Year End Results

Period Ended December 31, 2017





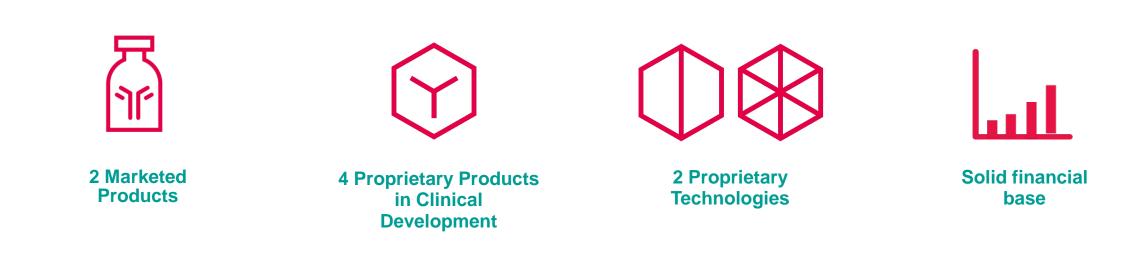
Forward Looking Statement

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.



Genmab At-A-Glance Solid Foundation

By 2025, our own product has transformed cancer treatment, and we have a pipeline of knock-your-socks-off antibodies





Building a Stronger Genmab For the Future



Doubled proprietary clinical pipeline in 2017

Continued investments in pre-clinical pipeline

Targeting 4 INDs over next 4 years

DARZALEX® royalties fund pipeline investments

Building commercialization and launch capabilities

RR) Increasing our staff to expand competencies

Growth will be carefully controlled



Key Achievements 2017

DARZALEX[®] (daratumumab)

- Approved in combination with len/dex and bort/dex for relapsed/refractory multiple myeloma (MM) in EU
- Approved in combination with pom/dex for relapsed-refractory MM in US
- Approved for relapsed or refractory MM in Japan
- Positive Phase III ALCYONE data in frontline MM – regulatory applications submitted in EU & US
- Multiple new studies announced
- DKK 1,109M in milestones from daratumumab collaboration with Janssen
- USD 1,242M net sales by Janssen in 2017
 - Resulting in DKK 1,013M in royalties

Pipeline Progress

- Tisotumab vedotin Phase I/II preliminary cervical cancer data
- DuoBody-CD3xCD20 IND
- HexaBody-DR5/DR5 IND
- New Janssen DuoBody program JNJ-64007957 Phase I study

Other Key Highlights

- New Chief Development Officer Judith Klimovsky, MD
- Seattle Genetics exercised option to codevelop tisotumab vedotin following positive cervical cancer data
- Improved revenue by DKK 549M vs. 2016



Income Statement: Year Ended December 31

	<u>2017</u> <u>2016</u> DKK millions Change		<u>2017</u> <u>2016</u> USD millions **		
Royalties Milestones Other Revenue Total Revenue	1,061 1,133 <u>171</u> 2,365	521 1,187 <u>108</u> 1,816	540 (54) 63 549	171 182 <u>28</u> 381	84 191 <u>18</u> 293
Operating Expenses	(1,021)	(763)	(258)	(165)	(122)
Operating Result	1,344	1,053	291	216	171
Net Financial Items Tax	(280) 40	77 57	(357) (17)	(45) 6	12 9
Net Result	1,104	1,187	(83)	177	192
Cash position increase* Cash position at end of period*	1,501 5,423	429 3,922		242 874	69 632

*Cash, cash equivalents, bank overdraft, and marketable securities

** USD 1.00 = DKK 6.2067 (Danish Central Bank spot rate on December 31, 2017)



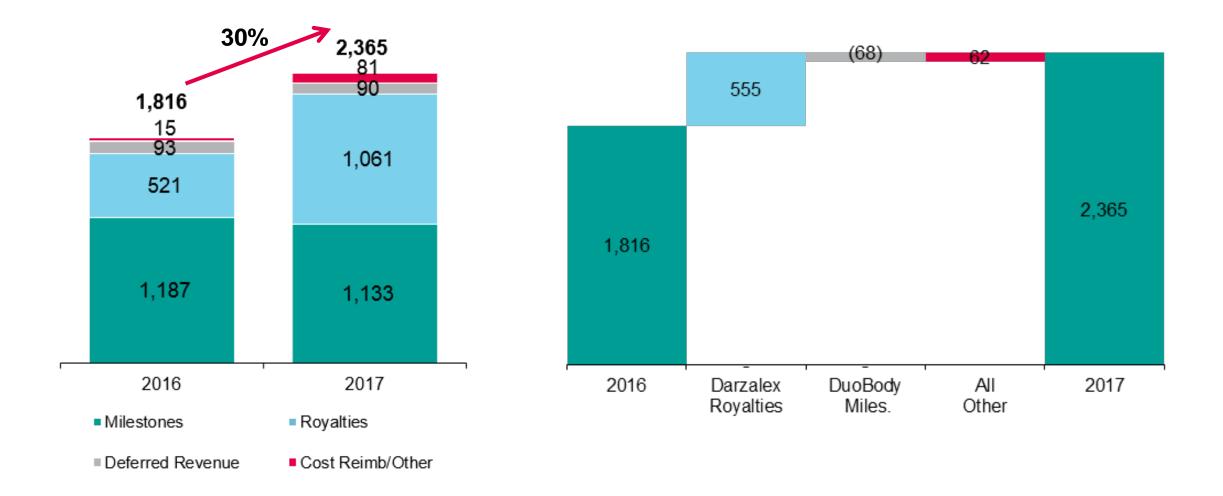
Overview – 2017 Guidance vs. Actual

DKK Millions	Original Guidance Feb 2017	Latest Guidance Nov 2017	2017 Actual
Revenue	1,950 – 2,150	2,240 - 2,440	2,365
Operating expenses	(1,000) – (1,100)	(1,000) – (1,100)	(1,021)
Operating income	900 - 1,100	1,190 – 1,390	1,344
Cash position at end of year*	> 4,500	> 4,900	5,423

*Cash, cash equivalents, and marketable securities



Revenue 2017 vs. 2016: Year Ended December 31

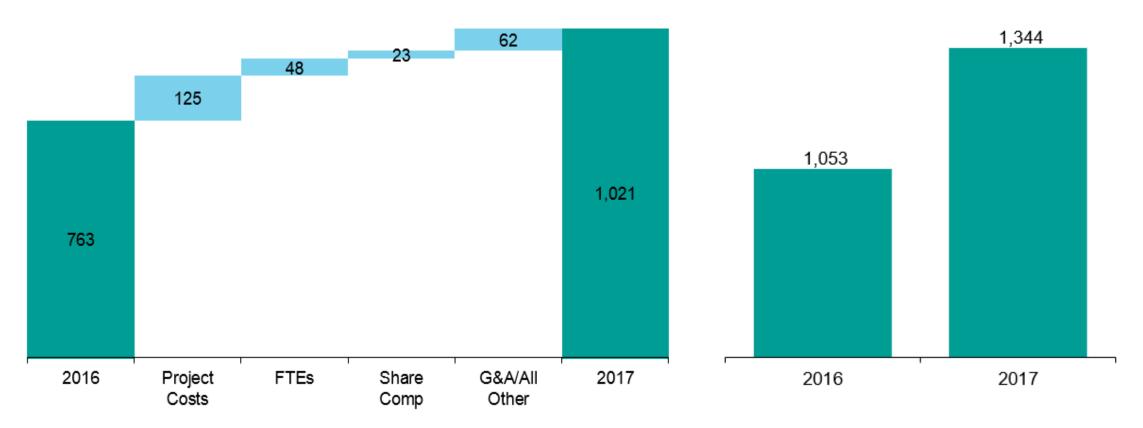




Operating Result: Investing in Our Pipeline

Operating Expenses increased 34% (+DKK 258M), driven by additional pipeline investment

Revenue growth outpaced expense increase - driving 28% (+DKK 291M) higher Operating Result

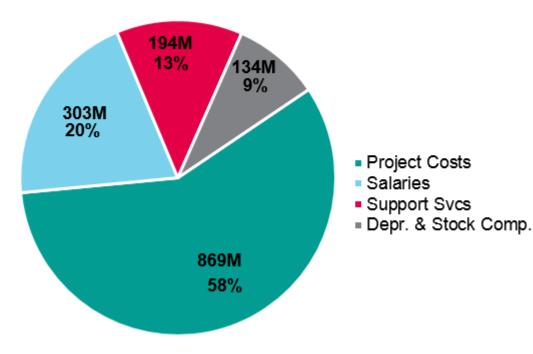




Overview – 2018 Guidance

DKK Millions	2018 Guidance	2017 Actual
Revenue	2,700 - 3,100	2,365
Operating expenses	(1,400) - (1,600)	(1,021)
Operating income	1,300 - 1,500	1,344

2018 Expense Base DKK 1,500M



DARZALEX sales mid-point USD 2.15 Bn

 Genmab's estimate of DARZALEX net sales USD 2.0 - 2.3 billion

Revenue mid-point DKK 2,900M

- DARZALEX royalties DKK 1,750M
- DARZALEX milestones DKK 550M
- Novartis one-time payment of DKK 300M

Expense mid-point DKK 1,500M

- Continued investment in our clinical & pre-clinical pipeline
- 10 pipeline projects drive ~DKK 765M, 51% of total expense



2018 Company Goals Maximizing Differentiated Product Portfolio Value

Priority	✓	Targeted Milestone
Maximize daratumumab progress		 » FDA and EMA decision on Phase III ALCYONE multiple myeloma (MM) submission » Start new Phase III MM study » Report early clinical data in solid tumors » Phase III MAIA MM efficacy analysis in frontline » Phase III CASSIOPEIA MM efficacy analysis in frontline
Optimize of atumumab value		» Complete recruitment Phase III subcutaneous of atumumab relapsing MS studies
Maximize tisotumab vedotin progress		 Start two Phase II studies cervical cancer (recurrent / metastatic & combination study in frontline) Start Phase II study in additional solid tumor indications
Strengthen differentiated product pipeline and technology partnership portfolio		 Start HuMax-AXL-ADC expansion phase in ongoing Phase I/II study Progress HexaBody-DR5/DR5 Phase I/II study Progress DuoBody-CD3xCD20 Phase I/II study Accelerate proprietary DuoBody Immuno-Oncology programs towards clinic Enter new technology or product collaborations
Disciplined financial management and building a commercial footprint		 » Execute controlled company growth with selective investments in product & technology pipeline » Continue investing in building commercialization and launch capabilities

Q&A

Upcoming Investor & Other Events Credit Suisse Healthcare Conference, February 27 Annual General Meeting 2018, April 10 Kempen Life Sciences Conference, April 18

