

- Generic API
- CDMO API
- CMO Drug product



## Orion at a glance (2020 figures)





Net sales 1,078 MEUR



Operating profit

**280** MEUR



Personnel

3,311



**R&D** investments

**123** MEUR



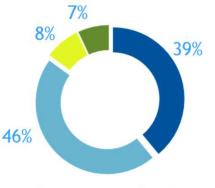
6 production sites in Finland



Own sales unit in **26** European countries, Singapore, Malesia and Thailand

Established in 1917

### Sales by business



■ Proprietary Products

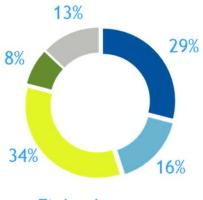
■ Specialty Products

Animal Health

■ Fermion & CM\*

\*) Contract manufacturing

#### Sales by market area



■ Finland

Scandinavia

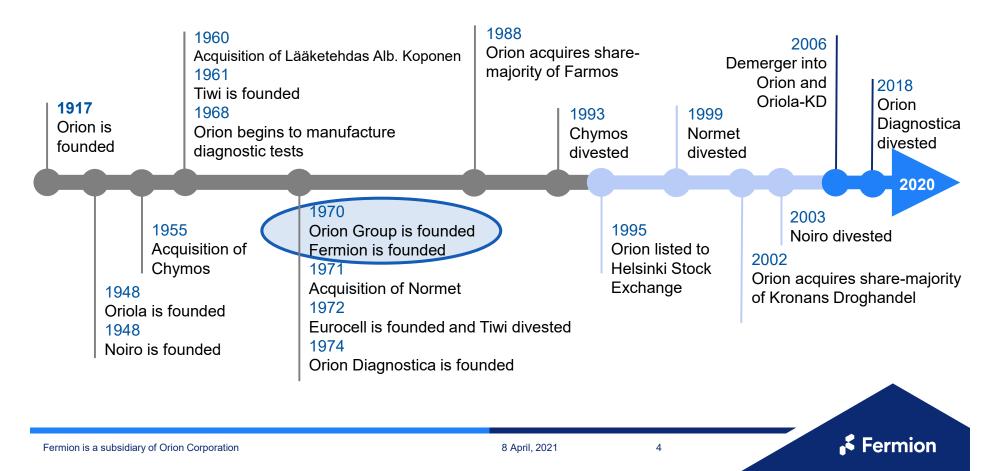
Other Europe

■ North America

■ ROW



## A century of Finnish industrial history



#### **Orion businesses**



Specialty Products 



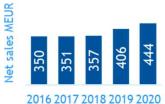
Fermion & Contract manufacturing

In-house developed drugs and other drugs with valid product protection for global markets. Own sales network in Europe. The most recent approved innovative drug is Nubeqa (NCE darolutamide)

Generic prescription drugs, OTC and non-medicinal products, biosimilars. Finland 56% Scandinavia 16% Eastern Europe 14% ROW 14%

Own animal drugs for global markets. Other drugs and well-being products. Own sales network in the Nordics and Eastern Europe.

Active pharmaceutical ingredients (APIs) for own proprietary products. CMO & CDMO services for other pharma companies.



39%



2016 2017 2018 2019 2020

46%



2016 2017 2018 2019 2020

8%



7%

5

= share of Group net sales in 2020



## Fermion Oy in 2020

- Develops, manufactures and sells active pharmaceutical ingredients (APIs)
- Markets drug product contract manufacturing services of Orion



Net sales 102 MEUR: External 58 %, Internal 42 %



Personnel ca. 370



Main office, S&M, R&D & Bench scale unit and Registration in Espoo



2 production sites: Hanko and Oulu Finland



Close to 230 customers, ca. 30 products

- 2020 Net sales 102 M€ (external 59 M€, internal 43 M€)
- Focus on APIs that are challenging to manufacture special expertise in the manufacturing of high potency APIs
- Leading global market share in some of its own products
- Main market areas are the United States, Europe, India, South Africa and Japan



## Fermion and CM strategic targets → Growth and profitability

Target to grow over 50% by 2025

Contribute to the growth and profitability of Orion Corporation Maximise the external generic API business Continue the renewal of generic API portfolio

Drive the growth of API CDMO business Develop
DP CMO business
building on Orion's
operational strengths
and API business
synergies



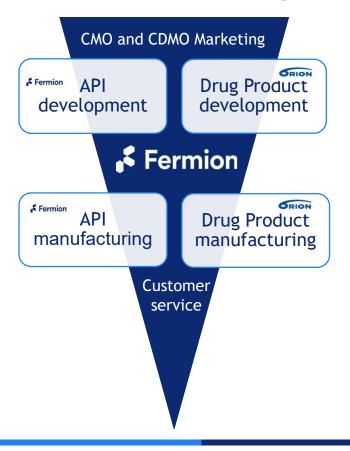
Improve productivity and ensure forecasted volume growth



#### Fermion's management **Fermion Board** Chairman Liisa Hurme Member Olli Huotari President Member Arto Toivonen ■ Management Team Members Jari Karlson Member **Executive Assistant** Virve Laitinen Solja Mannermaa **Finance Manager Tuomo Virolainen Head of EHS** Mervi Satta Operations **Marketing and Sales** R&D QA,QC,Regulatory Satu Vartiainen Marjaana Tapio Marko Salo **Arne Grumann Vice President Vice President Vice President Vice President**

8

## **Contract development and manufacturing at Orion Group**





## **API and Drug Product sites**





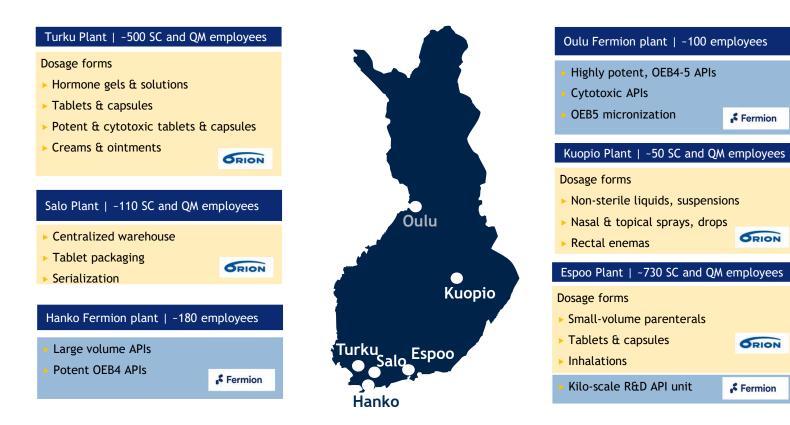








## **Locations and facility profiles**





## Orion has invested over 160 M€ to its manufacturing units during the last five years



2012-2015: Espoo unit expansion



2013-2014: New packaging and logistics center in Salo

27 Me



2011-2014: Fermion Oulu HPAPI unit expansion



2012-2014: Turku unit expansion



2015: Fermion kilo scale API unit expansion



2015-2019: Fermion Hanko, new large scale unit 4 40 Me



2019-> :
Continuous investements
to improve EHS,
automation, process
analytics and Al



2020-> :
New blister line for cytotoxic oral solids and renovation of ointment department at Turku 17 Me









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## Fermion has a strong position in global market with a number of its generic APIs

Product	US DMF	Europe CEP	Japan JMF	Specification
Alprazolam	✓	✓	✓	USP, Ph.Eur.
Azathioprine	✓	$\checkmark$	$\checkmark$	USP, Ph.Eur.
Benserazide		applied		Ph.Eur.
Budesonide		applied		
Buspirone HCI	✓	ASMF		USP, Ph.Eur.
Cabozantinib	✓			
Carbidopa	✓	✓		USP, Ph.Eur.
Dexmedetomidine	✓	ASMF	✓	
Diltiazem HCI	✓	ASMF		USP, Ph.Eur.
Entacapone	✓	ASMF		USP, Ph.Eur.
Fluticasone propionate		applied		
Flutamide	✓	✓		USP, Ph.Eur.
Formoterol fumarate	$\checkmark$	✓	✓	USP, Ph.Eur.
Glipizide	✓			USP
Hydroxychloroquine sulphate	✓	✓		USP, BP
Irinotecan HCI	✓	✓	✓	USP, Ph.Eur.
Levosimendan	✓	ASMF		

Product	US DMF	Europe CEP	Japan JMF	Specification
Mercaptopurine	✓	✓	✓	USP, Ph.Eur.
Methotrexate	✓	$\checkmark$	✓	USP, Ph.Eur.
Methotrexate disodium	✓	ASMF		USP, Ph.Eur.
Nadolol	✓	ASMF	✓	USP, Ph.Eur.
Nintedanib	✓	ASMF		
Nitrofurantoin Macrocrystals	✓	applied		
Nitrofurantoin Monohydrate	✓	ASMF		
Propafenone HCI	✓	$\checkmark$		USP, Ph.Eur.
Quetiapine fumarate	✓	✓		USP, Ph.Eur.
Salmeterol xinafoate		✓		Ph.Eur.
Sodium cromoglycate	✓	✓	✓	USP, Ph.Eur.
Tamsulosin HCI	✓		$\checkmark$	USP, Ph.Eur.
Tolnaftate	✓	ASMF		USP, Ph.Eur.
Toremifene Citrate	✓	ASMF	✓	
Trazodone HCI	✓	ASMF		USP, BP
Vemurafenib				









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## Fermion R&D at your disposal

Ability to take a molecule from pre-clinics all the way to commercial

Best from both worlds, proprietary and generics

Personnel of 50: 28 scientists, 29 % PhD, 20 technicians

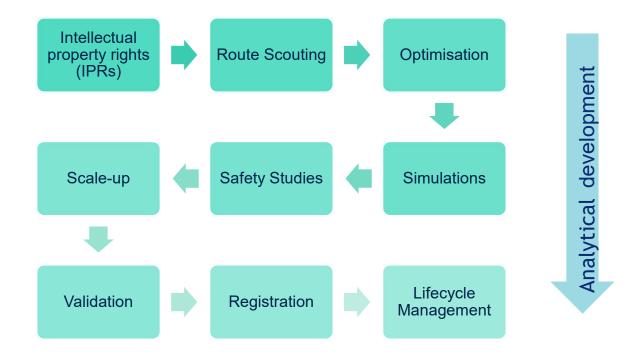
50 Years experience captured in organisational know-how

Dedicated engineer in every project involving upscaling

Automation in R&D and production

Strong particle engineering and physical properties control

## **R&D - Fully Integrated, Pick What You Need**



# Fermion has highly automated production units with recent expansions to highly potent compound (OEB4-5) manufacturing capabilities





- Small quantities for developmental purposes and clinical studies
- HPAPI handling capability up to OEB 5
- QC lab for development and PM



Oulu production incl. R&D pilot

- 75 m<sup>3</sup> reactor volume
- Small volume commercial products
- HPAPI handling capability, including micronisation, up to OEB 5
- Production QC lab



**Hanko production** 

- 250 m<sup>3</sup> reactor volume
- Large volume commercial products
- HPAPI handling capability up to OEB 4-5
- Production QC lab

## **Fermion Quality System**

Fermion's Quality System complies with the ICH Q7 GMP Guide for API's" and the EU GMP guideline "The Rules Governing Medicinal Products in the European Union – Part II: Basic requirements for Active Substances used as Starting Materials". The Quality System follows also the ICHQ9 guideline (Quality Risk Management), the ICHQ10 guideline (Pharmaceutical Quality System) and all other relevant guidelines for API manufacturing and GMP.



- The main Quality System elements are Change Control, Risk Management, CAPA, Deviation Management, OOS Management, SOP system and Internal Audits. The Management is committed to the continuous improvement of the Quality System. The QMS is reviewed annually in the Management Review.
- Quality Unit is involved in all GMP critical operations during the whole product lifecycle from the development phase to regulatory filing and market.



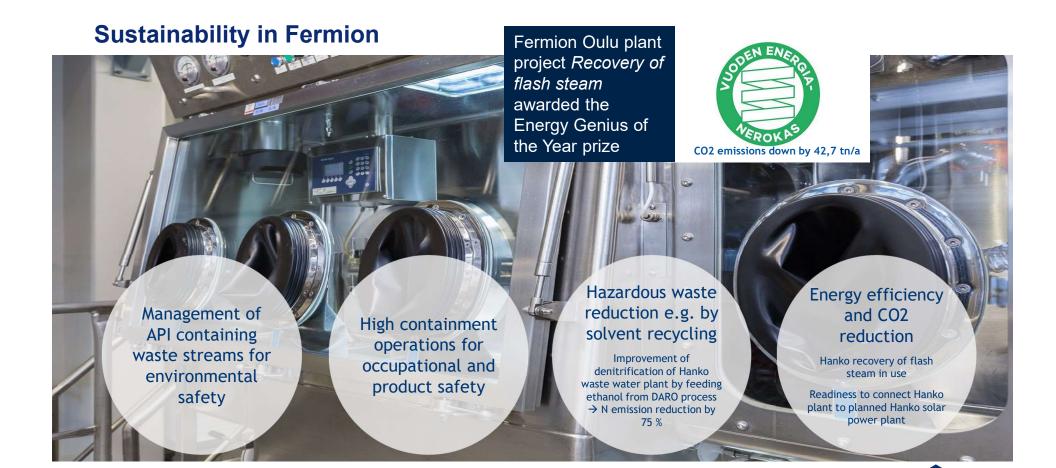
## Fermion is regularly inspected by health authorities

Fimea inspected Espoo and Oulu plants in Feb 2020.
 Based on Fimea inspection also FDA classified Oulu plant as acceptable (MRA applied).

CFDI = Center for Food and Drug Inspection

NMPA= National Medical Products Administration

	Espoo	Hanko	Oulu
FIMEA (* unit IV)	1995 1998 2002 2004 2007 2010 2013 2015 2017 2020	1995 2002 2005 2008 2011 2014 2017 2018*	1995 2002 2005 2008 2011 2014 2017 2020
US FDA	1993 1995 1997 1998 2001 2003 2011	1992 1993 1995 1997 1998 2001 2004 2007 2011 2014 2016 2018	1992 1995 1997 2001 2003 2006 2010 2013 2016 2017 2020
MOH (Mexico)			2010
KFDA (Korea)			2010
ANVISA (Brazil)			2011 2016
PMDA (Japan)		2006	











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## If you are looking for a partner who...



with high reliability throughout product life cycle

#### **WE OFFER**

- Established and highly developed processes and technologies
- Approvals of global regulatory bodies and long track record of inspections
- On-time delivery reliability
- Sustainability, CSR and EHS practices well in place
- Our experience to Customers



you to get your product from development to the market

#### **WE OFFER**

- Upscaling product to commercial stage
- Wide expertise throughout the whole value chain
- Diverse production and process capability as your needs grow
- Tailor-made complex solutions
- Our established processes to Customers

## **Espoo plant capabilities**



#### **Tablet production**

- •Direct mixing, high shear mixing, fluid bed drying
- Batch sizes of over 400kg
- •OEB4
- 1,5 billion tablets/capsules in 2020



#### **SVP** production

- •Vials (5-36ml) and ampoules (2-5ml)
- •20-600 litre batch sizes
- •OEB 4-5 and products for human and veterinary use
- Flammable material handling
- Solutions and suspensions
- •158 bulk batches and 2,5m packages with serialization in 2020



## Latest inspections by main authorities at Espoo site

Authority	Date	Area
FDA (USA)	May 24-Jun 1, 2018	Sterile and non-sterile production (GMP)
ANVISA (Brazil)	Apr 24-28, 2017	Sterile and non-sterile production
FIMEA (Finland)	Sep 14-17, 2020	Sterile and non-sterile production
INAME (Argentina)	Aug 31- Sep 4, 2009	Non sterile production
KFDA (Korea)	Nov 3-5, 2009	Non sterile production
MOH (Turkey)	Apr 18-22, 2016	Sterile and non-sterile production
The Turkish Ministry of Food, Agriculture and Livestock (MoA)	Jan 15-18, 2018	Sterile productions (Veterinary productions)
MoIT (Russia)	January 22-24, 2020	Non sterile production

## **Turku plant capabilities**



#### **Oral solids production**

- Direct mixing, high shear mixing, fluid bed drying
- Batch sizes starting from less than 100kg
- OEB5, cytotoxics
- 1 billion tablets / capsules in 2020
- 2,3 million packages of cytotoxic oral solids in 2020



#### Non-hormonal creams, ointments and liquids production

- One syringe line, two tube lines, one bottle line
- 150-1000 litre batch sizes
- OEB4-5, products for human and veterinary
- 811 tons of bulk product in 2020
- 4,8 million packages in 2020



#### Hormonal gels, creams and solutions production

- Single dose sachet line and three lines for bottles, all with serialization
- 150-200 litre batch sizes
- OEB4-5
- 99 tons of bulk product in 2020
- 2,7 million packages with serialization capability in 2020



## Latest inspections by main authorities at Turku site

Agency	Date	Subject
FDA (USA)	Sep 2-6, 2019	Non sterile production (all departments)
FIMEA (Finland)	May 8-10, 2019	Non sterile production (all departments)
MOH (Turkey)	Apr 16-20, 2012	Anti-Cancer Dept
ANVISA (Brazil)	Aug 13-17, 2018	Two products
KFDA (Korea)	May 6-8, 2013	Hormone Gel Dept
Taiwanese Authorities	Mar 6-8, 2013	Hormone Gel Dept
Gulf States (Saudi-Arabia, Bahrain, Kuwait)	Apr 14-16, 2014	Anti-Cancer Dept
MoIT (Russia)	Jun 20-21, 2018	One Product



## **Kuopio plant capabilities**

	2018
Bulk batches	184 pcs
Finished product batches	383 pcs
Solutions	254 000 I
Suppositories	0,3 Mpcs
Finished product packages	4,3 Mpcs



- Non-sterile liquids and suspensions
- Nasal and topical sprays, drops
- •Rectal enemas
- •200-6000 litre batch sizes
- •Four lines with different closure system capabilities
- •OEB4

The Plant is run mainly in one shift



## Latest inspections by main authorities at Kuopio site

Authority	Date	Subject
FIMEA (Finland)	Sep 3-5, 2013	Non sterile production
FIMEA (Finland)	Mar 2-3, 2016	Non sterile production
FIMEA (Finland)	Jan 29-31, 2019	Non sterile production



## Tablet packaging operations at a glance



All Orions' tablet packaging operations centralized to Salo site, excluding cytotoxics,  $\beta$ -lactames, cephalosporines and sex-hormones.

#### Facts:

- 7 packaging lines, 4 bottle line and 3 blister line
- More than 37 million sales packages per year
- 3 shifts, ~85 employees
- 10 000 m2 total, clean areas 5000 m2



## Latest inspections by authorities at Salo site

Authority	Date	Area
FIMEA (Finland)	Feb 27 – 28, 2018	Non sterile packaging and warehousing
FDA (USA)	Sep 28 – 29, 2015	Non sterile packaging and warehousing (PAI/GMP)
ANVISA (Brazil)	Dec 12-16, 2016	Non sterile packaging and warehousing
MOH (Turkey)	Oct 10-13, 2016	Non sterile packaging and warehousing
The Turkish Ministry of Food, Agriculture and Livestock (MoA)	Jan 17, 2018	Non sterile packaging and warehousing
MoIT (Russia)	Jan 20-21, 2020	Non sterile packaging and warehousing



## Orion's Sustainability Agenda and indicators 2020



Patient safety and ensuring reliable supply of medications



Responsibility for the environment



Responsibility for Orionees



Business ethics and transparency



Customer complaints (pharmaceuticals)

**/ O** Ppm (76)

0

GxP\* audits by Orion

141

Greenhouse gas emissions (scope 1&2)

18,611

tCO2e (20,123)

0

Energy savings target set for 2025 achieved

**53**%

(51%)

(BIII)

Injury rate

3.6

LTIF 1 (6.6)

Code of Conduct training, no. of participants

3 410



## **Contact us for CDMO API and CMO Drug Product services**



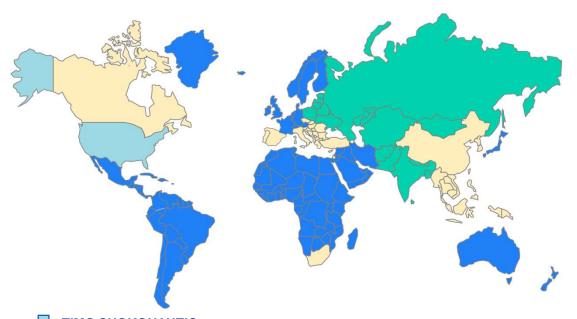
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