

March 2, 2016

MorphoSys AG

Year End Results 2015 & Outlook 2016



Today on the Call



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**Chief Executive
Officer**



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**Chief Financial
Officer**



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**Chief Scientific
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This presentation includes forward-looking statements.

Actual results could differ materially from those included in the forward-looking statements due to various risk factors and uncertainties including changes in business, economic competitive conditions, regulatory reforms, foreign exchange rate fluctuations and the availability of financing. These and other risks and uncertainties are detailed in the Company's Annual Report.

**Proprietary
Pipeline**

**Partner
Pipeline**

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Strong and Differentiated Biopharmaceutical Pipeline



Successful Transformation from Technology Provider to Drug Development Company

- First partnered programs approaching the market
- Proprietary portfolio is gaining momentum

Increasing Confidence in Proprietary Portfolio

- Additional resources
- Majority of spending for clinical programs

Long-term Strategy

- Growing portfolio of innovative therapeutics
- Creation of long-term, sustainable value



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MOR208

First- & Best-in Class Potential

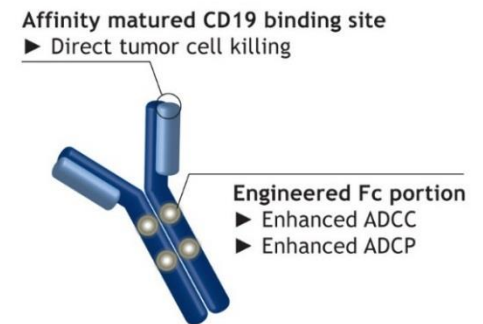


Updated phase 2 monotherapy data in NHL

First data from phase 2 combination trial MOR208 plus lenalidomide in CLL (IIT)

MOR208 - Differentiation

- Benign safety profile
- Long half-life in circulation
- Straightforward manufacturing
- Minor structural modification leads to significantly enhanced B cell depletion

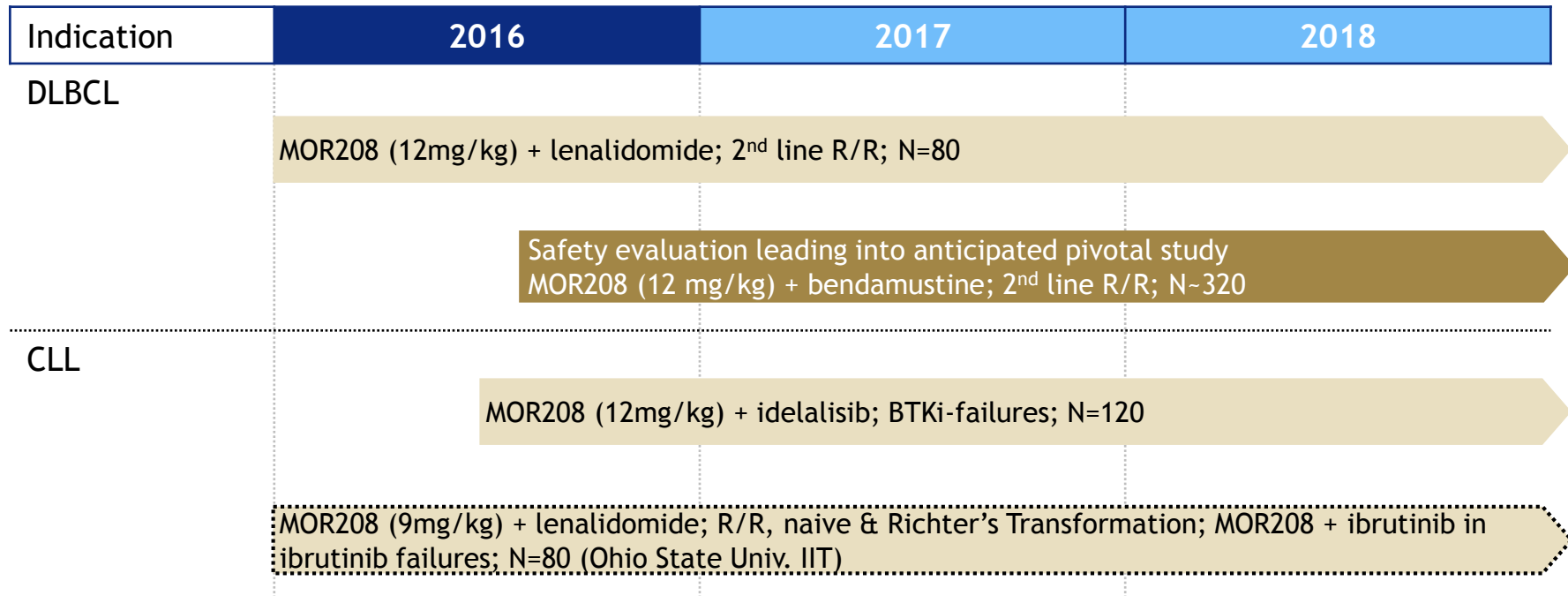


MOR208

MOR208 has the potential to become an important combination partner for the treatment of r/r DLBCL and Btk-inhibitor refractory CLL patients.

MOR208

Comprehensive Clinical Development Plan



- Phase 2
- Phase 2/3
- IIT: Investigator-initiated trial

MOR202

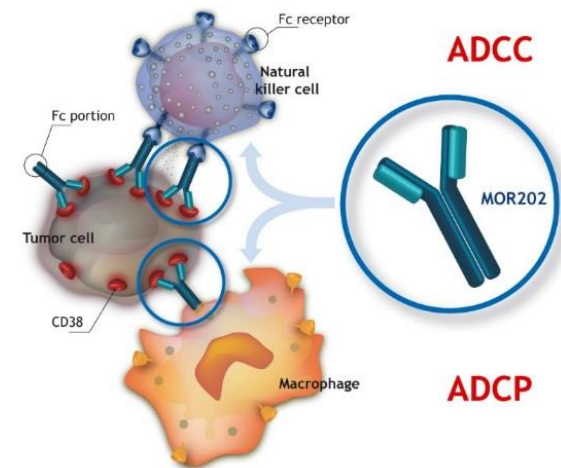
A Differentiated Antibody for Multiple Myeloma

Encouraging phase 1/2a data in r/r multiple myeloma

- MOR202
- MOR202 + lenalidomide
- MOR202 + pomalidomide

MOR202 - Differentiation

- Convenience: 2-hour infusion
- Best-in class infusion tolerability
- Preservation of NK cells



MOR202 is shaping up to be a genuine advance in the treatment of multiple myeloma, potentially offering patients superior outcomes and quality of life.



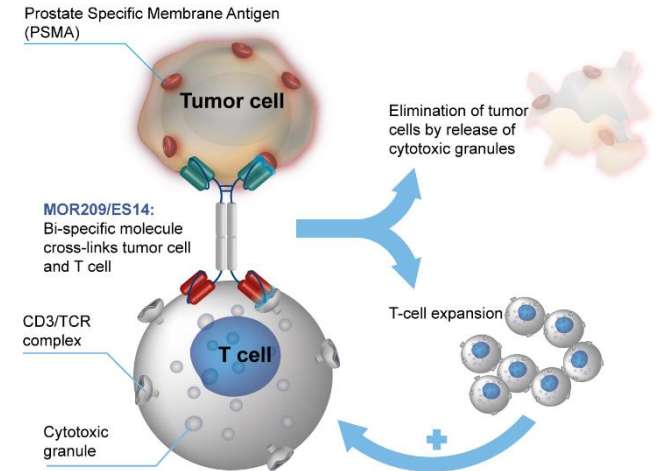
MOR209/ ES414

Co-development Agreement with Emergent BioSolutions

- Phase 1 clinical trial in mCRPC patients was started in March of 2015

Restructured Agreement with Emergent BioSolutions

- Adjustment of dosing regimen and administration
- Reduction of MorphoSys's cost sharing and reduced milestone payments



Clinical development will continue in 2016 under an adapted clinical development plan.



MOR106

Co-development with Galapagos

- Inflammatory program, novel mechanism-of-action
- Start of phase 1 in H1 of 2016
- Evaluation of safety and pharmacokinetics of the antibody in healthy volunteers and also in patients



MOR107

Lanthipeptide from the Lanthio Pharma acquisition

- Agonist of the angiotensin II type 2 receptor
- Strong anti-fibrotic efficacy in pre-clinical models
- Start of phase 1 expected in Q4 of 2016

MOR103/GSK3196165

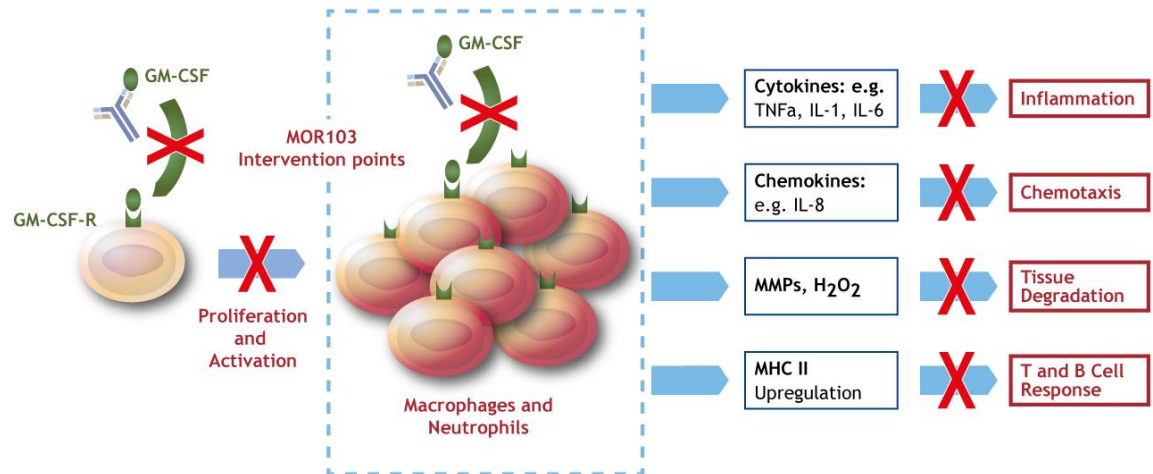
Anti-inflammatory Program Licensed to GSK



MOR103/ GSK3196165

License Agreement with GSK

- HuCAL antibody specific for GM-CSF
- Phase 2b study in RA (BAROQUE) ongoing
- Phase 2 in hand osteoarthritis to start in 2016



Powerful Technology Base Ensures Pipeline Sustainability

Innovative Targets

GPCRs, ion channels



Immune checkpoints



MHC-presented, tumor-associated peptides



Source of novel targets



Differentiated drug candidates

Proprietary Platforms

Antibody library



Protein optimization



Lantipeptides



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Bimagrumab (BYM338)

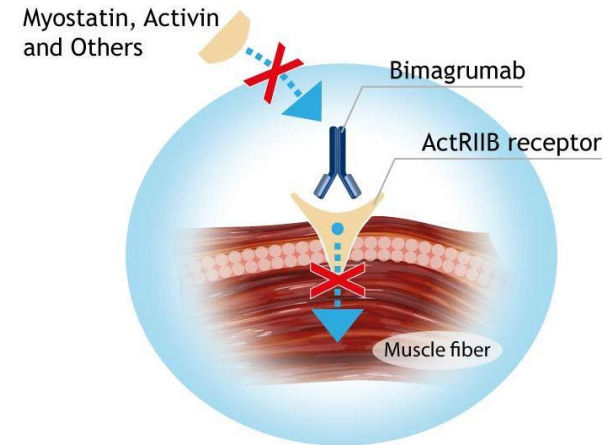
A Novartis Musculoskeletal Program

Bimagrumab

- HuCAL antibody specific for ActRIIB, antagonizes myostatin binding to muscle cells
- Development in muscle-wasting conditions
- FDA breakthrough therapy designation

Current Status

- Pivotal study in sIBM with 240 patients ongoing, phase 3 data expected in H1 2016
- Phase 2 studies in sarcopenia, cachexia and hip fracture surgery



Bimagrumab could become the first marketed product from our HuCAL technology platform

Guselkumab (CNTO1959)

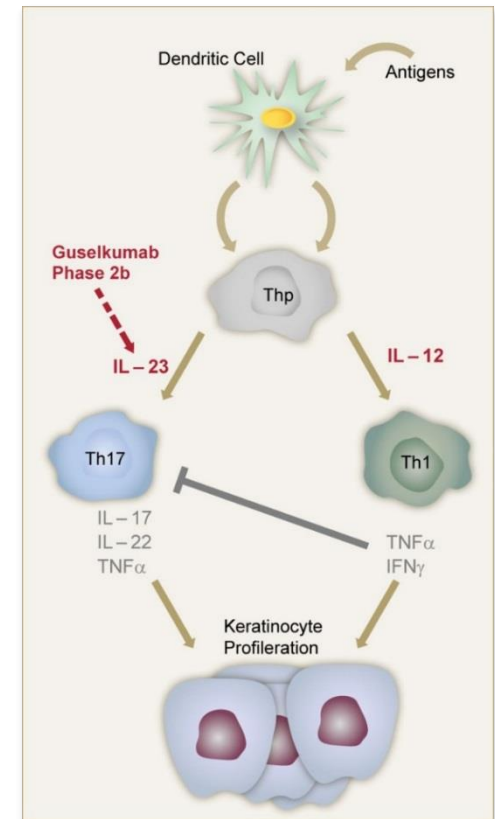
A Janssen Anti-Inflammatory Program

Guselkumab

- A HuCAL antibody specific for IL-23, does not bind IL-12
- Being developed in psoriasis and psoriatic arthritis

Current Status

- Six Phase 3 clinical trials ongoing
- First Phase 3 data expected in 2016
- Anticipated filing in 2016



Guselkumab would then be the second marketed product based on the HuCAL technology and the second source of royalty-based income



Gantenerumab

A Roche Alzheimer's Disease Program

- Higher doses will be tested
- Phase 2/3 (SCarlet RoAD) in prodromal Alzheimer's disease ongoing (open label extension)
- Phase 3 (Marguerite RoAD) in mild Alzheimer's disease ongoing (open label extension)



Anetumab ravtansine

A Bayer Cancer Program

- Start of global phase 2 clinical trial in mesothelioma; a study designed to support registration
- ADC program targeting mesothelin-expressing cancer cell

The MorphoSys Pipeline

25 Clinical Product Candidates, 103 Total



Most advanced development stage

Program	Partner	Target	Disease Area	Discovery	Preclinic	Phase 1	Phase 2	Phase 3	
Bimagrumab (BYM338)	Novartis	ActRIIB	sIBM (musculoskeletal)						Y
Guselkumab (CNTO1959)	Janssen	IL23p19	Psoriasis						Y
Gantenerumab	Roche	Amyloid-β	Alzheimer's disease						Y
MOR208	-	CD19	ALL, CLL, NHL						Y
MOR202	-	CD38	Multiple myeloma						Y
MOR103/GSK3196165	GSK	GM-CSF	Inflammation						Y
Anetumab Ravtansine (BAY94-9343)	Bayer	Mesothelin (ADC)	Solid tumors						Y
BHQ880	Novartis	DKK-1	Multiple myeloma						Y
BPS804	Mereo/Novartis	Sclerostin	Brittle bone syndrome						Y
CNTO3157	Janssen	-	Inflammation						Y
CNTO6785	Janssen	-	Inflammation						Y
LFG316	Novartis	C5	Eye diseases						Y
LJM716	Novartis	HER3	Cancer						Y
Tarextumab (OMP-59R5)	OncoMed	Notch 2	Solid tumors						Y
VAY736	Novartis	BAFF-R	Inflammation						Y
MOR209/ES414	Emergent	PSMA/CD3	Prostate cancer						Y
BAY1093884	Bayer	TFPI	Hemophilia						Y
BI-836845	BI	IGF-1	Solid tumors						Y
NOV-7	Novartis	-	Eye diseases						Y
NOV-8	Novartis	-	Inflammation						Y
NOV-9	Novartis	-	Diabetic eye diseases						Y
NOV-10	Novartis	-	Cancer						Y
NOV-11	Novartis	-	Blood disorders						Y
PF-05082566	Pfizer	4-1BB	Solid tumors						Y
Vantictumab (OMP-18R5)	OncoMed	Fzd 7	Solid tumors						Y
MOR106	Galapagos	-	Inflammation						Y
MOR107 (LP2)	-	AT2-R	Fibrosis						Y
Immuno-oncology program	Merck Serono	-	Cancer						Y
Immuno-oncology program	Immatics	-	Cancer						Y
6 MOR programs	-	-	Various						Y

89 Partnered Discovery Programs
 13 MOR Programs
 1 Outlicensed Program

In addition, 25 partnered programs in pre-clinic, and 43 partnered programs in discovery

Proprietary
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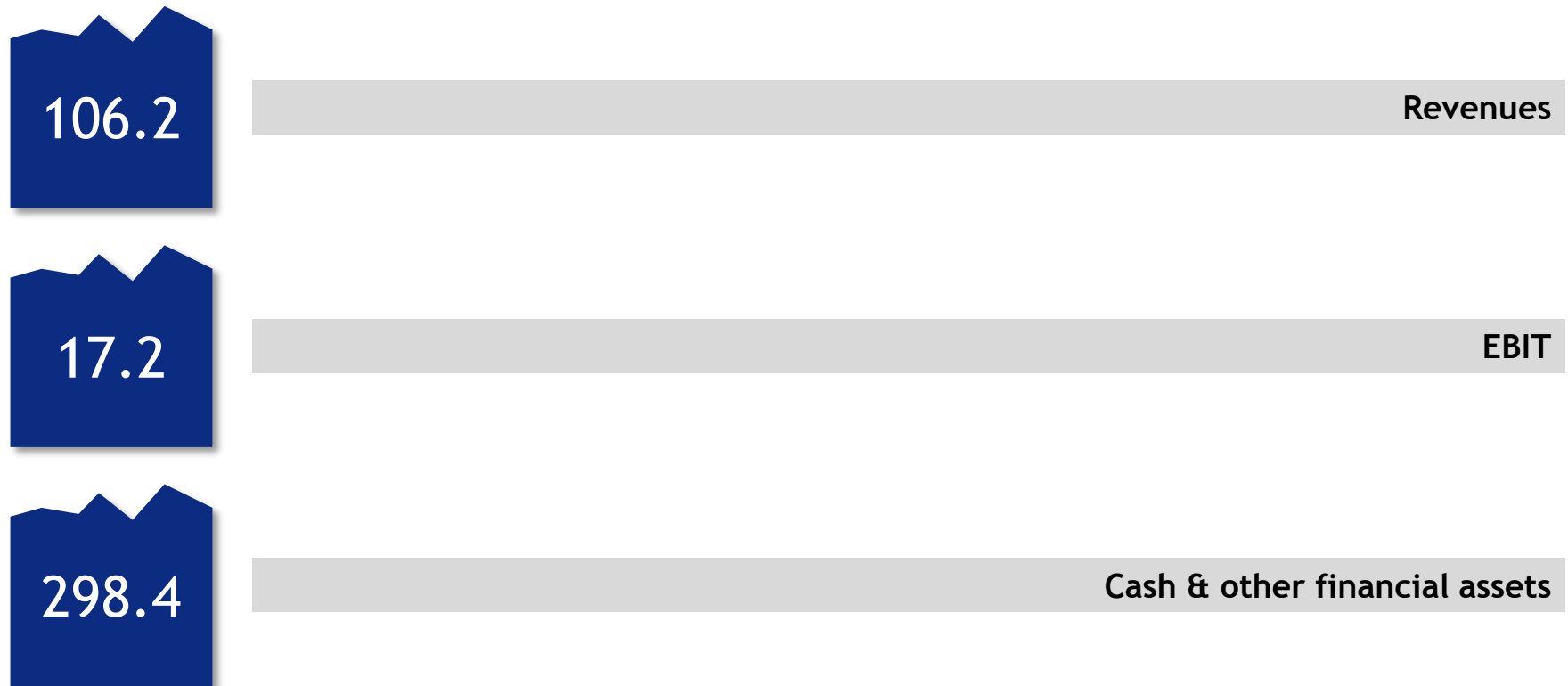
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Financial Highlights in 2015



in € million

Revenue and EBIT for 2015 contain a non-recurring contribution of approximately €59 million from the recognition of deferred revenues and a payment attributable to the early ending of the Celgene collaboration in March 2015.

FY2015: Income Statement

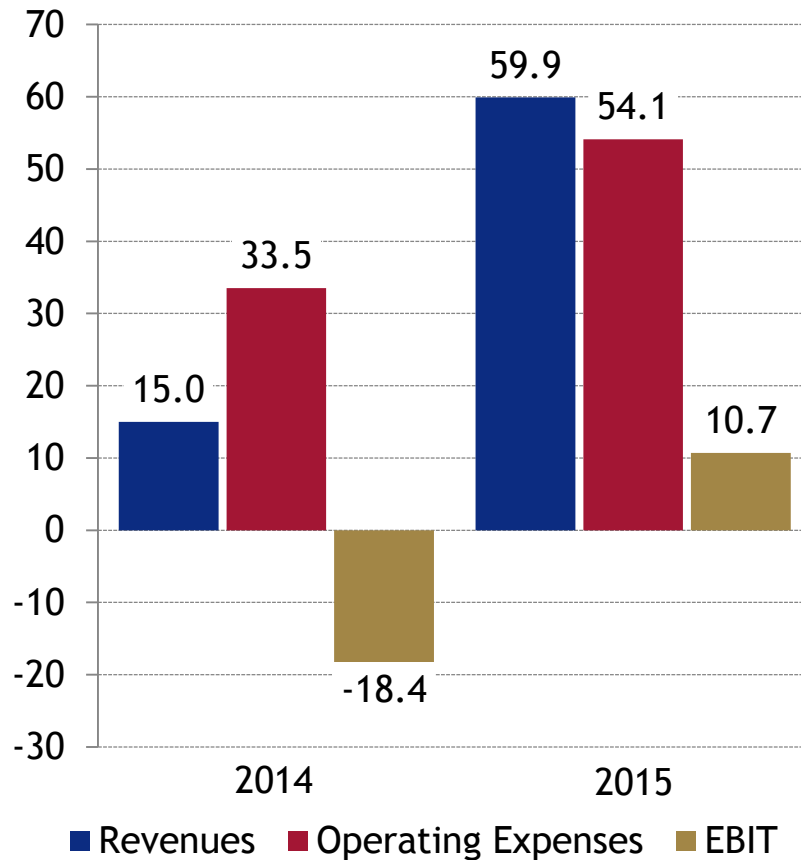


in € million*	2015	2014
Revenues	106.2	64.0
Research and Development Expenses	78.7	56.0
General & Administrative Expenses	15.1	14.1
Total Operating Expenses	(93.7)	(70.1)
Other Operating Income/(Expense)	4.7	0.2
EBIT	17.2	(5.9)
Finance Income	3.8	1.8
Finance Expenses	0.4	0.2
Income Tax (Expense)/Income	(5.7)	1.3
Consolidated Net Profit/(Loss)	14.9	(3.0)
Diluted Net Profit/(Loss) per Share (in €)	0.57	(0.12)

* Differences due to rounding

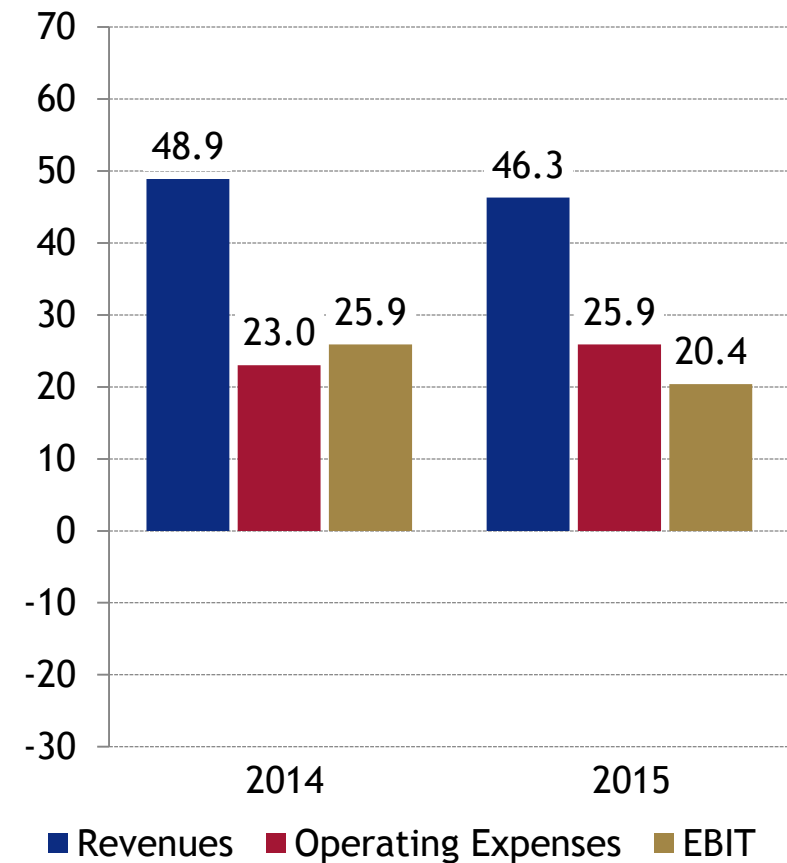
Proprietary Development

in € million



Partnered Discovery

in € million



FY2015: Balance Sheet



in € million*	Dec 31, 2015	Dec 31, 2014
Assets		
Cash and Cash Equivalents [#]	90.9	32.2
Available-for-sale Financial Assets [#]	64.3	106.1
Bonds, Available-for-sale [#]	33.1	7.5
Financial Assets classified as Loans & Receivables [#]	94.6	157.0
Other Current Assets	17.2	19.6
Total Current Assets	300.1	322.4
Financial Assets classified as Loans & Receivables, Net of Current Portion [#]	15.5	50.0
Total Non-current Assets	100.0	104.1
Total Assets	400.1	426.5
Liabilities & Stockholders' Equity		
Total Current Liabilities	27.5	32.7
Total Non-current Liabilities	9.9	45.0
Total Stockholders' Equity	362.7	348.8
Total Liabilities & Stockholders' Equity	400.1	426.5

[#] As of Dec. 31, 2015, MorphoSys held liquid funds and marketable securities as well as other short-term and long-term financial assets in the amount of €298.4 million (Dec. 31, 2014: € 352.8 million) * Differences due to rounding

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in € million	2015A	Guidance 2016
Group Revenues	106.2	47 to 52
Proprietary R&D Expenses (incl. Technology Development)	56.6	76 to 83
EBIT	17.2	-58 to -68

Cash, cash equivalents & marketable securities
as well as other short-term and long-term financial assets

298.4

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What to Expect?



Bimagrumab	sIBM	Data from pivotal trial and regulatory filing expected
Guselkumab	Psoriasis	Data from 3 pivotal trials and regulatory filing expected
MOR208	DLBCL	<ul style="list-style-type: none"> ■ Phase 2 lenalidomide combo trial to start in Q1 2016 ■ Phase 2 bendamustine combo safety evaluation to start mid 2016 ■ Phase 3 bendamustine combo pivotal study planned for 2017 ■ First data of combination trials in 2017
	CLL	<ul style="list-style-type: none"> ■ Phase 2 idelalisib combo trial to start in Q1 2016 ■ First data of combination trial in 2017
MOR202	MM	Updated data from phase 1/2a trial at ASCO 2016 and ASH 2016
MOR209	Prostate cancer	Continuation of trial under amended protocol, clinical data in 2017
MOR106	Inflammation	Start of phase 1 with Galapagos in H1 2016
MOR107	Fibrosis	Start of phase 1 in Q4 2016
MOR103	RA	<ul style="list-style-type: none"> ■ Start of phase 1b/2a in osteoarthritis of the hand
	Osteoarthritis	<ul style="list-style-type: none"> ■ Data from the phase 2b in RA in 2017
Pipeline		<ul style="list-style-type: none"> ■ Up to 5 new program starts ■ Around 5 clinical milestones

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Thank You

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