quality facility located in eastern Germany
- Long-established and stable team
- From cosmetics to specialised pharmaceutical products

## CERTIFICATIONS

- EU-GMP Certification (human & animal products)

# LIVRON, FRANCE

With over 110 years of experience, our facility in France focuses on **manufacturing sterile liquids and suppositories**, providing stability studies, narcotics handling and analytical testing services.



## OUR EXPERTISE

- Controlled drugs handling
- 100+ molecules handled
- Long-established and stable team

## CERTIFICATIONS

- EU-GMP
- France-ANSM
- France-ANSES
- Korea-KFDA

## OUR CAPABILITIES

### Sterile Liquids

- 4 production lines: 2 aseptic with RABS technology and non-aseptic
- Batches from 30 litres to 1,500 litres
- Ampoules from 1ml to 20ml

### Suppositories

- 2 filling machines and packaging line
- Batches from 100kg to 2,000kg
- Suppositories or ovules up to 4,6g



Gaël  
Le Saux

Site Head



# HALDEN, NORWAY

One of Northern Europe's largest production sites with experience specializing in the production of IV bags and ampoules (BFS). Halden operates as a partnership with the Prange Group, the owner and a significant shareholder of the Adragos group.



Ian  
Cooper

CEO Halden



Nina  
Gabrielsen

Business  
Development

## OUR CAPABILITIES

### IV Bags

- 1 new bag filling line (Q1-2026)
- IV Bags from 50ml to 1,000 ml

### Ampoules

- 3 BFS lines
- Ampoules from 10ml to 30ml

## OUR EXPERTISE

- 40+ years legacy
- Proven experience with Big Pharma
- Global supply to more than 75 countries
- 95% of products exported outside Norway

## CERTIFICATIONS

- EU-GMP
- US-FDA
- Brasil - ANVISA
- Japan - PMDA
- China - NMPA
- Turkey – MOH
- Australia - ARGPM
- Taiwan – TFDA
- UAE

# Our C-suite



Dr. Andreas **Raabe**

CEO

McKinsey  
&Company



**SANDOZ** A Novartis  
Division



Henny **Zijlstra**

CCO

**Lonza**



Dr. Daniel **Wigbers**

CFDO



Marco **Gorgas**

COO



Dr. Norbert **Kuebler**

CSO







## Contact Us

[Info@adragos.com](mailto:Info@adragos.com)

[www.adragos.com](http://www.adragos.com)

[www.linkedin.com/company/adragos-pharma](https://www.linkedin.com/company/adragos-pharma)

