

Bayer Pharmaceuticals

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39th Annual J.P. Morgan Healthcare Conference

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Cautionary Statements Regarding Forward-Looking Information

This presentation may contain forward-looking statements based on current assumptions and forecasts made by Bayer management.

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Guidance at constant currencies, not including portfolio divestitures if not mentioned differently.



Driving Performance and Delivering New Growth Opportunities



Maximize the value of the existing portfolio

Exploit full potential of Xarelto and Eylea



Deliver on key pipeline assets

Three new potential blockbusters in Oncology, Cardiovascular and Women's Healthcare



Take advantage of breakthrough technologies

Expand into Cell & Gene Therapy

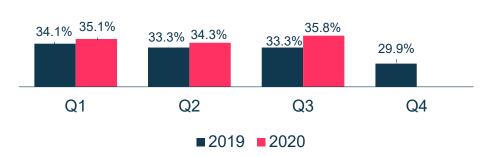


After a Challenging Q2, V-shaped Recovery in Q3

Sales Growth vs Prior Year (cpa) by Quarter



EBITDA margin before special items



* Sales related growth rates are reported as currency and portfolio adjusted, and compared to the previous year

Pharma business recovering in Q3 from Covid-19

impact

- Xarelto's growth trajectory intact (+13% YTD)
- Stringent cost management to protect the bottom-line
- Outlook for FY2020 confirmed on November 3, 2020:
 - Expected Sales growth: -1%
 - Expected underlying EBITDA margin: 34-35%



Increased Momentum to Deliver Innovation for Patients

Major achievements 2019/2020

Phase Transitions; New Trials¹



- Finerenone HFmr/pEF
- Aflibercept high dose nAMD & DME
- Aflibercept retinopathy of prematurity
- Regorafenib IO combi
- Regorafenib glioblastoma
- Factor XI portfolio
- P2X3 inhibitor
- PEG ADM inhale
- Runaciguat

Positive Phase III Trial Results



- Finerenone (FIDELIO-DKD)
- Vericiguat (VICTORIA)
- Copanlisib (CHRONOS-3)
- Darolutamide (ARAMIS OS-data)
- Rivaroxaban (PRONOMOS, VOYAGER PAD, EINSTEIN-Jr.)

Filings & Approvals



- Darolutamide
- Larotrectinib
- Vericiguat
- Finerenone
- Eylea PFS
- Rivaroxaban pediatric

External Innovation



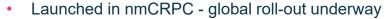
- BlueRock
- AskBio
- KaNDy
- Atara
- Daré Bioscience
- leaps[⊕] investments

¹ Key Phase II and III trials only



A Successfully Matured Late-stage Pipeline that May Deliver Three New Potential Blockbusters

Nubeqa¹ – AR-Antagonist





- Differentiated clinical profile 31% OS benefit & favourable safety profile (ARAMIS)
- Prim. completion of phase III trial in mHSPC mid-2021e (ARASENS)
- Peak sales potential ≥€1bn

KaNDy NT-814 – Dual NK 1,3 Receptor-Antagonist



- First-in-class oral, non-hormonal, once-daily neurokinin-1,3 receptor antagonist
- Positive phase IIb data for the treatment of frequent menopausal symptoms
- Phase III to be initiated in 2021
- Peak sales potential >€1bn

Finerenone – MR-Antagonist



- Significant reduction of renal and cardiovascular outcomes in patients with CKD and T2D
- Filed in key markets for CKD in T2D
- Phase III trials in CKD/T2D (FIGARO) and in HFm/pEF (FINEARTS-HF) ongoing
- Peak sales potential ≥€1bn

Vericiguat² – **sGC-Stimulator**



- Significant reduction of the combined risk of CVdeath or HF-hospitalization in HFrEF-patients (VICTORIA)
- Dosed on top of existing HF-therapies
- Filed FDA action date January 20, 2021
- Peak sales potential ~€500m

¹In collaboration with Orion Corporation; ²In collaboration with Merck & Co. Inc., Kenilworth, NJ, USA



We Provide the Resources Needed for a Successful Commercialization of our Late-stage Pipeline Assets

	Asset	Short- to mid-term Investment Areas
0	Nubeqa ¹	 Continued global rollout in additional markets Preparation for ARASENS readout Continue to leverage positive OS data and differentiated safety profile
	Finerenone	 Preparation for potential first launches in 2021 Investing to re-enter the US-market in CV diseases Preparation for FIGARO readout Continuation of the HFm/pEF development program
XO O	KaNDy NT-814	Initiation of pivotal Phase III trial in 2021
	Vericiguat ²	Potential launches in first markets in 2021

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Increased Focus on External Innovation to Accelerate Replenishment of Pipeline and Broaden Modalities

Selected high-level overview

Momentum Significantly Increased

- More than 20 Transactions signed in 2020
 - Deals covering the entire spectrum from equity investments (with LEAPS), over licensing agreements to acquisitions
 - Active portfolio management taking internal assets outside (e.g. Vincera Pharma)

Strategic Focus

- Venturing into new modalities (Cell & Gene Therapy)
- Broadening the Oncology pipeline (e.g. Systems Oncology, Atara)
- Commercial partnerships in China (e.g. Hua Medicine)
- Deals in the Digital Space (e.g. R&D: Schroedinger, Exscientia, Recursion; Commercial: One Drop)
- Continued augmentation of core TAs: (WHC: KaNDy)
- Strengthening the Cardiovascular pipeline (PeptiDream, Broad Institute)



Our Cell & Gene Therapy Strategy Builds on Four Integrated Platforms to Drive the Next Wave of Innovation at Pharma

Gene Augmentation Stem Cells Allogeneic Cell Therapy Collaboration with Atara *AskBio BlueRock **Biotherapeutics** Industry-leading AAV Creating induced Next-generation, mesothelin-directed pluripotent stem cells vector gene (iPSC) with broad augmentation CAR-T cell therapies platform differentiation Focus on potential Monogenic & Create an entirely new allogeneic, "off the pathway diseases generation of cellular shelf" tumor therapies medicines **CDMO** business (Viralgen) already IND for lead program in Parkinson's disease generates revenues Gene Editing as cross-functional enabling technology



Comprehensive Cell & Gene Therapy Pipeline that has Proven Ability to Yield Commercial Stage Assets

Project	Discovery	Preclinical	Phase I/II
Pompe Disease - Gene Therapy			
Parkinson's Disease - Gene Therapy			
Congestive Heart Failure - Gene Therapy			
Factor VIII - Gene Therapy ¹			
MSLN CAR-T Therapy (ATA2271) ²			
Parkinson's Disease - Cell Therapy			
MSLN CAR-T Therapy (ATA3271) ²			
> 15 preclinical assets			

- Comprehensive Cell & Gene Therapy pipeline established
- Pipeline already comprises 6 clinical assets and multiple IND generating opportunities
- AAV technology included in commercialized assets

¹ In collaboration with Ultragenyx; ² In collaboration with Atara Biotherapeutics



Key Takeaways

- We continue to maximize the value of the existing portfolio
- Increased momentum to deliver innovation late-stage pipeline may deliver three new potential blockbuster products
- We provide the resources needed for a successful commercialization of our late-stage pipeline assets
- Cell & Gene Therapy platform in place to drive the next wave of breakthrough innovation
- C>-Pipeline established 6 clinical assets and multiple IND generating opportunities



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Abbreviations

AAV	Adeno-associated virus	Ю	Immuno-oncology
AR	Androgen receptor	iPSC	Induced pluripotent stem cells
CAR-T	Chimeric antigen receptor modified T cells	mHSPC	Metastatic hormone sensitive prostate cancer
CDMO	Contract development and manufacturing organization	MRA	Mineralocorticoid receptor antagonist
CKD	Chronic kidney disease	nAMD	Neovascular age-related macular degeneration
C>	Cell and gene therapy	NK	Neurokinin
сра	Currency and portfolio adjusted	nmCRPC	Non-metastatic castration resistant prostate cancer
CV	Cardiovascular	OS	Overall survival
DKD	Diabetic kidney disease	PAD	Peripheral artery disease
DME	Diabetic macula edema	PFS	Pre-filled syringe
EBITDA	Earnings before interest, tax, depreciation, and amortization	R&D	Research & Development
FDA	U.S. Food and drug administration	sGC	Soluble guanylate cyclase
HF	Heart failure	T2D	Type 2 diabetes mellitus
HFmrEF	Heart failure with mid-range ejection fraction	YTD	Year to date
HFpEF	Heart failure with preserved ejection fraction		

HFrEF IND Heart failure with reduced ejection fraction

Investigational New Drug