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CROSSJECT: DEVELOP SELF-ADMINISTERED DRUGS FOR EMERGENCY SITUATIONS



- Emergency
- Outside a medical area

- Self-administered
- Fast and easy to use

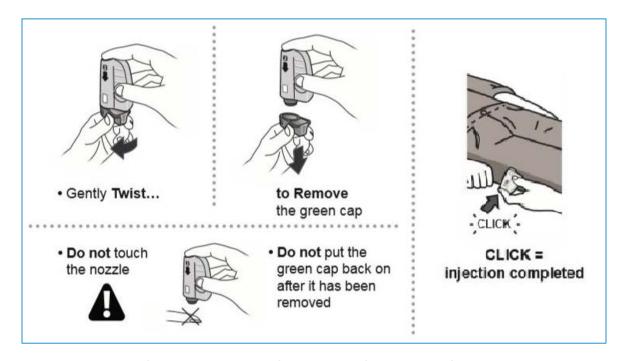


ZENEO®: self-administered medication in an emergency

- The patient is diagnosed as prone to attacks
- An emergency dose is prescribed
 - The patient keeps the dose with him/her constantly
 - The patient has an attack
 - The patient or someone else injects the emergency dose

ZENEO®, A BREAKTHROUGH TECHNOLOGY

- ZENEO®, an innovative, fast and safe needle-free self-injector, allowing patients to receive timely emergency therapy
 - o Pre-filled, single-use, 2-step auto-injector
 - o Intuitive, easy and quick administration
 - o Dose pre-set in factory for subcutaneous (SC) or intramuscular (IM) injection



Innovative therapeutic solutions, closer to the emergency

PIPELINE

Indication	Life-threathening emergency *** Emergency ** Other *	Therapeutic entities
Opioid overdose	***	ZENEO® Naloxone
Epileptic seizure	***	ZENEO® Midazolam
Allergic shock	***	ZENEO® Adrénaline
Acute migraine / Cluster headache	**	ZENEO® Sumatriptan
Asthma attack	***	ZENEO [®] Terbutaline
Acute adrenal insufficiency	***	ZENEO® Hydrocortisone
Rheumatoid Arthritis	*	ZENEO® Methotrexate
Parkinson	**	ZENEO® Apomorphine





RESULTS 2018

€ thousand	2018	2017	Impact of the production issues in the second half 2018
Operating income	3,524	4,142	
Operating expenses	(15,080)	(12,763)	
Other purchases and external expenses	(7,659)	(7,371)	Limited increase : tight rein on expenses in a context of development
Personnel expenses	(3,972)	(3,059)	expenses in a context of development
Taxes and duties	(126)	(59)	Hiring in 2018 in line with the needs at this
Depreciation, amortisation and provisions	(3,324)	(2,274)	stage of development. 75 at the end of
Operating profit/(loss)	(11,556)	(8,621)	2017 vs 58 at the end of 2018
Financial income/(expense)	(737)	159	Mainly impairment of treasury shares
Exceptional income/(expense)	(10)	(278)	
Income tax	1,592	1,129	Research tax credit
Net profit/(loss)	(10,711)	(7,611)	

2018 CASH FLOW STATEMENT

In € thousand	2018	2017		
Net Profit/(loss)	(10 711)	(7 611)	_	
Depreciation, amortisation and provisions	3 838	2 054		
Other comprehensive income and expenses	(120)			
Cash flow from operations	(6 993)	(5 557)		Increasing of the gross value of
Change in working capital requirements	(1024)	583		inventories illustrating the
(1) Net cash generated by (used in)				company's further development
operating activities	(8 017)	(4 974)		
Acquisition of fixed assets	(3 273)	(4 248)	_	
(2) Net cash generated by (used in)				
investing activities	(3 273)	(4 248)		Consolidation of Crossject's financial structure: capital
Capital increase	4 005	7 412		increase (4 M€) and convertible
Bond	7 750			bonds (nearly 7,8 M€)
Commercial paper		(755)		
Debts on fixed assets	100	(720)		
Repayable advances	1 448	3 457	_	
(3) Cash flow financing operations	13 303	9 394		
Variation cash (1)+(2)+(3)	2 013	172	_	
Opening cash positions	2 806	2 634		
Closing cash position	4 819	2 806	•	Marked improvement of the cash position

BALANCE SHEET 2018

€ thousand, as of 31 st december

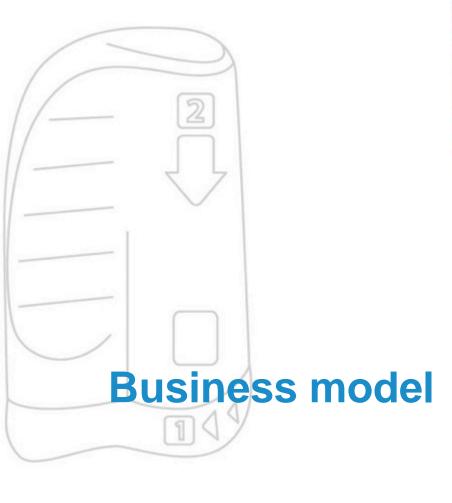
	2018	2017
Fixed assets	11 327	11 418
Of which R&D	4 615	4 031
Of which industrial plant and equipment	5 903	5 395
Current assets	8 832	6 298
Of which available cash	4 760	2 641
Total Assets	20 159	17 716

Shareholder's equity	1 655	6 086
Conditionnal advances	5 195	3 747
Provisions for risks ans expenses	125	93
Borrowings and debts	13 184	7 790
Of which bonds	5 475	
Of which debts on fixed assets	2 614	2 514
Total Liabilities	20 159	17 716

CASH CONTRIBUTIONS SINCE 31 DECEMBRE 2018

- January 2019 : 1,1 M€ loan
 - Granted jointly by Bpifrance and the Bourgogne-Franche-Comté region
 - 8 years, grace period of 3 years
- Expected additional cash contributions in 2019 :
 - Research tax credit + already agreed aid : 3 M€
 - Commercial revenues from licensing agreements

- Funding of
 Crossject's priority
 developments in
 2019
- Other aid currently under preparation or development
- Non-dilutive contributions currently under discussion
- Strenthening Crossject's long-term financing





BUSINESS MODEL



- ZENEO®, an innovative, fast and safe needle-free self injector
- New therapeutic entities: combination of ZENEO® with recognized molecules in targeted indications



- High-capacity production lines
- Manufacturing and supply organisation in place



- Sales of exclusive licences by product and by country to pharma companies operating in the relevant therapeutic area
- Take-up by physicians facilitated: molecules already known to practitioners

TYPICAL DEVELOPMENT PLAN



A simple pharmaceutical development process

1

Drug-system combination of a formulation of the molecule injectable by ZENEO® 2

Pharmaceutical development: galenic, industrial transposition, clinical and approval batches 3

Completion of a comparative bioavailability study in healthy volunteers to demonstrate good delivery in the body

4

Verification of the drug's stability over time



Filing of marketing authorisation (MA) requests, then receipt of MA roughly 1 year later 10 clinical studies including

- 1 published MRI study presenting the intramuscular performances of ZENEO®
- 1 published clinical trial presenting the subcutaneous performance of ZENEO®

20 consultations with regulatory authorities



- Easy and fast duplication of ZENEO® for new pathologies once the industrial process is finalised
- For each new drug, cost estimated at €2m-3m and development time at 2 to 3 years

Manufacturing process in place



- O Insourcing of strategic items in the supply chain
 - PARC® production unit (for Prêt À Remplir Crossject) in Dijon
 - Gray site: 3 production lines
 - Quality control and metrology
- Partnership with Cenexi, a recognised manufacturer
 - Production line dedicated to Crossject fill/finish operation
- End-2018: production capacity of 500,000 units per year
 - Sufficient space available to duplicate facilities
- O An acknowledged Quality approach: pharmaceutical establishment status obtained in early 2019



Easy-to-duplicate industrial equipment



DEAL-MAKING PROCESS



- Selection of possible partners
 - Direct and via brokers
 - By product/country
 - Focused on therapy area



4 deals signed (regional and worldwide deals)

- Negotiation of non-binding offers received
- Signing of licensing agreements
 - Publication of a press release

2018



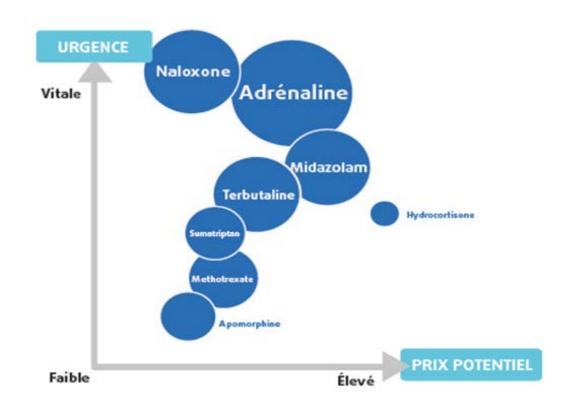
5 times as many NDAs with US companies than 3 years ago

PRICING POWER DRIVEN BY VALUE ADDED



Valuation criteria

- Save lives
- Reduced hospitalisation time and mobilisation of medical staff
- O Better compliance



€639M estimated IN-MARKET PEAK SALES*

Indication	Therapeutic entities	Key facts	Peak sales*	Priority
Opioid overdose	ZENEO® Naloxone	49 000 deaths/year in the US 97.3m patients with at least 1 opioid prescription	€64m	1
Epileptic seizure	ZENEO® Midazolam	6 ooo deaths/year in Europe and in the US	€88m	1
Allergic shock	ZENEO® Adrénaline	2.3m auto-injectors sold per year in the UK 4.5m units sold per year in the US	€158m	0
Acute migraine Cluster headache	ZENEO® Sumatriptan	+7m sumatriptan injection/year	€65m	2
Asthma attack	ZENEO [®] Terbutaline	7 ooo deaths/year in Europe ⁹	€107M	2
Acute adrenal insufficiency	ZENEO® Hydrocortisone	No easy to use solution adapted for emergency situations	€18m	2
Rheumatoid Arthritis	ZENEO® Methotrexate	Disabled patients	€86m	3
Parkinson	ZENEO® Apomorphine	Disabled patients	€53M	3

^{* :} Peak sales by Crossject distributors. Source: Average of financial analysts estimates / Crossject

PRIORITISATION OF MA FILINGS FOR 3 LIFE-SAVING DRUGS







OPIOID OVERDOSE: ZENEO® NALOXONE

Market potential

An epidemic in the United States...

49,000 deaths a year in the US

... that is creating additional commercial potential

FDA seeking to develop:

- Co-prescription (48.5 M doses per year¹)
- Over-the-counter sales (without a medical prescription)

Strong interest from pharmas

Recent deal Emergent
Biosolutions/Narcan® (\$625m)

Needle free

No risk of accidental contamination (e.g. HCV or HIV)

Superior early exposure

Early exposure @5 min : intramuscular = 2.7 x intranasal

Universal

12% nasal obstruction among US overdose cases 3





Narcan® Nasal spray

Competitive advantages

EPILEPTIC SEIZURE: ZENEO® MIDAZOLAM

Market potential

A crisis that can create severe neurological lesions resulting in death 50m people worldwide 6,000 deaths / year in Europe

Targeted indication

- US: finalisation of a development strategy, no expected overlap with existing orphan indications
- Europe: regulatory pathway confirmed for prolonged acute seizures, no orphan exclusivity on epilepsy

Route of administration

Intramuscular Midazolam (recommended by the AES guidelines)

Fast administration without injury on a convulsing patient

Absolute biovailability of intramuscular injection > 90% ⁴









Buccolam® (Europe) Oromucosal solution Infants and adolescents only

Competitive advantages

ALLERGIC SHOCK: ZENEO® ADRENALINE

Market potential

A widespread and growing pathology

2.3m auto-injectors sold per year in the UK¹⁰ 4.5m units sold per year in the US¹¹

Worldwide licence signed in 2013

€8m milestone payments upon drug approval

True intramuscular injection

MRI study - Advances in Therapy, 2017 ⁶

Universal injection into the muscle

including overweight people



Auvi-q®



Epipen®
1.1bn sales in 2016 ⁷
~16 mm needle length ⁸



Adrenaclick®



Symjepi®

Competitive advantages





ROLLING OUT OUR MODEL: ALL CLEAR



Performance: 10 clinical studies

- An innovative intuitive auto-injector
- Regular exchanges with health authorities



Production capacity of 500ku/y and control of the supply chain



- A resolute business model
- An acknowledged quality approach

ON TRACK FOR THE FIRST MA FILINGS

2021 2020 Marketing 2019 of ZENEO® First MA application filings Pharmaceutical establishment status Extension of the Gray site Production of clinical batches in the spring Launch of bioequivalence studies Signature of licensing agreements



SHAREHOLDER INFORMATION

MARKET DATA AS OF 28 FEBRUARY 2019

Price: €2.53

Number of shares: 17,428,074

Market cap: €44.09m

SHARES

Business sector: Specialty Pharma Main index: EnterNext © PEA-PME 150

Market: Euronext Growth Venue: Euronext Paris (France)

Ticker: ALCJ

ISIN code: FR0011716265

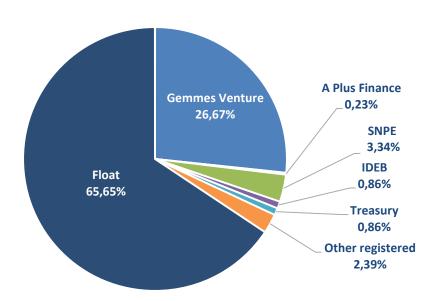
Crossject is eligible for the PEA/PME



FINANCIAL ANALYSTS

- ALPHA VALUE
- CM CIC MARKET SOLUTIONS
- INVEST SECURITIES
- KEPLER CHEUVREUX

SHAREHOLDING STRUCTURE 28 FEBRUARY 2019



CROSSJECT: KEY FIGURES



8 médicaments en portefeuille

dédiés à l'urgence dont 5 en urgence vitale

dédié à une maladie chronique





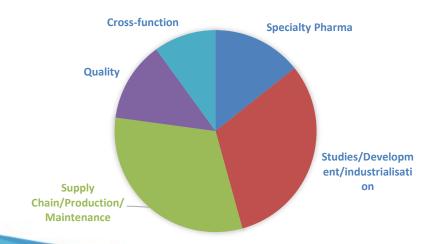






COMMITTED MANAGERS AND TEAMS

- Experienced managers
- More than 70 employees
 - Internalization of key functions
 - Experienced industrial and pharmaceutical profiles
 - Cross-cutting collaboration and longterm operational support





Patrick ALEXANDRE
Founder of Crossject, Chairman of the executive Board, CEO
Supelec Engeneering School
Ex Arcelor, Fournier labs.
Founded Crossject in 2001



Olivier GIRÉ COO Specialty Pharma EDHEC Business School Ex Ipsen, Amdipharm, Exeltis Joined in 2016



Isabelle LIEBSCHÜTZ Chief Quality and Regulatory Officer Doctor in Pharmacy Ex Fournier labs, Solvay, Plasto Santé Joined in 2013

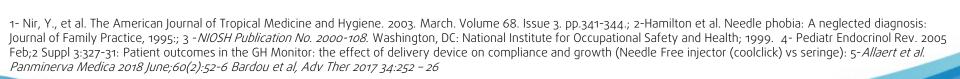


Henri DE PARSEVAL COO Engineering & Industries Ecole Nationale Supérieure d'Arts et Métiers Paris Tech Ex HP, Michelin, Ansaldo. Joined in 2014

KEY BENEFIT FOR BIOTECH PRODUCTS NEXT CROSSJECT GROWTH DRIVERS

ZENEO® Technological breakthrough Pre-filled needle-free single-use auto-injector

- Patient compliance & injectable biologics.
 - Phobia or fear of needles: 8-21% of patients (1);(2)
 - Risk of needle-stick injury or contamination (3)
 - Usability
- Needle free injection: demonstrated benefits for patients:
 - Longer adherence to treatment (4)
 - Better compliance
- Biologics needle-free injection with ZENEO®
 - Easy to use for 97% of people (5)
 - No button: several years of design and test with arthritis sufferers
 - Delivers the full dose in less than 1/10th of a second
 - Successful clinical trials demonstrating ZENEO® bioequivalence (6)
 - Life Cycle Management oriented towards patients and competitive edge





FOCUS ON THE INDUSTRIAL PROCESS



SOURCES

- (1) FDA
- (2) https://www.accessdata.fda.gov/drugsatfda_docs/nda/2014/2057870rig1s000SumR.pdf accessed on 16 march 2018
- (3) Weiner SG, Joyce AR, Thomson HN. Journal of Opioid Management. 2017;13(2):69 7
- (4) http://www.medsafe.govt.nz/profs/datasheet/m/MidazolaminjPfizer.pdf
- (5) http://base-donnees publique.medicaments.gouv.fr/extrait.php?specid=60034085 (3) Brown et al, Annals of Emergency Medicine Vol 67 No.3 2015 (4) Umasunthar T et al. Allergy 2015
- (6) Advances in Therapy, 2017; Bardou et al, Adv Ther 2017 34:252 26
- (7) https://www.reuters.com/article/idUSL8N1C24SS last accessed March 2018
- (8) Tsai G et al. Allergy Asthma Clin Immunol. 2014; 10
- (9) Eurostat. https://ec.europa.eu/eurostat/documents/2995521/6980739/3-10092015-AP-EN.pdf/bc1e347e-9895-4131-9972-4ef718869c22
- (10) http://www.mhra.gov.uk/home/groups/comms-ic/documents/websiteresources/con423091.pdf
- (11) https://seekingalpha.com/article/4234265-patience-will-required-sandoz-adamis-phased-launch-symjepi

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