

COMPANY PROFILE

PARTNER FOR COOPERATION

VUOS

Where you can find us













VUOS ownership







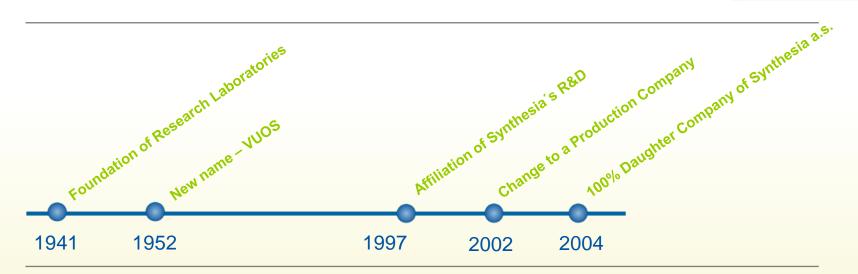
History









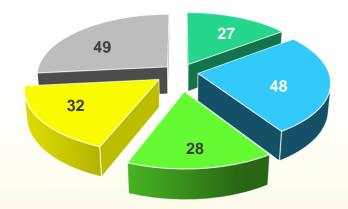




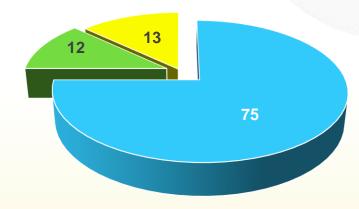
VUOS main businesses

- 1. R&D and Production of Fine Chemicals
- 2. Toxicology and Analytics









Sales in %



The Certificates of VUOS

- Certificate ČSN EN ISO 9001: 2016 (Organic chemicals syntheses development and manufacture)
- Certificate of Good Manufacturing Practice (GMP)
- Certificates of Good Laboratory Practice in Test Facility (OECD GLP Principles)

CERTIFICATE OF GMP COMPLIANCE CERTIFIKÁT SPRÁVNÉ VÝROBNÍ PRAXE

Part I / Část I

Institute for the State Control of Veterinary Biologicals and Medicines as national competent authority of the Czech Republic issues according to Section 16(2) letter a) item 3 of the Act No. 378/2007 Coll., on Pharmaceuticals and Amendments to Several Related Laws in current wording (hereinafter referred to as "Act on Pharmaceuticals No. 378/2007 Coll.") and in accordance with Art. 80(5) of Directive 2001/82/EC as amended, this certificate that to confirm that manufacturer

Ústav pro státní kontrolu veterinárních biopreparátů a léčiv se sídlem v Brně jako příslušný úřad České republiky vydává podle § 16 odst. 2 písm. a) bod 3. zákona č. 378/2007 Sb., o léčivech a o změnách některých souvisejících zákonů (dále jen zákon č. 378/2007 Sb., o léčivech) a v souladu s článkem 80(5) Směrnice 2001/82/EC, ve znění pozdějších předpisů, tento certifikát, kterým potvrzuje, že výrobce

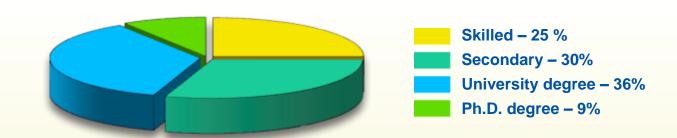
Výzkumný ústav organických syntéz a.s.



Strong Points

- Education structure of employees
- VUOS is located in area of long-lasting chemical tradition and complete infrastructure
- Cooperation with University of Pardubice









R&D, upscaling



- Pharmaceutical intermediates also under GMP standard
- Intermediates for microelectronics
- Dyes and pigments for special application
- Custom syntheses of chemical specialities





R&D, upscaling





Grams	-	in laboratory
Kilograms	-	in kilolab unit
Metric tons	-	in VUOS production plant

Hundreds of MT -





VUOS key chemistry



Adamantane derivatives



N-Heterocycles



API Building Blocks



VUOS 's key technology



Hydrogenation



Phosgenation



VUOS technology/manufacture expertise includes:

Organometallic and Cryogenic Chemistry

- Organolithium Chemistry
- Organozinc Chemistry
- Grignards

Transition-Metal Catalysis

- Homogeneous and Heterogeneous Transition-Metal Catalysis
- Cross Couplings (Suzuki, Kumada, Heck, etc.)

Hazardous and Unpleasant Chemistry

- Phosgenation
- Halogenation (PCI₃, POCI₃, SOCI₂, BCI₃, BBr₃, etc.)
- · Rections with Nitroalkanes
- Nitration (in small scale)
- Sulfonation

Reduction and High-Pressure Hydrogenations

- Reduction with Sodium borohydride, Red-Al, Superhydride, etc.
- Reductive Alkylation
- Reductive Amination
- Hydrogenolysis
- Dehalogenation





Kilolab and Production Unit

Phosgenation

Reactors up to 1 600 I

2x High performance rectification columns

Laboratory phosgenation units from 1 to 30 I





Kilolab and Semi-Pilot Unit

Hydrogenation



3x autoclaves 100 l each, 60, 120 and 130 bar, up to 200 °C, Hastelloy C clad

1x autoclave 300 l, 120 bar, 140°C, Hastelloy C clad

1x autoclave 700 l, 100 bar, 200 °C, stainless steel

2x autoclaves 1 000 l, 10 and 60 bar, 140 °C, stainless steel

many small autoclaves up to 10 l

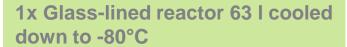






Kilolab and Production Unit

Low temperature, RLi, RMgX, R₂Zn



1x Glass-lined reactor 400 I cooled down to -80°C

1x Glass reactors 630 I cooled down to – 60°

3x Glass reactors 100 I cooled down to – 40°C







Kilolab and Semi-Pilot Unit

Multipurpose chemistry



4x glass reactors, 100 l each

6x glass lined reactors (200-1 500 l)

2x evaporators 50 l/h

4x stainless steel reactors (300–1 500 l) sufficient filtration and distillation capacity





GMP Production Line

Production of API, advanced intermediates, controlled finalization



Glass-lined vessels 400 and 630 l

Stainless steel vessels 400 and 630 l

Filtration, Centrifuging and Drying





Production Unit

Production Equipment

52x glass lined reactors, total vol. 85 000 l (500 – 6 000 l)

16x stainless steel reactors, total vol. 29 000 I (1 000–6 000 I)

8x High performance rectification columns

sufficient filtration capacity (nutsches, centrifuges)

computer controlled







New

Pharma Intermediates Production Unit

Purpose



5 reactors, total volume 10 000 l, stainless steel and glass lined

Filtration, distillation, drying

In operation from March 2021

Investment cost 4 mil. EUR





















New Final Operations Unit dedicated for grinding, drying and blending of dry materials

Purpose

Dedicated for the for grinding, blending and drying of pharma intermediates under GMP standard

3 tray dryers in separate rooms with separate ventilation

Equipment for grinding and blending

Investment cost 1.5 mil. EUR







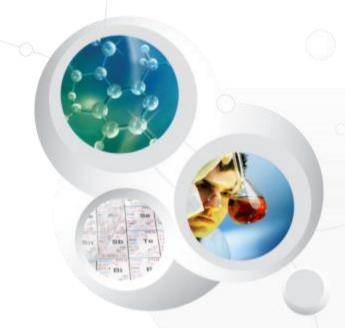
New Final Operations Unit will be in operation in

4Q. 2021















REACH and CLP Services

- Literature search
- Preparation of dossiers in IUCLID 5
- Preparation of Chemical Safety Report
- Non-testing approaches
 - (Q)SAR
 - Grouping/Read-across
- Preparation of SDS
- Classification of substances and mixtures







Analytical methods

- NMR Spectrometry, Mass Spectrometry (LC/GC-MS)
- ICP-AES, IR Spectrometry, Chromatography (HPLC, GC), UV-VIS

GMP control laboratory

Servis for Customers: Validation, IPC, Quality Control









Toxicological tests

In vivo and in vitro tests according to the OECD or EU methods:

- Acute toxicity
- Skin and Eye irritation
- Sensibilisation
- Repeated dose toxicity
- Reproductive toxicity
- Carcinogenicity
- Mutagenicity





Ecotoxicological tests

- Aquatic toxicity
- Biotic degradation
- Abiotic degradation
- Adsorption/desorption

Physico-chemical tests

 Information on physicochemical properties of substance





Contacts

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