Strategy Day Ipsen

June 9, 2011





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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.

Today's agenda

M. de Garidel Chairman and Chief Executive Officer





Agenda for the day

Introduction	Marc de Garidel	13:30
Endocrinology/ Somatuline®	Christophe Jean	14:00
Field of NET	Pr. Wouter de Herder	14:20
Neurology/ Dysport®	Christophe Jean	14:45
Field of toxins	Pr. Pierre Denys	15:05
R&D	Claude Bertrand	15:25
Q&A session		15:45
Break - 20 minutes		16:05
Uro-oncology/ Decapeptyl®	Christophe Jean	16:25
Hemophilia	Marc de Garidel	16:40
Field of Hemophilia	Pr. Claude Négrier	16:50
USA	Christophe Jean	17:10
China	Eric Bouteiller	17:30
Primary care	Marc de Garidel	17:50
Conclusion	Marc de Garidel	18:00
Q&A session		18:10

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

M. de Garidel Chairman and Chief Executive Officer



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



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

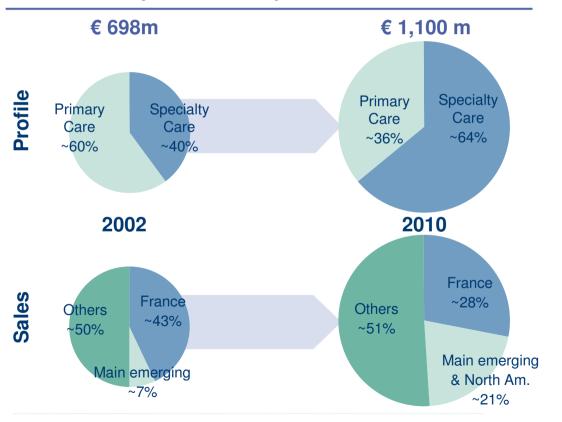
First assessment





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

Ipsen's key strengths

Innovative & differentiated platforms

- Peptides
- Toxins

Open model culture

Partnerships from research to marketing

Comparatively low exposure to patent expiry

- Specialty Care
- Life cycle management

International Footprint

Pharmerging & US

Geographical reach

13 Ipsen - Strategy Day - June 9, 2011



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

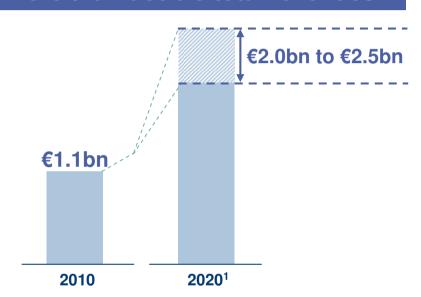
"To become a global leader in targeted debilitating diseases"



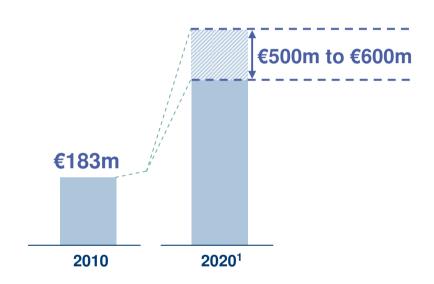


Ipsen's 2020 ambitions

More than double total revenues...



..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



To achieve our ambition, we need to...

... Increase focus

- 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

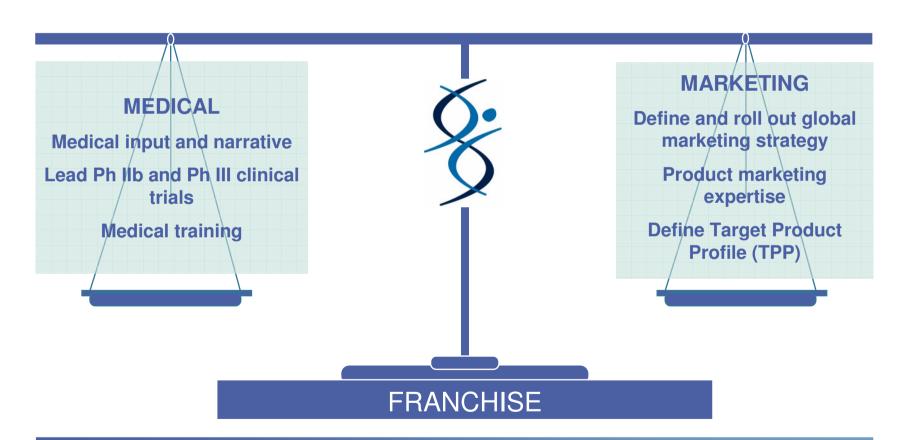
... Leverage footprint

- 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

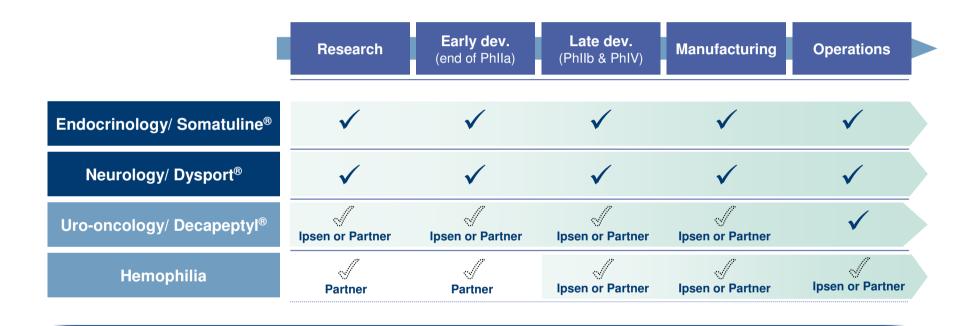


Countries are responsible for P&L performance





Increase market focus on 4 franchises, driving innovation

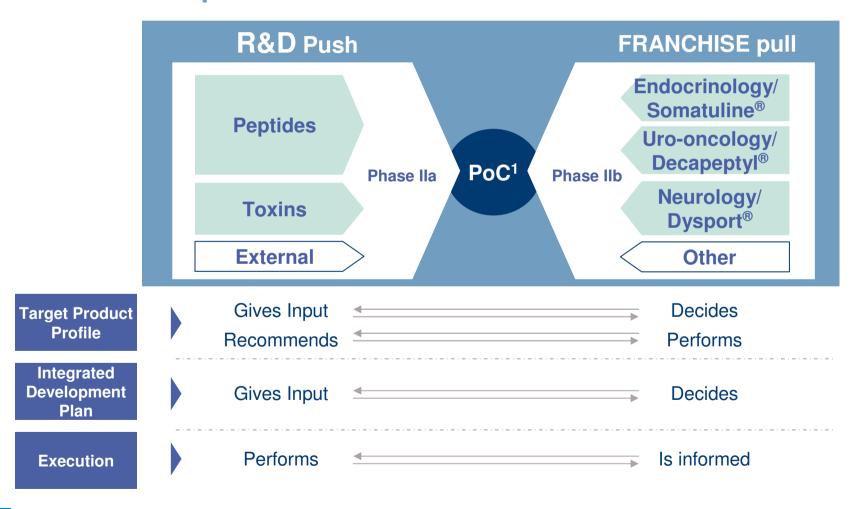


Primary care and Short Stature in a commercial optimization strategy





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

Market driven R&D

- 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





INVEST TO GROW: Ipsen's way forward

Invest to grow over 2011-2015

Focused investment effort to further develop Somatuline® and Dysport®

- Dysport® Ph III in spasticity
- Dysport® Ph II in urology
- Dysport® Ph III Next Generation

- Somatuline® Ph III in NET
 - Functioning in the US
 - Non functioning WW

Increase investment effort in growth markets: pharmerging, US

- Somatuline® and Dysport® full label in the US
 - Local/ regional partnering
- Decapeptyl® 6 month in the emerging markets
- Increase sales force in China to benefit from existing positions
- Reinforce GI portfolio around Smecta® in emerging markets

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

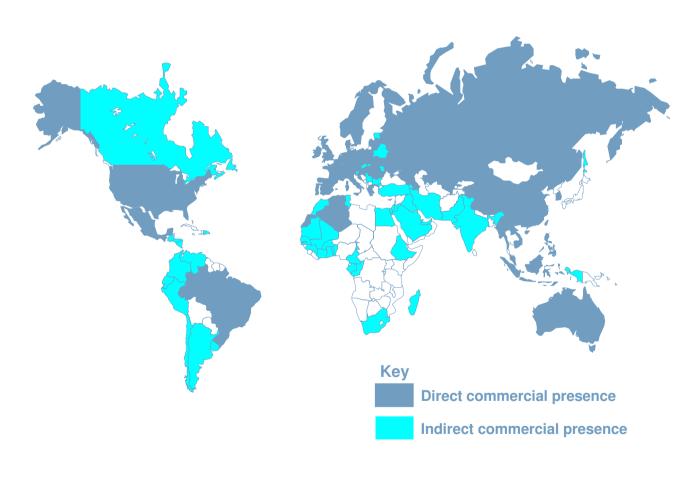
Enhance leadership in technological platforms
Grow market share in selected disease areas







An already existing extensive commercial reach...



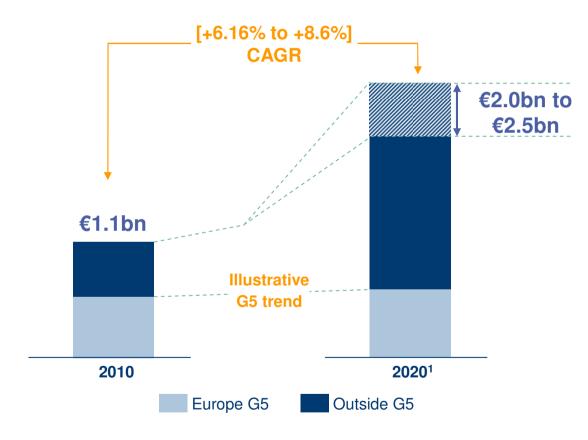
Sales recorded in 115 countries:

- 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





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

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®

Pharmerging

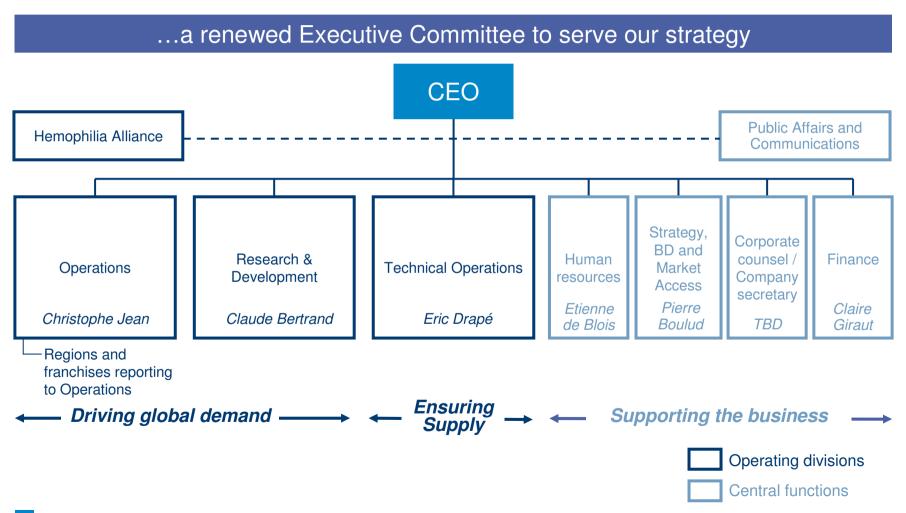




INVEST TO GROW

LEVERAGE FOOTPRINT

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







INVEST TO GROW

Enhance Ipsen's corporate culture



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®

C. Jean EVP, Chief Operating Officer





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

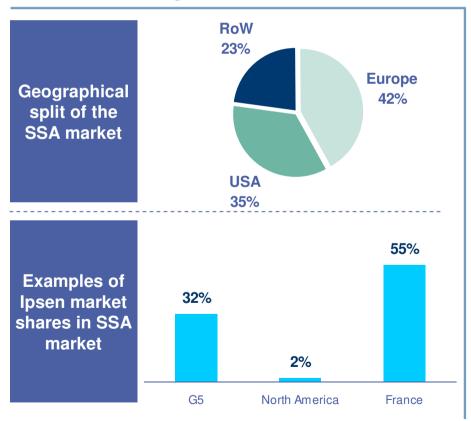
Franchise territory : Adult endocrinology, NET





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

Q4, 2010 market figures



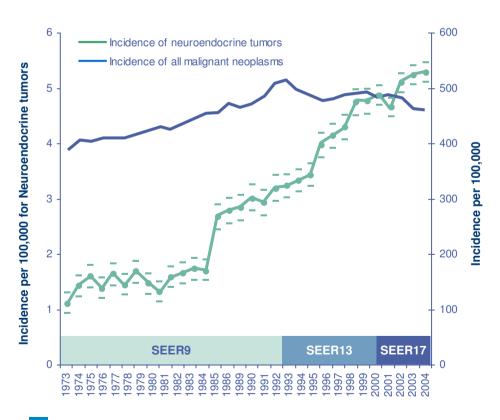
- 2010 SSA market: ~€ 1.1bn
- Solid SSA market growth (+9%¹ in 2009 and +18%¹ in 2010)
- A fairly balanced geographical split between Europe (42% of total sales), the US (35%) and the RoW (23%)
- Somatuline[®], an established product in Europe both in Acromegaly and in NET with 55% SSA market share in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



... 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

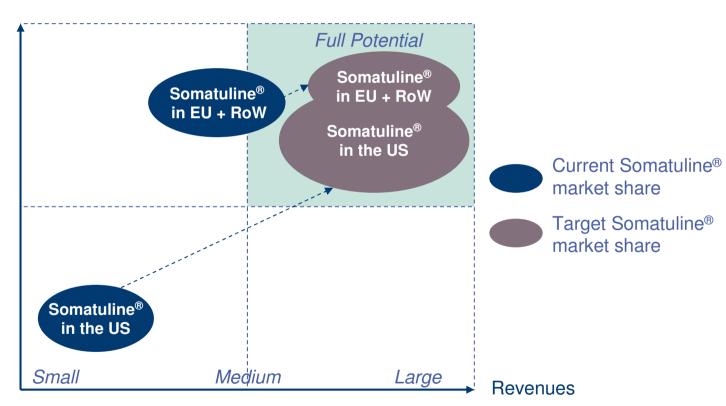
The US

 +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

Technological platform

Barrier to entry

Competition

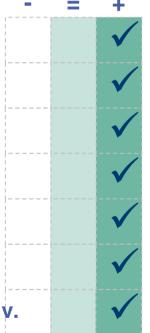
Market size/ Growth

Market share

Geographies

Somatuline® competitive adv.

Somatuline® growth potential



Core: peptides

High – pre filled syringe & long acting peptide

Well characterized – 1 major competitor, targeted combo therapies emerging

Large/ **Solid** – 3.8% CAGR until 2020 (€1.59bn)

Room to grow: Somatuline® 2nd player

US + **Pharmerging** potential – solid growth in Europe

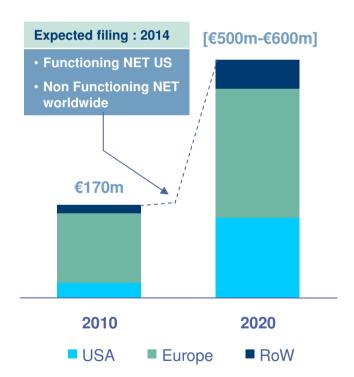
Significant – elements of differentiation to be further enhanced

Strong - 2 ongoing PhIII in NET (non functioning WW + functioning US)...

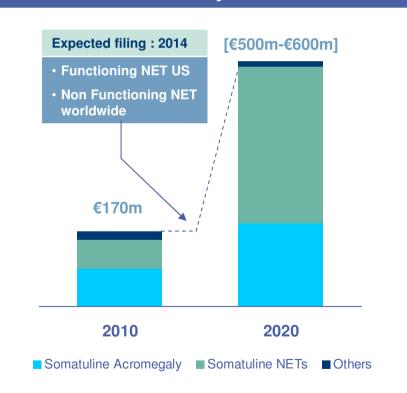


Somatuline® ambition: 10% to 15% CAGR* until 2020

Balanced geographical growth...

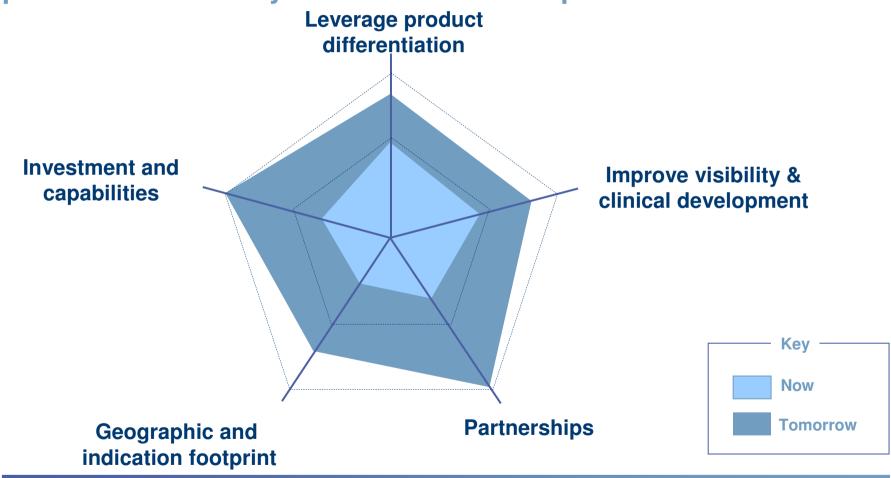


...driven by NET





Ipsen to work on key levers to reach full potential



NET and the US: two main growth drivers



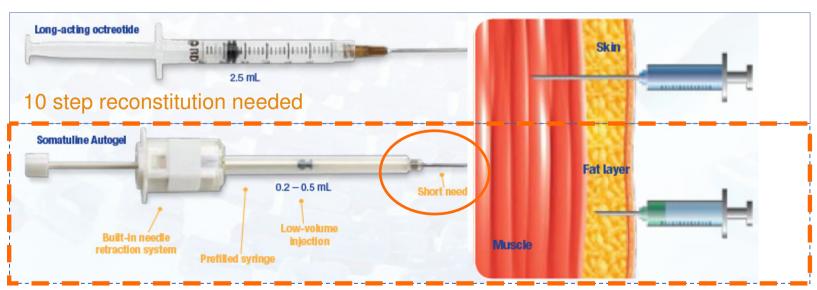
New additional elements of differentiation

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

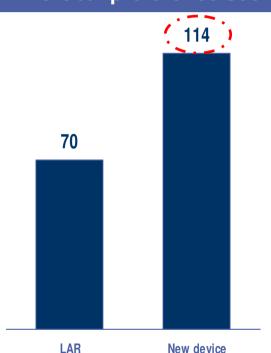




Somatuline® New Device: preferred by Nurses

Somatostatin Analog Nurse Preference Study - 1st publication

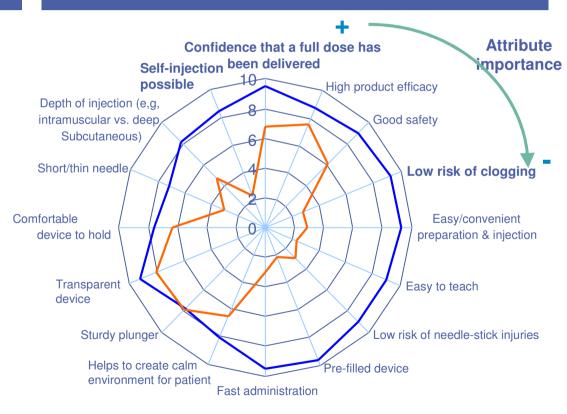
Global preference score



Global preference score = Added sum of each attribute's importance score/10 * Product Evaluation score

= significant difference compared to Ocreotide LAR (p< 0.05)

New device better on all attributes



— Lanreotide Autogel® new device

Octreotide LAR



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



SANOE

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



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
 Brazil Croatia Czech Rep. India Latvia Poland 	RussiaSerbiaSouth AfricaTurkeyUkraine

Non Functioning NET worldwide - CLARINET

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

Austria	• Italy
 Belgium 	Poland
 Czech Rep. 	Slovakia
Denmark	Spain
France	Sweden
Germany	• UK
India	• US

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



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	•	-	•	_	





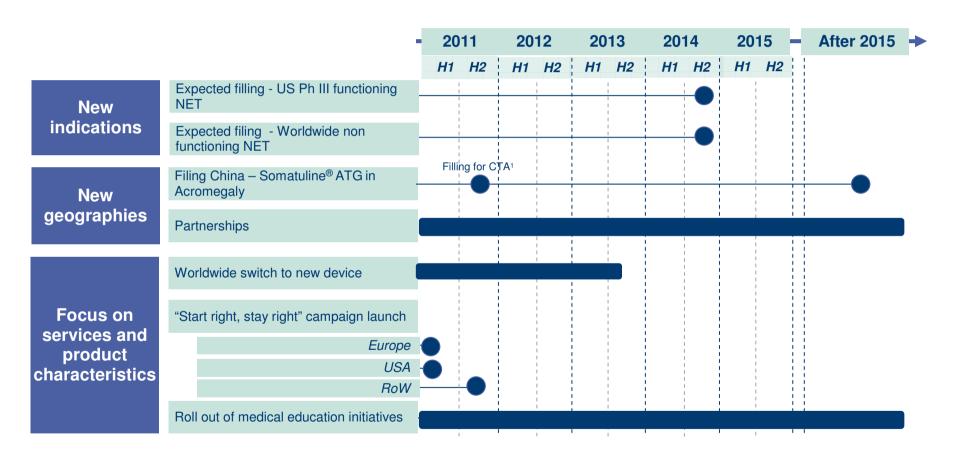
Somatuline® Autogel 2020: a globalized reach

Geography/ Indication	Europe	US	China	Brazil	Russia
Acromegaly	√	√	√	√	√
Functioning NET	✓	√		✓	✓
Non functioning NET	√	√		√	√
				✓ lp	sen presence

LatAm and Asia covered through partnerships



Somatuline[®] milestones



Professor Wouter de Herder

Professor of endocrine oncology at the Erasmus University, Rotterdam, Netherlands

Chairman and Vice-Chairman of ENETS (European Neuroendocrine Tumour Society)

Member of the advisory boards of ENETS and NANETS (North American Neuroendocrine Tumor Society)



Neurology/ Dysport®

C. Jean EVP, Chief Operating Officer





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)

2010 BonTA market figures ← Therapeutic → ← Aesthetics → ~€370m ~€420m ~€310m ~€240m **USA RoW USA RoW** Botox Dysport Others



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

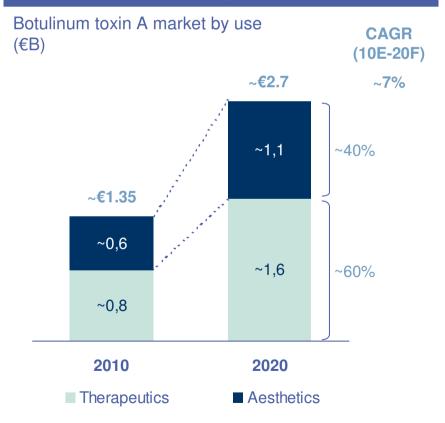
US to remain half of 2020 WW market ...

Botulinum toxin A market by geography (€Bn)

CAGR (10E-20F)



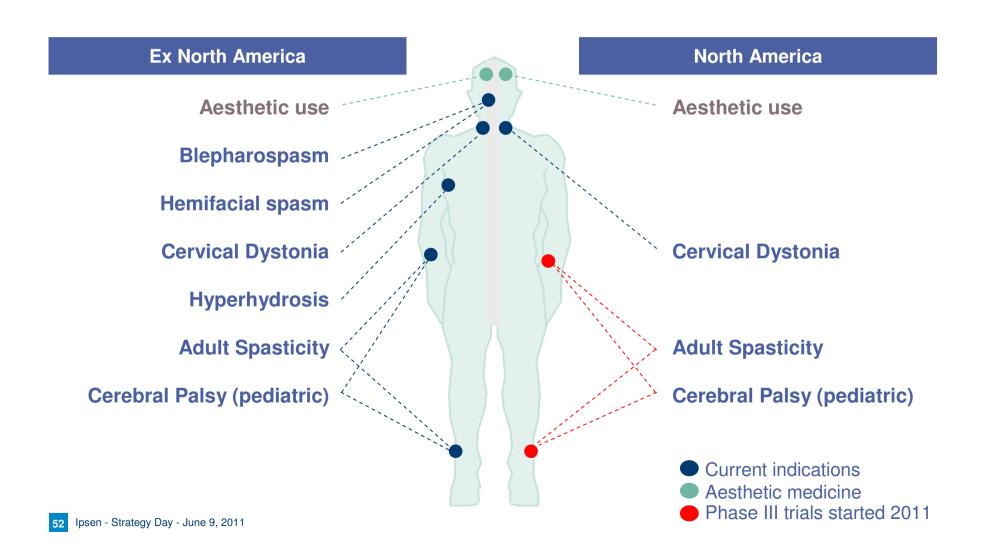
...with split between therapeutics and aesthetics remaining stable



Ipsen - Strategy Day - June 9, 2011 Source: Ipsen analysis



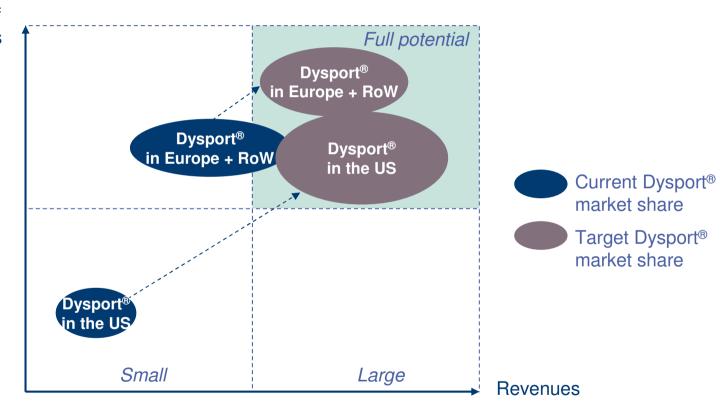
Room for new indications in North America





Full potential of Dysport® lies ahead...

Number of indications



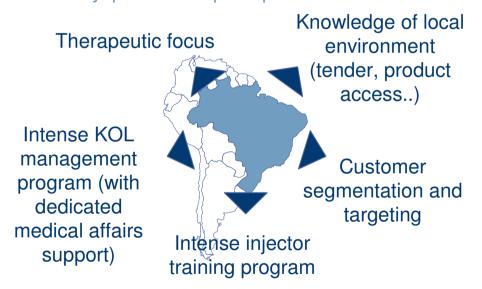
... and BonTA market is expected to grow 7% CAGR until 2020



Brazil, success built on focus and strong selling fundamentals

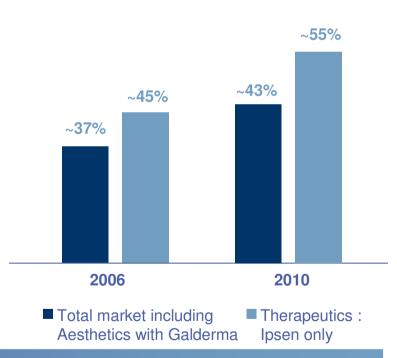
Focus and strong selling fundamentals...

Dysport®: The Ipsen product in Brazil



... to become market leader

Dysport®'s market share in Brazil1



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®)

Ipsen - Strategy Day - June 9, 2011

Note 1: Internal data



Commitment across the full value chain

Competitive landscape

Toxin Technological platform **Barrier to entry**

Competition

Market size/ Growth

Dysport® Market share

Geographies

Dysport® competitive adv.

Dysport® NG competitive adv.

Dysport[®] growth potential



Core to Ipsen

High

Well characterized – 2 major competitors. more to come – strong Aesthetics alliances

Large/ Solid – 7% CAGR until 2020

2nd after Botox® – room to grow

US and emerging markets potential

At par with competitors

Further differentiation with Next Generation

4 ongoing PhIII in spasticity...

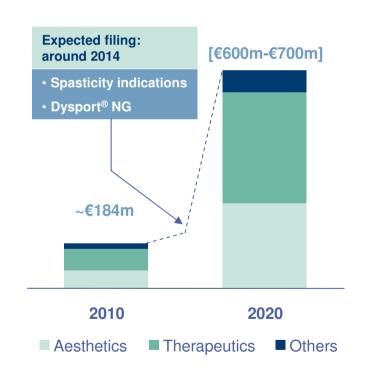


Dysport® ambition: 10% to 13% CAGR* until 2020

Driven mainly by US...

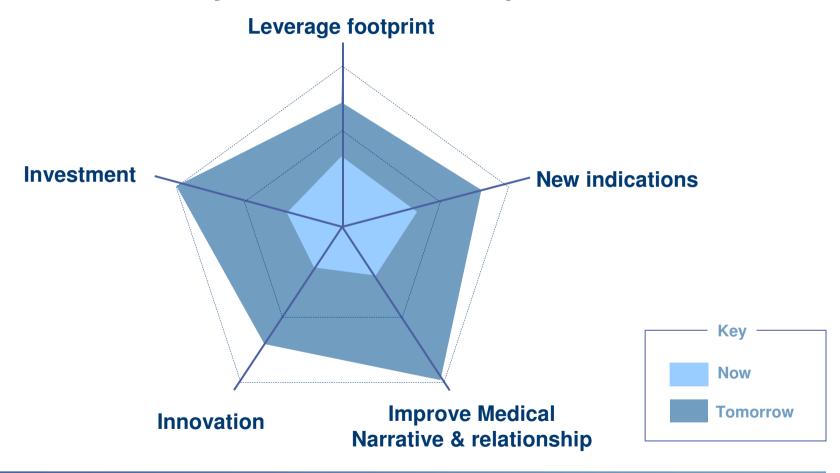
Expected filing: [€600m-€700m] around 2014 Spasticity indications Dysport® NG ~€184m 2010 2020 USA Europe RoW

...and therapeutics





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 Over- activity	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 Over- activity	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



Ipsen to improve medical narrative and prescriber relationship

Leverage unique clinical Dysport® value experience **Enhance collaborations with** medical and scientific **Enhance services to physicians** communities 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

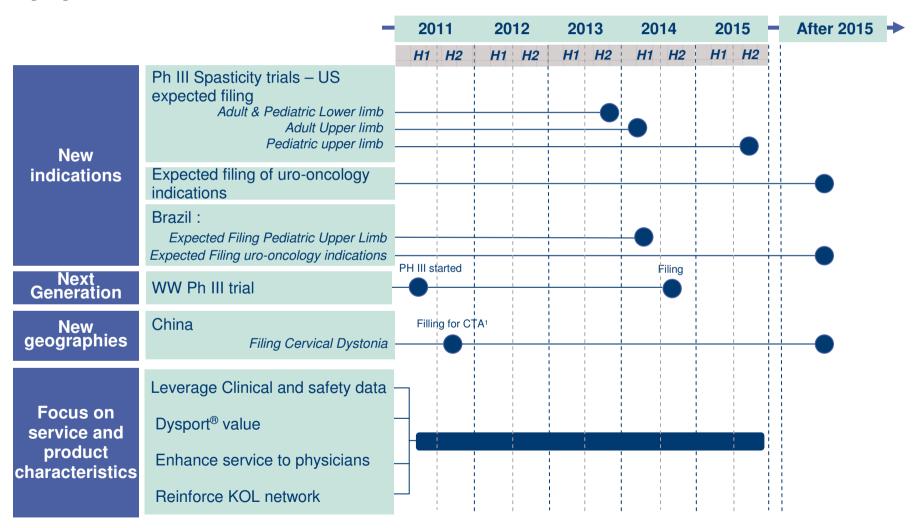
A potentially transforming project

- 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



Dysport® milestones:



Professor Pierre Denys

Professor of Physical Medicine and Rehabilitation at the Medicine University, Paris West, France

Hospital Physician in the Neuro-Urology Unit, Raymond Poincaré Hospital, Garches, France



Research & Development

C. Bertrand EVP, Chief Scientific Officer





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

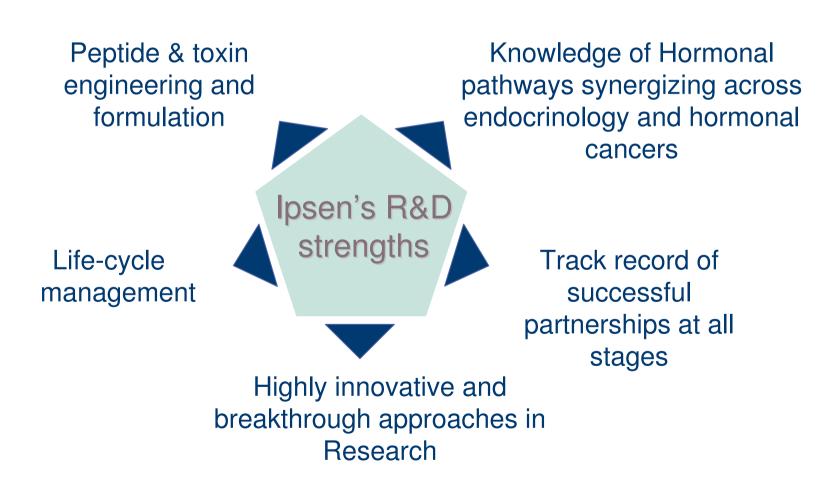
Focus on patients (Patient centric)

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

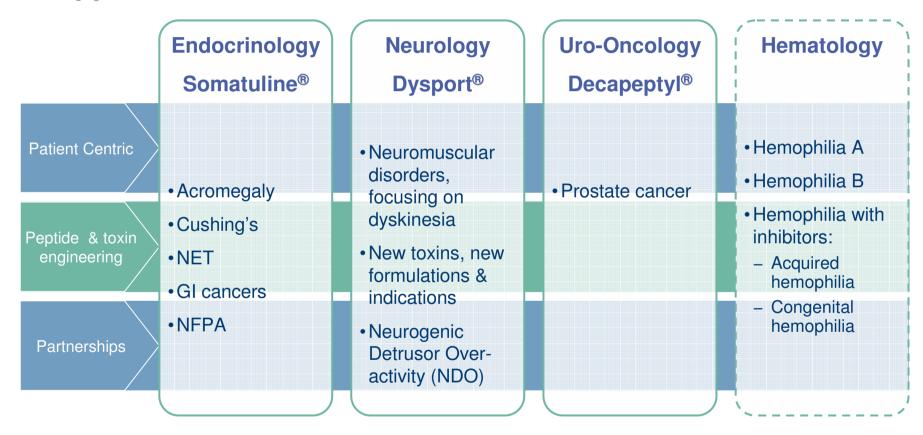
- Highly focused in targeted debilitating diseases
- Core Translational Sciences/
 Medicine capabilities
- Integrated R&D and lean organization
- Internal resources and competencies focused on peptides and toxins
 - Open Innovation & built from our strong partnerships

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





... applied to well defined disease areas and indications

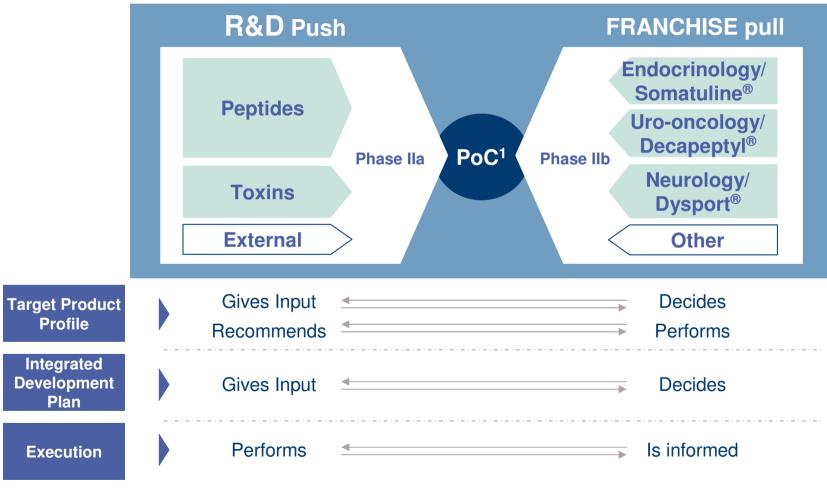


Focused. Aligned. Synergies across therapeutic areas





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

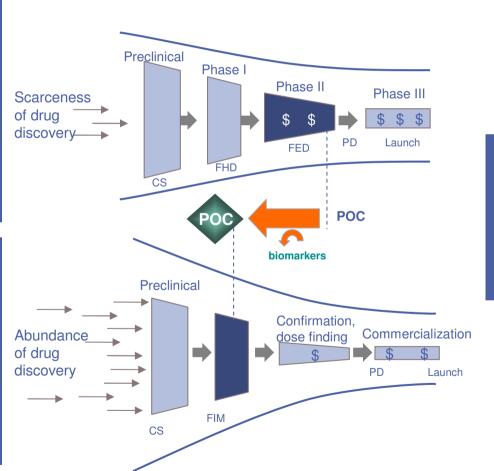






Biomarkers help compounds move to POC quicker while reducing failure rate in late development stages

Traditional Development approach



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

Proiect killing

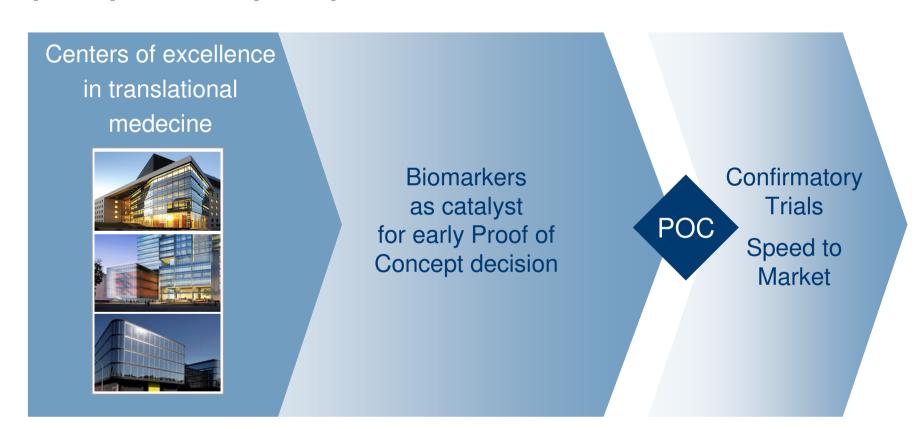
CS: Candidate Selection FED: First efficacy dose FIM: First in Man PD: Product decision POC: Proof of Concept

New approach





Ipsen partnerships in personalized medicine



Current partnerships : Centers of Translational Medicine & BioMérieux

Focus on two core technologies...





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

Enhance potency & efficacy

Improve target selectivity

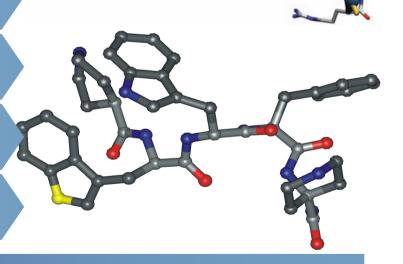
State-of-theart scientific expertise and technology

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

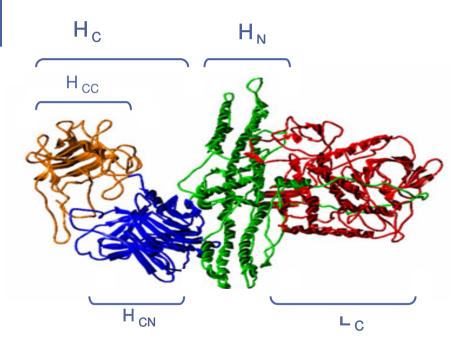




Botulinum toxin: Recombinant technology offers promise of development well beyond current marketed products

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



Hematology







2003 - 2005

2006-2011



Our key drivers entail major decisions

Summary
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



Key decisions on Ipsen's Ph II pipeline

Assets no longer in strategic focus

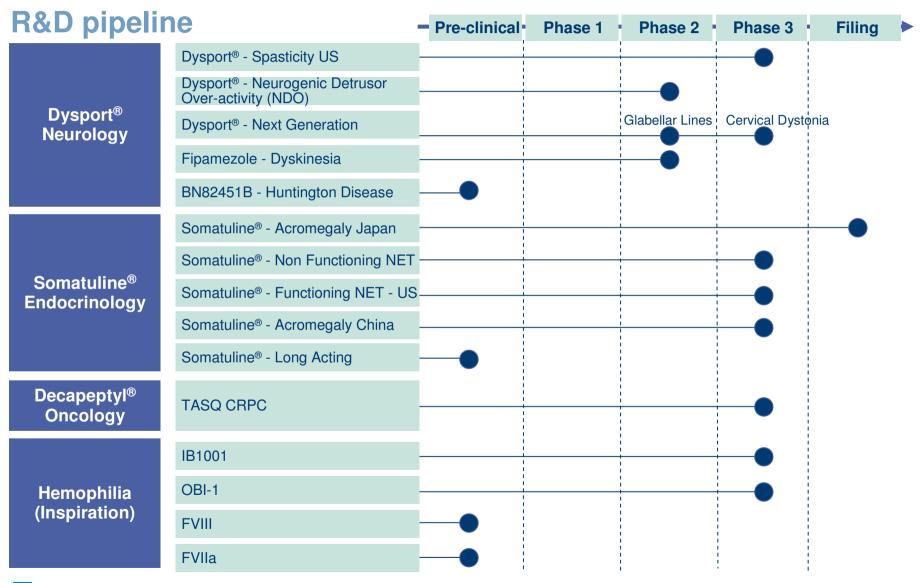
Oncology

- 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
 Ipsen 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

Q&A



Break

Presentation resumes in 20 minutes



Uro-Oncology/ Decapeptyl®

C. Jean EVP, Chief Operating Officer





A franchise with renewed growth opportunities

Tasquinimod

for castrate resistant tumors

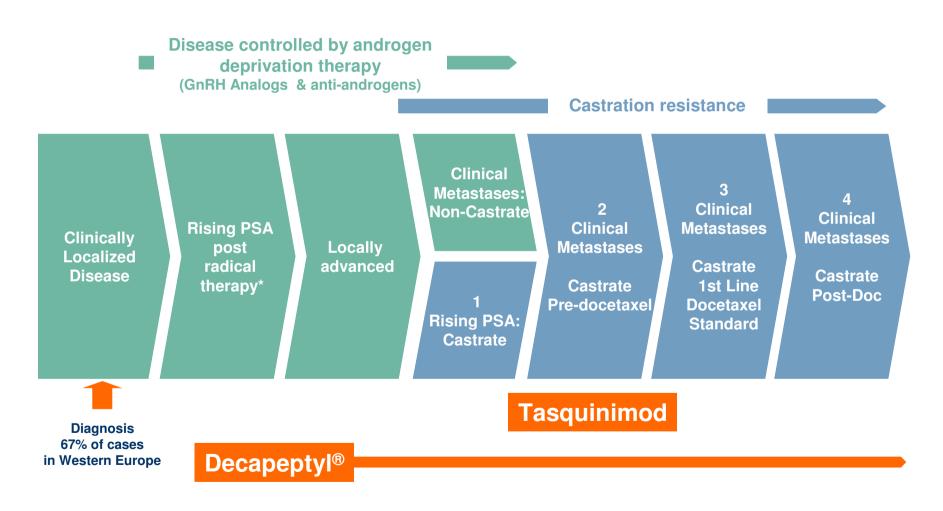
Once a day oral formulation in PhIII

Decapeptyl®

for hormone-sensitive tumors



Prostate cancer: disease evolution towards castration resistance





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

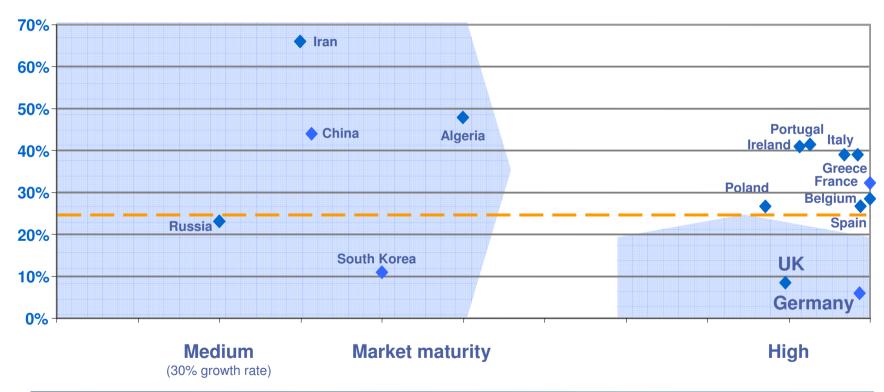
... and still poised to grow

- 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)

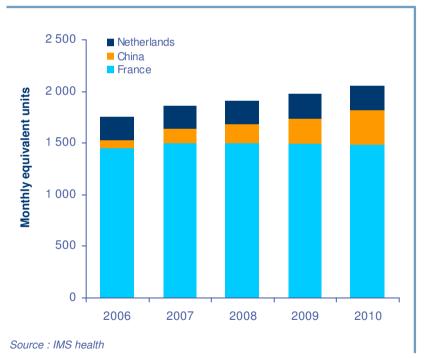


Market share in the UK and Germany (~40% of G5 GnRH Analogs sales) has doubled in the past 3 years

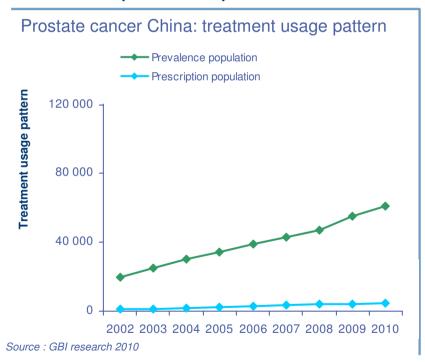


Emerging countries, and in particular China, as growth engine

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**

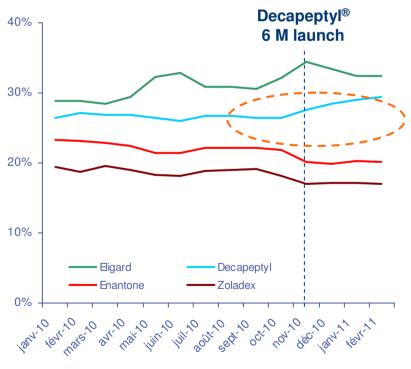
Local Tolerance

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

Formulation reconstitution

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

e.g. Impact on market shares in Spain



Source: IMS sales data

^{*}Lundstrom E & Al, , ClunDrug Invest 2009; 29(12):757-765

^{**}Mounedji N & al, J clin Oncol 29: 2011 (suppl 7; abstr 162)

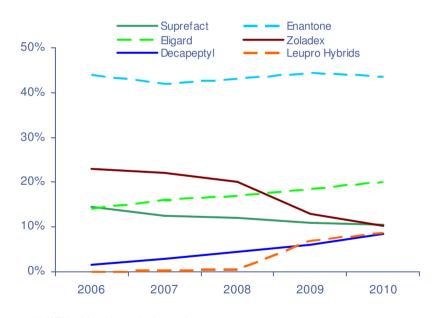


No true generics of GnRH analogs anticipated, only Hybrids

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

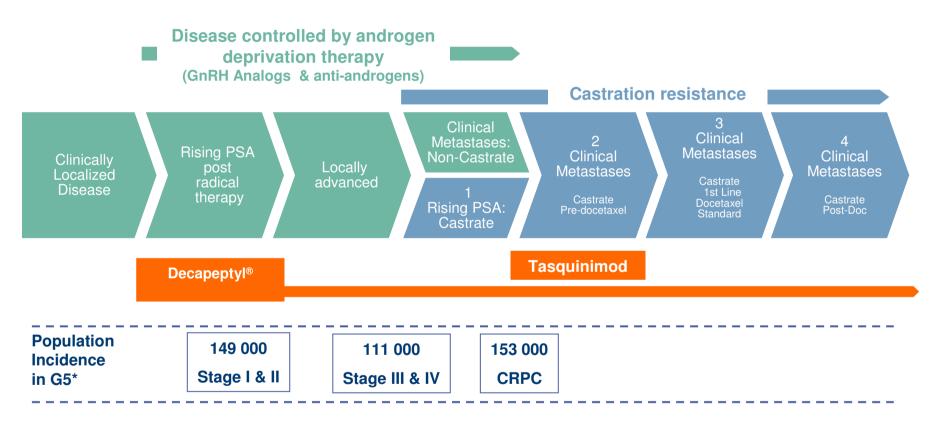


(*) MEU = Monthly equivalent units Source: Insight Health, OdV data - Germany

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



Tasquinimod: a perfect strategic fit



- 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)

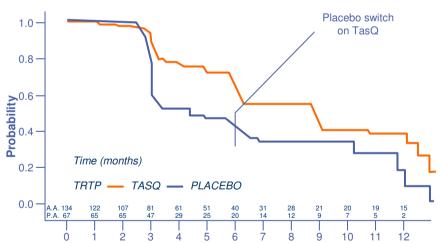


Tasquinimod, promising phase II results

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

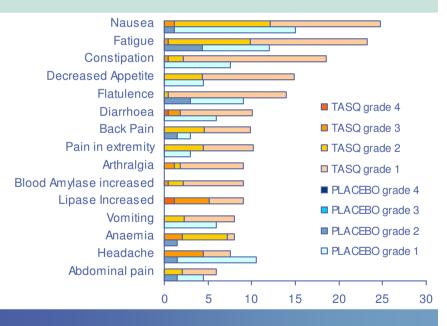
Primary end point Proportion of patients with progression at 6 months: n=134/67 31% in Taquinimod group vs. 66% in placebo group





Tasquinimod improves Radiographic Progression Free survival vs. placebo (8.8 months vs. 4.4 months)

Most common AE-s and percent of patients with grade 1-4 in Double-blind phase



Side effects are manageable

^{*} ASCO-GU, 2011, J. Armstrong¹, M. Haggman², W. M. Stadler³, J. R. Gingrich⁴, V. J. Assikis⁵, O. Nordle⁶, G.Forsberg⁶, M. A. Carducci⁷, R. Pili⁸



Tasquinimod, Phase III program ongoing

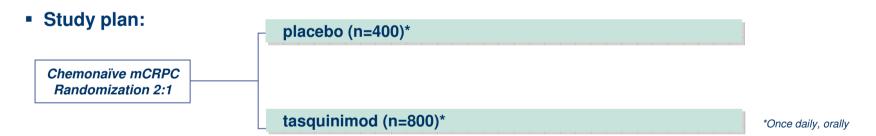
A Phase III randomized, double-blind, placebo-controlled study of Tasquinimod in men with asymptomatic/mildly asymptomatic Metastatic Castrate Resistant Prostate Cancer

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

M. de Garidel Chairman and Chief Executive Officer





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

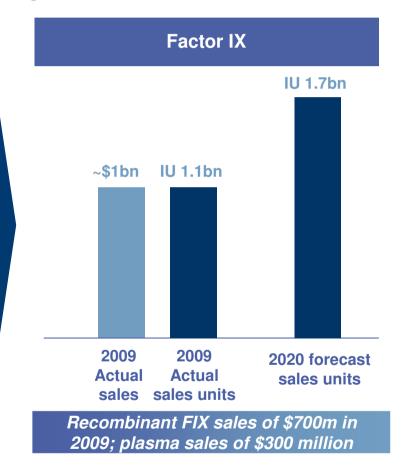
- FIX market overlooked primary focus FVIII
- 90% recombinant in developed markets
- 40% recombinant and growing in underserved markets

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



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



Historical and projected hemophilia market growth drivers

Hemophilia population growth

Diagnosis & treatment rate

Prophylaxis penetration

Conversion to recombinant

Increased competition

Increasing weight of demographics

Limited, longer-term opportunity

Market opportunity

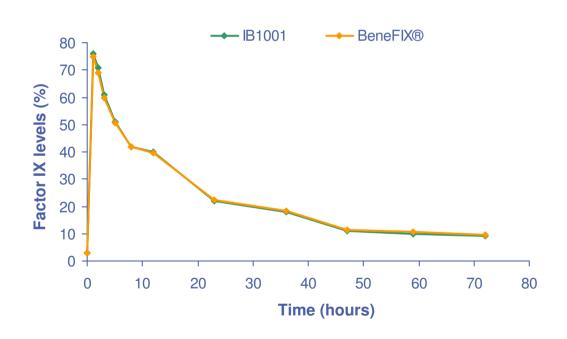
Market opportunity

Opportunity to enhance prophylaxis 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

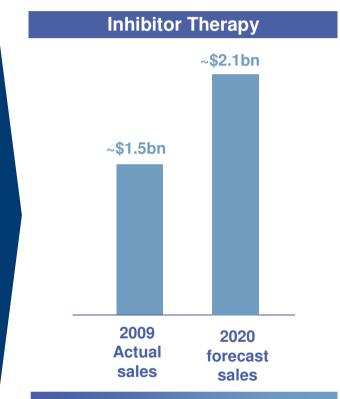
- 1/3 of hemophilia A patients will require inhibitor therapy during their lifetime:
 - Inhibitor therapy (IT) 70% recombinant in developed markets
 - IT <10% recombinant in underserved markets
 - Current therapies are FVIIa bypassing agents

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

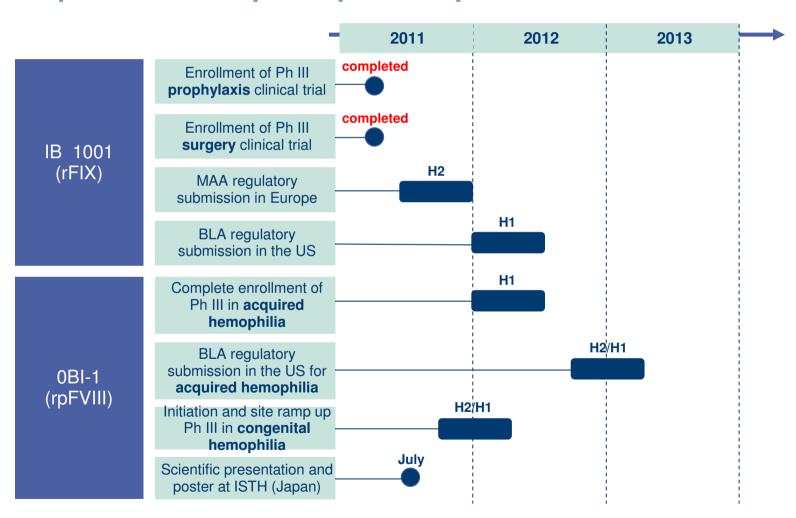


Recombinant FVIIa sales of \$1.1bn in 2009; plasma sales of \$400m

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

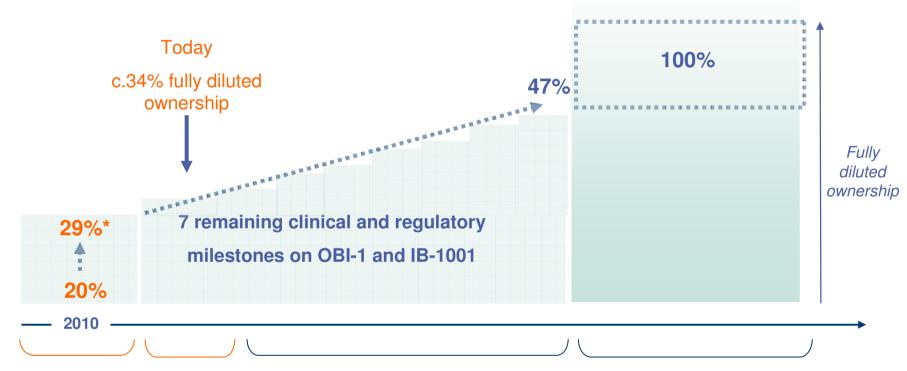


Inspiration hemophilia product portfolio – short term timeline





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



Initial equity **OBI-1 PhIII** stake: \$85 m initiation + OBI-1 \$50 m paid upfront: \$50 m by Ipsen in + 27.5% royalty exchange for rate on OBI-1 convertible bonds

Total development funding of \$124m in exchange for convertible bonds maturing the later of 7 years or the end of the call exercise period

Call at market value exercisable on triggering events expiring at the latest in 2019

Professor Claude Négrier

Head of the Hematology Department at Edouard Herriot University Hospital in Lyon, France

Professor of Hematology at the Lyon School of Medicine, France

Adjunct Professor in the Division of Hematology at the University of North Carolina, Chapel Hill, USA



US platform

C. Jean EVP, Chief Operating Officer





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

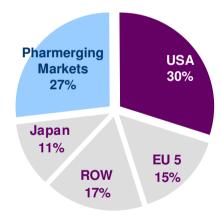




The U.S. will continue to lead the global marketplace and remains an important pillar in Ipsen's global strategy

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









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)

8 on-going phase IIIs in the US

- 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

- 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 NFT
- 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

E. Bouteiller General Manager, China







China amongst top pharma markets...

	2010 rank			2015 rank				2016 rank	
1	United States		1	United States			1	United States	
2	Japan		2	Japan		\longrightarrow	2	China	*1
3	China	*1	3	China	*)		3	Japan	

Region contribution to global growth

Region market share of global sales 2015



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

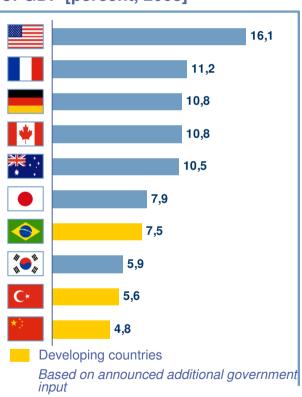
In 2010, China became the world's second largest economy ahead of Japan





...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

115 Ipsen - Strategy Day - June 9, 2011 Source: McKinsey analysis





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

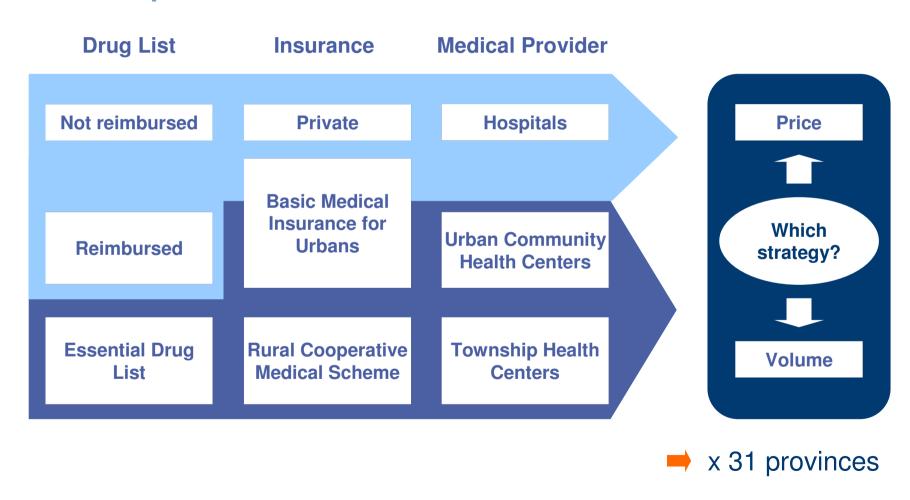
Healthcare Reforms:

- 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

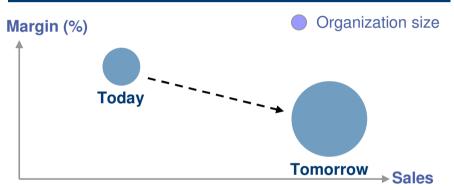






China today, a tougher place for new comers...

Increasing cost of doing business



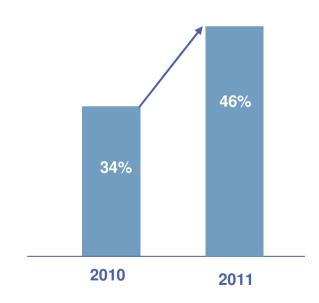
China is more and more costly:

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

with price cuts

China is increasingly discriminant

Perception governmental policies 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



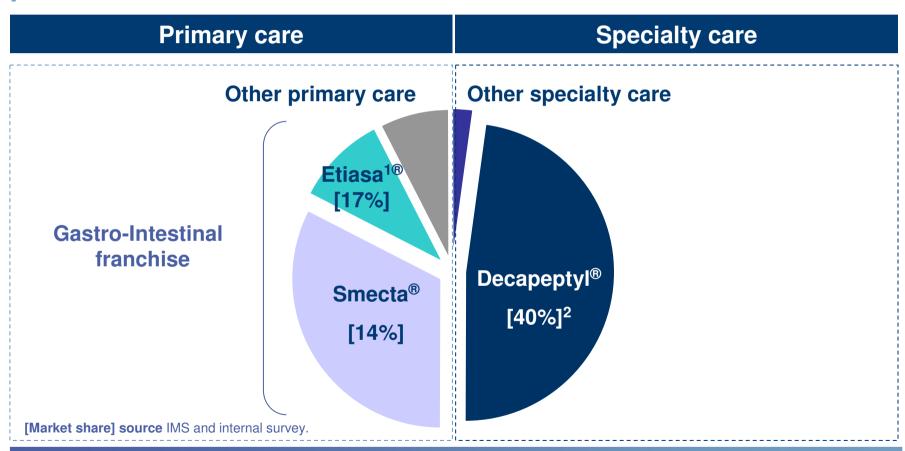
- Regional sales force teams focused on key coastal cities, now spreading towards the west into lower tier cities.
- Excellent National & Provincial market access team (patent, pricing, reimbursement, bidding ,etc.)
- Experienced medico-regulatory team to conduct local clinical and registration trials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



2005 - 2010 CAGR: 22% at constant exchange rate







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





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

2020 ambition

- Introduction of 2 major lpsen 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 and Trade representatives
- Highly 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

Extend and deepen coverage

- Triple sales force team by 2020

2020 ambition



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

...committed...

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

Primary Care France

M. de Garidel Chairman and Chief Executive Officer





Contrasted primary care dynamics



800 | 650 -500 -350 -

2007

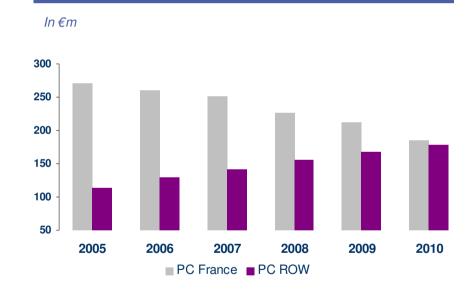
■ Total Specialty care ■ Total Primary care

2008

2009

2010

French vs. RoW primary care sales



Pressure on Ipsen's French primary care

2005

2006



A European-wide pharmaceutical industry situation

Large, mid and small Pharma situation in Europe

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

A growth potential to materialize

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

Conclusion

M. de Garidel Chairman and Chief Executive Officer





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



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



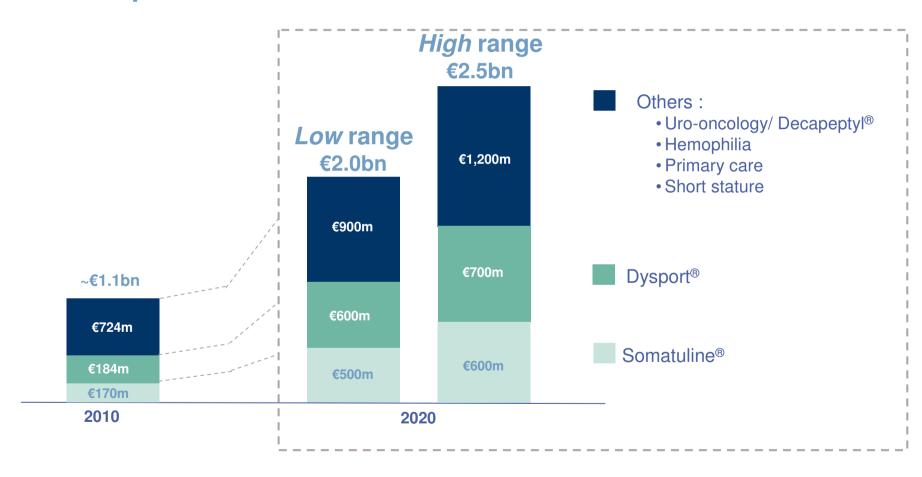
Closing of R&D activities of Barcelona site



 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

1

2011 - 2015 Invest to grow

Clinical trials:

- Dysport® in spasticity
- Dysport® in urology
- Dysport® Next Generation
- Somatuline® in NET:
 - Functioning
 - Non functioning

Leveraged geographies:

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

Overall efforts

Stronger commercial push on key products

2

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



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
V	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
V	Dysport® CD CTA¹ 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
	CTA ¹ 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 staffed		OBI-1 PhIII (Acquired H) enrollment completed				Dysport® P.U.L filedin Brazil	
	Franchise org. implemented		OB-1 PhIII Congenital H.			Inspiration option assessment Smecta® EDL		
	IB1001 filed in Europe		US platform reorganized				assessment (China)	

Thank you



Q&A

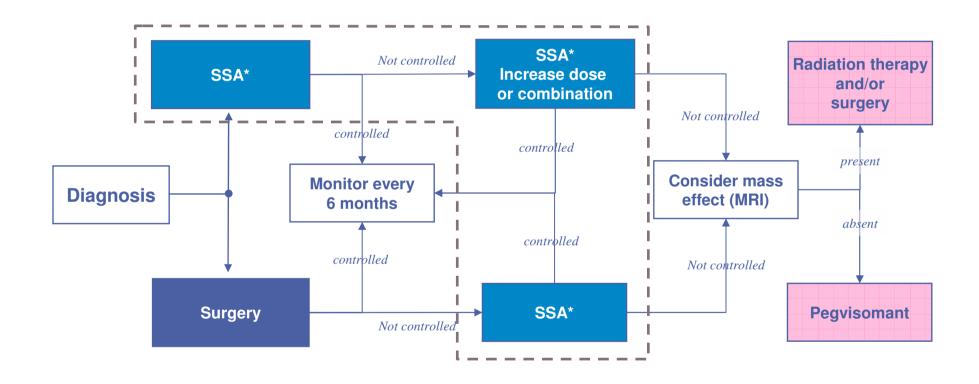


Appendices





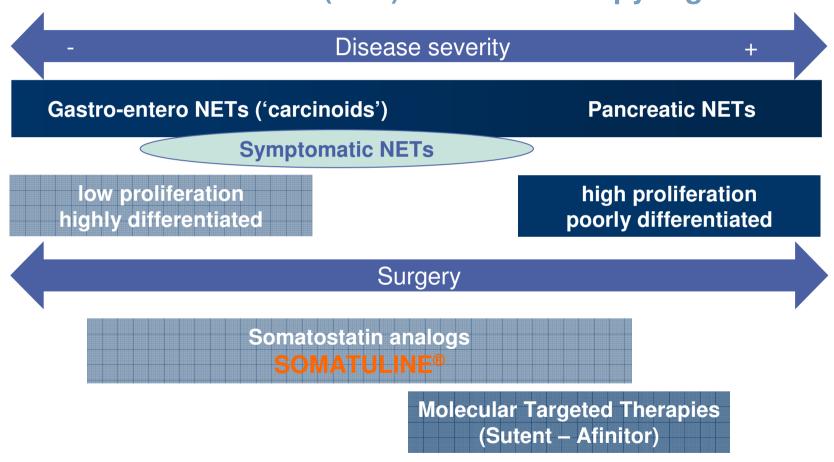
Acromegaly: current therapy algorithm



*SSA = Somatostatin Analogs



Neuro Endocrine tumors (NET): Current Therapy Algorithm



Cytotoxics