Ipsen

Strategy Roadshow - Société Générale

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The implementation of the strategy has to be submitted to the relevant staff representation authorities in each country concerned, in compliance with the specific procedures, terms and conditions set forth by each national legislation.

Executive summary





Ipsen's 2020 aspiration: Become a global leader in targeted debilitating diseases

Increase Focus

Invest to Grow

Leverage Footprint

A market-oriented franchise model...

...driving an R&D patient centric organization focused on core platforms, peptides and toxins.

More than double revenues¹

...and more than triple EBIT2

Introduction



Quick overview of the pharma context





Over the past decade, the pharmaceutical environment has significantly toughened

Cross industry topline pressure...

- Increased primary care pressure
- Intensifying competition in specialty care globally
- Patent cliff 2010-2014

...fewer NMEs and higher R&D costs...

- R&D spend more than doubled since 1997² with lower R&D productivity
- 50% less New Molecular Entities approved per year vs.1997²
- Decrease to 50% from 70% of Ph III success rate¹

...and more complex market access

- Overall pressure on new drug prices
- Increasing hurdles
- In EU, regional decision making, tenders...

Increasing pressure across P&L

7

1 Nature Review Drugs Discovery, Feb 2011



Growth potential lies in specialty care and global footprint

Rise of specialty care

- High unmet medical needs addressed by Specialty Care provide growth:
 - Global 2010 growth of 6.3%1 (vs. 2.3%1 for global primary care growth)
- Lower exposure to substitutable generics

Rise of emerging markets

- Steady development of emerging countries :
 - Improving healthcare coverage
 - Increasing drug purchasing power
- Pharmerging markets to grow ~15%¹ CAGR (2010-2015)

US to remain N°1 market

- US market to remain by far the largest market (with ~30%¹ of WW market in 2015)
- Expected contribution to global growth to remain important: 11%¹ between 2010 and 2015

First assessment





Over the last decade, Ipsen has succeeded in adapting to a fast changing environment...

Evolution of Ipsen's sales profile



Ipsen is ideally positioned to benefit from current market trends



... building key assets, creating a unique profile in 2011

Specialty care

Valuable specialty care portfolio

- Growth potential with Somatuline®, Dysport® & Decapeptyl®
- New products: FIX¹, OBI-1¹, Tasquinimod

Talents

- Strong top and middle management
- Entrepreneurial subsidiaries

Innovative & differentiated platforms

- Peptides
- Toxins

Open model culture

 Partnerships from research to marketing

Strengths Comparatively low exposure to patent expiry

- Specialty Care
- Life cycle management

Partnerships from research to

International Footprint

Pharmerging & US

Ipsen's key

Geographical reach

11

NOTE 1 : Inspiration Biopharmceutical



During the strategic review, several key questions were identified...

Are we focused enough?

Have we fully leveraged our current portfolio's potential?

Is our early stage compound portfolio sufficient to sustain long term growth?

How do we improve return in the US?

How do we address the primary care situation?



... and answered

| FOCUS | There is still significant scope to enhance focus |
|-------------------|---|
| FULL POTENTIAL | We can further leverage the potential that lies in Dysport [®] and Somatuline [®] |
| PIPELINE | We need to replenish our early stage pipeline for the out years |
| US | Focus on our core products Dysport® and Somatuline® |
| FRENCH PC | All partnering opportunities are being assessed |



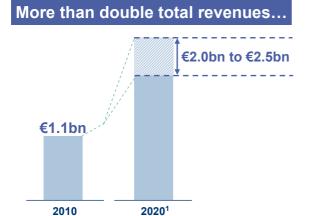
Long term Ambition

"To become a global leader in targeted debilitating diseases"

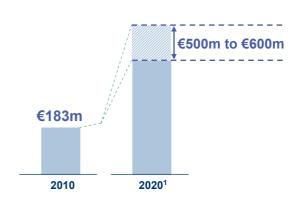




Ipsen's 2020 ambitions



...and more than triple EBIT²...



... over 2 periods:



2011 - 2015 Investment

- Label extension
- Leveraged geographies
- Increased commercial investments
- Inspiration option



2016 - 2020 Solid growth

- Dysport® and Somatuline® on track for full potential
 New products: Inspiration option, tasquinimod, others...
 R&D productivity/ pipeline delivering
- US platform to materialize potential

15

NOTE 1 : 2020 figures include contribution of Inspiration portfolio and are set at constant FX
NOTE 2: prior to purchase accounting recordings and non recurring elements



To achieve our ambition, we need to...



- Capture Ipsen's assets full potential, on a limited number of:
 - technological platforms (R&D focus)
 - Disease areas (commercial focus)

... Invest to grow

- Enhance leadership in technological platforms (R&D and manufacturing)
- Grow market share in selected disease areas
- Build pipeline
- R&D productivity

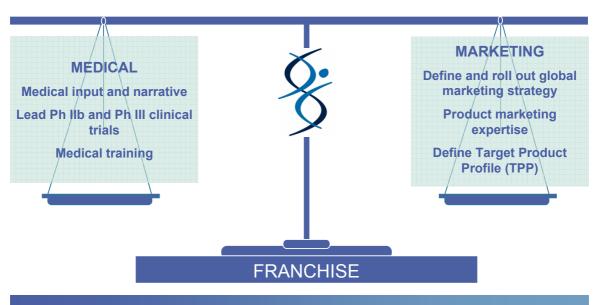
... Leverage footprint

- Leverage Ipsen's presence to broaden access to:
 - US specialty care growth reservoir
 - Accelerate Pharmerging market penetration in both primary and specialty care





Franchise will bring commercial reality at the center of drug development



Countries are responsible for P&L performance







Increase market focus on 4 franchises, driving innovation

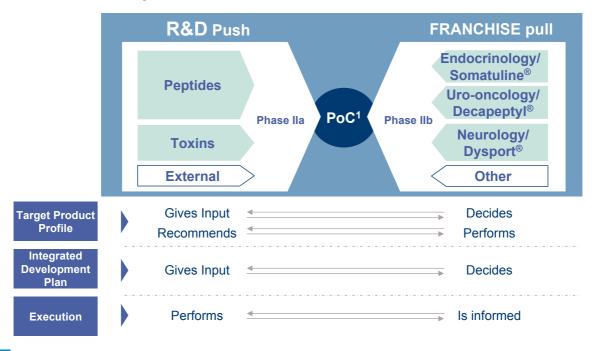
| | Research | Early dev. (end of Phlla) | Late dev. (Phllb & PhIV) | Manufacturing | Operations |
|----------------------------|------------------|------------------------------|-----------------------------|------------------|------------------|
| Endocrinology/ Somatuline® | \checkmark | ✓ | ✓ | ✓ | ✓ |
| Neurology/ Dysport® | ✓ | ✓ | ✓ | ✓ | ✓ |
| Uro-oncology/ Decapeptyl® | Ipsen or Partner | Ipsen or Partner | Ipsen or Partner | Ipsen or Partner | ✓ |
| Hemophilia | Partner | Partner | Ipsen or Partner | Ipsen or Partner | Ipsen or Partner |

Primary care and Short Stature in a commercial optimization strategy





An integrated R&D "push-pull" model to fulfill patient/commercial requirements











Ipsen's path to INCREASED FOCUS

Key projected priorities

Marketfocused franchise model

- Implementation of four market-driven franchises:
 - 2 franchises covering full value chain supporting Somatuline[®] and Dysport[®]
 - 2 franchises primarily focused on late stage development and commercial performance around Decapeptyl[®] and Hemophilia

Projected deprioritizations

 Short Stature in commercial optimization perspective. Ipsen will explore all options to maximize value while meeting its obligations to patients and partners. It will be managed directly by regions and countries

- Focus R&D on core platforms, peptides and toxins
- Merge "R" and "D" departments, increase productivity and improve time to market
- Streamline R&D footprint

- Proteins and small molecules
- Short Stature and non-prostate cancer oncology R&D programs are no longer in Ipsen's strategic priorities
- R&D activities at Barcelona site to close

Market driven R&D





INVEST TO GROW: Ipsen's way forward

Invest to grow over 2011-2015

Focused investment effort to further develop Somatuline® and Dysport®

- Dysport® Ph II in urologyDysport® Ph III Next Generation
- Somatuline® Ph III in NET

 Functioning in the US

 Non functioning WW

Increase investment effort in growth markets: pharmerging, US

Stronger commercial push on key products to catch up vs. competition

Allocate reprioritized R&D resources to support highly differentiated peptide and toxin technological platforms

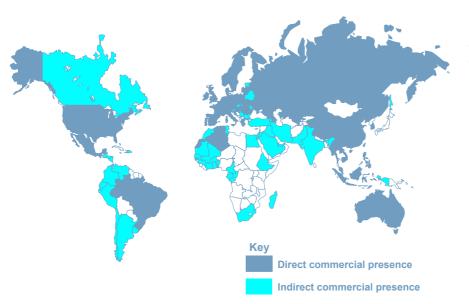
> Enhance leadership in technological platforms Grow market share in selected disease areas







An already existing extensive commercial reach...



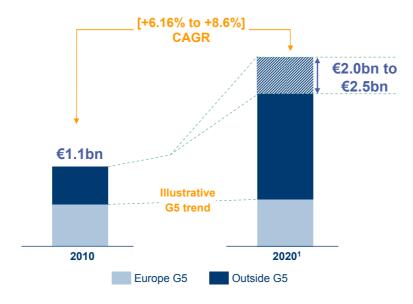
Sales recorded in 115 countries:

- Direct commercial presence in 49 countries
- Commercial presence through partners in 66 countries





... that will provide most of the growth in the future



Evolution of Group sales outside G5

23

NOTE 1: 2020 figures include contribution of Inspiration portfolio and are set at constant FX





LEVERAGE FOOTPRINT as a major growth driver

Leverage Footprint

- Allocate additional commercial efforts to Somatuline® and Dysport® to capture sales full potential
- Inspiration opportunity
- Move teams to the East Coast

Projected deprioritizations

 Apokyn® and Increlex® are no longer in Ipsen's strategic priorities. The Group will explore all options to maximize value while meeting its obligations to patients and partners

Pharmerging

US

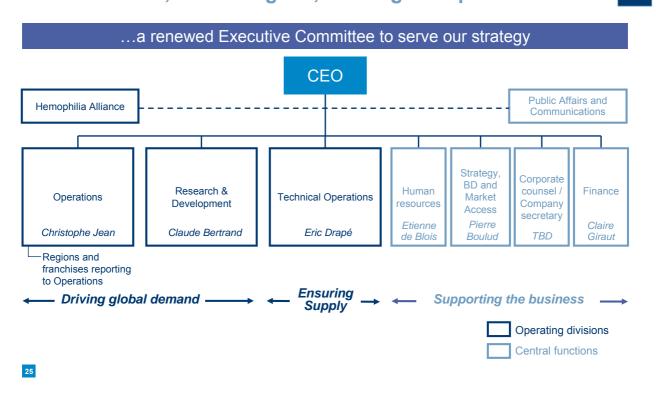
- Further leverage lpsen's profitable emerging market base:
 - Leverage primary care portfolio
 - Expand specialty care portfolio to most existing countries
- China:
 - Support strong Decapeptyl® growth in China
 - Register Dysport[®] and Somatuline[®]

24





Increase focus, Invest to grow, Leverage footprint...







In short...





Become a global leader in targeted debilitating diseases

Increase Focus

Invest to Grow

Leverage Footprint

A market-oriented franchise model...

...driving an R&D patient centric organization focused on core platforms, peptides and toxins.

2020 ambition

More than double revenues¹

...and more than triple EBIT²

Endocrinology/ Somatuline®





Endocrinology/ Somatuline® franchise

Acromegaly

- Pituitary disorder triggering excess GH secretion and leading to gigantism and growth of soft tissues
- Prevalence: 60 per 1 million
- 5-10 years less life expectancy ¹
- North America: ~15,000 patients
- Europe: ~15,000 patients
- Ipsen geographies: Europe, RoW and North America



Neuro Endocrine Tumors (NET)

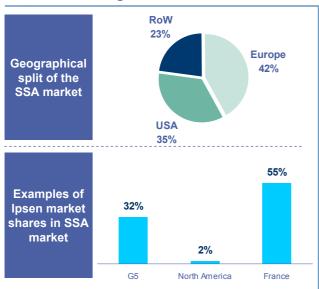
- Arise from cells with both neuronal and endocrine origins¹
- Prevalence: 25 to 50 per 1 million
- Prevalence x5 in 30 years² in the US
- Can arise from almost any organ, most commonly GI tract, pancreas, and lung
- 5 years of survival for carcinoid tumors in 60% of cases³
- Ipsen geographies : Europe, RoW ex North America

Franchise territory: Adult endocrinology, NET



Global Somatostatin Analog (SSA) market in 2010 : ~ 1.1 billion euros...

Q4, 2010 market figures



- 2010 SSA market: ~€ 1.1bn
- Solid SSA market growth (+9%¹ in 2009 and +18%¹ in 2010)
- A fairly balanced geographical split between Europe (42% of total sales), the US (35%) and the RoW (23%)
- Somatuline[®], an established product in Europe both in Acromegaly and in NET with 55% SSA market share in France and 32% SSA market Share in G5
- Ramping up acromegaly sales in the US with only 2.4% SSA long acting market share in 2010

Note 1: Actual (Somatuline® + Sandostatin) reported sales Others : based on company reported sales ; IMS MIDAS MAT Q4 2010

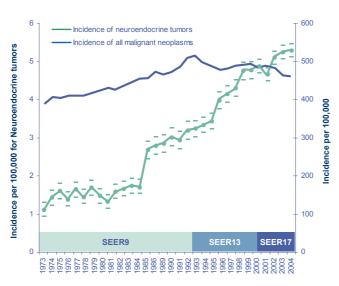




... exceeding 1.6 billion euros in 2020, driven by NET

NET incidence over 30 years

Incidence per 100,000 for NET between 1973 - 2004²



Steady 3.8%¹ CAGR until 2020

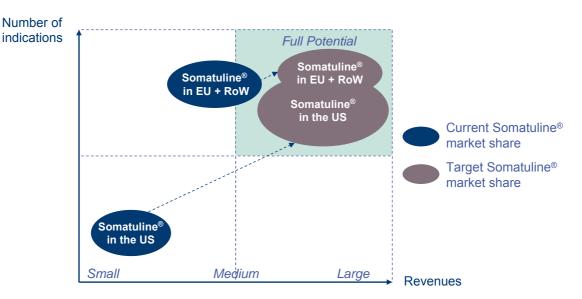
2020 SSA market: ~€1.6bn¹ (+ 45% or 3.8% CAGR)

Growth in the SSA market mainly driven by:

- NET
 - Studies suggest that NET incidence has been growing rapidly over the past several decades, particularly in the US
 - Increased awareness of NETs results in a wider availability of improved diagnostic techniques
- The US
 - +4.6%¹ expected market growth in the US between 2010-2020 (world most solid growth)



Great potential lies ahead for Somatuline[®]...



... while SSA market is expected to grow 3.8% CAGR until 2020



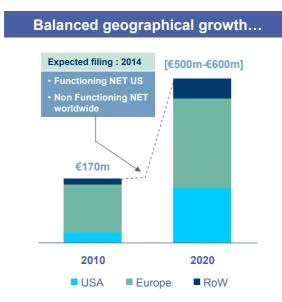


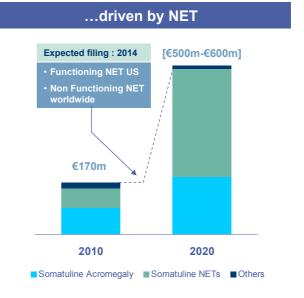
Commitment across the full value chain

Competitive landscape Core: peptides **Technological platform Barrier to entry High** – pre filled syringe & long acting peptide Well characterized - 1 major competitor, targeted Competition combo therapies emerging Market size/ Growth **Large/ Solid** – 3.8% CAGR until 2020 (€1.59bn) **Market share** Room to grow: Somatuline® 2nd player **Geographies US** + **Pharmerging** potential – solid growth in Europe Significant – elements of differentiation to be further Somatuline® competitive adv. Somatuline® growth Strong - 2 ongoing PhIII in NET (non functioning WW + functioning US)... potential



Somatuline® ambition: 10% to 15% CAGR* until 2020



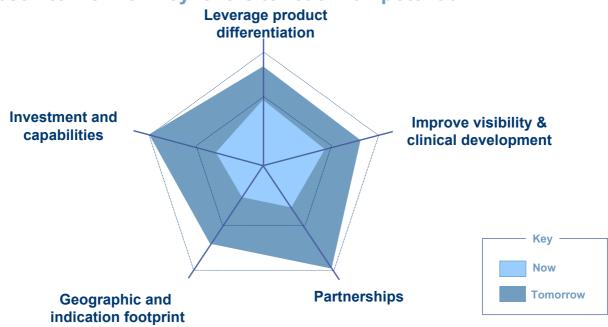


35

* two sequences : one invest; two return



Ipsen to work on key levers to reach full potential



NET and the US: two main growth drivers



New additional elements of differentiation

Increased extended dosing interval worldwide

- Approved in the US in March 2011
- From one injection every 4 weeks (60-90mg) to every 6-8 weeks (120mg)
- Increased comfort for the patients
- Economic benefit

New device

- Retractable needle to ensure full dose release
- Optimal safety for hospital care practitioners/ patients
- Health economic benefits related to absence of clogging and no need for reconstitution

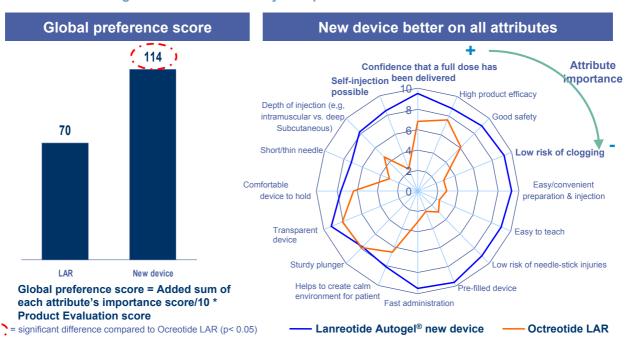






Somatuline® New Device: preferred by Nurses

Somatostatin Analog Nurse Preference Study - 1st publication





Improve share of voice & clinical development



Leverage clinical and safety data

Enhance differentiation elements

New campaign:

"Start right, stay right"

Enhance collaborations with medical and scientific communities

Enhance services to physicians, nurses, patients and payers





Partnerships to explore new treatment paradigm

Innovative partnership with Pfizer Europe in Neuro Endocrine Tumors (NET)

Medical education initiative kicked off at ENETS (joint symposium on March 11th 2011 in Lisbon)



Build upon respective best-in-class position to develop medical education on gastro-entero-pancreatic NET (GEP NET) management

Drive guidance on patients profiles who would benefit most from both agents



Partnerships to increase penetration in emerging markets

Promotion agreement with Sanofi in Latin America

IPSEN Innovation for patient care

SANOF

Working together for patients

- Long term agreement between Sanofi and Ipsen in emerging markets focused on Latin America
- Started in January 2009
- Potential extension in other geographies

Promotion agreement with Invida¹ in Asia







- New Geographical footprint for Ipsen's Specialty Care Portfolio in Endocrinology (and Oncology)
- Started in April 2010

- Accelerates penetration of Ipsen's products
- Leaves
 Ipsen's
 options
 open for the
 future



Note 1: JV between Zuellig - Quintiles and Temasek



New indications: Functioning NET in the US and Non Functioning NET worldwide

Functioning NET for US label

- Recruitment target: 100 patients
- Global recruitment status on target for completion end of 2012
- Carcinoid syndrome initially slow to recruit due to trial design and ongoing competitive trials
- 12 countries planned (US + 11 ROW countries), 66 sites (56 Row + 10 US)

| • USA | | | | | |
|---|---|--|--|--|--|
| BrazilCroatiaCzech Rep.IndiaLatviaPoland | RussiaSerbiaSouth AfricaTurkeyUkraine | | | | |

Non Functioning NET worldwide - CLARINET

- RECRUITMENT COMPLETED end of April 2011
- 200 patients accrued (45 centers in 14 countries)
 - Austria

Denmark

France

India

Germany

- Italy
- BelgiumCzech Rep.S
- PolandSlovakia
 - Spain
 - Sweden
 - UK • US

Somatuline®, potentially the only SSA with functioning and non-functioning NET label

SIPSEN Innovation for patient care

Somatuline[®] Autogel 2010 footprint and indications

| Geography/ Indication | Europe | US | China | Brazil | Russia |
|---------------------------|----------|----|-------|-------------------------------|-------------------------------|
| Acromegaly | ✓ | ✓ | - | But no national reimbursement | But no national reimbursement |
| Functioning NET | √ | - | - | But no national reimbursement | But no national reimbursement |
| Non functioning NET | _ | • | _ | • | • |







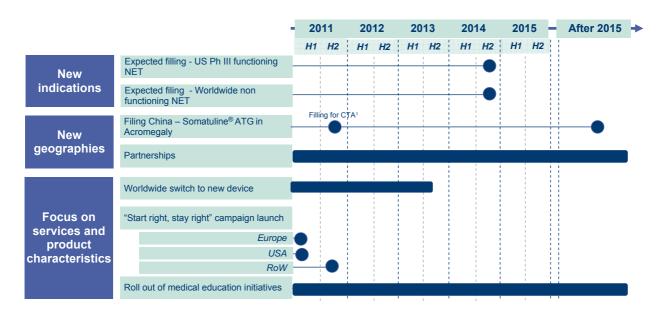
Somatuline® Autogel 2020: a globalized reach

| Geography/ Indication | Europe | US | China | Brazil | Russia |
|---------------------------|--------------|----|----------|-------------|--------------|
| Acromegaly | \checkmark | ✓ | √ | √ | √ |
| Functioning NET | √ | ✓ | • | ✓ | ✓ |
| Non functioning NET | √ | ✓ | - | √ | √ |
| ' | | | | √ Ip | sen presence |

LatAm and Asia covered through partnerships



Somatuline[®] milestones



45

NOTE 1: CTA or filing for Clinical Trial Authorization

Neurology/ Dysport®

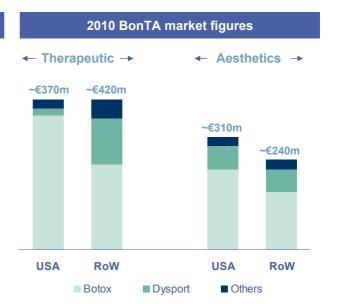




A 2010 botulinum toxin market in excess of 1.3 billion euros

Dysport® market metrics

- 2010 Botulinum toxin market : ~€1.35bn¹
- The US represent north of 50% of the market
- Therapeutic indications represent 58% of the market
- Dysport[®], a solid second player
- Dysport® recently launched by Ipsen in the USA (November 2009) with a single medical indication (cervical dystonia) and by Medecis in aesthetics (Glabellar lines)



Note 1 : Internal company data

Source: Ipsen analysis



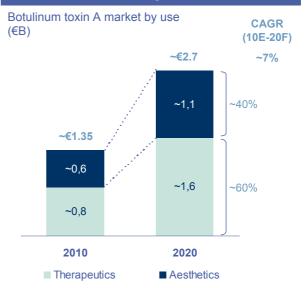


Botulinum toxin market expected to grow by ~7% p.a. to 2.7 billion euros in ten years

US to remain half of 2020 WW market ...

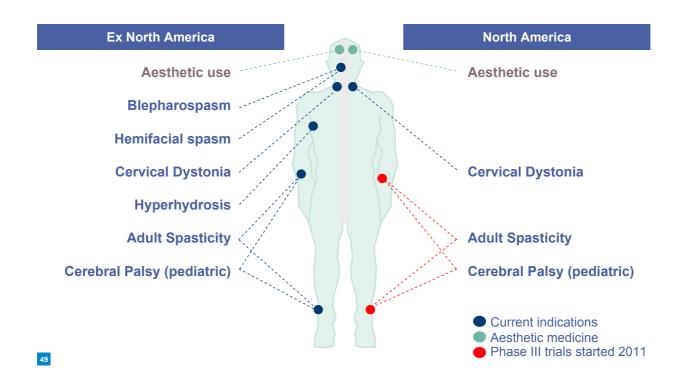
Botulinum toxin A market by geography CAGR (€Bn) (10E-20F) ~€2.7 ~7% ~1,3 ~50% ~€1.35 ~0,7 ~50% ~1,4 ~0,7 2010 2020 **US** RoW

...with split between therapeutics and aesthetics remaining stable



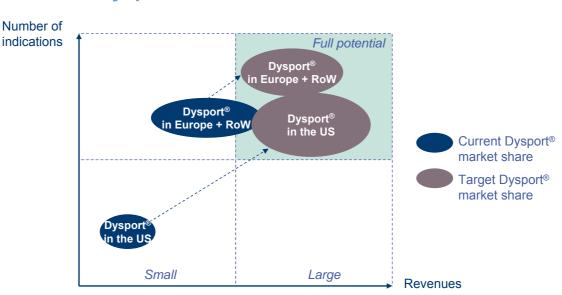


Room for new indications in North America





Full potential of Dysport® lies ahead...

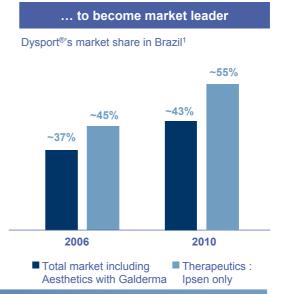


... and BonTA market is expected to grow 7% CAGR until 2020



Brazil, success built on focus and strong selling fundamentals





In Brazil, Dysport® gains market share and expands the market with a 2006-10 CAGR of 13% vs. 10% for the market (Botox®, Xeomin® and Prosigne®)



Note 1: Internal data



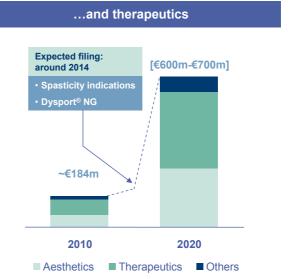
Commitment across the full value chain

Competitive landscape Toxin Technological Core to Ipsen platform **Barrier to entry** High Well characterized - 2 major competitors, Competition more to come – strong Aesthetics alliances Market size/ Growth Large/ Solid - 7% CAGR until 2020 **Dysport® Market share** 2nd after Botox® - room to grow Geographies US and emerging markets potential **Dysport® competitive adv.** At par with competitors Dysport® NG competitive adv. **Further differentiation with Next Generation** Dysport® growth potential 4 ongoing PhIII in spasticity...



Dysport® ambition: 10% to 13% CAGR* until 2020



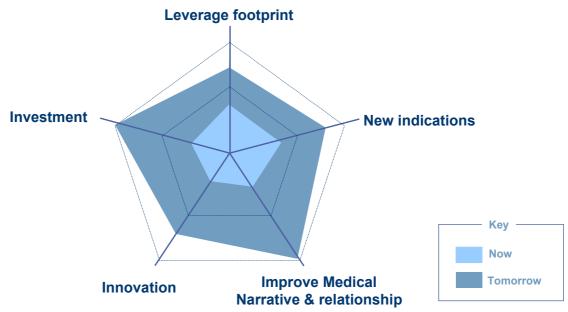


53

* two sequences : one invest; two return



Ipsen to work on key levers to reach full potential



Spasticity and the US: two main growth drivers



Dysport® 2020 footprint aspiration : More geographies, more indications

| Geography/ Therapeutic area | Europe | US | China | Brazil | Russia |
|--------------------------------|---|--|--|---|---|
| Therapeutic | 1. Cervical Dystonia 2. Adult arm spasticity 3. Blepharospasm 4. Hemifacialspasm 5. Paediatric per equinus spasticity (Cerebral palsy) 6. Adult leg spasticity (in three EU markets) 7. Hyperhidrosis 8. Pediatric arm spasticity 9. Neurogenic Detrusor Overactivity | 1.Cervical Dystonia 2.Adult Upper Limb 3.Adult Lower Limb 4.Pediatric Lower Limb 5.Pediatric Upper Limb 6.Neurogenic Detrusor Overactivity | 1.Cervical Dystonia 2.Other indications under assessment | 1. Cervical Dystonia 2. Adult arm spasticity 3. Blepharospasm 4. Hemifacialspasm 5. Paediatric per equinus spasticity (Cerebral palsy) 6. Adult leg spasticity 7. Hyperhidrosis 8. Pediatric Upper Limb 9. Neurogenic Detrusor Overactivity | 1. Cervical Dystonia 2. Adult arm spasticity 3. Blepharospasm 4. Hemifacial spasm 5. Paediatric per equinus spasticiy (Cerebral palsy) 6. Hyperhidrosis 7. Pediatric Lower Limb 8. Pediatric Upper Limb 9. Neurogenic Detrusor Overactivity |
| Aesthetic | 1.Glabellar Lines 2.Canthal Lines | 1.Glabellar Lines 2.Canthal Lines | 1.Glabellar Lines | 1.Glabellar Lines | 1.Glabellar Lines |





Dysport® in aesthetics

Through partnerships...

- Rationale: access the aesthetic market potential with partners offering a range of aesthetics products
- Successful partnerships with :
 - Medicis in North America



 Galderma in Europe and other selected geographies (e.g. Brazil)



... or solely as Ipsen

- In geographies not covered by partnerships...
- ... when it makes business and economic sense



New indications: Focus on spasticity and urology indications

Focus on spasticity in the short term...

- Current spasticity indications:
 - Adult upper (ex-US) and lower limb (limited markets)
 - Pediatric lower limb (ex-US)
- Spasticity, a major short-term growth opportunity:
 - Stroke: 15 million people worldwide every year.
 5 million are left permanently disabled¹
- World-wide Adult and Pediatric Ph III program (4 trials):
 - 4 new indications in the US
 - New and/ or Improved labeling ex-US

... and in urology in the longer term

- Leverage current access to prescriber base:
 - Clear synergies with Uro-oncology franchise in Europe
 - Clear WW synergies with neuro-rehabilitation environment
- Neurogenic Detrusor Overactivity: Ph IIa started (NCT01357980):
 - First patient screened in May 2011
 - Limited cost and high probability of success
- Urology indications, a significant mid term growth potential



Note 1: WHO The Atlas of Heart Disease and Stroke, Dr Judith MacKay and Dr George A. Mensah



Ipsen to improve medical narrative and prescriber relationship

| Leverage unique clinical experience | Dysport [®] value |
|--|---|
| Enhance collaborations with medical and scientific communities | Enhance services to physicians and payors |



Dysport® Next Generation: a potential new exciting opportunity

The first ready-to-use toxin A...

- ...is a breakthrough innovation bringing clear differentiation vs. competitors
- ...saves time by avoiding reconstitution
- ...improves safety (dilution/dosage, reconstitution, single use product ...)
- ...has very positive qualitative and quantitative market research results¹ (c. 500 participants):
 - 83% of potential adopters on time saving and improved safety grounds

A potentially transforming project

- A WW Ph III program to assess safety and efficacy:
 - -Indication: Cervical Dystonia
 - -350 patients
 - -71 sites (42 in Europe, 29 in the US)
 - -First patients recruited in Europe
 - -US recruitment pending feedback from FDA in Q3 2011
- A complex manufacturing process with technical hurdles to be addressed
- Ipsen team fully mobilized to bring R&D project to fruition

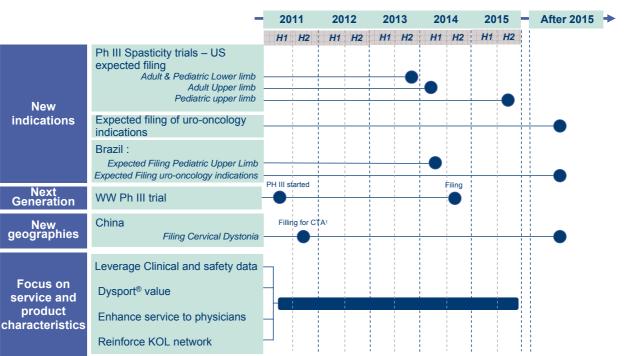
Potentially, a major change in market paradigm

59

Note 1: with Neurologists and Neuro-rehabilitators



Dysport® milestones:



Research & Development





A changing environment that calls for a dramatic change in the way of doing R&D

Tougher times for the Pharma Industry...

- Increased competition
- Generic/hybrid entry
- Persistent attrition rate
- Pressure of payers

... calls for

Scientific & medical excellence

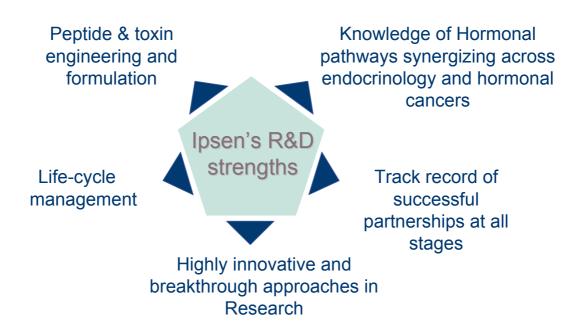
Focus on patients (Patient centric)

Speed of execution across the value chain

Collaborative innovation



Ipsen's R&D has recognized strengths...







... but still has room for improvement...

Ipsen's R&D has experienced setbacks...

- BIM23A760 (Phase IIb)
- Taspoglutide (Late Phase III)1
- Difficulties to properly position GH-IGF-1 combo program in short stature
- Difficulty to derive value out of promising research compounds (CDC25, diflomotecan, elomotecan, angiomate...)
- Irosustat in monotherapy (Phase IIa)

... push for greater focus across R&D

- Increase alignment between science & business analysis
- Focus resources on key projects and technological platforms
- Increase alignment in decision making
- Implement culture of decision and speed of execution



... calling for an overhaul of our R&D engine...

Success
to come from
highly
differentiated
healthcare
solutions with
demonstrated
patient
benefits

- Focus & align priorities between R&D and Commercial Operations in expert mid-size niches in specialty care
- Patient-centric driven R&D
- Continue to foster partnerships
- Speed of execution & Operational Excellence
- Highly differentiated technological platforms

Highly focused in targeted debilitating diseases

Core Translational Sciences/
Medicine capabilities

Integrated R&D and lean organization

Internal resources and competencies focused on peptides and toxins

Open Innovation & built from our strong partnerships

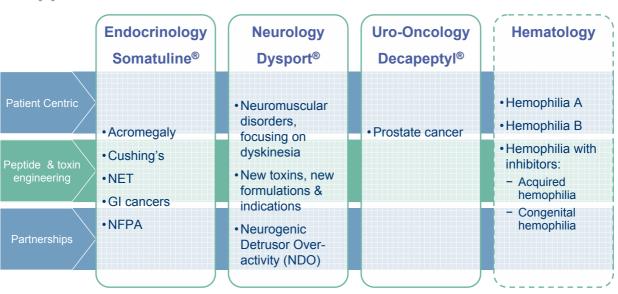
R&D to focus on short, medium and long term deliveries







... applied to well defined disease areas and indications

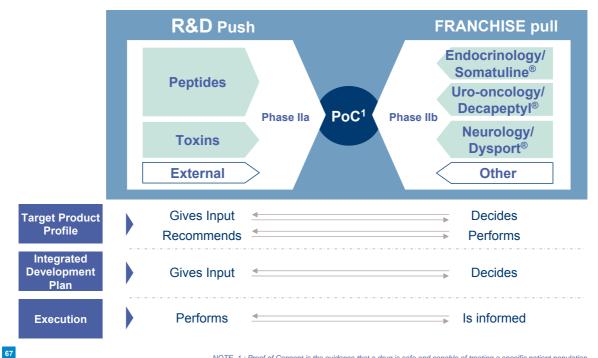


Focused. Aligned. Synergies across therapeutic areas





An integrated R&D "push-pull" model to fulfill patient/ commercial requirements

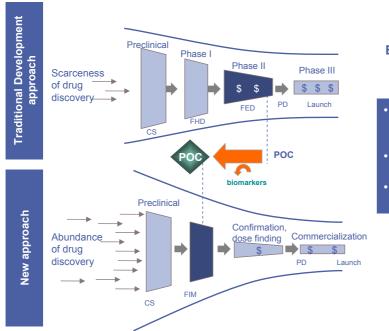


NOTE 1: Proof of Concept is the evidence that a drug is safe and capable of treating a specific patient population





Biomarkers help compounds move to POC quicker while reducing failure rate in late development stages



Benefits from the new approach

- Anticipation of Clinical plan at early Research stage
- Early recycling for some projects
- · Higher Probability of Success for projects entering Phase II-b

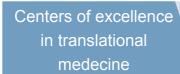
Project killing

CS : Candidate Selection FED : First efficacy dose FIM : First in Man PD : Product decision POC: Proof of Concept





Ipsen partnerships in personalized medicine





Biomarkers as catalyst for early Proof of Concept decision Confirmatory
Trials
Speed to
Market

Current partnerships : Centers of Translational Medicine & BioMérieux

69

Focus on two core technologies..





Ipsen has built a state-of-the-art peptide engineering platform...

State-of-theart scientific expertise and technology Enhance potency & efficacy

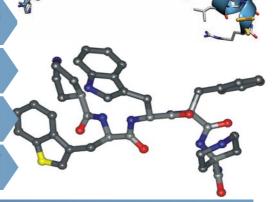
Improve target selectivity

Increase enzymatic stability and prolong duration of action

Target specific tissues, cells and tumors

Synergize activities at multiple targets with chimeric peptides

Combine sustained release expertise with chemical engineering to enhance compatibility with novel formulations



Ipsen has a proven peptide track record with marketed drugs and drug candidates





... and will continue to invest in emerging technologies

Emerging peptide-based technologies have potential to further expand applications of peptide-based agents

Current
state-of-theart peptide
platform

Cell-penetrating peptides, "stapled" peptides

- Peptides used as targeting agents to deliver therapeutic "war heads"
- Oral delivery technologies
- Chimeric peptides
- Phage display technology
- Cytotoxic-peptide and toxin-peptide conjugates, peptide-siRNA chimeras
- Peptide toxins & protein mimetics, inhibiting proteinprotein interactions

Ipsen to expand and reinforce its peptide platform, in-house and with selected partners



Focus on two core technologies...

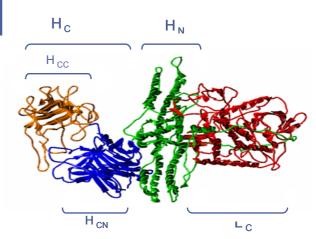




Botulinum toxin: Recombinant technology offers promise of development well beyond current marketed products

New opportunities with recombinant technology

- Design of novel targeted toxins
- Design of toxins with different characteristics (onset of action, duration)
- Platform versatility based on feasible modification(s) of the functional domains of native Botulinum toxin :
 - Binding domain
 - Translocation domain
 - SNARE cleavage domain
- Possibilities to improve the properties of the current available toxin formulation
- Clear and measurable clinical effects







Ipsen is ideally placed to become a key player in "the toxin of the future"

Track record expertise in botulinum toxin with Dysport®

Pharmacological, preclinical and clinical expertise in Botulinum Toxin in neuromuscular diseases

State-of-the-art facilities: research, scale up, development and manufacturing



Valuable partnerships synergize with Ipsen

Established network of Toxin experts







Partnerships at the Heart of Ipsen's Innovation







2003 - 2005

2006-2011



Our key drivers entail major decisions

of major R&D decisions

- Merge Research and Development departments
- Close R&D activities at Barcelona site
- Stop internal non toxin and non peptide research*
- Build project management excellence



* The Group keeps the OBI-1 development platform in Milford, MA, USA



Key decisions on Ipsen's Ph II pipeline

Assets no longer in strategic focus

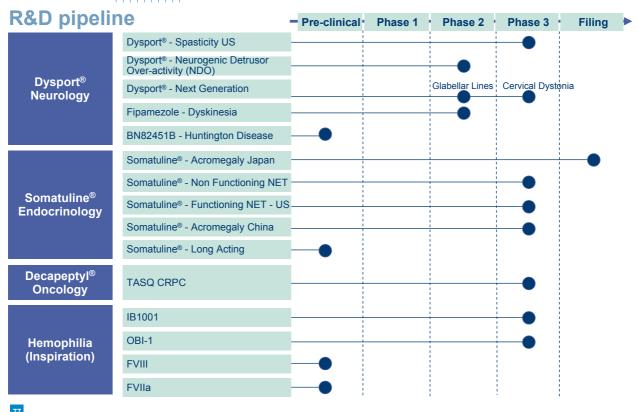
Oncology

- Discontinuation of the development of Irosustat as a single agent (monotherapy) in all indications
- Seeking partnership

Endocrinology

 GH- IGF-1 Combo and all IGF-1 programs are deprioritized lpsen to explore all options to maximize value while meeting its obligations to patients and partners







Ipsen "new" R&D ambition

5 novel pre-clinical candidates...

...out of which 3 will reach POC decision by end of 2015...

...while all life cycle initiatives are achieved on time

Uro-Oncology/ Decapeptyl®





A franchise with renewed growth opportunities

Tasquinimod

for castrate resistant tumors

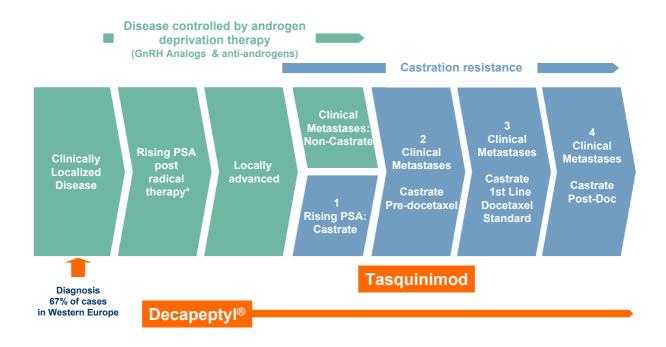
Once a day oral formulation in PhIII

Decapeptyl®

for hormone-sensitive tumors



Prostate cancer: disease evolution towards castration resistance





*Radical treatment : prostatectomy or radiotherapy



Decapeptyl®: a solid basis to develop a Uro-Oncology franchise

A 20-year growth story...

- Marketed in over 60 countries (Excl. the Americas and Japan)
- Indications:
 - Prostate Cancer (more than 70% of sales)
 - Gynaecology
 - · Precocious puberty
 - IVF
- Formulations: Daily, 1 month, 3 months and 6 months
- No true generics of GnRH analogs anticipated

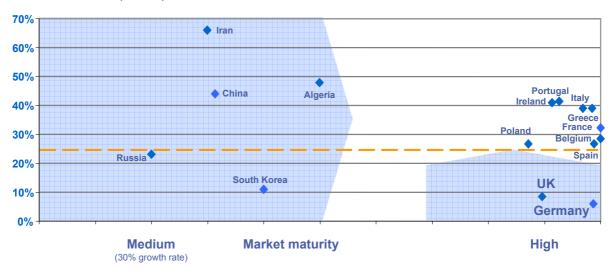
... and still poised to grow

- GnRH analogs remain mainstay of first line hormonal manipulation in PCa
- 6 month formulation enables market share gain in key EU countries
- Emerging countries, and in particular China, provide a long term growth
 - In 2010, emerging markets contributed to ~65% of Decapeptyl[®]'s growth
- European sales affected by increased competition and price pressure



Decapeptyl®: strong market shares throughout the world, and room to grow

Current market share (in units)



Market share in the UK and Germany (~40% of G5 GnRH Analogs sales) has doubled in the past 3 years



SOURCE: IMS – Year 2010 100 indicates most mature markets

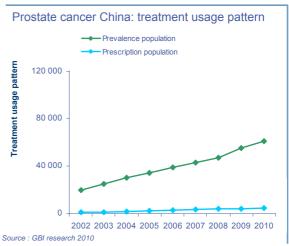


Emerging countries, and in particular China, as growth engine

China, limited GnRh analog market but poised to grow (2006-2010)



Increasing access of patients to medicines (2002-2017)



China expected to become 2nd country in terms of Decapeptyl® sales in 2011



Decapeptyl® 6 month formulation: a differentiated product profile, enabling market share gain

Decapeptyl 6 month formulation

Efficacy

- Sustained low level of testosterone, without breakthrough*
- Similar efficacy on PSA control and Testosterone across all formulations**

Local Tolerance

- Limited local side effects (6.7% of patients)
- No nodule or abscess at site of injection

Formulation reconstitution

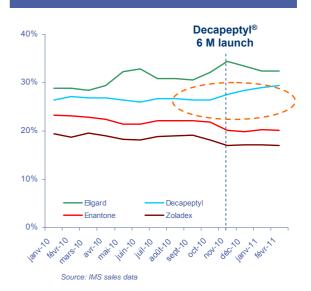
FIPSEN

- Lyophilized slow release formulation
- Easy to reconstitute with no product loss

*Lundstrom E & Al, , ClunDrug Invest 2009; 29(12):757-765 **Mounedji N & al, J clin Oncol 29: 2011 (suppl 7; abstr 162)



e.g. Impact on market shares in Spain

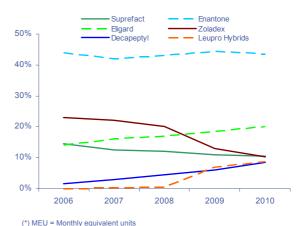


No true generics of GnRH analogs anticipated, only Hybrids

Hybrids rather than true generics

- Risk of true <u>long acting</u> GnRH analogs generics entry expected to be low
- Only hybrids of leuprorelin are available today¹
- Hybrids are currently not substitutable and priced 20-25% below original products
- In Germany, the 2 leuprorelin hybrids have reached less than 10% MS in 3 years² with no impact on class price yet

Impact of hybrids on market shares MEU (*) in Germany



Source: Insight Health, OdV data - Germany

Hybrids represent a moderate threat to GnRHa established brands compared to true generics



Tasquinimod: a perfect strategic fit

Disease controlled by androgen deprivation therapy (GnRH Analogs & anti-androgens) **Castration resistance** Clinical Metastases: Non-Castrate Rising PSA post radical therapy Clinical Metastases Clinical Metastases Locally advanced **Tasquinimod** Decapeptyl® **Population** 149 000 111 000 153 000 Incidence Stage I & II **CRPC** in G5* Stage III & IV

- Leverage the Group's current leadership position in prostate cancer
 - Expand to medical oncology
 - Access to significant sales potential
- Beyond prostate, tasquinimod has potential in other cancers (such as GI)

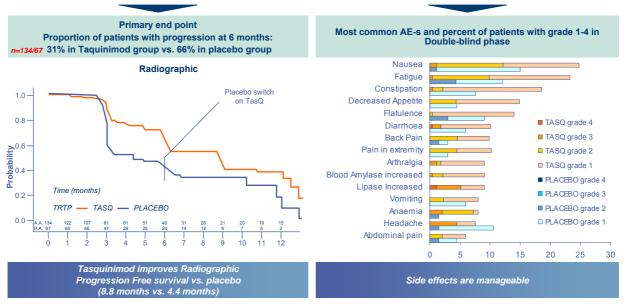


* Oncos Da Vinci, 2008



Tasquinimod, promising phase II results

Safety and efficacy analysis* of Phase II study of Tasquinimod in chemotherapy naïve patients with asymptomatic metastatic castrate-resistant prostate cancer (CRPC) (n=201)

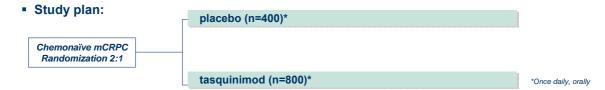




Tasquinimod, Phase III program ongoing

A Phase III randomized, double-blind, placebo-controlled study of Tasquinimod in men with asymptomatic/mildly asymptomatic Metastatic Castrate Resistant Prostate Cancer

- Objectives
 - TASQ in chemonaïve patients with metastatic castrate-resistant prostate cancer
 - Effect of Tasquinimod on delaying disease progression compared with placebo
- Endpoints
 - Primary: Radiological progression-free survival (PFS)
 - · Secondary Endpoint: Overall Survival (OS) Study powered for OS



- Principal investigators:
 - America : Michael A Carducci, Johns Hopkins Kimmel Cancer Center, Baltimore, USA
 - Europe: Cora N Sternberg, San Camillo and Forlanini Hospitals Rome, Italy

International Pivotal Phase III opened 1Q 2011...

... filing expected in 2014





Tasquinimod, deal terms for Ipsen

Geographies

World excluding Japan and the Americas

Execution

- Active Biotech: Pivotal registration PhIII
- Ipsen: Supportive study

Financials

- Milestones :
 - Upfront payment of €25 million
 - Additional payments of €175 million contingent upon progress/ achievement of clinical, regulatory and commercial milestones
- Royalty rate: progressive on the level of sales starting in the low teens

Expected peak sales: in excess of €250m

Hemophilia





Ipsen and Inspiration are aiming at all levels of the coagulation cascade for the treatment of hemophilia

A full fledged hemophilia franchise, with potentially 4 products ...with a broad potential inhibitor therapy offering (OBI-1, FVIIa)...

...and the first recombinant competitor in hemophilia B therapy, IB1001 ...differentiated with
OBI-1, the only recombinant porcine FVIII product...

- → An \$8bn market
- → A high margin market
- → 2 products in Ph III:
 - OBI-1: a highly innovative porcine recombinant FactorVIII (orphan drug)
 - IB1001: first rFIX biosimilar in an underserved, fast growing market



Growing Market Opportunity in Hemophilia B

Current market

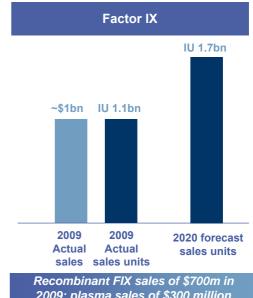
- FIX market overlooked primary focus FVIII
- 90% recombinant in developed markets
- 40% recombinant and growing in underserved markets

Long-term growth prospects

- 2% to 6% CAGR until 2020
- Broadening access to care grows overall market:
 - Driven by prophylaxis in developed markets
 - Driven by more patients treated in underserved markets

Critical unmet medical need: access to treatment for more patients

 Hemophilia B market potential of 6.5 billion International units (IU), based on population



2009; plasma sales of \$300 million

Significant market opportunity for IB1001, a recombinant FIX currently in Phase III





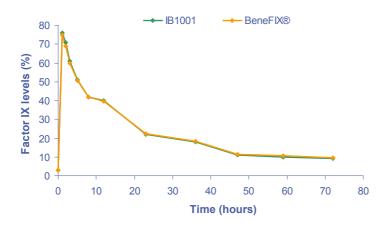
Historical and projected hemophilia market growth drivers

Hemophilia **Prophylaxis** Conversion Increased Diagnosis & population treatment penetration to competition recombinant growth rate **Opportunity to** Increasing Limited, enhance Market Market weight of prophylaxis longer-term opportunity opportunity demographics opportunity and conversion to rFIX



IB 1001 demonstrated non-inferiority to BeneFIX®

Mean FIX activity by time and treatment



The preliminary safety data collected during the PK study phase indicate that IB1001 has an acceptable safety profile and is well tolerated

Study IB1001-01 is ongoing and further analyses on safety and efficacy will be available in 2011.





Growing Market for More Effective Inhibitor Treatments

Current market

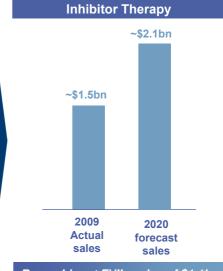
- 1/3 of hemophilia A patients will require inhibitor therapy during their lifetime:
 - Inhibitor therapy (IT) 70% recombinant in developed markets
 - IT <10% recombinant in underserved markets
 - Current therapies are FVIIa bypassing agents

Long-term growth prospects

- 3% to 6% CAGR until 2016, 0-1% thereafter (due to increased competition, downward pricing pressure):
 - Driven by prophylaxis in developed markets
 - Driven by more patients treated in underserved markets
- Opportunities in treating inhibitors to human FVIII and acquired hemophilia

Most important unmet need: Additional inhibitor therapy options

- Current inhibitor therapies are expensive; not always effective
- Lack of biomarkers to predict efficacy

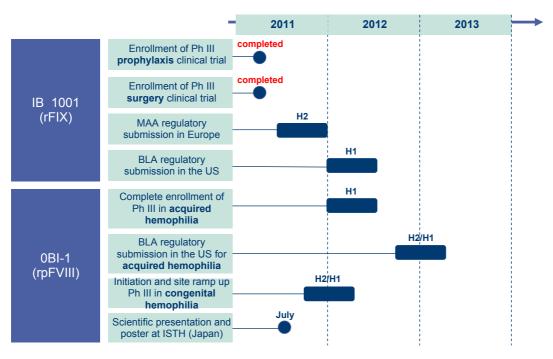


Recombinant FVIIa sales of \$1.1bn in 2009; plasma sales of \$400m

Unique positioning for OBI-1, a differentiated porcine recombinant FVIII currently in Phase III



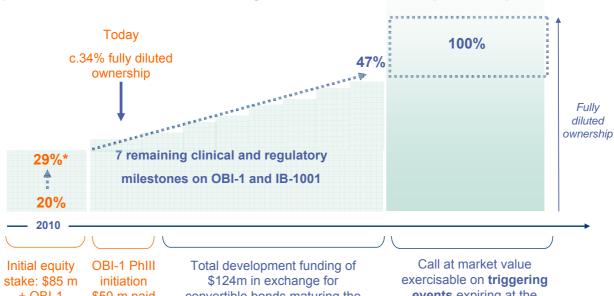
Inspiration hemophilia product portfolio – short term timeline







Ipsen now has ~34% of fully diluted ownership of Inspiration



+ OBI-1 \$50 m paid upfront: \$50 m by Ipsen in + 27.5% royalty exchange for rate on OBI-1 convertible bonds

convertible bonds maturing the later of 7 years or the end of the call exercise period

events expiring at the latest in 2019

US platform





Ipsen has completed a comprehensive assessment of global operations and reaffirms its commitment to the US market

Significant upside potential for new indications for Dysport® and Somatuline Depot®

Significant market opportunity for hemophilia portfolio

Significant value in further developing Ipsen's presence and insight into the US market

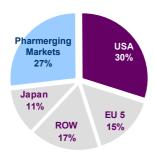




The U.S. will continue to lead the global marketplace and remains an important pillar in Ipsen's global strategy

In 2015, the U.S. market will represent ~30% of the global pharmaceutical market...

2015 Global Pharmaceutical Market Share (Sales)1



The US will contribute to 11%¹ of the 2010-15 global pharma market growth and specialty products are anticipated to grow faster than the overall market...

...with significant opportunity for Dysport® and Somatuline®

Dysport®

- Global therapeutic botulinum toxin market expected to grow at 7.4%² CAGR to €1.6bn in 2020
- US botulinum toxin market to grow at 7.4%² CAGR to €1.4bn in 2020 (50% of total market)

Somatuline[®]

- Global SSA³ market expected to grow at 3.8% CAGR to ~€1.6bn²
- World most solid growth in the US with a SSA³ market expected market growth 4.6%² CACG in the US

SOURCE: (1) IMS Health 2011 (2) Company estimates (3) SSA: Somatostatin Analogs







To ensure we maximize the opportunity, there will be four key changes to the U.S. Organization

Focus investment and resources to drive Dysport[®] and Somatuline Depot[®] growth in current and future indications

Restructure the US organization to increase focus and align US operations with the new global franchise structure

Better integrate the US business with the global organization and move US commercial operations to the East Coast

Allocate internal effort to activities that drive differentiation and focus and form outsourcing partnerships to support other activities





U.S. growth also will be fueled by life-cycle management and new products

8 on-going phase IIIs in the US

4 Dysport® (Spasticity)

- Adult upper limb spasticity
- Adult lower limb spasticity
- Pediatric upper limb spasticity
- Pediatric lower limb spasticity

Expected filing 2014 - 15

2 Somatuline® (NET)

- Functioning NET
- Non Functioning NET

2 Hemophilia (Inspiration

- IB 1001 (rFIX)
- OBI-1 (rpFVIII)

Expected filing 2014

Expected filing H1 2012 for IB1001 Expected filings 2012/ 2013

Opportunistic Business Development will be considered







A two-phased US strategy supports short- and long-term growth objectives

— 2011 — — 2014 - 2015 — — 2020 →

- Invest in current indications for Dysport® and Somatuline® Depot to gain share in cervical dystonia and acromegaly markets respectively
- Invest in clinical trials to support life cycle management of Dysport[®] and Somatuline[®] Depot
 - Dysport®: 4 Ph III in spasticity
 - Somatuline[®]: 2 PhIII in functioning and non functioning NET
- Prepare the organization for the launch of new indications

Assess the option to enter US hemophilia market

- Launch and grow Dysport[®] in spasticity and Somatuline[®] Depot in NET
- Hemophilia franchise option
- Become a significant sales and profit contributor to the Ipsen Group



A new President and General Manager for Ipsen North America

Sean McKercher

- 30 years of experience in the healthcare industry, working in many different locations including Canada, Asia, Africa, US and most recently Europe.
- **Joined Ipsen in 2007**: instrumental in the 2008 acquisitions of Vernalis US, Tercica and the product OBI-1.
- Since 2009, head of Ipsen's business development and alliance management organization completing over 20 different transactions including the creative relationship with Inspiration.
- In 2010, named the Business Development Executive of the Year by the UK Pharmaceutical Licensing Group.
- Recently named President and General Manager for Ipsen North America



China focus:

Invest to accelerate value growth to harvest the full benefit of 20 years experience







China amongst top pharma markets...

| | 2010 rank | | | 2015 rank | | 2016 rank | | |
|---|---------------|---|---|---------------|---|-----------|---------------|----|
| 1 | United States | | 1 | United States | | 1 | United States | |
| 2 | Japan |] | 2 | Japan | | 2 | China | *) |
| 3 | China | | 3 | China | * | 3 | Japan | • |

Region contribution to global growth

Region market share of global sales 2015



 $^{^{\}star}$ 2015 sales estimated using 2011-2014 CAGR sept. 2010 for Egypt & Ukraine in Tier 3 and RoW Source : IMS Health Market Prognosis, March 2011

In 2010, China became the world's second largest economy ahead of Japan

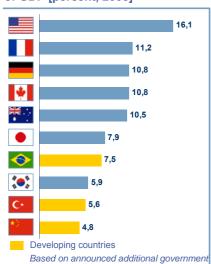






...and more Healthcare potential to come

Healthcare expenditure as percent of GDP [percent, 2008]



China healthcare expenditure - projected spend [USD billions]



A US\$ 600 billion market by 2015, trending towards \$1 trillion by 2020





China a bursting market place

Competitors

- International competitors: "Arms race" on coverage, investment, products, talents, etc...
- Local competitors: strong development & ambitions, with strong cash reserves (subsidies for State-owned enterprises, IPO for private companies...) that start to venture abroad

Patients

- Looking for quality
- Paying out of pocket
- Better informed

Chinese Healthcare

Authorities

- More balanced development
- Support national champions
- Healthcare as a political challenge

Healthcare Reforms:

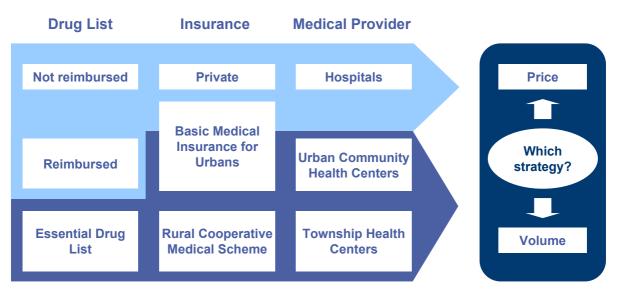
- Expand basic medical insurance programs
- Establish national essential drug system
- Develop primary healthcare services system
- Provide equal access to urban and rural residents
- · Accelerate public hospital reform







Essential Drug System may reconfigure Chinese healthcare modus operandi



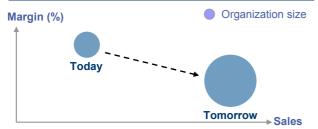
x 31 provinces





China today, a tougher place for new comers...

Increasing cost of doing business



China is more and more costly:

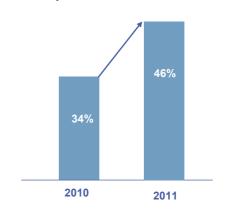
- Salaries
- Taxes and employer contributions
- Regulatory requirements
- Inflation
- X Strong headcounts increase for better coverage
- Decreased overall profitability

with price cuts

111

China is increasingly discriminant

Perception of governmental policies as discriminating against Foreign-Invested Companies for the next 2 years



Source : European Union Chamber of Commerce, Business Confidence Survey, May 2011





Ipsen benefits from a longstanding presence in China, now its 2nd affiliate

- Established in 1992
- HQ in Tianjin. Total staff of ~ 500 employees
- ~€90m 2010 sales
- A truly Chinese organisation with a local JV partner and 2 expatriates



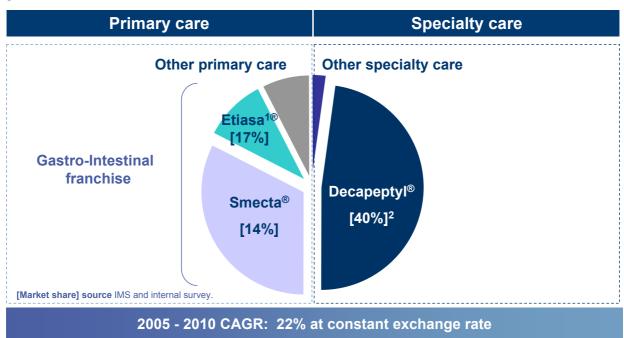
- Regional sales force teams focused on key coastal cities, now spreading towards the
 west into lower tier cities
- Excellent National & Provincial market access team (patent, pricing, reimbursement, bidding ,etc.)
- Experienced medico-regulatory team to conduct local clinical and registration trials
- World class manufacturing site since 2000

Proven track record of building brand value and leadership in targeted therapeutic areas





Solidly anchored on two pillars, with three market leader products



113

NOTE 1:Etiasa® in-licensed from Ethypharm NOTE 2 : 40% market share over all indications (Gynecology, prostate cancer, IVF, ...)

Primary care





Strengthening our Gastro Intestinal Franchise

Currently promoted portfolio

- -Smecta®
- –Etiasa®
- –Fortrans®
- -Forlax®
- _Meteospasmyl[®]
- -Tanakan®

Strong Alliance Management

- Currently: 2 in-licensed products: Etiasa[®] in 1999; Meteospasmyl[®] in 2011
- Continuing local business development efforts

Life Cycle Management

2020 ambition

Current coverage and sales force

- -180 cities
- –250 Medical and Trade representatives

Expand coverage

- Cover more than 400 cities by 2020
- Double sales force team by 2020

Assess EDL opportunity regularly

2020 ambition





2020

ambition

Developing our Specialty care franchise

Currently promoted portfolio

Decapeptyl® with a complete range of formulations:

- -Prostate Cancer
- -Gynecology
- Consolidate gynecology leadership position while developing prostate cancer indication with launch 3M formulation
- Prostate Cancer extension
 - Tasquinimod
 - Additional licensing projects under discussion

Introduction of 2 major Ipsen products

- Somatuline® CTA¹ filling 2011; regulatory approval expected in 2016
- Dysport[®] in Cervical Dystonia and aesthetic indications: CTA¹ filling 2011; regulatory approval expected in 2016 and 2017 respectively

Current coverage and sales force

- 120 cities
- 120 Medical andTrade representativesHighly focused and

trained teams

- Increase medicalization :
 - Already ongoing with 4 studies in 60 centers and more than 1500 patients in 2011 (against none in 2009)
 - 2012: Creation of a Clinical trial platform in Beijing

2020 ambition

Extend and deepen coverage

Triple sales force team by 2020



NOTE1: CTA filing for Clinical Trial Authorization



Focusing to secure long term growth

Shorten timeto-market

- Strengthen medical development capabilities
- Creation of a clinical trial platform in Beijing in 2012
- Earlier integration of China in global product development

Anticipate healthcare reform and implementation

- Constantly adapt to the changing market conditions
- In every province: Local Government Affairs to be close to local decisions makers

Execute the plan

- High turnover in whole pharmaceutical industry with new competitors local / state companies
- Recruit / retain / develop in a talent war context





In summary, Ipsen in China is...

...well positioned for its size...

...in this market of unique magnitude and growth opportunities, with a solid portfolio and a highly competitive infrastructure

...well aware and vigilant...

...about market fragmentation, dynamics and complexity and has become a "market mover" in its selected therapeutic areas



...with a professional and motivated team dedicated to gaining market share and delivering sales and profit growth

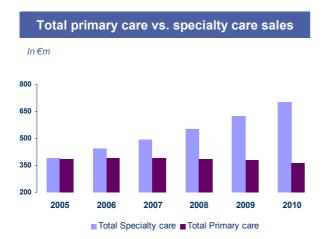


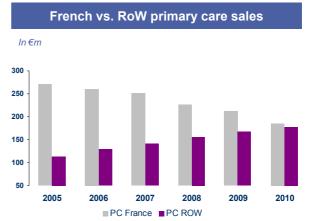
Primary Care France





Contrasted primary care dynamics





Pressure on Ipsen's French primary care





A European-wide pharmaceutical industry situation

Average price for daily treatment cost has declined and will continue to decline R&D productivity lower in GP products Increased restriction on promotion

Mature products are still responsive to promotion but may not justify a large dedicated sales network

Potential solutions: add products and/or work the cost base



Ipsen's French PC remains an efficient platform ready for further commercial leverage



Adenuric[®], Exforge[®] and perspectives A potential leverage in the OTX market with established brands

Solid foundations

An established portfolio of primary care products

A recognized and respected sales and marketing organization





Today, optimization is no longer sufficient for French PC

Straight forward co-marketing and co-promotion optimization deals will not remain sufficiently financially attractive

We need to explore more engaging partnership models



- Enlarge the portfolio of promoted products :
 - Usual co-promotion and co-marketing in-licensing agreement
 - Combine Ipsen's portfolio with other companies' primary care products, including "mature" products
- Have access to OTC/ OTX know-how and capabilities, including pharmacy sales force
- Mutually optimize the cost bases



Primary care France – Manufacturing

A well established manufacturing facility in Dreux...

- Drug manufacturing and packaging activity for finished products including Smecta[®], Forlax[®], Tanakan[®]
- Specific expertise and high volumes in Sachet manufacturing
- While French related volumes are expected to decline, international activity will provide a favorable basis for partnering discussions

Other industry actors like CMOs...

...could ensure a sustainable future and add value to Dreux manufacturing site...

...better than Ipsen

123

Conclusion





Become a global leader in targeted debilitating diseases

Increase Focus

Invest to Grow

Leverage Footprint

A market-oriented franchise model...

...driving an R&D patient centric organization focused on core platforms, peptides and toxins

2020 ambition

More than double revenues¹

...and more than triple EBIT2

125

NOTE 1: 2020 figures include contribution of Inspiration portfolio and are set at constant foreign exchange rate

NOTE 2: prior to purchase accounting recordings and non recurring elements



One-off costs associated with the new organization

Expected one-off costs of 80 to 100 million euros before tax over 2011 and 2012



USA transfer costs to east coast



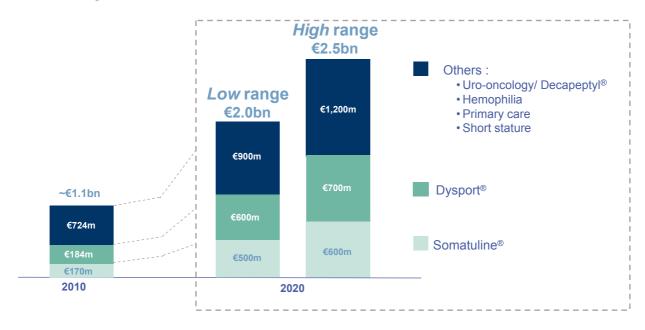
Closing of R&D activities of Barcelona site



 Other one-off costs related to the implementation of the strategy and of new organization



A significant contribution of Dysport® and Somatuline® to 2020 sales aspiration







Requiring an initial investment phase

2011 – 2015 Invest to grow

Clinical trials:

Dysport® in spasticity
Dysport® in urology
Dysport® Next Generation
Somatuline® in NET:
Functioning

2016 - 2020 Solid growth

Dysport® and Somatuline® on track for full potential:

- US
- Launch in China/ Russia/ Brazil
- Dysport[®] Next Generation

Global Decapeptyl 6 month supported by tasquinimod

Leveraged geographies:

Non functioning

- New geographies for Dysport[®] and Somatuline[®]
- New geographies for Decapeptyl® 6 months

Overall efforts

Stronger commercial push on key products



Implementation: main milestones to success

| | 2011 —— | 2012 —— | 2013 | 2014 —— | | 2015 |
|--------------|--|---|--|------------------------------------|----------|-------------------------------------|
| ✓ | Define strategy Merge R&D | R&D « PoC » machine implemented | Somatuline® New device rolled out globally | TASQ filed in Europe | | Inspiration option assessment |
| \checkmark | Reinforce Uro- oncology franchise (TASQ) | Barcelona R&D site closed | Dysport® A.& P. L.L spasticity filed | Somatuline® F. NET filed in the US | | Dysport® P.U.L filed in the US |
| \checkmark | Dysport [®] CD CTA ¹ filing in China | French primary care commercial activities | Dysport® NDO Ph III initiated | Somatuline® NF NET filed WW | | 5 new Pre clinical candidates (vs. |
| V | Somatuline® Acromegaly | partnered | Smecta® EDL assessment (China) | Dysport® A.U.L filed | | June 2011) O/W 3 reach POC |
| | CTA¹ filing in China | IB1001 filed in the USA | OBI-1 Acquired H. filed in the US | Dysport® NG filed | | Smecta® EDL assessment (China) |
| | New extended Executive Committee | OBI-1 PhIII (Acquired H) enrollment completed | | Dysport® P.U.L filedin Brazil | | |
| | staffed | oomplotod | | Inspiration option assessment | | |
| | Franchise org. implemented | OB-1 PhIII Congenital H. | | Smecta® EDL | | |
| | IB1001 filed in | initiated | | assessment (China) | | |
| | Europe | US platform reorganized | | | | |
| 129 | | | | NOTE 1: CTA | or filii | ng for Clinical Trial Authorization |

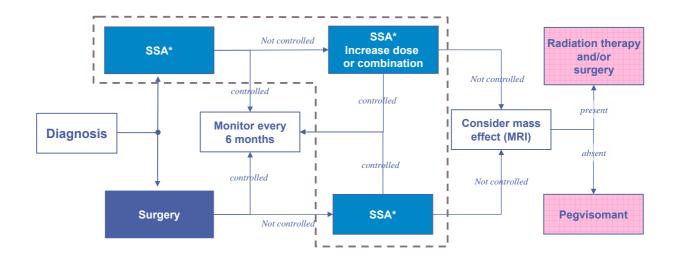
Subject to the relevant staff representation authorities in each country concerned, in compliance with the specific procedures, terms and conditions set forth by each national legislation

Appendices





Acromegaly: current therapy algorithm



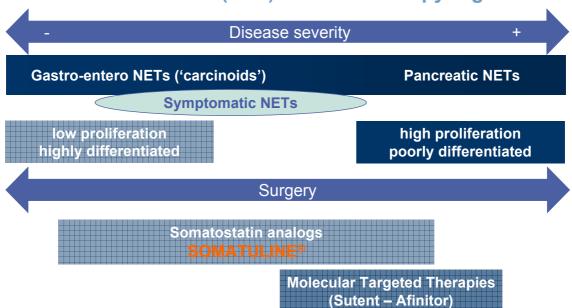
*SSA = Somatostatin Analogs



Adapted from Melmed; J Clin Endocrinol Metab, May 2009, 94(5):1509-1517



Neuro Endocrine tumors (NET): Current Therapy Algorithm



Cytotoxics