

Investor Handout

March 2015



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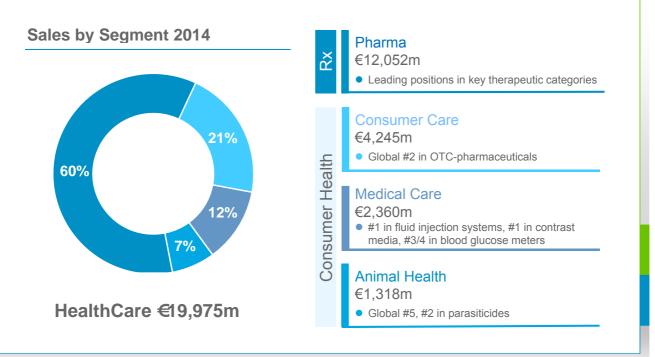
Figures for 2012 have been restated due to changes in accounting policies relating to the accounting standards IAS 19R ("Employee Benefits") and IFRS 11 ("Joint Arrangements"). In addition, Bayer changed accounting for the stock-based compensation program.



Disclaimer

Leading Positions in Key Healthcare Markets

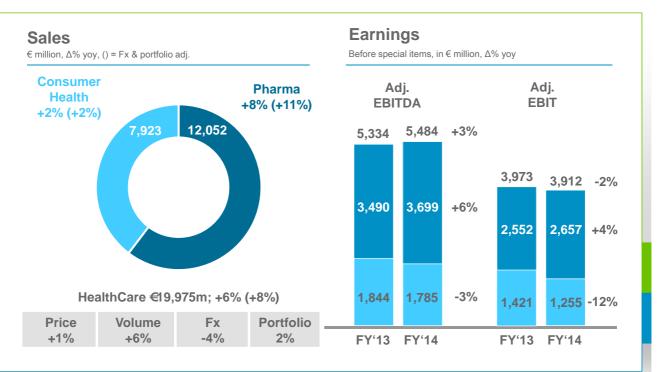




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FY 2014 - Dynamic Growth of HealthCare Driven by Pharma







HealthCare: 2015 Financial Outlook

Sales Δ% Fx & portfolio adj.

		Sales*	EBITDA before special items*
H	HealthCare	Mid-single-digit % increase (Fx +3%)	Mid-teens % increase
	Pharma	Mid-to-high-single-digit % increase (Fx +2%) Launch product sales: increase towards €4bn	Low-teens % increase Slight margin improvement
	Consumer Health	Mid-single-digit % increase (Fx +3%)	Mid-to-high twenties % increase

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*Assuming spot Fx rates as of Dec 31, 2014 for FY'15 Outlook depends on specific planning assumptions as detailed in the Annual Report

HealthCare: Aspirations 2017



	Sales CAGR (2014-2017)	(before special items)
HealthCare*	~6% to above €25bn	29 - 31%
Pharma	~7% to above €15bn	32 - 34%
Consumer Health*	~4% to above €10bn	24 - 26%



Strategic Priorities for Pharma



Maximize the value of recently launched products



Achieve phase III readiness for key phase II assets by mid-2016

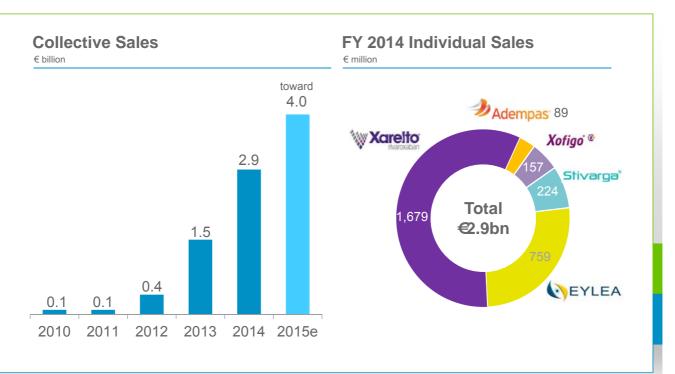


Advance early and mid-stage pipeline

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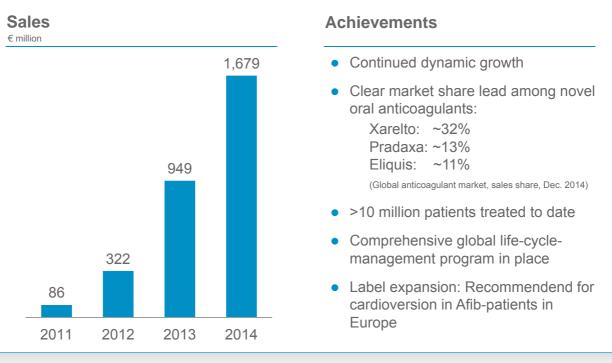
Pharma Launch Products Drive Growth: Combined Peak Sales Potential of ≥€7.5bn





Xarelto: The Leading Novel Oral Anticoagulant

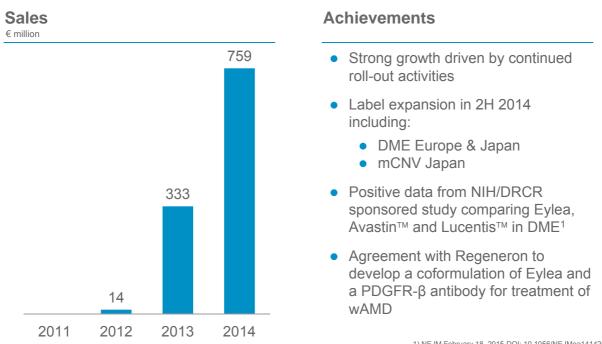




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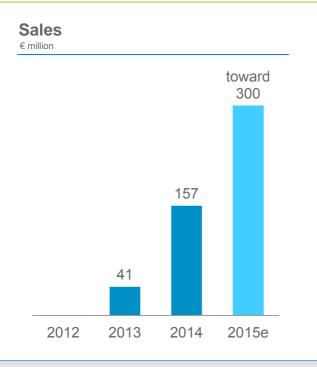
Eylea: Label Expansion to Support Further Growth







Xofigo: Significant Growth Expected



Achievements

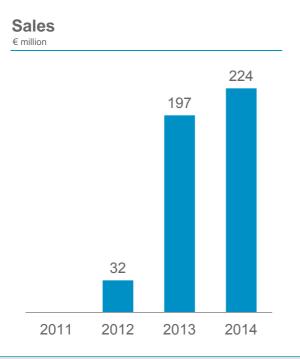
- Launch activities ongoing
 - Approved in 41 countries¹
 - Launched in 27 countries¹
- Q4 2014 affected by temporary drug shortage
- Supply of Xofigo resumed in November 2014
- Life-cycle management program targeting label expansions in CRPC and additional cancer indications underway

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CRPC: Castration resistant prostate cancer
1) As of January 2015

Stivarga: Leading 3L+ Treatment of Metastatic Colorectal Cancer





Achievements

- Most prescribed agent in 3rd-line plus treatment of mCRC in the US and in Japan
- Roll-out activities in mCRC and GIST continuing
- Targeting increased use in 3Ltreatment of mCRC
- ~40,000 patients treated to date

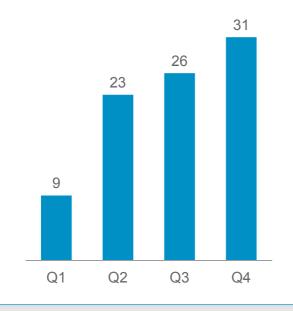
mCRC: Metastatic colorectal cancer; GIST: Gastrointestinal stromal tumor

Adempas: Encouraging Early Launch Experience



Sales 2014

€ million



Achievements

Launched in PAH and CTEPH in first countries

	PAH	CTEPH
Approved ¹	45	48
Launched ¹	24	26

- ~3,000 patients treated to date²
- Life-cycle management trials in systemic sclerosis, cystic fibrosis and in PH associated with idiopathic interstitial pneumonias initiated
- Agreement with Merck & Co. for joint development and commercialization of sGC-modulators in place

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1) as of January 2015; 2) as of January 2015, PH: Pulmonary hypertension
 PAH: Pulmonary arterial hypertension, CTEPH: Chronic thrombo-embolic pulmonary
 hypertension, sGC: Soluble guanylate cyclase

Increasing R&D-investment to Support Short-, Mid- and Long-term Growth at Pharma



Life-cycle Management

Projects from Midand Late-stage Pipeline

Opportunities in Research and Early Pipeline

- Xarelto (Rivaroxaban)
- Eylea (Aflibercept)
- Xofigo (Radium-223 dichloride)
- Stivarga (Regorafenib)
- Adempas (Riociguat)
- Finerenone
- Copanlisib
- Molidustat
- Vericiguat
- Vilaprisan
- Damoctocog alfa pegol
- ODM-201

- Cardiology
- Hematology
- Oncology
- Gynecological Therapies
- Ophthalmology
- Inflammation

Rivaroxaban - Comprehensive Life-cycle Management Program Underway

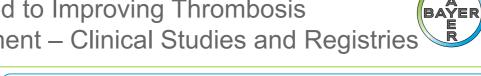


Indication	Trial	End date [*]
Areas with patients at risk of arterial thromboembolism:		
Chronic heart failure and significant coronary artery disease	COMMANDER-HF	Apr 2017
Major cardiovascular events in coronary or peripheral artery disease	COMPASS	Jan 2018
Embolic stroke of undetermined source (ESUS)	NAVIGATE ESUS	Jan 2018
Peripheral artery disease undergoing peripheral artery interventions	VOYAGER PAD	Yet to start
Acute coronary syndrome (ACS)	GEMINI ACS 1	Jan 2017 (PH II)
Specific AF-patient populations:		
Non-valvular atrial fibrillation + catheder ablation	VENTURE-AF	completed
Non-valvular atrial fibrillation + cardioversion	X-VeRT	published
Non-valvular atrial fibrillation + left atrial/left atrial appendage thrombus	X-TRA	completed
Non-valvular atrial fibrillation + percutaneous coronary intervention	PIONEER AF-PCI	Aug 2016
DVT and PE in specific patient populations:		
Long-term prevention of symptomatic recurrent DVT and/or PE	EINSTEIN CHOICE	Nov 2016
Symptomatic DVT and/or PE due to concurrent medical illness	MARINER	Feb 2017
Treatment and sec. prevention of DVT and/or PE in pediatric patients	EINSTEIN JUNIOR	Nov 2018

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Rivaroxaban is being developed in cooperation with Janssen Research & Development LLC, a Johnson & Johnson company *) Primary trial completion dates as of February 20, 2015

Committed to Improving Thrombosis Management - Clinical Studies and Registries



Registries N≈97,000

Gärfield **ORBIT** AFF

Gärfield **CLOT-**AF



Phase IV/NIS N≈47,000

XCIMOS

XONTUS

XOLIA

Phase IIIb N≈2,400

VENTURE AF

X-TRO



Phase II/III N≈51,000

COMPASS ** VOYCIGER PAD KNOVIGATE ESUS ** MARINER K EINSTEIN CHOICE EINSTEIN JUNIOR COMMANDER HES GEMINI ACS 1 15

Completed

ROCKET AF 5 J-ROCKET AF 5 ATLOS MAGELLON*

Over 275,000 patients are expected to be included in both clinical trials and real world settings.

Regorafenib – Potential for Significant Advances in Treating wAMD



Current Standard intravitreal injection

Today's standard of care is administered by injection into the eye

Project Goal topical treatment (dro

topical treatment (drops) as additional option

Eye drops could reduce efforts, costs, logistics and would be more convenient

- Regorafenib inhibits VEGF receptor signaling, a well established principle to treat wAMD
- Regorafenib eye drops may offer topical treatment of wAMD - targeting superior convenience but non-inferior efficacy
- Phase I completed
- Phase IIa/b initiated
 - Comparator: Lucentis
 - Target enrollment: N=540 patients
 - Completion expected April 2016







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wAMD: Wet age-related macular degeneratio; Primary completion dates as published at clinicaltrials.gov, status February, 2015

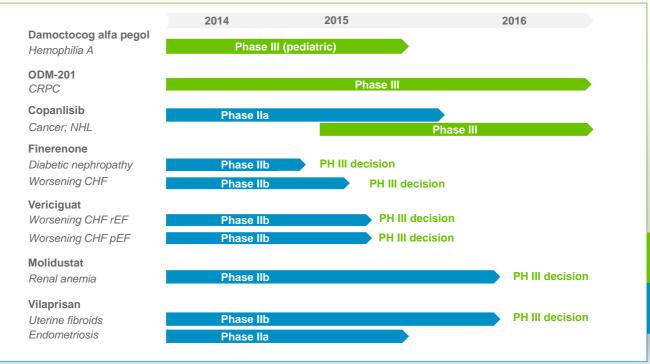
Key Mid- and Late-Stage Pharma Pipeline Assets



Project	Mechanism	Indication	
Damoctocog alfa pegol	Damoctocog alfa pegol Long-acting rFVIII		
ODM-201 Androgen Receptor Inhibitor		Castration Resistant Prostate Cancer	
Copanlisib	PI3-Kinase Inhibitor	Cancer (NHL)	
Finerenone	MR Antagonist	Chronic heart failure (CHF) Diabetic nephropathy (DN)	
Molidustat	HIF-PH Inhibitor	Anemia	
Vericiguat	sGC Stimulator	Worsening chronic heart failure	
Vilaprisan (sPRM)	Progesterone Receptor Antagonist	Symptomatic uterine fibroids Endometriosis	

Mid- and Late-Stage Pharma Pipeline Progressing – Current Timelines





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CHF: Chronic heart failure; CRPC: Castration-resistant prostate cancer; NHL: Non-Hodgkin's lymphoma; r/pEF: Reduced/preserved ejection fraction

Attractive Early Pipeline Projects*



Project	Mechanism	Target Indications	Status
BAY 1067197	Non-nucleosidic partial A1 receptor agonist	Worsening clinical heart failure	Phase IIa
BAY 1213790	FXIa antibody	Anti-coagulation therapy	Phase I
BAY 1142524	Chymase inhibitor	Cardiovascular diseases	Phase I
Plasminogen- Inhibitor	Inhibition of fibronolysis	Various bleeding disorders incl. hemophilia	Phase I start Q1 2015e
BAY 1000394	Pan-CDK-inhibitor	Small-cell lung cancer	Phase II
BAY 94-9343	Mesothelin antibody-drug- conjugate	Solid tumors	Phase lb
BAY 1179470	FGFR2 antibody	Solid tumors	Phase I
BAY 2010112	PSMA-BiTE	Prostate cancer	Phase I
BAY 1128688	AKR1C3 inhibitor	Endometriosis	Phase I



Well-positioned for Growth

- One of the fastest growing global Pharma companies
- Pharma launch products drive growth Combined peak sales potential of ≥€7.5bn
- Comprehensive life-cycle management program to fully exploit potential of launch products
- Mid- and late-stage pharma pipeline progressing Expect increasing visibility in 2015

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Date	Event	Publication
Thursday, April 30, 2015	Investor Conference Call	First Quarter 2015 Results Stockholders' Newsletter
Wednesday, May 27, 2015	Annual General Meeting	
Thursday, June 25, 2015	Meet Management in New York	Investor Conference
Wednesday, July 29, 2015	Investor Conference Call	Second Quarter 2015 Results Stockholders' Newsletter
Thursday, October 29, 2015	Investor Conference Call	Third Quarter 2015 Results Stockholders' Newsletter



Reporting Events and AGM



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