

Year End Results

Period Ended December 31, 2018



Forward Looking Statement

This presentation contains forward looking statements. The words “believe”, “expect”, “anticipate”, “intend” and “plan” and similar expressions identify forward looking statements. All statements other than statements of historical facts included in this presentation, including, without limitation, those regarding our financial position, business strategy, plans and objectives of management for future operations (including development plans and objectives relating to our products), are forward looking statements. Such forward looking statements involve known and unknown risks, uncertainties and other factors which may cause our actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by such forward looking statements. Such forward looking statements are based on numerous assumptions regarding our present and future business strategies and the environment in which we will operate in the future. The important factors that could cause our actual results, performance or achievements to differ materially from those in the forward looking statements include, among others, risks associated with product discovery and development, uncertainties related to the outcome of clinical trials, slower than expected rates of patient recruitment, unforeseen safety issues resulting from the administration of our products in patients, uncertainties related to product manufacturing, the lack of market acceptance of our products, our inability to manage growth, the competitive environment in relation to our business area and markets, our inability to attract and retain suitably qualified personnel, the unenforceability or lack of protection of our patents and proprietary rights, our relationships with affiliated entities, changes and developments in technology which may render our products obsolete, and other factors. Further, certain forward looking statements are based upon assumptions of future events which may not prove to be accurate. The forward looking statements in this document speak only as at the date of this presentation.

Key Achievements 2018

DARZALEX® (daratumumab)

Regulatory approvals

- US & Europe in combination with bortezomib, melphalan and prednisone (VMP) in frontline multiple myeloma (MM)
- Europe for split dosing regimen and in US in 2019

Regulatory submissions

- China for relapsed/refractory MM
- Japan in combination with VMP in frontline MM
- US in combination with lenalidomide & dexamethasone (DRd) in frontline ASCT-ineligible MM (2019)

Positive topline Phase III results in frontline MM

- MAIA study combining daratumumab with lenalidomide & dexamethasone (Rd)
- CASSIOPEIA study combining daratumumab with bortezomib, thalidomide and dexamethasone (VTd)

USD 75M milestone from Janssen for \$2 billion in sales in calendar year

USD 2,025M net sales by Janssen in 2018 - resulting in DKK 1,708M in royalties

Key Achievements 2018

Additional Highlights

Pipeline Progress

- Recruitment completed in Phase III relapsing MS studies of subcutaneous ofatumumab
- 4 new studies with tisotumab vedotin including potential registration Phase II study in cervical cancer
- Expansion phase started in enapotamab vedotin Phase I/II study in solid tumors
- First patients dosed in HexaBody[®]-DR5/DR5 Phase I/II study in solid tumors
- First patients dosed in DuoBody[®]-CD3xCD20 Phase I/II study in B-cell malignancies

Other Key Highlights

- Strategic partnership with Immatics
- Successful Capital Markets Day
- HexElect[™] antibody platform introduced
- Improved revenue by DKK 660M vs. 2017

Income Statement: Year Ended December 31

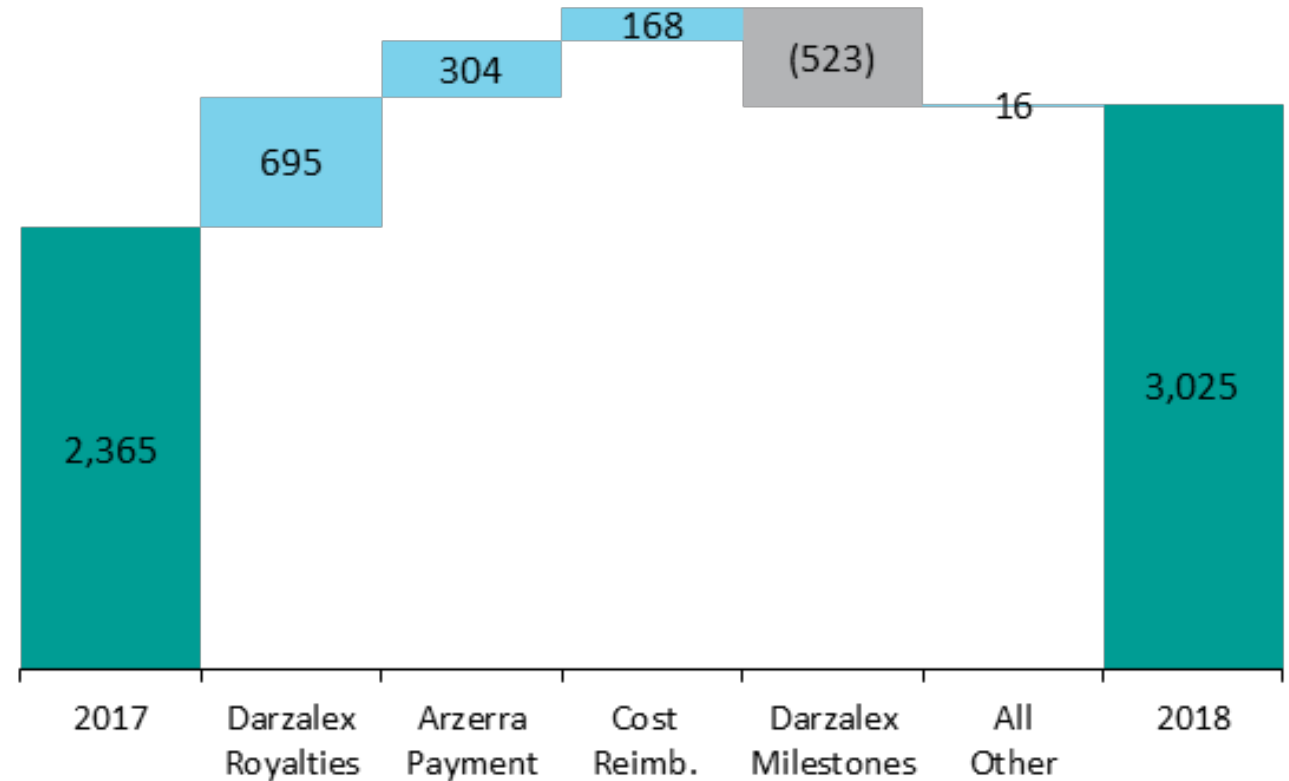
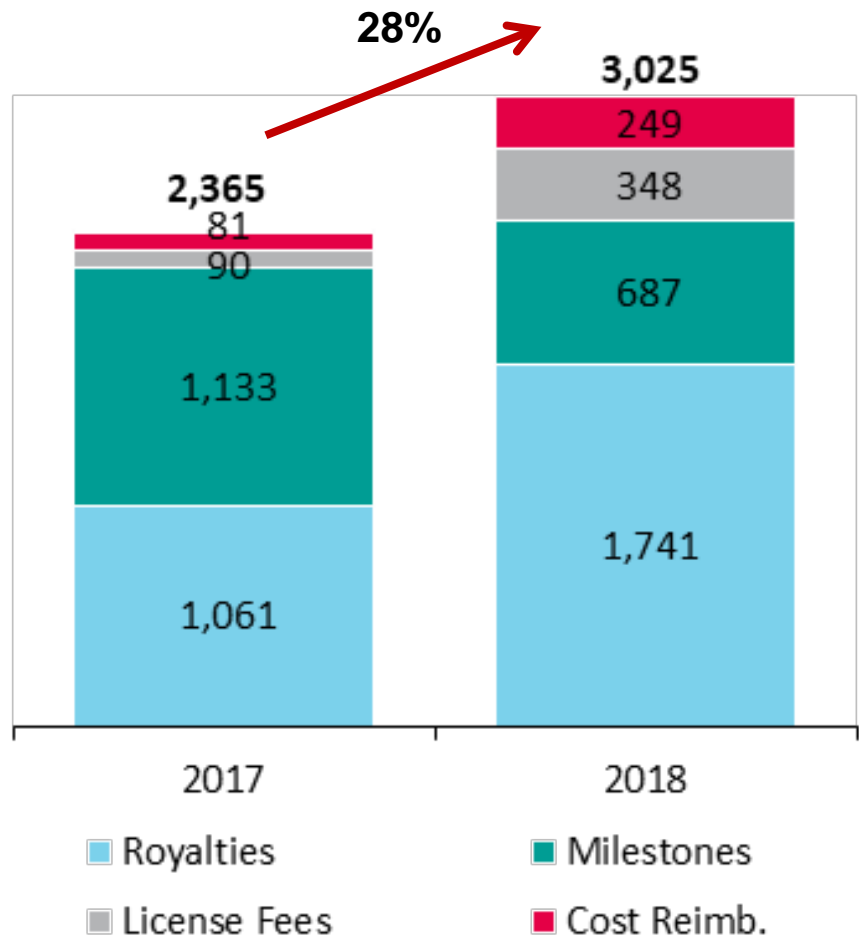
	<u>2018</u>	<u>2017</u>		<u>2018</u>	<u>2017</u>
	DKK millions		Change	USD millions *	
Darzalex Royalties	1,708	1,013	695	262	155
Reimbursement Income	249	81	168	38	12
Other Revenue	1,068	1,271	(203)	164	195
Total Revenue	3,025	2,365	660	464	362
R&D Costs	(1,431)	(874)	(557)	(219)	(134)
G&A Expenses	(214)	(147)	(67)	(33)	(23)
Operating Expenses	(1,645)	(1,021)	(624)	(252)	(157)
Operating Result	1,380	1,344	36	212	205
Net Financial Items	232	(280)	512	36	(43)
Tax	(140)	40	(180)	(21)	6
Net Result	1,472	1,104	368	227	168

* USD 1.00 = DKK 6.5213 (Danish Central Bank spot rate on December 31, 2018)

Overview – 2018 Guidance vs. Actual

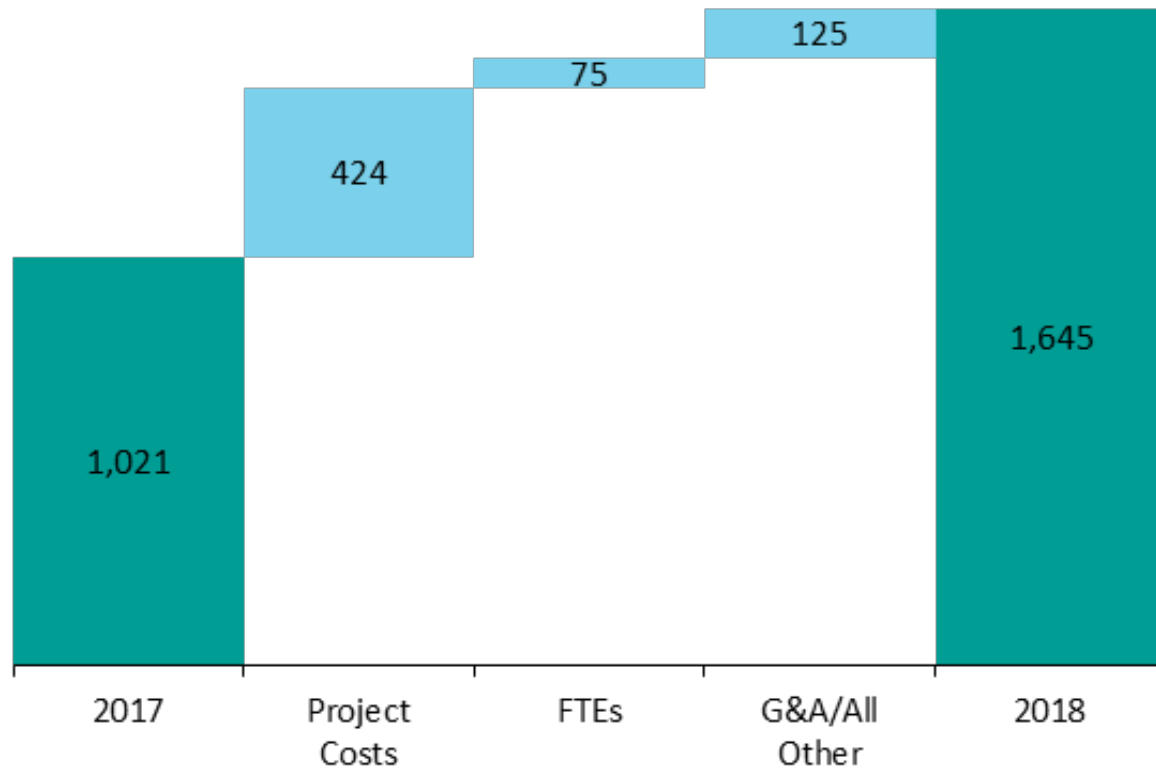
DKK Millions	2018 Guidance	2018 Actual
Revenue	2,700 - 3,100	3,025
Operating expenses	(1,400) - (1,600)	(1,645)
Operating income	1,300 - 1,500	1,380

Revenue 2018 vs. 2017: Year Ended December 31

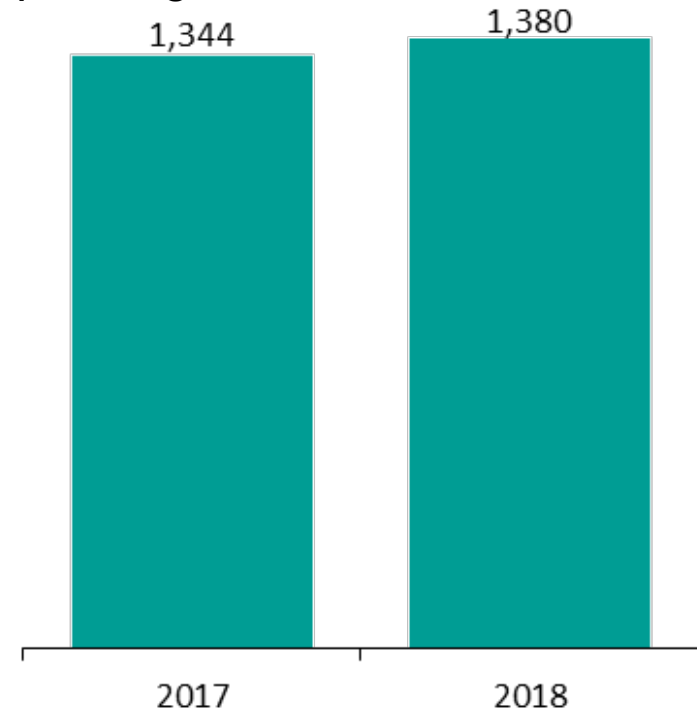


Operating Result: Investing in Our Pipeline

Operating Expenses increased 61% (+DKK 624M), driven by additional pipeline investment



Revenue growth outpaced expense increase - driving DKK 36M higher Operating Result



Overview - 2019 Guidance

Income Statement

DKK Millions	2018 Actual	2019 Guidance	Change	%
Revenue	3,025	4,600	1,575	52%
Operating Expenses	(1,645)	(2,600)	(955)	58%
Operating Income	1,380	2,000	620	45%

Revenue Detail

DKK Millions	2018 Actual	2019 Guidance	Comments
Darzalex Royalties	1,708	2,685	Darzalex Net Sales USD 3.0 billion
Darzalex Milestones	586	1,500	Milestone payment of USD 150 million (DKK 900 million) from Darzalex Net Sales of USD 3.0 billion
All Other	731	415	2018 includes Novartis one-time payment of USD 50 million (DKK 304 million)
Total Revenue	3,025	4,600	

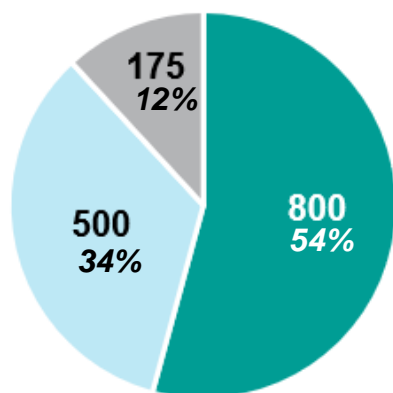
Overview - 2019 Guidance – Pipeline Investment

Expense Detail

DKK Millions	2018 Actual	2019 Guidance	Change	%	Comments
Project Investment	985	1,475	490	50%	Driven by Top 10 Projects
Personnel Costs	365	650	285	78%	Increase in 2019 by 180 FTEs
Business Support	295	475	180	61%	Including Technologies & Systems, Commercial & Medical Affairs
Total Operating Expenses	1,645	2,600	955	58%	

Total Project Investment

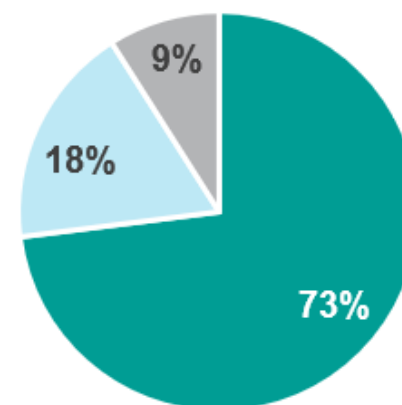
1,475 Top 10 = 1,300



- Tisotumab Vedotin & Enapotamab Vedotin
- Next 8
- All Other

2019 FTE Growth

180 FTEs



- R&D
- Support Functions
- Medical Affairs/Commercial

Key 2019 Priorities - External

Building a Robust Differentiated Product Portfolio

Priority	✓	Targeted Milestones
Daratumumab		<ul style="list-style-type: none"> » FDA decision on Phase III MAIA multiple myeloma (MM) submission » FDA decision on Phase III CASSIOPEIA MM submission » Phase III COLUMBA MM subcutaneous (SC) daratumumab safety & efficacy analysis
Ofatumumab		<ul style="list-style-type: none"> » Phase III ASCLEPIOS I & II relapsing multiple sclerosis SC ofatumumab study completion and reporting
Tisotumab vedotin		<ul style="list-style-type: none"> » Phase II innovaTV 204 tisotumab vedotin recurrent / metastatic cervical cancer study enrollment complete by mid year
Innovative pipeline		<ul style="list-style-type: none"> » Phase II enapotamab vedotin expansion cohort efficacy analysis » Phase I/II HexaBody-DR5/DR5 initial clinical data » Phase I/II DuoBody-CD3xCD20 clinical data dose escalation cohorts » File INDs or CTAs for 3 new products

Q&A

Upcoming Investor & Other Events

Carnegie Healthcare Seminar 2019, March 5

Genmab Annual General Meeting, March 29

H.C. Wainright Global Life Science Conference, April 9

Kempen Life Sciences Conference, April 16-17

