Confirming Ipsen's biopharmaceuticals profile

Exane 12th Healthcare Conference *Paris, May 10, 2010*



Jean-Luc Bélingard – Chairman and Chief Executive Officer Pierre Kemula – Investor Relations Officer



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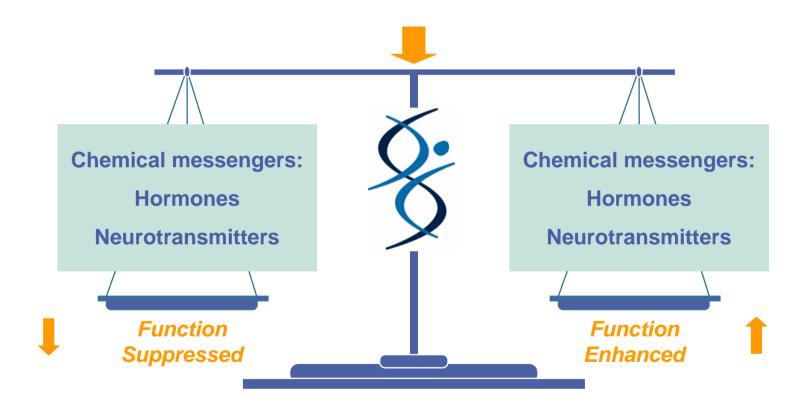
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Quick overview of some fundamentals





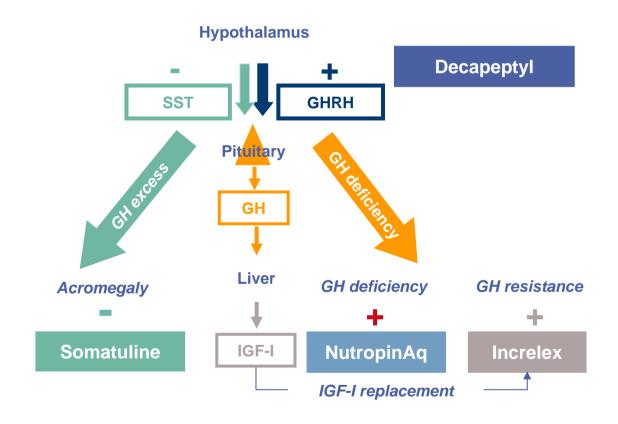
The fundamentals: restoring physiological balance



Therapeutic corollary: restore physiological levels, no more, no less



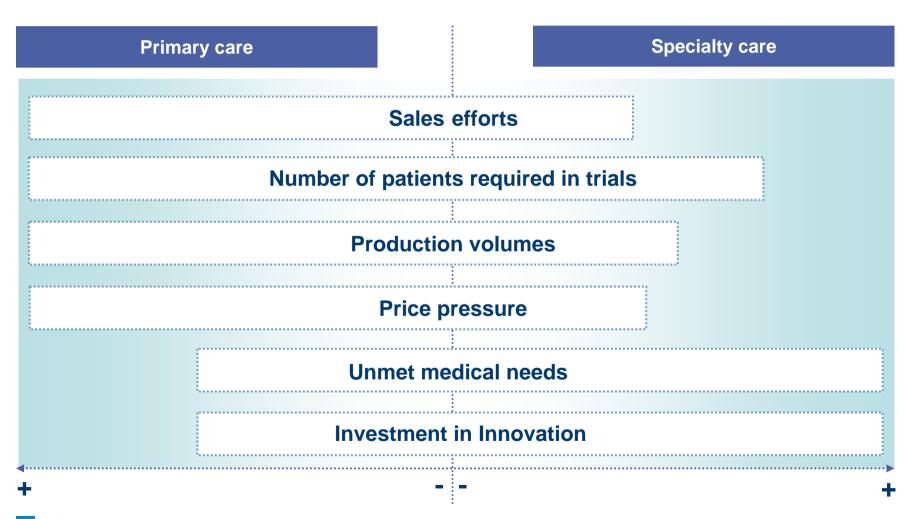
The fundamentals: the hormonal pathway and cascade



Ipsen has a specific knowledge of pituitary disorders



The fundamentals: Ipsen's specialty care vs. primary care businesses

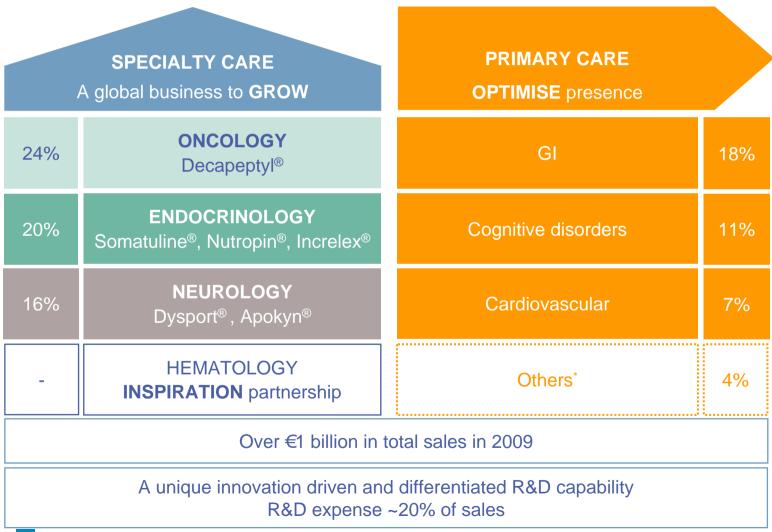


Ipsen in short

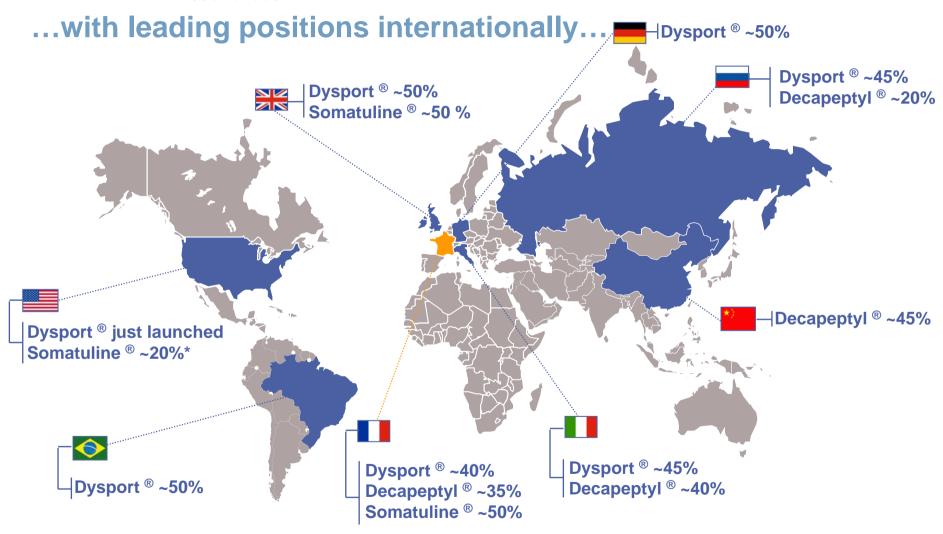




Ipsen today: a global biotech specialty pharma...







2008 rounded Market shares, except for Somatuline® in the US: 4Q09

Market shares are for (i) Dysport® in medical indications (ii) Decapeptyl in prostate cancer (China gynecology only) and (iii) Somatuline in Acromegaly only.

* In the US, market share in patients treated by endocrinologists and in pituitary centers

Sources: IMS, Insight Health/ODV, Ipsen estimates, Strategix



...continuously outgrowing its main competitors

2009 sales of Dysport® +18.0% y-o-y at constant currency

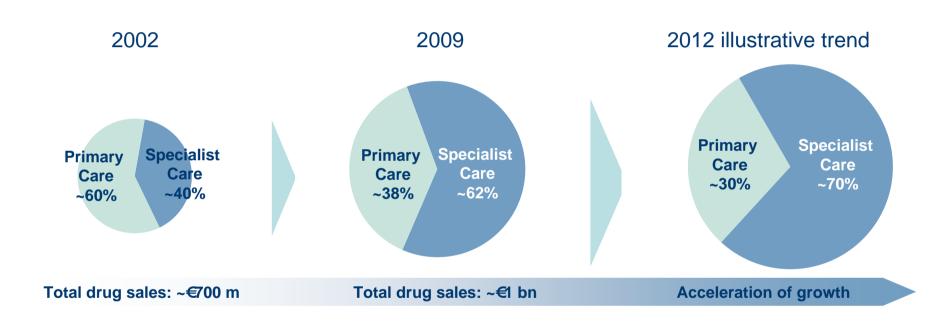
2009 sales of Somatuline® +18.2% y-o-y at constant currency

2009 sales of NutropinAq[®] +26.5% y-o-y at constant currency

Strong and continued specialty care dynamics +13.9% y-o-y at constant currency



Today, a reinforced profile...



Confirming Ipsen's biotech specialty care profile

Truly Differentiated R&D Capabilities

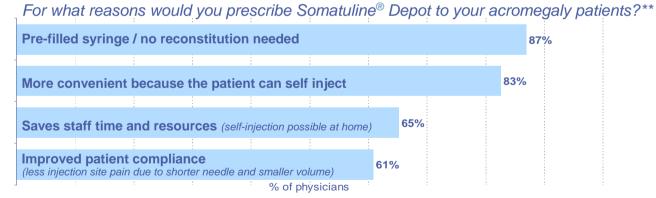




Example 1 : Somatuline® Depot

		Sandostatin LAR®	Somatuline® Autogel®		
Administration		2.0 ml Intramuscular	0.3 ml – 0.5 ml Subcutaneous		
Presentation		Powder vial + solvent filled syringe + 2 needles	Pre-filled syringe		
Injection technique		10 steps needed to reconstitute	Ready to use Self administration*		





Somatuline® Depot: A self administration syringe, an improved quality of life

* In selected countries



Example 2 : a unique technology convergence, taspoglutide

		Liraglutide Flex Pen	Byetta LAR*	Taspoglutide PFS**	
Administration		Daily	Weekly	Weekly	
Device		Multiple use Needle: 32 gauge	Single-use, syringe & vial Needle : 23 gauge	Single-use, staked-in needle Needle: 29 gauge	
Injection technique		1.Attach needle 2.Prime pen 3.Set dose 4.Inject	1.Re-constitute 2.Draw into syringe 3.Inject	1.Remove cap 2.Inject	







Taspoglutide: 50 to 300 µl of highly concentrated aqueous solution devoid of excipient

Selected potential future growth drivers





Snapshot

Internationalization

Fast growing presence in Russia, China, Brazil and emerging markets

Entering the US

Life cycle management

Somatuline IGF-I+GH coadministration Decapeptyl 6-month

Hematology

Capitalizing on OBI-1 to build a unique portfolio of compounds in hemophilia

Rich R&D pipeline

Oristusane BIM 23A760

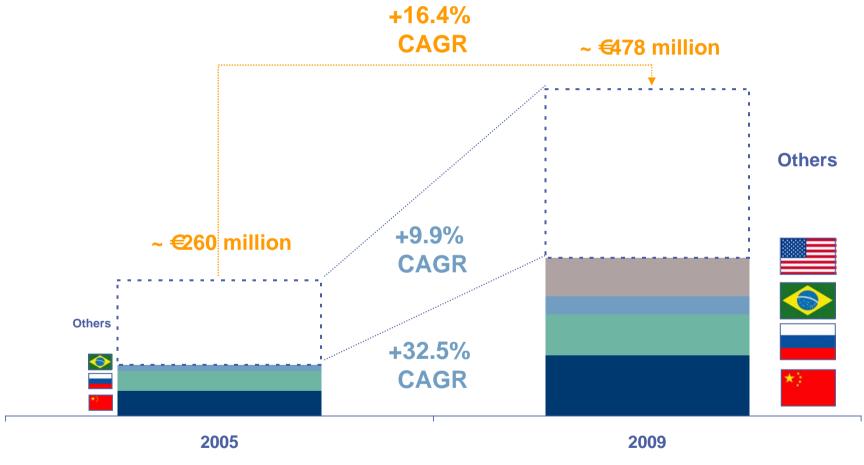




Growth and globalization beyond G5... A diversified portfolio in the fastest growing Eastern European pharma market **Entering the world's largest pharma market** with 4 specialty care products Dysport® Decapeptyl® Dysport® Tanakan[®] Somatuline® Smecta[®] Increlex® **Ipsen's G5** Apokyń **Dynamic specialty care** presence in the largest South **American pharma market** Longstanding and strong positions in one Dysport[®] of the world's largest pharma market Partnership with Galderma for aesthetics Decapeptyl® Smecta[®]



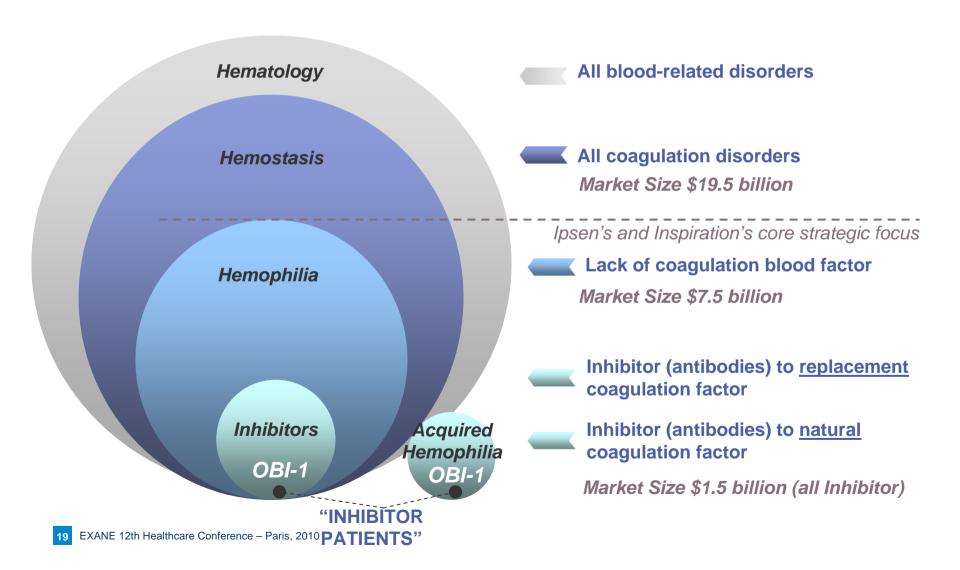
...translating into rapid expansion



Evolution of Group sales outside the G5

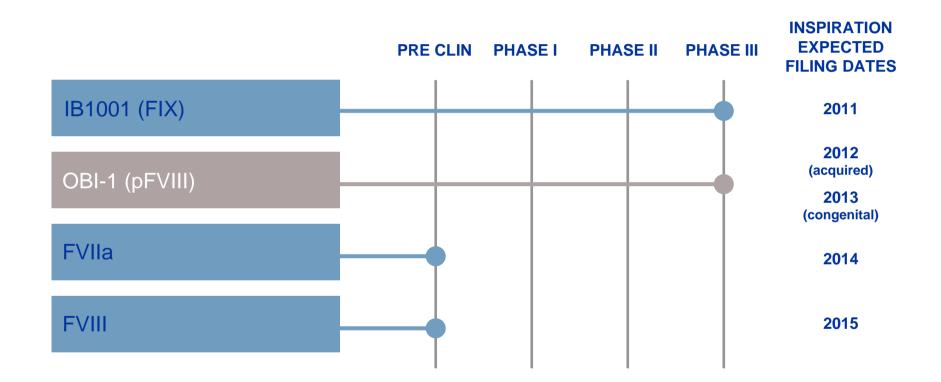


Capitalizing on OBI-1 with the Inspiration partnership...





....to build a unique franchise covering all needs in hemophilia...



A recombinant product in each segment of the hemophilia market



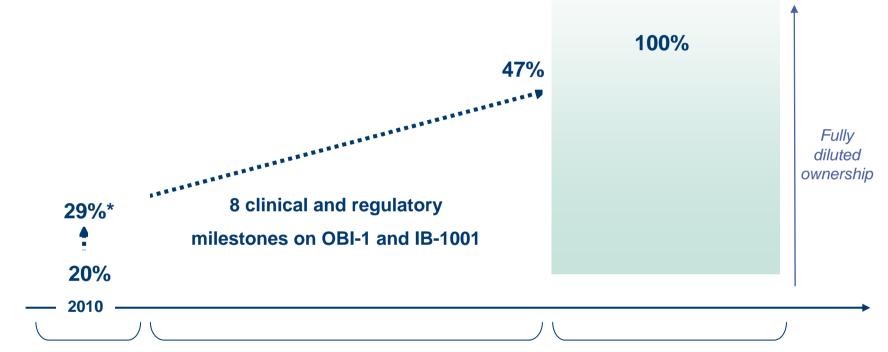
... with no equivalent currently on the market

	Inspiration	Wyeth	Baxter	CSL Behring	Bayer	Novo Nordisk	Biogen Idec and Biovitrum
FIX	Phase III	BeneFIX	Phase I	Mononine plasma derived	-	Phase I	Phase III (fusion protein)
OBI-1	Phase III	-	-	-	-	-	-
FVIIa	Preclinical	Preclinical	-	Preclinical	Phase I	Novoseven	-
FVIII	Preclinical	Refacto Xyntha	Advate Recombinate	Helixate	Kogenate	Phase III	-

Inspiration will be the company with the most comprehensive portfolio of hemophilia solutions



A progressive path of control of Inspiration



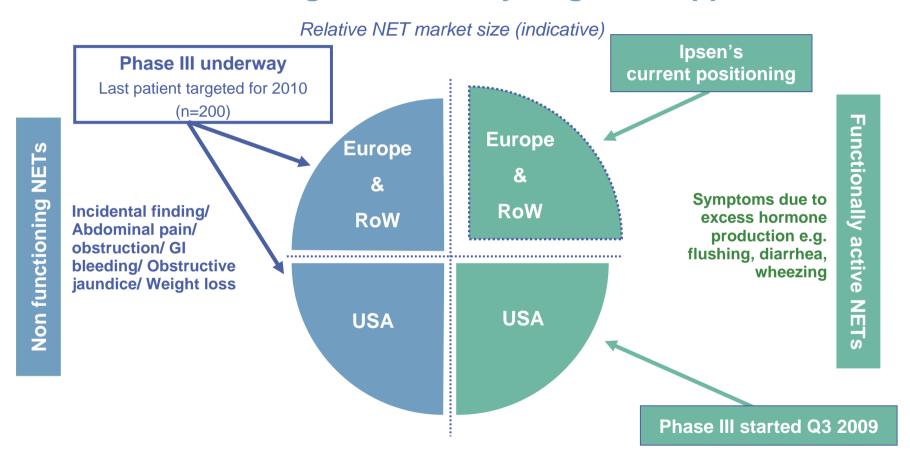
Initial equity stake: \$85 m + OBI-1 upfront: \$50 m + 27.5% royalty rate on OBI-1

Total development funding of \$174m 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



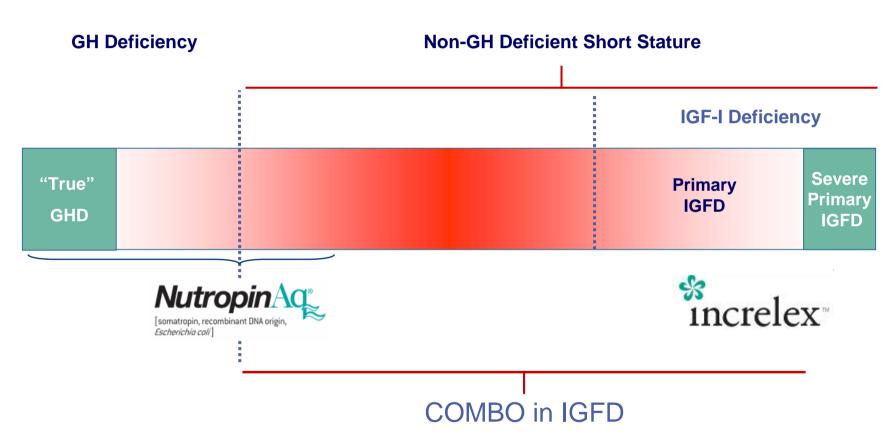
Somatuline® offers significant life cycle growth opportunities



Significant scope for expansion



Redefining the treatment of short stature

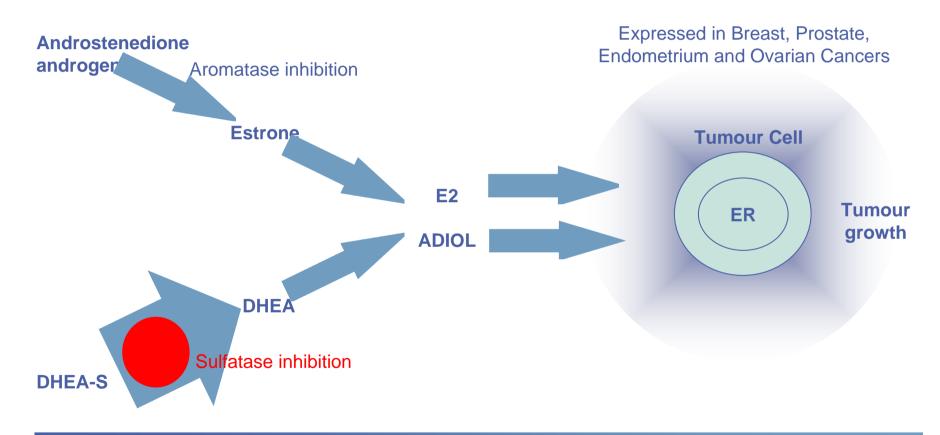


- MS316: Ph II dose titration study recruitment completed in Q2 '09
 - Interim results announced in September 09
 - Complete results available in H2 '10



Rationale for Sulfatase inhibitor development

Inhibition of Androstenediol synthesis from DHEA-S



After encouraging phase I results, Oristusane is now in phase II clinical trial





Phase I Phase II

ENDOMETRIAL CANCER

Phase II initiated - 80 patients to be treated

First patient dosed on November 25, 2009

METASTATIC BREAST CANCER (ER)

Phase I/II on going - 50 patients

Preliminary results presented at the SABCS (dec. 09).

OBD determined: 40 mg 1 per day

Additional 15 patients included to study target enzyme (STS) inhibition in cancerous cell

PROSTATE CANCER

CASTRATE RESISTANT

Phase I/II on going

Dose escalation

Progress and Outlook





2009: major initiatives, in a rigorous execution of the Group's strategy

Grow and Globalize Ipsen's specialty care business

4 products in the US, 3 global

6 products in launch phase

Decapeptyl® 6M approved in Europe

Rich phase II/ III programmes

Out licensing of none core compounds



Optimize Ipsen's *primary care* business

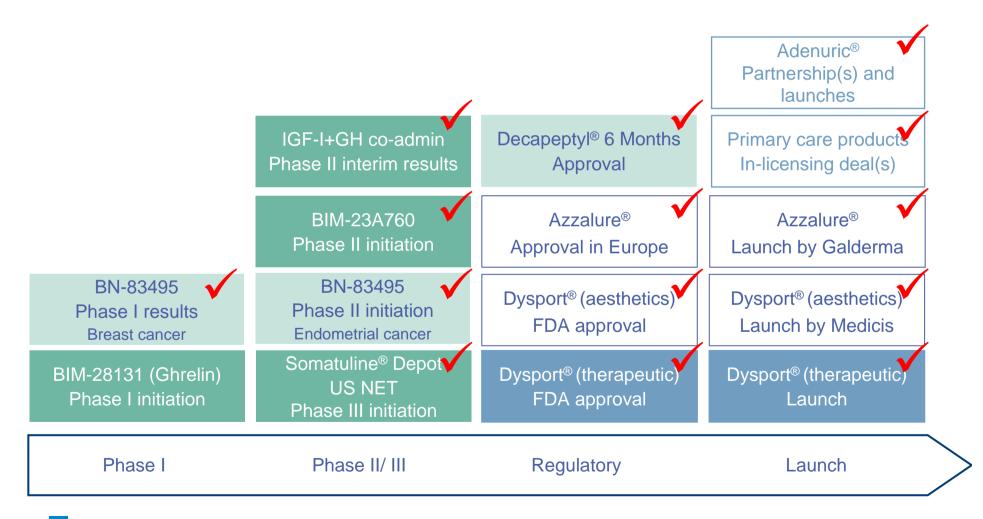
Rich deal with Menarini on Adenuric®

Promising headline results for taspoglutide (Roche)

BLI-800 (Braintree)

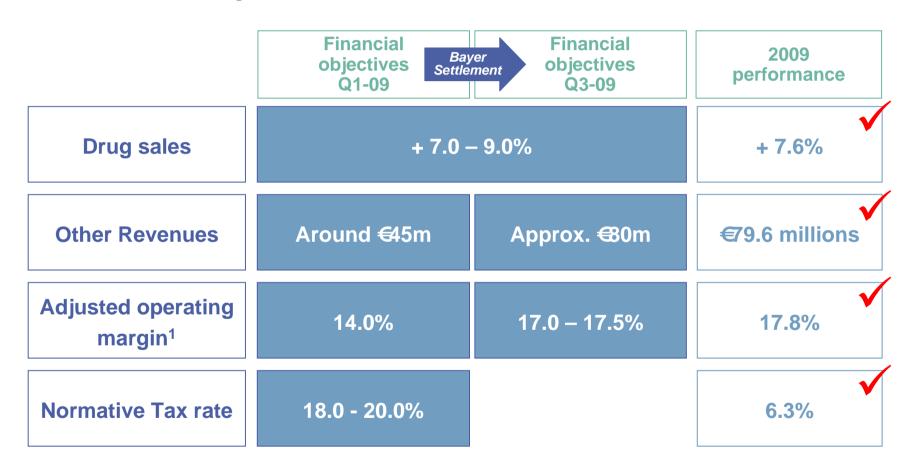


All key milestones delivered in 2009





All financial objectives have been met in 2009





A rich newsflow in 2010, already initiated

BIM-28131 (Ghrelin)
Partnered

IGF-I+GH co-admin
Phase II
Data available

BIM 23A760
Phase II initiation (WW)
Neuro Endocrine Tumors (NET)

Taspoglutide
Phase III results
Diabetes

OBI-1
Phase III initiation

Dysport®
Phase III initiation (US)
Adult+children lower limb spasticity

Tanakan® Phase III data available (GuidAge®) Somatuline® Depot US filing Extended Dosing Interval

Somatuline® Autogel® Launch in Russia Acromegaly + NET

Decapeptyl® 3 Months Launch in China Prostate cancer



Adenuric[®]
Launch



Clinical development

Regulatory / Launch

Continuous and rigorous execution of Ipsen's strategy

- → Specialty care growth & globalization, and increase in R&D efficacy
 - → Primary care contribution optimization



All financial objectives for 2010

Specialist Care - Drug sales

Primary Care - Drug sales

Other Revenues

Recurring Adjusted operating margin**

Recurring Adjusted EPS**

Financial objectives 2010

Close to double digit growth

(5)% to (7)% decrease*

3% to 5% growth*

Close to 50€million

Approx. 15% growth*

Relative stability with 2009

The above objectives are set at constant currency 2010 objective excludes any potential non recurring items

* Margins expressed in % of sales

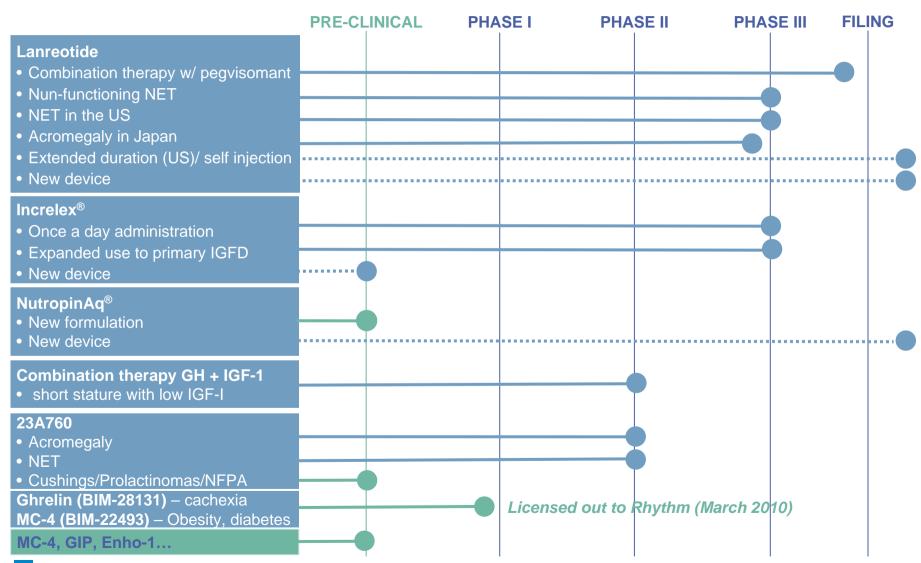
^{**} Prior to any accounting implications in connection with the purchase accounting of its acquisitions in North America and non recurring elements

APPENDIX R&D Pipeline



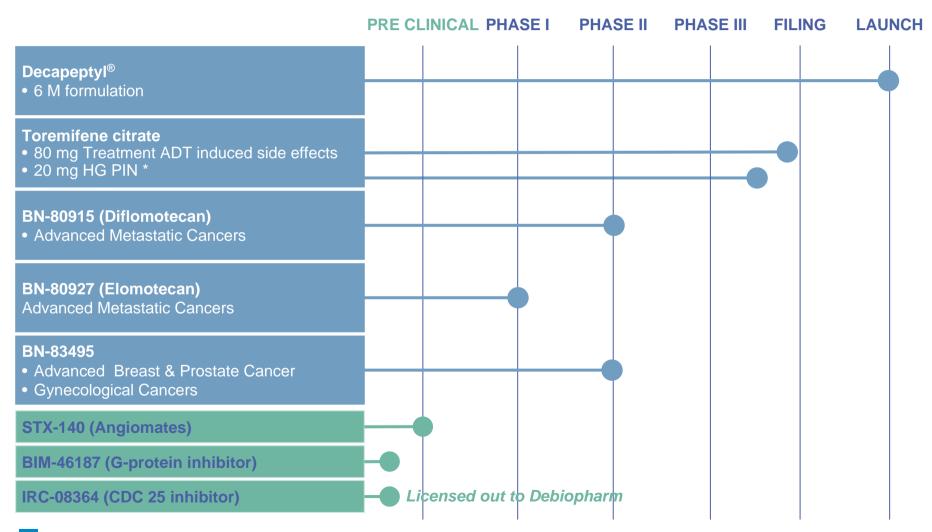


A rich endocrinology pipeline





A promising Oncology pipeline





Taspoglutide: Potential Best-in-class profile

Reported

T-emerge programme:

- -T-E 1: Taspoglutide vs. placebo
- -T-E 2: Taspoglutide QW vs. exenatide BID
- -T-E 4: Taspoglutide vs. sitagliptin vs. placebo
- -**T-E 5:** Taspoglutide vs. insulin glargine
- -T-E 7: Taspoglutide vs. placebo (metformin failures [high BMI])

Toxicology data:

- -Results of carcinogenicity studies support the ongoing clinical development of taspoglutide;
- -These results apply to mice as well

Filling during 2011

To be reported

T-emerge programme:

- -T-E 3: Taspoglutide vs. placebo (pioglitazone + metformin failures)
- -T-E 6: Taspoglutide vs. pioglitazone

Full body of data (efficacy, safety, potential side effects):

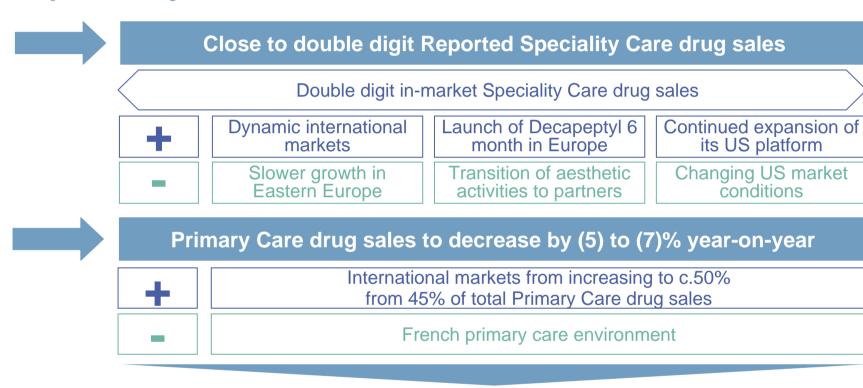
> -American Diabetes Association, June 26-29 2010

APPENDIX Financial objectives





Top line objectives for 2010



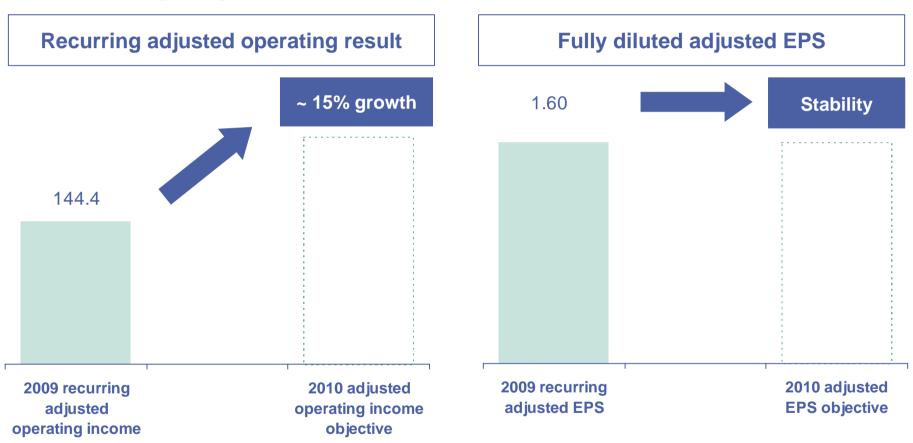
Group Drug Sales growth between 3.0 and 5.0% year-on-year



Other Revenues close to €50 million depending on the performance of the Group's partners



Profitability objectives for 2010



The Group targets an increase of its adjusted operating result and a relative stability of its consolidated income in a context of a significantly expanded R&D footprint



In the longer term...

July 2008 2009 2010 2011 Longer term

High US double-digit growth

coupled with significant profitability improvements

Continued US penetration

with 4 marketed specialty care products, of which Dysport® just recently launched

Financial crisis

profound changes in global equilibrium and macroeconomic conditions

Increased primary care competitive environment

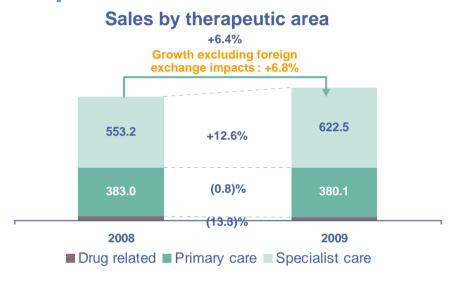
The Group today cannot confirm its 2011 and 2012 perspectives, or at least their timeframe

APPENDIX Full year 2009 detailed financial performance





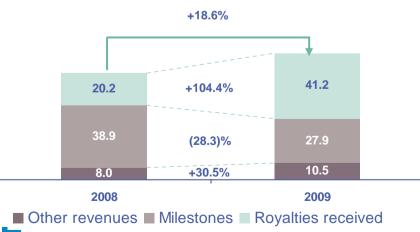
Top line evolution



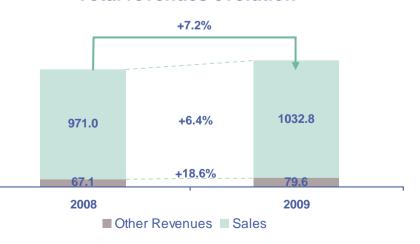
Sales by region +6.4% Growth excluding foreign exchange impacts: +6.8% 559.5 (0.9)% 554.7 236.2 (0.8)% 234.3 45.7 164.1 +20.8% 198.2

■ ROW □ North America ■ Other European Countries ■ European G5

Other revenues evolution

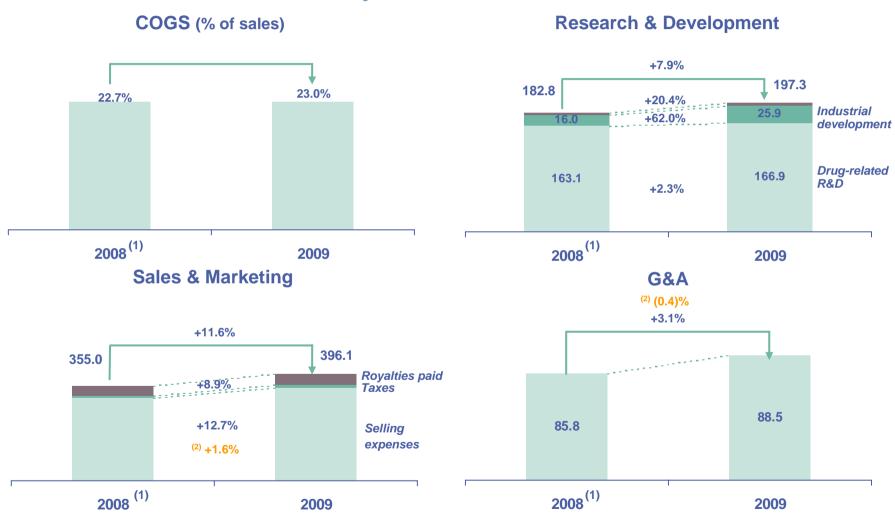


Total revenues evolution



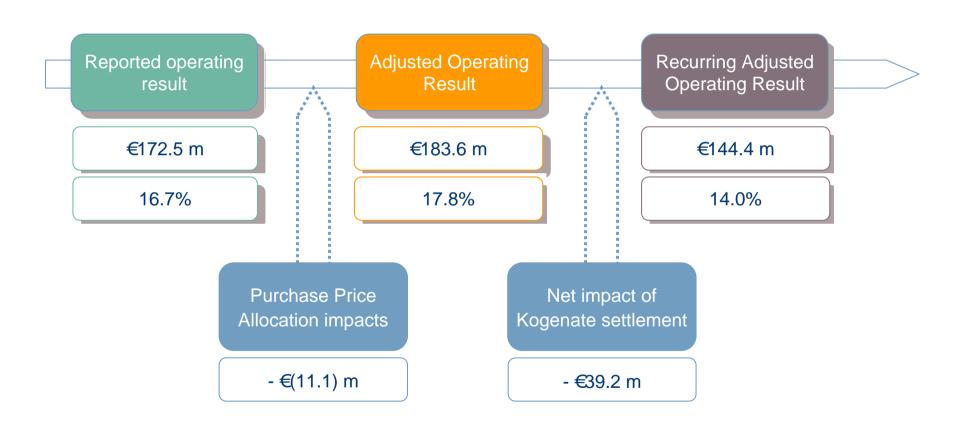


Evolution of main P&L expenses





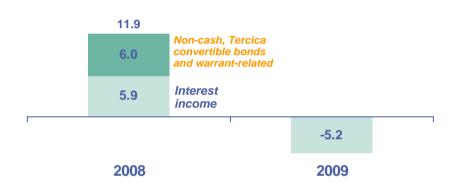
P&L – 2009 operating result and margin



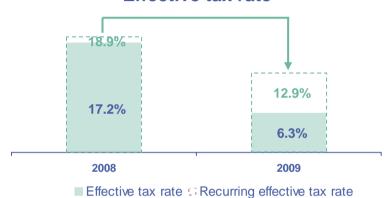


P&L - below EBIT

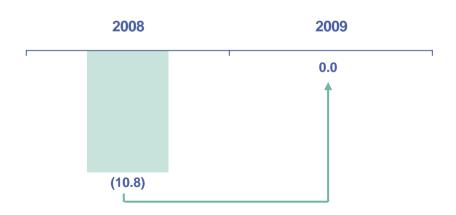
Financial result (€m)



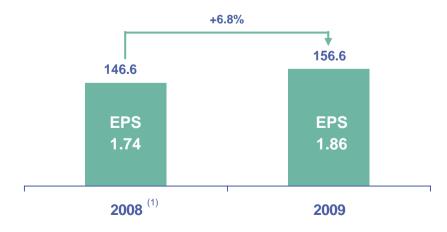
Effective tax rate



Income from Associates (€m)



Consolidated result (€m - group share)





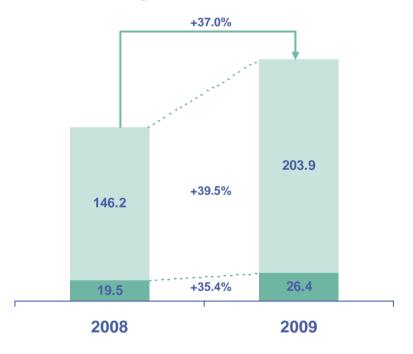
Balance Sheet evolution

(€m) Assets			Liabilities		
	31 Dec 08	31 Dec 09		31 Dec 08	31 Dec 09
Goodwill	290.8	290.2	Equity	885.0	982.6
Property. plans & equipments	237.9	251.8	Minority interests	1.6	1.7
Intangible assets	232.9	237.0	Total equity	886.6	984.3
Other non-current assets	112.9	145.5	Long-term financial debts	162.7	12.2
Total non-current assets	874.5	924.5	Other non-current liabilities	194.2	270.3
Total current assets	688.6	652.4	Short-term debts	10.6	21.4
Incl. cash and cash equivalents	239.6	218.6	Other current liabilities	305.4	286.7
Discontinued operations	1.3	0.0	Discontinued operations	4.9	2.0
Total assets	1,564.4	1,576.9	Total Liabilities	1,564.4	1,576.9
Net Cash	66.2	185.6			



Significant increase of partnership related deferred revenues

Total Milestones cashed-in but not yet recognised as revenues



- Payments recognised as revenues in n+2 and beyond
- Payments recognised as revenues in n+1

Main milestones cashed-in in 2009

May

Medicis: \$75m upon approval of Dysport®

March - September

Galderma: €20m upon approval and
launches of Azzalure®

October

Menarini: €20m upon signing of partnership for Adenuric®



Cash flow statement

€m	31 Dec 08	31 Dec 09	
Cash Flow before change in working capital	196.3	192.7	■ Net inc
- Increase/ Decrease in working capital	7.4	64.9	partner
Net cash flow generated by operating activities	203.7	257.6	
Investment in tangible assets	(61.4)	(40.3)	
Investment in Intangible assets	(33.8)	(24.7)	
Others	(190.3)	(6.3)	
Net cash flow used in investing activities	(285.5)	(71.3)	
Net change in borrowings	141.0	(151.3)	■ Reimbu
Dividends paid	(55.0)	(58.0)	credit fa
Others	(7.0)	(5.4)	
Net cash flow used in financing activities	79.0	(214.8)	
Discontinued operations	0.7	(1.0)	
Change in cash and cash equivalent	(2.1)	(29.5)	
Impact of exchange rate fluctuations	(1.5)	(2.4)	
Closing cash & cash equivalents	237.3	205.4	
Closing Net Cash	66.2	185.6	

ncrease in deferred revenues from erships: +€73.9m

bursement of syndicated t facility €(150) m

Appendix

Focus on the performance of Ipsen's US franchise





US platform integrated and fully operational

Strong and experienced management team :

New President and General Manager (May 2009)

New leadership team in key positions (Clinical & Medical Affairs, legal, HR, ...)

Fully operational managed care organisation (22 FTEs), including Payer Relation Management

Customer support programs in place, essential to US success Implementing PACE program (Patient assistance, Access to services, Continuity of care, Education) for each product

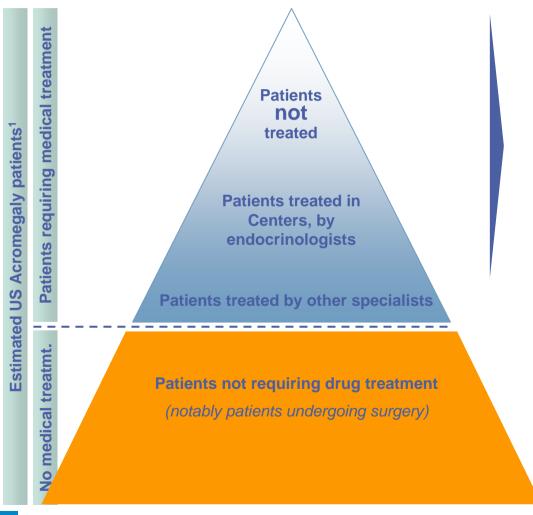
Full clinical development and regulatory capability in the US allowing for global developments of key programs

4 products now marketed, promoted by a Sales Force of 75



Somatuline® Depot market in the US

US Market structure - 15 000 to 18 000 patients



- Good penetration of Somatuline® Depot
 - 20% share² of endocrinologists prescriptions
 - 40% market share² in pituitary centers
- Significant pool of untreated patients
- Relatively low compliance compared to Europe

Drive Somatuline® Depot as first line recommendation in Pituitary centers and Endocrinologists

Drive Somatuline® Depot access and persistence



Snapshot on Increlex® and Dysport®

Increlex®

Established treatment option: Number of SMN¹s up more than 31% year-on year

Establishment of specialized Reimbursement & Endocrinologist dedicated support teams

+40% increase in sales in 2009

30% decrease in patient drop out rate in 2009

Dysport®

Great interest from commercial payors

Dysport® enjoys an 85% coverage rate (commercial) and a 100% coverage rate (government)

Excellent brand awareness

Dysport® known by more than 80% of target prescribing population at launch Fully integrated in US reimbursement system

Dysport®'s J-Code secured for March 2010

Higher-than-expected success of sampling campaign

Dysport® requested by twice as many neurologists as originally anticipated



A change in US context

Difficult economic situation impacting finances of patients

Increased pressure from commercial payers with tougher reimbursement criteria

Enforcement of strict compliance environment

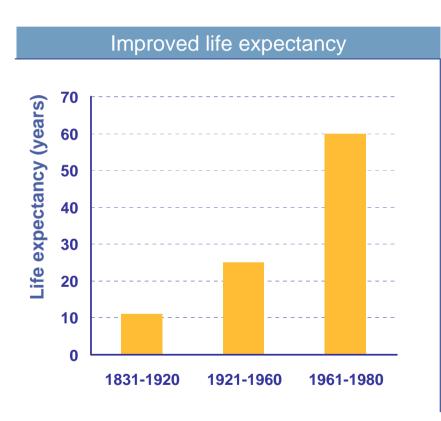
Appendix

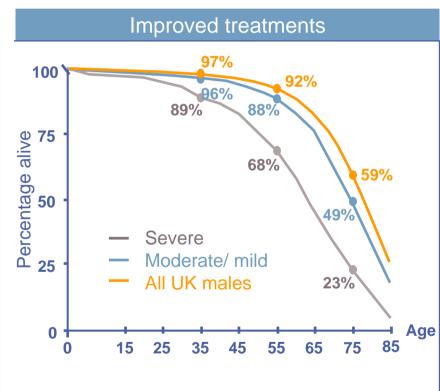
Partnership with Inspiration in Hematology





Despite improved life expectancy and treatment of patients with hemophilia...



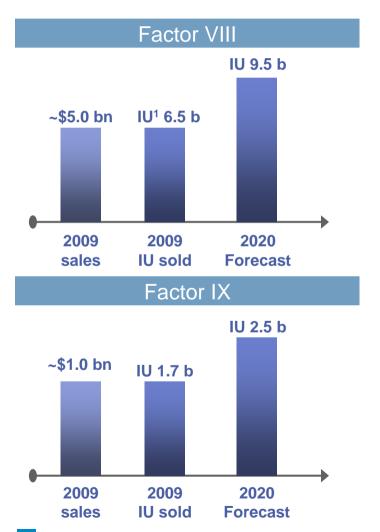


SOURCE: Larsson SA, et al. Br J Haematol 1985;59:593-602

Survival in men in the UK with hemophilia who were not infected with HIV and in the general male population of the UK in 1999 SOURCE: Darby SC. et al. Blood 2007;110:815–825



...FVIII & FIX market are still underserved globally...



Low penetration

 70% of hemophilia patients worldwide do not have access to factors

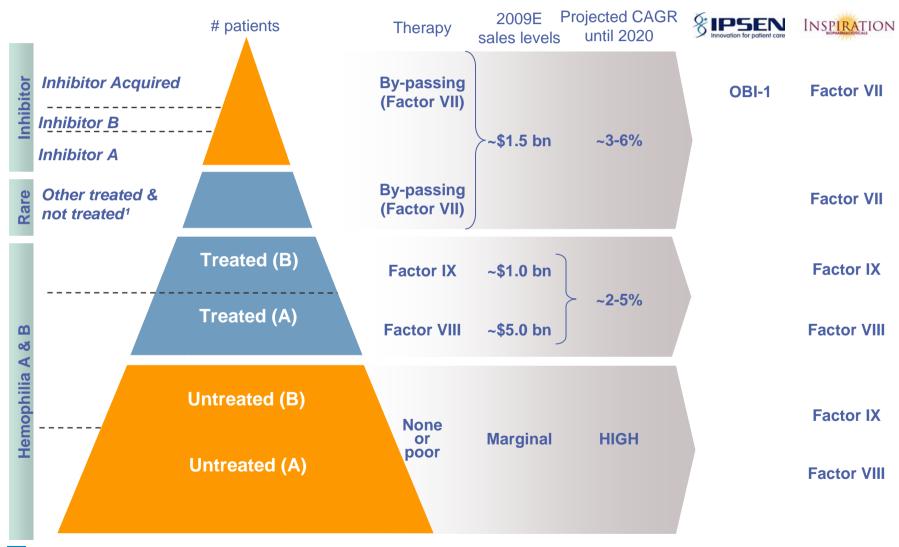
Long term growth prospects

- 4% to 10% CAGR until 2020
- Driven by prophylaxis in developed world
- Driven by more patients treated in RoW
- 90% recombinant in developed world
- 40% and growing in RoW

Most important unmet need today: Enable access to treatment to more patients



...resulting in high unmet medical needs in all segment of hemophilia



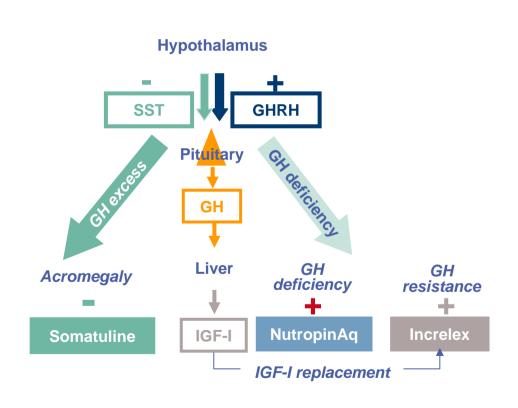
BACK UP SLIDES

Endocrinology





A unique focus on pituitary disorders and hormone dependent diseases

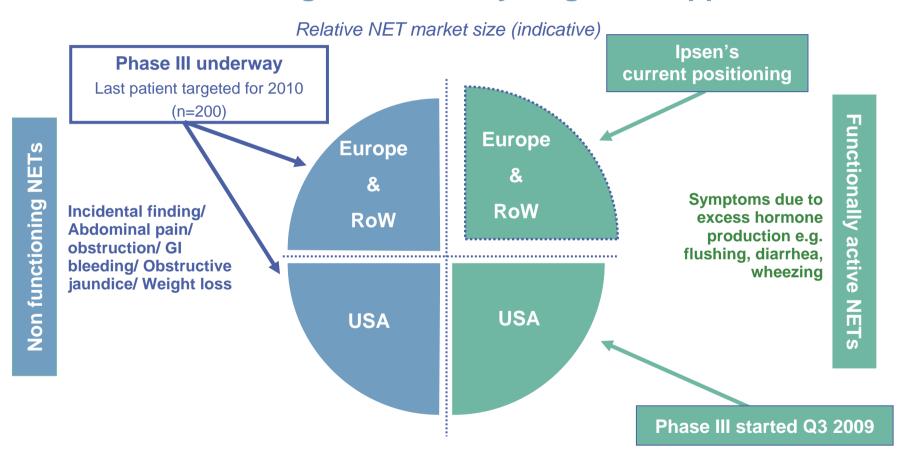


A strong franchise

- A range of products addressing today
 Short Stature, Acromegaly and NET
 - High morbi-mortality
 - Debilitating pathologies
 - High unmet medical needs
- Somatuline[®], NutropinAq[®] and Increlex[®] contributed to ~16 % of 2008 Group sales, ie. ~ €158 million.
- A fast growing franchise: sales doubled in the past 3 years



Somatuline® offers significant life cycle growth opportunities



Significant scope for expansion

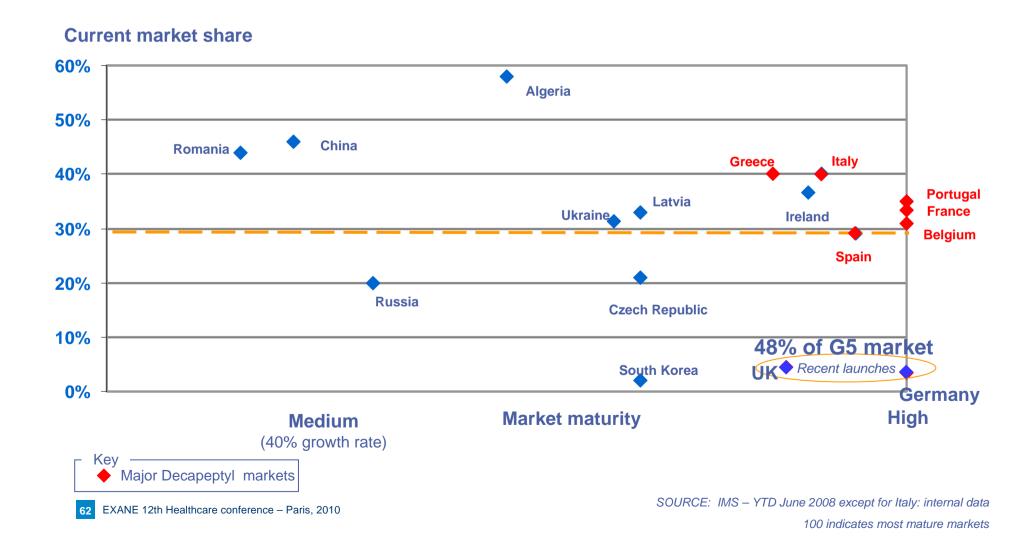
BACK UP SLIDES

Oncology





Decapeptyl®: strong positions, and poised to grow





Decapeptyl® 6 month formulation: a more differentiated product profile

Efficacy

- Comparable efficacy to 1 and 3 months formulation
 - Castration levels (testosterone)
 - Disease control (PSA)

Local Tolerance

Limited local side effects (6.7% of patients)

Storage and reconstitution

- Storage at room temperature (no need to heat up before reconstitution)
- 5 Steps to reconstitute, change needle, and inject IM route

Formulation/ Efficacy

6 month competitor 1

- 80% of patients castrated after 6M²
- Testosterone to be tested every 6M* 1
- Formation of Nodules or abscess ¹

6 month competitor 2

- Slow release formulation dependent on manual 60 mixture¹ step
- Storage at 2-4°: need to heat up for reconstitution ¹

BACK UP SLIDES

Neurology





A good track record at catching-up market shares...

Market share in medical indications Russia Botox launched in '94, now 56% market share **Dysport®:** Launched 5 years later, now 44% market share South Korea Botox launched in '95, now has 40% market share Dysport®: Launched 4 years later, now 23% market share Botox launched in '90, now has 40% market share **Dysport®:** Launched 10 years later, now 51% market share



An increasingly transactional model

ENDOCRINOLOGY













ONCOLOGY















NEUROLOGY









PRIMARY CARE

















2003 - 2005

2006-2009