Full Year Results 2011

Geoffrey McDonough (CEO) and Lars Sandström (CFO)



Stockholm, 23 February, 2012



Who We Are

Sobi is an innovative bio-pharmaceutical company with a leading position in the commercialization of niche and rare disease products.



Our operations are driven by a diversified and **growth-oriented portfolio** in niche and rare disease therapies.



We have a **late-stage pipeline** with substantial commercial potential.



We are a differentiated partner with world-class capabilities in protein biochemistry and biologics manufacturing development for **rare diseases**.



Actions to Reach Our Goals

We intend to earn our way into our future based on operational performance.

- 1. Revenue growth through focus on key products
- 2. Ongoing cost discipline
- 3. Gross margin improvement
- Improving cash flow from operations and working capital



Highlights

Financial

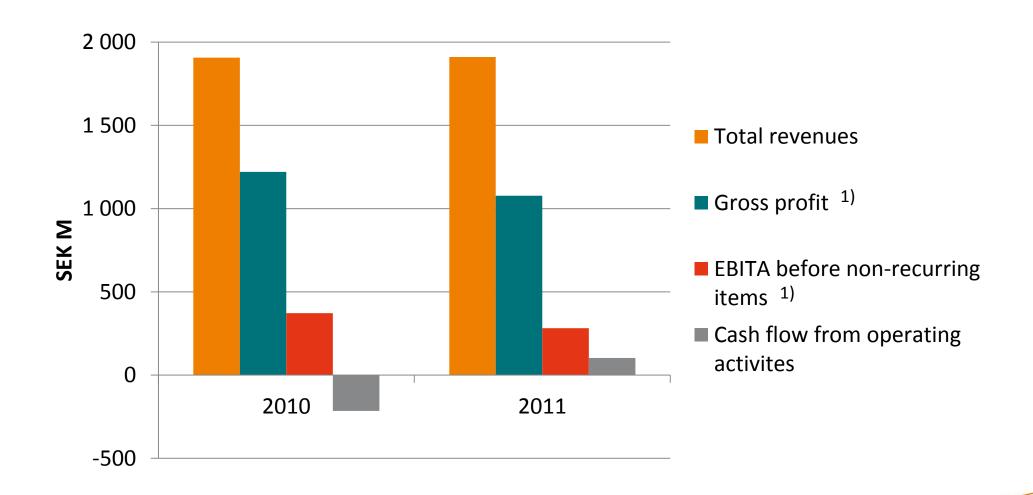
- Total revenues SEK 1 911 M (1 907)
 - Unchanged vs. last year
 - Increase by 9% adjusted for currency and discontinued products
- Gross margin lower due to reduced margin for manufacturing and currency effects
- Operating expenses reduced reflecting ongoing streamlining of operations
- Financial position strengthened through a rights issue
- Positive cash flow in Q4 2011

Business

- Strengthened management team
- New organizational focus in three business lines
- US subsidiary in place
- Phase III programs advanced according to plan
 - Kiobrina FPI July
 - Enrollment substantially completed in Factor programs with Biogen Idec
- Extension of ReFacto manufacturing agreement



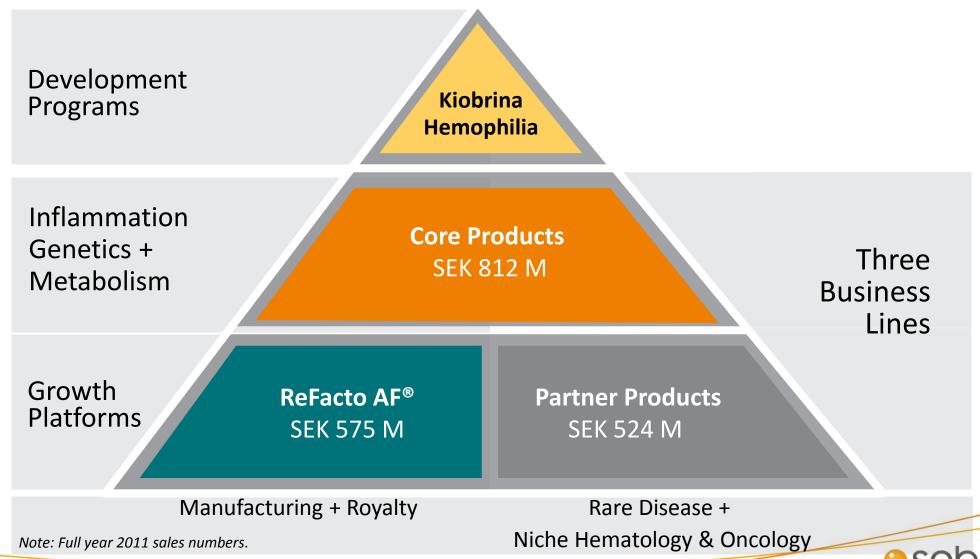
FY 2011 Consolidated Results



¹⁾ Adjusted for balance sheet write-downs in Q4-11.



FY 2011 Revenues by Business Lines



2011 Product Cross-Walk

Core Products (SEK 812 M)

Inflammation

- Kineret
- Ruconest

Genetics + Metabolism

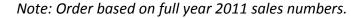
- Orfadin
- Ammonaps
- Ammonul

Partner Products (SEK 524 M)

- ReFacto AF® Co-promotion
- Kepivance
- Yondelis
- Ferriprox
- Betapred
- Buronil
- Aloxi
- Willfact
- > 40 products (each with annual sales < SEK 15 M)

ReFacto AF® (SEK 575 M)

- Manufacturing
- Royalty





Recent Agreements with Pfizer

Supply Agreement

Facts

- Extended supply agreement for ReFacto AF®/XYNTHA® until 31 December 2020 (previously 2015)
- Sobi continues to be the global supplier of the drug substance for ReFacto AF®/XYNTHA®

Financial Implications

 Secures long-term base business and associated cash flows

Divestment of Co-promotion Rights

Facts

- Divestment of co-promotion rights for ReFacto® and BeneFIX® in the Nordic region as of 15 February 2012 (previously 2016)
- Proceeds of USD 47.5 M (SEK 320 M)

Financial Implications

- Lower revenues and profits (approx. SEK 100 M annually)
- Strengthens balance sheet



2011 Pro Forma (Adj. for Divestment of Co-promotion Rights)

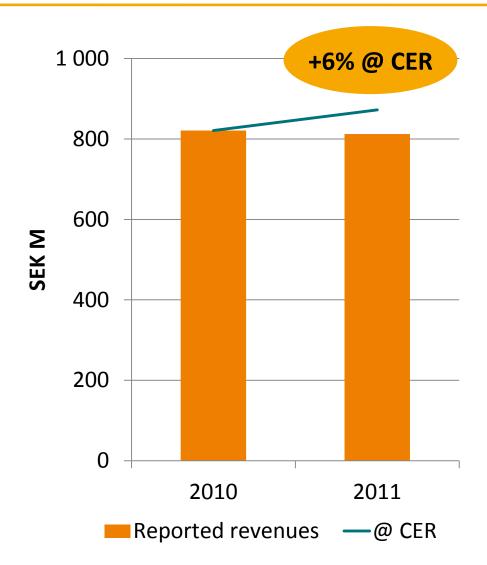
SEK M	2011	Co-promotion	Pro Forma 2011
Total revenues	1 911	105	1 806
Gross profit 1) Gross margin 1)	1 078 56,4%	105 100,0%	973 53,9%
EBITA before non-recurring items 1)	282	105	177

¹⁾ Adjusted for balance sheet write-downs in Q4-11.



Revenues From Core Products





- Substantial shift in resources H2 2011 to focus on Core Products
- Building broader commercial capabilities
- Life cycle management a key focus in the medium term

Note: Core Products include Kineret, Orfadin, Ammonaps, Ammonul and Ruconest

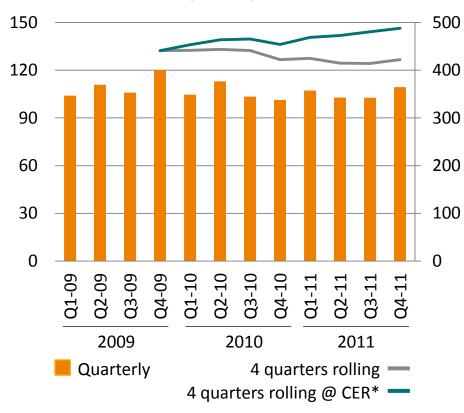


Kineret® – Steady Growth









^{* 4} quarters rolling using average exchange rates for 2009 as base.

- Awareness in Rheumatoid Arthritis space increasing, as is interest in broader inflammation
- A NIH study¹ shows that Kineret
 (anakinra) effectively stop progression of
 organ damage in neonatal-onset
 multisystem inflammatory disease
 (NOMID)
- Expect EMA response on pediatric investigation plan for CAPS (pediatric inflammation) in H1 2012
- Sobi will file for CAPS in EU 2012
- Sobi will file for NOMID indication in US 2012

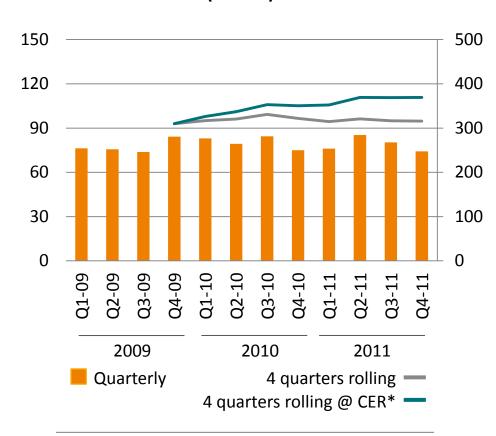
¹ Sibley et al., Sustained Response and Prevention of Damage Progression in Patients with Neonatal-<u>Onset Multisystem Inflammatory</u> <u>Disease (NOMID) Treated with Anakinra; Arthritis & Rheumatism.</u> <u>January 2012</u>



Orfadin® – Understanding the Growth Drivers





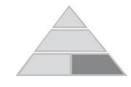


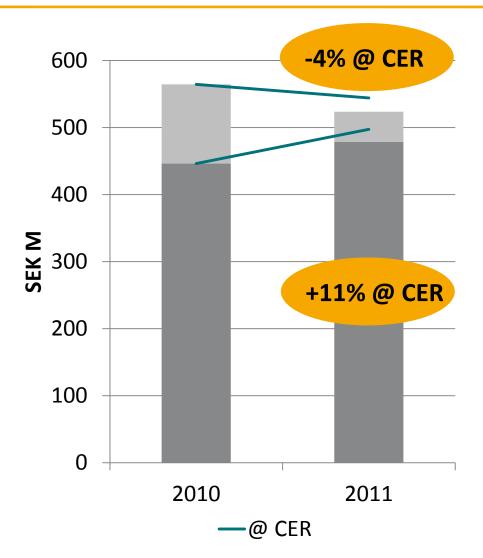
^{* 4} quarters rolling using average exchange rates for 2009 as base.

- Strong growth in US due to new born screening, balanced by Obama healthcare Medicaid rebates
- Strong growth in Russia and South America
- Europe focus on dosing and compliance
- Expect EMA response on our pediatric investigation plan submission for liquid formulation in H1 2012
- Liquid formulation in Europe opens opportunity to extend Orphan Drug exclusivity through 2017



Revenues From Partner Products





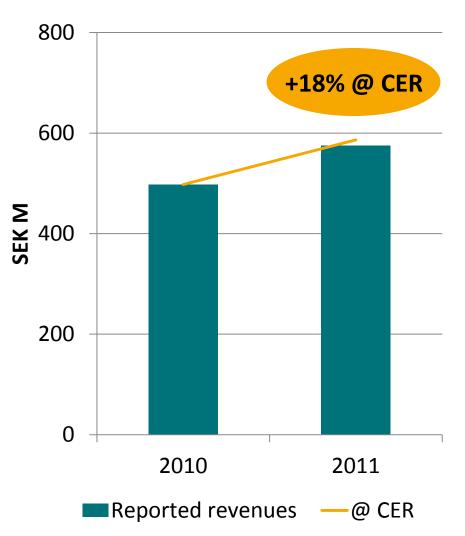
- Dedicated business unit focus in order to drive momentum and deal flow in Nordic and CEE
- Continued strong underlying growth
- 5 discontinued products (SEK 45 M)
 - Shire products: Equasym, Fosrenol and Xagrid
 - Mimpara
 - Tracleer
- New agreement signed
 - Gentium / Defibrotide
 - Discontinued products
 - Current portfolio, includingCo-promotion revenues

Note: Partner Products includes ReFacto® Co-promotion, Kepivance, Yondelis, Ferriprox, Betapred, Buronil, Aloxi, Willfact and additional >40 products.



Revenues From ReFacto AF®



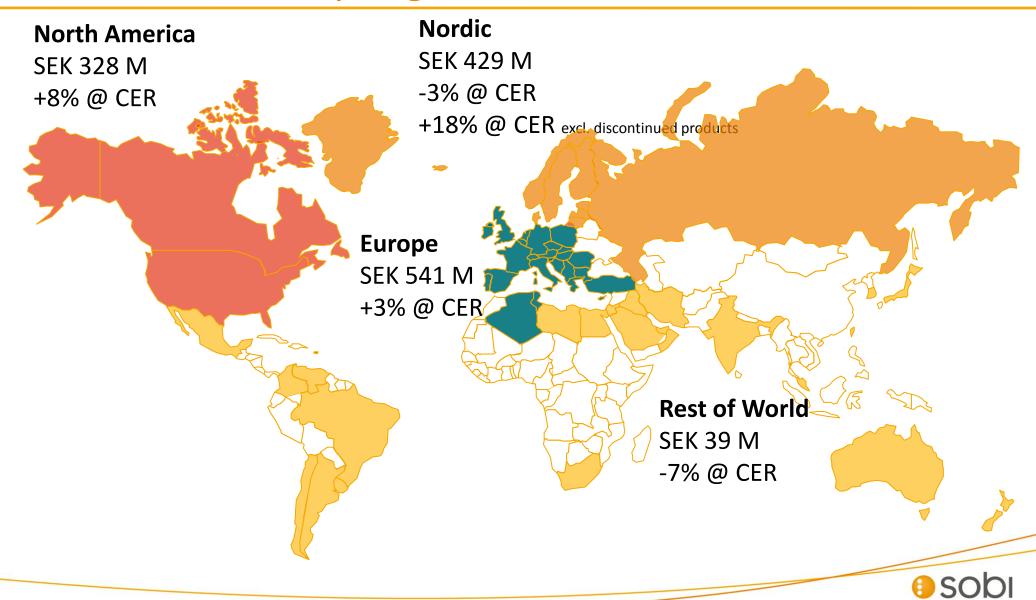


- Steady growth in volumes
- Validation batches contributed SEK 42 M to sales in 2011
 - Underlying growth +9% @ CER adjusted for validation batches
- Supply agreement extended from 2015 to 2020, with option to renew
- Due to process improvements, capacity will increase in H2 2012

Note: ReFacto AF® includes manufacturing and royalty revenues



2011 Revenues by Region



Financials

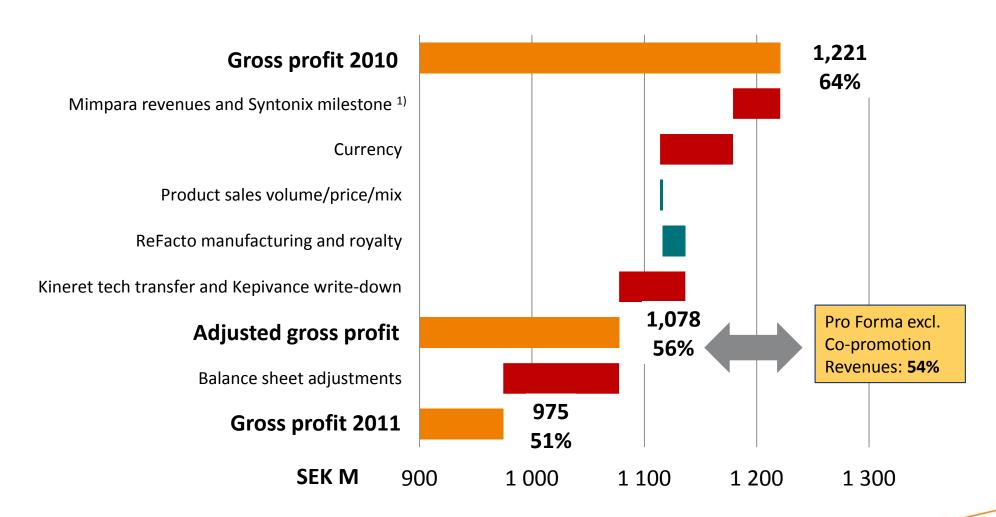
Lars Sandström (CFO)



We provide valuable medicines to patients with rare diseases



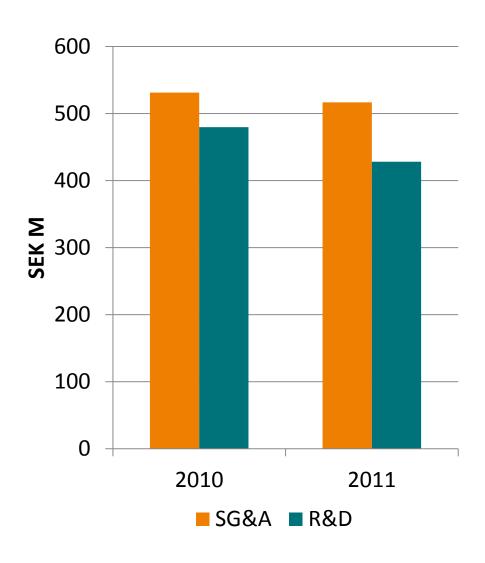
Gross Profit



¹⁾ Co-promotion revenues from Mimpara (SEK 22.7 M) and Milestone revenues (SEK 23.5) from Syntonix related to factor programs.



Operating Expenses



- Investments in sales organization finalized
- Cost savings within Research & Development offset by increased Kiobrina phase three investment
- Substantial General & Administration cost reductions



Balance Sheet Adjustments

Asset	Product	Amount (SEK M)
Inventory	Kineret [®] Kepivance [®]	103.2
Trade receivables		14.8
Intangibles	Leptin modulation program	126.9
Real estate		46.5
Total		291.4



Income Statement – Cross-Walk for Write Downs

	Reported	Adjusted	Balance Sheet	Reported
SEK M	2010	2011	Adjustments	2011
Total revenues	1 907	1 911	0	1 911
Gross profit	1 221	1 078	-103	975
Gross margin	64,0%	56,4%	-	51,0%
OPEX	-1 011	-945	-50	-995
Other operating revenues/expenses	162	149	-2	147
EBITA before non-recurring items	372	282	-154	127
Non-recurring items	-88	-78	0	-78
Amortizations	-294	-231	-137	-368
EBIT	-10	-27	-291	-319
Net financial items	-82	-52	0	-52
Tax	-12	389	0	389
Profit/loss	-104	309	-291	18

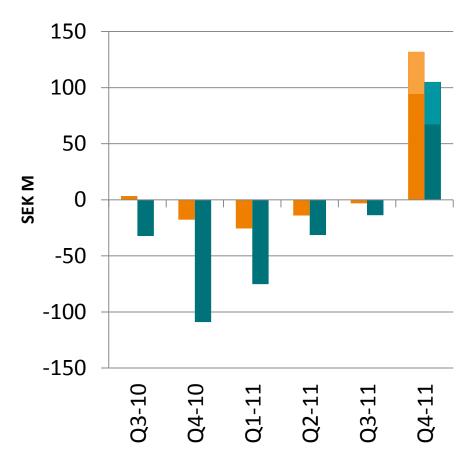


Cash Flow

SEK M	Reported 2010	Reported 2011
Cash flow from operations before change in working capital	250	118
Change in working capital	-465	-15
Cash flow from operating activites	-215	103
Cash flow from investing activites	-1 884	-44
Change in external debt	468	-472
Change in equity	1 414	594
Cash flow from financing activities	1 882	122
Period cash flow	-218	181



Cash Flow

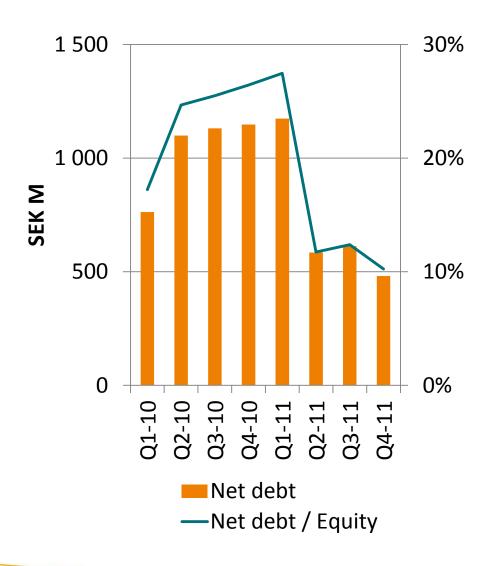


- Cash flow before financing (excl. acq.)
- Whereof working capital

- Underlying profitability reflected in cash flow from operations
- Build-up of inventory due to Kineret tech transfer finalized in Q3 2011
- Large payments from Pfizer in Q4 2011 impact working capital positively
 - Lighter colors in Q4 2011 indicate onetime payment (approx. SEK 40 M) to be passed through to Pfizer in Q1 2012



Net Debt



- Rights issue in June substantially improved our financial position
- Divestment of the co-promotion rights for ReFacto® and BeneFIX® (USD 47.5 M / SEK 320 M)
 - Pay down debt
 - Strengthen our cash position



Outlook

Geoffrey McDonough (CEO)



We provide valuable medicines to patients with rare diseases



Outlook 2012

- Market conditions are expected to continue to be challenging but do not significantly deteriorate
- The current SEK/USD and SEK/EUR exchange rates prevail during the year
- Total revenues for the full year 2012 are expected to be approximately SEK 100 M lower than 2011, reflecting the divestment of the ReFacto® co-promotion rights
 - Revenues for Core Products and Partner Products are expected to show mid to high single digit growth, and revenues for ReFacto® manufacturing and royalty are expected to show low single digit growth
 - Revenues in 2011 from validation batches (SEK 42 M) and discontinued products (SEK 45 M) will not recur in 2012

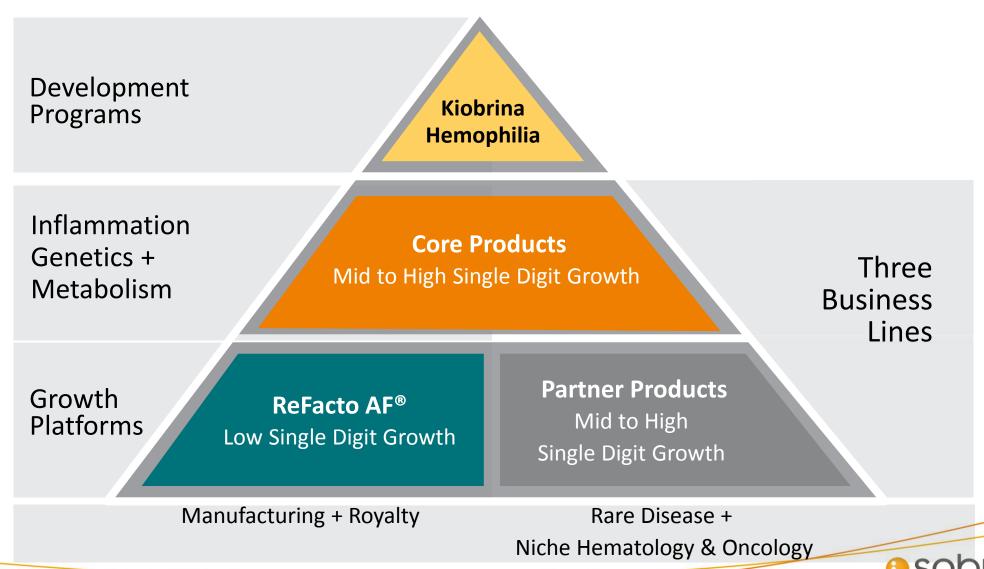


Outlook 2012 cont.

- The gross margin for the full year 2012 is expected to be in line with the 2011 margin, which was 54% after adjustment for the balance sheet write-downs and the divestment of co-promotion rights
- Costs in 2012 related to the transfer of Kineret® production are estimated at SEK 60 M and are expected to impact the gross margin primarily in the first half of the year
- As a result of the renegotiated ReFacto[®] manufacturing agreement with Pfizer, reduced quarterly variations in the ReFacto[®] manufacturing gross margin are expected
- Milestone payment to Amgen of USD 55 M expected in Q4 2012 or in Q1 2013
- Operating expenses excluding amortizations are estimated at or below SEK 950 M

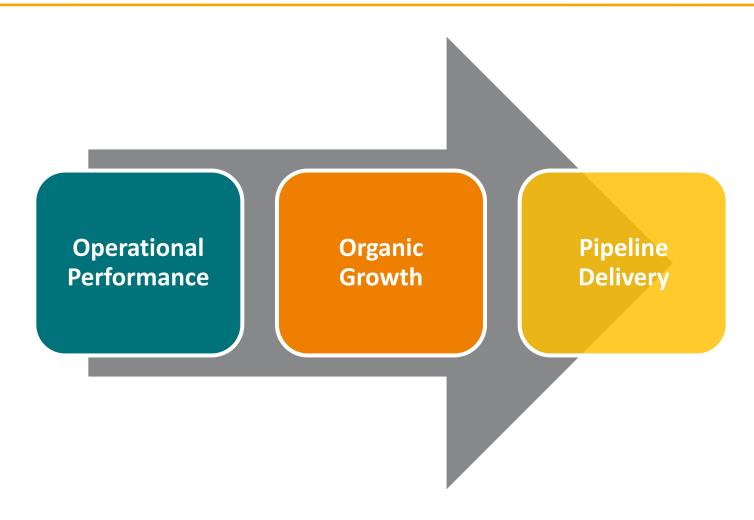


Outlook by Business Lines



Focus on Delivery







2012 Calendar Highlights

Event	H1 2012	H2 2012
Orfadin Liquid Formulation PIP Response		
Kineret CAPS PIP Response		
Complete Tech Transfer Kineret Manufacturing		
FPI Pediatric Trials rFVIIIFc + rFIXFc Programs (BIIB)		
Kineret NOMID Filing FDA		
Kineret CAPS Filing EMA		
Data for rFVIIIFc + rFIXFc Programs (BIIB)		
Kiobrina Complete Phase 3 Enrollment		



Summary

- Diversified commercial portfolio focused on improving cash flow and profitability
- 2. Working to efficiently commercialize our proprietary innovative medicines for rare disease patients globally
- 3. Business model oriented to **building**value through partnerships from global early stage biologics development to late stage specialty distribution in Europe







