



SHIFTING THE LINES AGAINST CENTRAL NERVOUS SYSTEM DISORDERS

**EURONEXT GROWTH** 



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A detailed description of the Company's business, financial situation and risk factors relating to the Company and the initial public offering is included in the prospectus of Theranexus (the "Prospectus") which received the approval of the Autorité des marchés financiers (the "AMF") under n°17-545 on October 10, 2017, comprised of the registration document (document de base) registered by the AMF on Septembber 27, 2017 under n°1.17-068 and a securities note (note d'opération) dated October 10, 2017 (which contains, in particular, the summary of the Prospectus) to which you are invited to refer to. Copies of the Prospectus are available on the AMF website (www.amf-france.org) as well as on the Company's website (www.theranexus.com).

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This document contains certain forward-looking statements. These statements are not guarantees of the Company's future performance. These forward-looking statements relate to the Company's future prospects, developments and marketing strategy and are based on analyses of earnings forecasts and estimates of amounts not yet determinable. Forward-looking statements are subject to a variety of risks and uncertainties as they relate to future events and are dependent on circumstances that may or may not materialize in the future. Forward-looking statements cannot, under any circumstance, be construed as a guarantee of the Company's future performance and the Company's actual financial position, results and cash flow, as well as the trends in the sector in which the Company operates, may differ materially from those proposed or reflected in the forward-looking statements contained in this document. Even if the Company's financial position, results, cash-flows and developments in the sector in which the Company operates were to conform to the forward-looking statements contained in this document, such results or developments cannot be construed as a reliable indication of the Company's future results or developments.

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#### Your contacts



Franck MOUTHON

## Co-founder and Chairman and CEO

- Franck Mouthon holds a degree in life sciences from the École Normale Supérieure
- Joined the Life Sciences Department of the French Alternative Energies and Atomic Energy Commission (CEA) in 1995
- Founded CEA spin-off Theranexus in March 2013 with Mathieu Charvériat
- Board member of France Biotech



Mathieu CHARVÉRIAT

#### Deputy CEO and Cofounder

- Mathieu Charveriat holds degrees from the Ecole Polytechnique and Mines ParisTech
- PhD in neurosciences from the Université Pierre & Marie Curie
- Joined the neurosciences research laboratory of CEA in 2009
- Founded CEA spin-off Theranexus in March 2013 with Franck Mouthon



Julien VEYS

**CBDO** 

- Julien Veys holds a master's degree in biochemistry and in information and intelligence management from Aix-Marseille University
- CBDO at Trophos where he played a key role in its sale to Roche laboratories for €470 million
- Joined Theranexus in 2016



Thierry LAMBERT

**CFO** 

- Thierry Lambert holds a degree in business administration from Birmingham University and an MBA from INSEAD
- 4 years of experience in syndicated and corporate finance
- 5 years as Chief Financial Officer for listed companies Naturex and then Safe Orthopaedics
- Joined Theranexus in 2017





- 1 THERANEXUS: SHIFTING THE LINES AGAINST CENTRAL NERVOUS SYSTEM DISORDERS
- A DISTRUPTIVE APPROACH TO THE CENTRAL NERVOUS SYSTEM
- THE CAPACITY TO GENERATE DRUG CANDIDATES TO RESPOND TO UNMET NEEDS WITH THREE DEVELOPMENTS ALREADY UNDERWAY, A BLOCKBUSTER POTENTIAL
- 4 AN AGILE ORGANIZATION WITH CAREFULLY-MANAGED FINANCING NEEDS
- 5 SHORT AND MID-TERM VALUE CREATION BACKED BY AN INNOVATIVE PLATFORM
- 6 TERMS AND CONDITIONS OF THE OFFERING



A French biotech that specializes in the central nervous system

#### A DISTINCTIVE BUSINESS MODEL

- AN ATTRACTIVE PROFILE within the Biotech sector
- Higher likelihood of SUCCESS
- FASTER access to the market...
- ... at LOWER COSTS

#### **AN ESTABLISHED PORTOFOLIO**

- 3 DRUG CANDIDATES\*
- BLOCKBUSTER POTENTIAL
- STRATEGIC MARKETS for pharmaceutical laboratories

<sup>\* 1</sup> in phase II and 2 ready to enter into clinical development

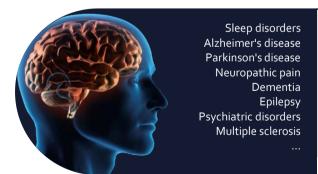




Central Nervous System disorders are one of the 1st causes of disability around the world

More than **one billion** [1] **people** are affected, i.e. **nearly 1 person** in 5

The cost in treating these disorders around the world is estimated at **more** than €2,000 billion per year, i.e. the equivalent of the gross domestic product of a country like France [1]; [2]



Unmet medical needs for many major conditions

- [1] WHO / Neurological Disorders: Public Health Challenges 2015
- [2] Source: Gustavsson et al., Eur Neuropsychophamacology 2011



## INDUSTRY INNOVATION FALTERS WHILE MEDICAL NEEDS ARE DRAMATICALLY GROWING

Growing demand for treatment

Prevalence is correlated to to AGING

UNMEET NEEDS are still very high (non-curative treatments, not all symptoms are relieved)



Value-added offers are increasingly rare

### AGING ARSENAL OF TREATMENTS

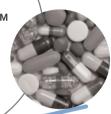
42 generic CNS drugs between now and 2030

### HIGH EXPECTATIONS FROM INDUSTRY

to see medical portfolios renewed

#### LOW SUCCESS RATE

in the development of new CNS drugs







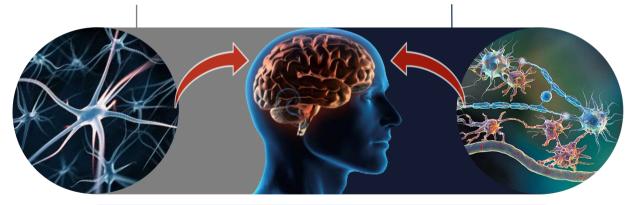
#### CNS DRUGS:

**1** molecule for **1** action on **1** family of cells (**neurons**)



#### THERANEXUS DRUG CANDIDATES:

2 separate molecules combined for 2 actions on 2 families of cells (neurons + glial cells)



**INNOVATION:** COMBINATIONS OF MOLECULES TO OPTIMIZE THE EFFICACY OF STANDARD OF CARE TREATMENTS





#### THERANEXUS PLATFORM: PROPRIETARY, SCALABLE & VERSATILE

#### **GLIAL CELL MODULATOR** CNS DRUGS **DRUG SEEN AS DRUG REPOSITIONED** THE 1ST LINE-TREATMENT **AS A MODULATOR** Condition with a strong unmet need for improved efficacy (with the current arsenal of therapeutics) Theranexus Action Optimization library of CNS drugs on the of the glial 27 glial cell 1<sup>st</sup> line- treatment neuron network modulators for CNS\* conditions **THN** XXX

#### 3 major advantages



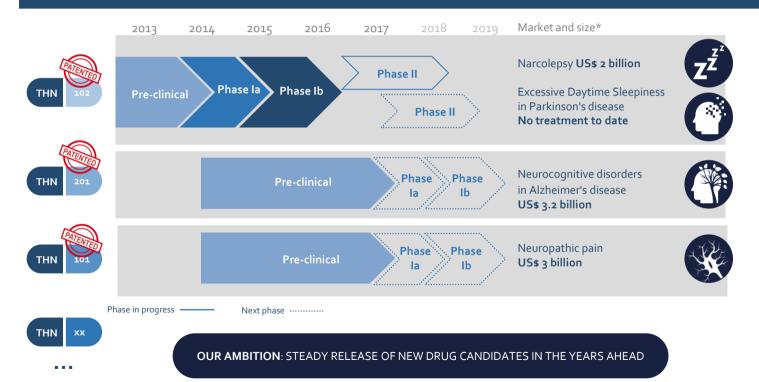




Higher probability of success, greater flexibility and shorter time-to-market



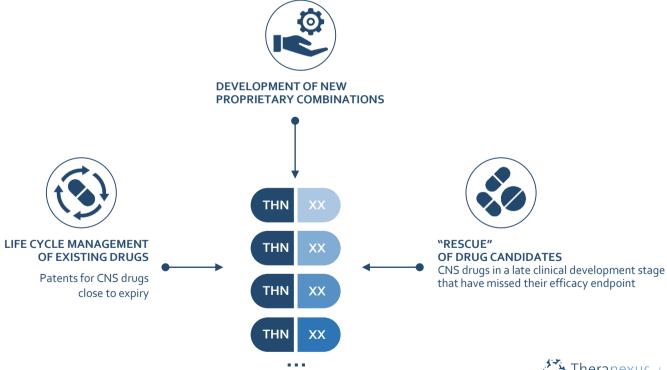
### 3 DRUG CANDIDATES IN JUST 4 YEARS







#### A GLOBAL STRATEGY ADAPTED TO THE NEEDS OF PHARMACEUTICAL COMPANIES





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#### TWO MAJOR FAMILIES OF CELLS IN THE CENTRAL NERVOUS SYSTEM (CNS)

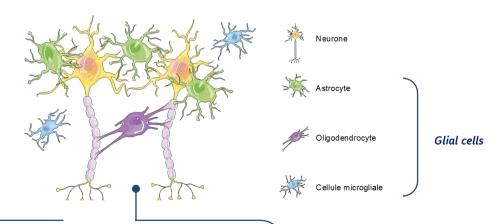
#### **Neurons:**

Cellular components that control our emotions, our mental activity, our memory, our senses, the way we feel pain, and even our motricity, etc.

#### Glial cells:

Able to respond quickly to neuron needs by providing the molecules needed for their metabolism

**Astrocytes** play a key role in neuronal communication



NEURONS DO NOT WORK INDEPENDENTLY BUT AS PARTS OF A CELLULAR CONTEXT





#### A DISCOVERY STEMMING FROM 10 YEARS OF RESEARCH

#### **NEURON CENTRIC APPROACH**

Suboptimal glial network

Limits the efficacy of the CNS drug



Neuronal and glial network with outside stimulation (psychotropic drug) leading to the overdevelopment of the glial network (suboptimal) which limits the efficacy of the drug



Optimal glial network



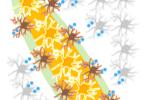
Astrocyte at rest



Neuron activated activated



CNS drug



Theranexus

Glial network close to natural state

Improves the efficacy of the CNS drug

Neuronal and glial network with outside stimulation and connexin modulator (drug candidate)



Zone activated by the psychotropic drug





Connexin modulator

OPTIMIZED SIZE FOR THE GLIAL NETWORK (ASTROGLIAL CELLS) WHICH IS FUNDAMENTAL FOR REGULAR NEURONAL ACTIVITY





Action on neurotransmitter systems

#### TRANSFORMING RESEARCH TO INNOVATION

#### **PRINCIPLE:**

Enhance neuron action with the modulation of glial cells

#### **APPLICATION:**

Combine medication that targets neurons with a medication that optimizes neuroglial interaction

Connexin modulator

CNS drug

(Psychostimulant, antidepressant, anxiolytic, etc.)

Prugs affecting the CNS

Non-neural networks

Astrocyte

Non-neural networks

Neural networks

The modulation of glial connexins optimizes the neuroglial interface to improve the way in which neurons react to CNS drugs

Giaume et al., Nat Rev Neurosci, 2010 Rouach et al., Science, 2008

Picoli et al., J Biomol Screen, 2012 Duchêne et al., Sleep, 2016 Charvériat et al. Front Cell Neuro, 2017

THE CHALLENGE: MAXIMISE NEURON RESPONSE TO EXISTING
DRUGS BY TARGETING THE ENVIRONMENT

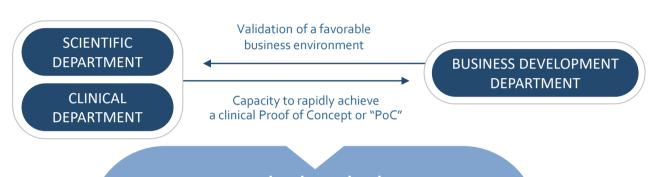




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#### DRUG CANDIDATES SELECTED FOR THEIR CLINICAL AND ECONOMIC VALUE



#### 4 selection criteria

- Patent-free CNS drug as the 1st-line treatment
- Demonstrated efficacy
- Clear room for improvement
- PoC within reach

**IN VIVO SELECTION OF THE BEST COMBINATION** of CNS drugs selected with a glial cell modulator





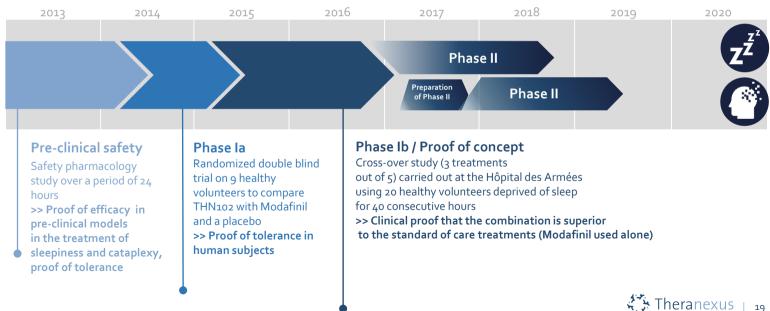
### THN102: A DRUG CANDIDATE FOR 2 CONDITIONS

Modafin	nil FI	ecainide
ZZ Narcolepsy	)	Parkinson's disease
Excessive daytime sleepiness ± cataplexy	SYMPTOMS	Excessive daytime sleepiness
Orphan disease: 300,000+ patients (France, Germany, United Kingdom, Italy, Spain, United States)	PREVALENCE	Close to <b>1 million patients</b> (G7) <b>30 to 50%</b> of patients diagnosed with Parkinson's
Modafinil 4 drugs on the market, none of which fully address both symptoms	STANDARD OF CARE TREATMENT	NONE  No approved treatment to date
US\$ 2 billion  (annual treatment cost/ patient of around US\$ 20k)	MARKET	-
7 drug candidates undergoing clinical trials None of which aim to prove their superiority over the standard of care treatments	RESEARCH	4 drug candidates undergoing clinical trials, all of which only target neurons.



#### THN102: THE MOST ADVANCED COMBINATION OF THE PORTFOLIO



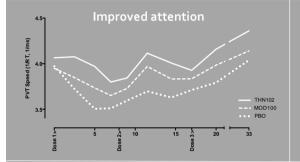




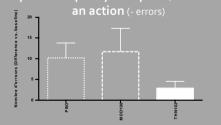
## THN102: SUPERIOR RESULTS OF THE COMBINATION OVER MODAFINIL ALONE AT THE END OF PHASE IB

#### **EFFICACY**

(sleep deprivation) vs placebo and Modafinil



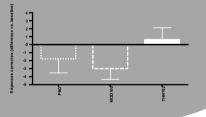
#### Improved capacity to repress/moderate



#### Improved mental flexibility



#### Improved working memory



## TOLERANCE vs Modafinil

	MOD 100 (n=12)	THN102 (n=35)	
Fatigue	83%	70%	
Headache	50%	23%	
Nausea	33%	14%	

SIGNIFICANT IMPROVEMENT IN
AWARENESS AND ATTENTION AND
GOOD TOLERANCE OF THE PRODUCT





#### THN102: 1ST DRUG CANDIDATE IN PHASE II

BEST IN CLASS

Modafinil

Flecainide

IN CLASS

ZŽ

Narcolepsy

#### Launch of the study Phase IIa in 2016

#### Double blind trial to compare 3 treatments:

Modafinil 300 mg/day alone or combined with two doses of FLECAINIDE, 3 and 27 mg/day

Cross-over study over three periods: each patient is randomly given each of the three treatments over three periods of two weeks each

Primary efficacy endpoint: ESS (Epworth Sleepiness Scale)

Trial carried out on 42 narcoleptic patients 20 patients already recruited on 3 sites

Results expected in Q<sub>3</sub> 2018 (Narcolepsy)



Excessive daytime sleepiness in patients diagnosed with **Parkinson's** 

## Regulatory package ready for a start to Phase IIa in Q4 2017

### Double blind trial to compare 2 doses of THN102 to the placebo

Cross-over study over three periods: each patient is randomly given THN102 or the placebo over three periods of two weeks each

Primary efficacy endpoint : ESS (Epworth Sleepiness Scale)

Study carried out on **60 patients** diagnosed with Parkinson's (some of whom possibly in the United States)

Results expected in Q2 2019 (Parkinson's)





#### THN102: DRUG CANDIDATE AS A FIRST-LINE TREATMENT

#### 4 drugs on the market:











**⊅MOD** 

 $\rightarrow$ SOX/PIT. 1 dose / day N/A

Yes

Yes

		Provigil® Modafinil	Nuvigil® ArModafinil	<b>Xyrem®</b> SOX	<b>Wakix®</b> Pitolisant
	Sleepiness	Yes	Yes	Yes	Yes
Marketing authorization label (1) (2) (3) (4)	Cataplexy	No	No	Yes	Yes
	Administration	2 doses / day	1 dose / day	2 doses / night	1 dose / day
	ANSM <sup>(5)</sup> drug database	N/A	- (	Class III	N/A
HAS* efficacy/safety ratio (1) (3) (4)		High	-	High	Moderate
Price in the EU (US\$/year) <sup>(6)</sup> (average in 5 countries)		2,600	00 - 11,850		12,250
Price in the US (US\$/year) (7)		36,000	8,600	120,500	-
Sales peak (US\$ million) <sup>(8)</sup>		2,100		1,108	ND

High **BLOCKBUSTER** POTENTIAL, **EFFECTIVE ON THE TWO MAIN SYMPTOMS** 

Estimate of benefits/risks of products currently on the market compared with the target profile for THN102 and the annual cost of treatments on the market (US\$ - rounded figures) \* French National Authority for Health

- (1) Transparency Commission Recommendation, CT-4626
- (2) FDA Label
- (3) Transparency Commission Recommendation, CT-2921
- (4) Transparency Commission Recommendation, CT-14970
- (5) ANSM drug database at 15/06/2017, includes GHB and its salts
- (6) France: CNAMTS; UK: BNF; Italy: AIFA; Spain: MSSSI; Germany: Apoteke
- (7) US Rx List internet drug index
- (8) Jazz Pharmaceuticals Investor Presentation of o6/o6/2017





## THN201 & THN101: TWO NEW MAJOR CONDITIONS TARGETED WITH VERY HIGH INDUSTRIAL STAKES

THN 201		THN 101
Neurocognitive disorders linked to Alzheimer's disease		Neuropathic pain
Impairment of memory, judgment, orientation	SYMPTOMS	Permanent background pain, with occasional stabbing pains, burning sensation and twinges
15 million patients in 2015 (G7) 19 million between now and 2030 45% undiagnosed patients	PREVALENCE	<b>70 million patients</b> (Europe, US, Japan)
DONEPEZIL	STANDARD OF CARE TREATMENT	AMITRIPTYLINE
US\$3.2 billion (annual cost of treatment/patient US\$4-5k)	MARKET	US\$3 billion (annual cost of treatment/ patient US\$3-4k)
23 drug candidates in clinical trials	RESEARCH	32 drug candidates in clinical trials



#### THN201 & THN101: TWO NEW MAJOR CONDITIONS TARGETED WITH VERY HIGH INDUSTRIAL STAKES

#### **DONEPEZIL** | MEFLOQUINE



#### Target profile:

Label for neurocognitive disorders linked to Alzheimer's

Performance target: THN201 versus DONEPEZIL:

- Improved cognitive function
- Delayed need for institutionalization

#### AMITRIPTYLINE | MFFLOOUINE



#### Target profile:

Label for neuropathic pain

Performance target: THN101 versus AMITRIPTYLINE:

- Reduction in pain intensity
- Increase in the number of patients experiencing a 50% reduction in pain
- Better tolerance profile

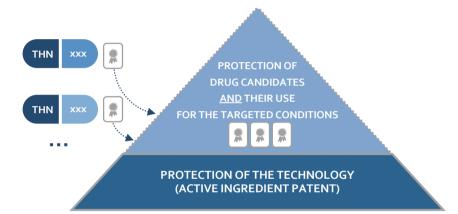
PROOF OF SUPERIORITY IN PRE-CLINICAL MODEL AND IN TERMS OF **TOLERANCE** 

2017 2018 2019 2020 Pre-clinical Phase Ia Phase Ib / PoC

CLINICAL PROOF THAT THE COMBINATION IS SUPERIOR TO THE STANDARD OF CARE TREATMENT TARGETED IN Q2 2019



## INNOVATION THAT IS FIRMLY PROTECTED BY AN INTELLECTUAL PROPERTY STRATEGY



	Family of patents 1 (platform patent)	Family of patents 2	Family of patents 3	Family of patents 4
Products	Anti-connexin agent + psychotropic molecule	THN201 Dementia	THN102 Narcolepsy / Parkinson's	THN101 Neuropathic pain
Expiry date	2029	2032	2034	2036
Geographic regions targeted		<b>○</b> • ■ *	<u> </u>	

FREEDOM TO EXPLOIT DRUG CANDIDATES
FREEDON TO DEVELOP NEW COMBINATIONS





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### A COMPLEMENTARY TEAM IN AN AGILE ORGANIZATION



Franck Mouthon CHAIRMAN & CEO









Werner Rein CMO









Mathieu Charvériat CSO









Julien Veys CBDO





Thierry Lambert CFO







### High profile partners



















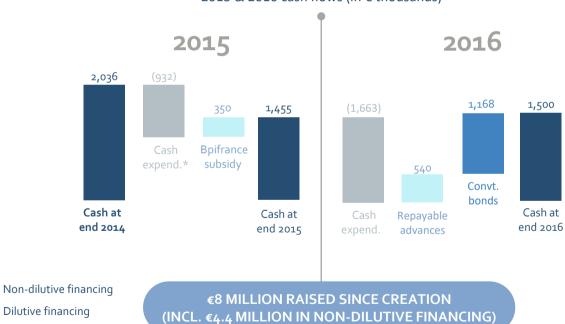
11 EMPLOYEES DEDICATED TO KEY FUNCTIONS





#### **CONTROLLED CASH EXPENDITURE**

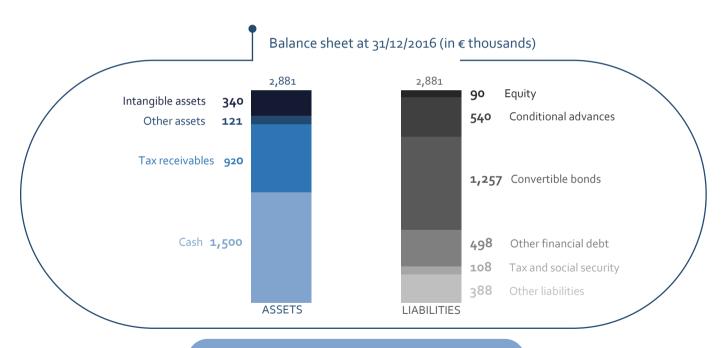
#### 2015 & 2016 cash flows (in € thousands)



<sup>\*</sup> Net flows from operations + Net flows from investments + Financial interest paid + Repayments of loans



#### **BALANCE SHEET STRUCTURE**



A HEALTHY AND ROBUST FINANCIAL STRUCTURE





#### **GOVERNANCE & SHAREHOLDERS**

#### **BOARD OF DIRECTORS**



Franck Mouthon
Theranexus, Chairman and CEO
Theranexus



Mathieu Charvériat
Theranexus, Deputy CEO





**Dominique Costantini** Independent director





Luc-André Granier Independent director





Auriga, represented by Florian Denis



CEA-Investissement, represented by Celia Hart



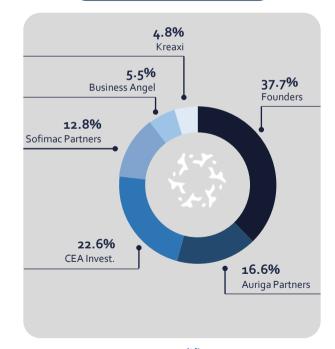
SOFIMAC partners capital investissement

**Sofimac Partner**, represented by **François Miceli** 



★ kreaxi
Kreaxi, represented
by Gwenaël Hamon
(non-voting member)

#### **SHAREHOLDERS**







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## STRONG INTEREST AMONG INDUSTRY PLAYERS FOR THE FIRST 3 CONDITIONS TARGETED

CONDITION	DATE	SELLER	BUYER	PROFILE	DEVELOPMENT STAGE	UP FRONT (US\$ m)	MILESTONES (US\$ m)	ROYALTIES (US\$ m)
Narcolepsy	2014 2013	Aerial Concert	Jazz Jazz	NCE <sup>[1]</sup> LCM <sup>[2]</sup>	Phase II Pre-clinical	125 5	272 115	NC NC
Neuropathic pain	2015 2015 2012	Convergence Spinifex Concert	Biogen Novartis Avanir	NCE NCE LCM	Phase II Phase II Phase I	200 200 NC	475 500 200	NC NC NC
Alzheimer's disease	2016 2013 2012	Chase Pharma Lundbeck Adamas	Allergan Otsuka Forest	Combination NCE Combination	Phase I/II Phase II Phase II	125 150 60	875 675 95	NC NC NC
Other neurological disorders	2014	Avanir	Otsuka	Combination	Market	3,500		

TURNING POINT IN VALUE AT THE END OF PHASE II (BETTER RATIO OF DEVELOPMENT COSTS TO IMMEDIATE AND SUBSEQUENT REVENUES)



<sup>[1]</sup> New Chemical Entity

<sup>[2]</sup> Life Cycle Management



#### THN102: A FIRST SOURCE OF VALUE CREATION





A DRUG CANDIDATE FOR 2 CONDITIONS

- + POTENTIAL PARTNERS
- + PROBABILITY OF SIGNING AN
- + PROBABILITY OF MAXIMISING VALUE

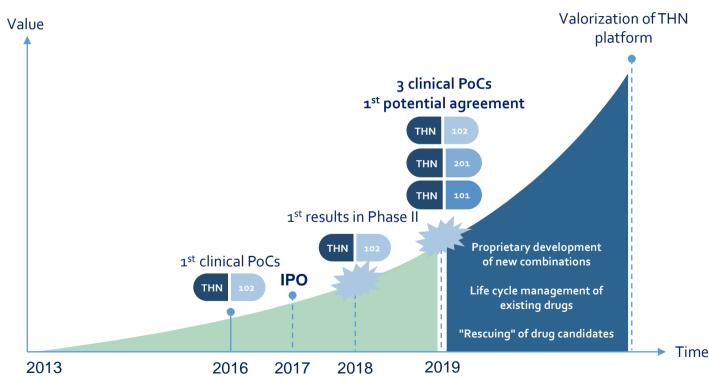
OPPORTUNITY FOR STRONG VALUE **CREATION BETWEEN NOW AND 2019** 

Boehringer Ingelheim





## A DUAL SOURCE OF VALUE CREATION IN THE SHORT AND MEDIUM TERM





#### **5 REASONS TO BECOME A SHAREHOLDER**

- A PATENTED unique approach to the CENTRAL NERVOUS SYSTEM in the treatment of CNS disorders
- A TECHNOLOGICAL PLATFORM capable of delivering several "best in class" candidates
- A clinical development profile to optimize the COMBINATION OF SPEED / PROBABILITY OF SIGNIFICANT SUCCESS
- A 1<sup>st</sup> drug candidate (**THN102**) with BLOCKBUSTER market potential
- A company structured around EXPERT PROFILES and supported by TIER ONE SHAREHOLDERS





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## KEY CHARACTERISTICS OF THE OFFERING

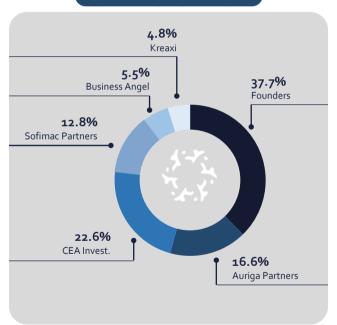
MARKET / CODES	Euronext Growth (Paris)
	Codes: FR0013286259 - ALTHX
PRICE RANGE	Between €14.00 and €18.80 per share
TERMS OF THE OFFERING	Public placement in France (OPO) & Global Placement reserved for institutional investors  1,097,560 new shares to be issued as part of a cash capital increase  1,262,194 new shares in the event of the full exercise of the Extension Clause  1,451,523 new shares in the event of the full exercise of the Extension Clause and Over-allotment Option
SIZE OF THE OPERATION	€18 million capital increase based on the offer's midpoint price of €16.40 per share, which may be increased to €23.8 million in the event of the full exercise of the Extension Clause and Over-allotment Option
SUBSCRIPTION COMMITMENTS	<ul> <li>€10.9 million in subscription commitments:</li> <li>€5.4 million in subscription commitments at any price from historical shareholders</li> <li>€4.0 million in subscription commitments from Financière Arbevel capped at a maximum price of €18.80 per share</li> <li>€1.5 million in subscription commitments from Alto Invest capped at a maximum price of €15.52 per share</li> </ul>
ABSTENTION, OWNERSHIP AND HOLDING COMMITMENTS	Franck Mouthon and Mathieu Charvériat, co-founders: holding commitment of 365 days  Historical shareholders: holding commitment of 365 days  Theranexus: abstention commitment of 180 days



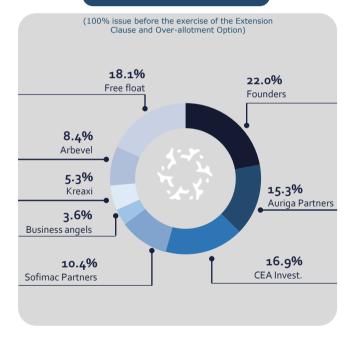
Approval of the Prospectus by the AMF	Tuesday 10 October 2017
Opening of subscription period	Wednesday 11 October 2017
Closing of subscription period	Tuesday 24 October 2017 at 5pm
Closing of Global Placement Setting of price and allocation of shares	Wednesday 25 October 2017
Settlement-delivery	Friday 27 October 2017
Start of trading on Euronext Growth	Monday 30 October 2017
Expiry of exercise period for Over-allotment Option / End of the stabilization period (where applicable)	Friday 24 November 2017



#### CAPITAL BEFORE OPERATION\*



#### **CAPITAL AFTER OPERATION**





### FINANCIAL INTERMEDIARIES AND ADVISORS

Consulting firm

Global Coordinator, Lead Manager, Book Runner

Co-Lead Manager, Book Runner







**Auditors** 



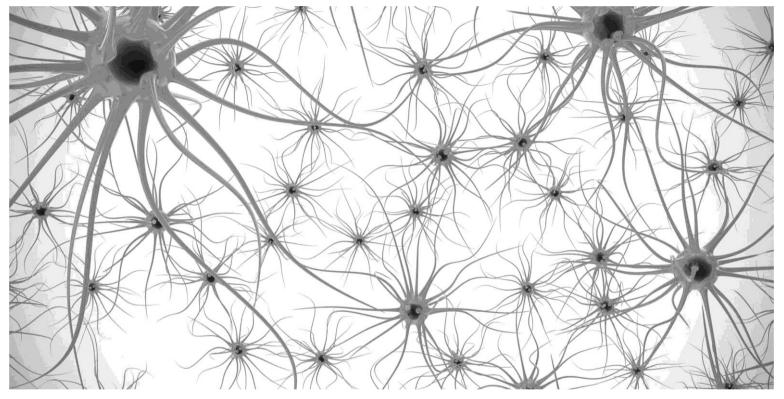














IPO ON EURONEXT GROWTH



### **Epworth Sleepiness Scale (ESS)**

Situation Please tick box	No chance of dozing	1 Slight chance	2 Moderate chance	3 Definitely would doze
Sitting and reading		F	F	
Watching TV				
Sitting inactive in a public place (e.g. Theatre or a meeting)			<b>S</b>	
As a passenger in a car for an hour without a break				
Lying down to rest in the afternoon when circumstances permit				
Sitting and talking to someone				
Sitting quietly after lunch without alcohol		- ATT	- ACT	
In a car, while stopped for a few minutes in traffic				

- Scored from o (no sleepiness) to 24 (highly severe sleepiness)
- Below 8: you have a healthy level of daytime sleepiness.
- From 9 to 14: you have a sleep debt, you need to improve your sleep hygiene.
- More than 15: you have a high level of excessive daytime sleepiness. You need to improve your sleep hygiene and consult your doctor for further medical help
- The average score for an untreated patient with narcolepsy is 18



## THN102 competitive landscape: 7 drug candidates for the treatment of narcolepsy undergoing clinical trials

<b>→</b> Ź				
Company	Molecule	Brand	Dev. stage	Mechanism of action
Jazz pharma	JZP-110	-	P <sub>3</sub>	NA / DA recapture inhibitor
Jazz pharma	JZP-258	-	P <sub>3</sub>	Xyrem® with reduced sodium content
Avadel	FT218	-	P <sub>3</sub>	Xyrem with sustained release
Taisho pharma	TS-091	-	P <sub>2</sub>	HIS H <sub>3</sub> receptor inverse agonist
Balance Tptx	BTD-001	-	P <sub>2</sub>	GABA-A blocker
Jazz pharma	JZP-507	-	P1	Xyrem® with reduced sodium content
Jazz pharma	JZP-386	-	P1	Deuterated Xyrem

Principal drugs and drug candidates indicated for the treatment of narcolepsy (source: informa Medtrack – June 2017) – Com: marketed; GABA: qamma-aminobutyric acid; HIS: histamine; NA: noradrenaline; DA: dopamine; 5HT: serotonin.

Theranexus



# THN102 competitive landscape: 4 drug candidates for the treatment of excessive daytime sleepiness in Parkinson's disease undergoing clinical trials

Company	Molecule	Brand	Dev. stage	Mechanism of action
Jazz pharma	JZP-110	-	P <sub>2</sub>	NA / DA recapture inhibitor
Benevolent Al	Bavisant	-	P <sub>2</sub>	HIS H <sub>3</sub> receptor agonist
Novartis	LML134	-	P1	HIS H <sub>3</sub> receptor inverse agonist
Eli Lilly	LY3154207	-	P1	RD1 allosteric modulator

Principal drugs and drug candidates indicated for the treatment of excessive daytime sleepiness in Parkinson's disease (informa Medtrack – clinicaltrials.gov July 2017); HIS: histamine; NA: noradrenaline; DA: dopamine; D1R: dopamine receptor D1.

NO PROJECTS AT A LATER STAGE THAN THN102 COMBINATIONS AND DRUG CANDIDATES THAT ONLY TARGET NEURONS

