To improve the lives of patients by creating & developing innovative antibody products





Approved Partnered Products

Solid Financial Base

DARZALEX® (daratumumab) / DARZALEX FASPRO™ (daratumumab & hyaluronidase-fihj)¹, Kesimpta® (ofatumumab)², TEPEZZA® (teprotumumab)³ Arzerra® (ofatumumab)⁴



Our Own Clinical Pipeline

Potential first-in-class / Best-in-class

Tisotumab vedotin⁵, enapotamab vedotin, HexaBody[®]-DR5/DR5, epcoritamab⁶, DuoBody-PD-L1x4-1BB (GEN1046)⁷, DuoBody-CD40x4-1BB (GEN1042)⁷ DuoHexaBody[®]-CD37 (GEN309)⁶ & DuoBody[®]-CD3x5T4 (GEN1044)⁶



Partner-owned Programs in the Clinic

Programs Built on Genmab's Innovation

15 in clinical development with partners with royalty streams for Genmab should they come to market. Includes 7 DuoBody products with Janssen



Technologies & Preclinical

R&D Engine: Powerhouse of Innovation

DuoBody tech., HexaBody tech., DuoHexaBody tech., HexElect® tech. & Rich preclinical pipeline incl. HexaBody-CD38 (GEN3014)⁸



Our Three-pronged Strategy

- · Focus on core competence
- · Turn science into medicine
- · Build a profitable & successful biotech



Our Vision

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



Proprietary technologies allow us to build a world-class pipeline



Match in-house expertise with strategic partnerships



Strong pipeline of first-in-class / best-in-class products



Deep insight into antibody biology & disease targets

2020 Guidance

Income Statement	DKKM	USDM ⁹
Revenue	9,250 – 9,850	1,423 – 1,515
Operating expenses	(3,850) – (3,950)	(592) – (608)
Operating income	5,350 - 5,950	823 - 915

⁹USD 1.00 = DKK 6.50

DKK

96B

~USD 14B¹⁰

2019 year-end market cap

2,728M

¹⁰USD 1.00 = DKK 6.6759

2019 operating expenses 87% invested in R&D

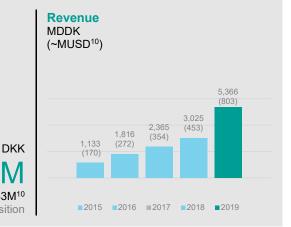
DKK

5,366M

~USD 803M¹⁰

2019 revenue 77% increase vs 2018

> 10,971M ~USD 1,643M¹⁰ 2019 year end cash position



com

In dev. by Janssen; In dev. by Novartis; In dev. by Horizon; "Commercialized by Novartis, no longer in active clinical development, "50:50 w/ Seattle Genetics; "50:50 w/ AbbVie; "50:50 w/ BioNTech; "Genmab is developing HexaBody-CD38 in an exclusive worldwide license and option agreement with Janssen Biotech, Inc This document contains forward looking statements that involve significant risks and uncertainties. For a further discussion of these risks, please refer to the risk management sections in Genmab's most recent financial reports, which are available on www.genmab.com and the risk factors included in Genmab's most recent Annual Report on form 20-7 and other filings with the U.S. Securities and Exchange Commission (SEC), which are available at www.ges.cgw. Nov 4, 2020

Innovation Powerhouse

Genmab Proprietary¹¹ Products

Product	Target	Rights	Indications	Most Advanced Dev. Phase					
				Pre-clin.	ı	1/11	II III	Appr	
Tisotumab vedotin	TF	50:50 Genmab Seagen	Cervical cancer						
			Ovarian cancer						
			Solid tumors						
Enapotamab vedotin	AXL	Genmab	Solid tumors						
Epcoritamab	CD3,	50:50 Genmab	Hem. malig.						
	CD20	AbbVie							
DuoBody-PD-L1x4-1BB	PD-L1, 4-1BB	50:50 Genmab BioNTech	Solid tumors						
DuoBody-CD40x4-1BB	CD40, 4-1BB	50:50 Genmab BioNTech	Solid tumors						
HexaBody-DR5/DR5	DR5	Genmab	Solid tumors						
DuoHexaBody-CD37	CD37	50:50 Genmab AbbVie	Hem. malig.						
DuoBody-CD3x5T4	5T4	50:50 Genmab AbbVie	Solid tumors					•	
IND/CTAs in 2020: HexaBoo	dy-CD38 ¹²								
Products Created by Ger	nmab ¹³								
Daratumumab	CD38	Janssen ¹⁴	MM						
			AL Amyloidosis						
			Non-MM blood						
			cancers						
Ofatumumab	CD20	Novartis ^{14,15}	RMS						
Teprotumumab	IGF-1R	Horizon ¹⁴	TED						
			dcSSc						
Partner-owned products	incorporating Ge	enmab's Innova	ation ¹⁶						
Amivantamab	EGFR, cMet	Janssen	NSCLC						
Teclistamab	BCMA, CD3	Janssen	R/R MM						
Camidanlumab tesirine	CD25	ADC Therapeutics	R/R Hodgkin						
			lymphoma						
	11 45		Solid tumors					 	
PRV-015 (AMG 714)	IL-15		Celiac disease					-	
Mim8	FIX(a), FX	Novo Nordisk	Hemophilia A						
Talquetamab	GPRC5D, CD3		R/R MM						
JNJ-63709178	CD123, CD3	Janssen	AML						
JNJ-63898081	PSMA, CD3	Janssen	Solid tumors					-	
JNJ-67571244	CD33, CD3	Janssen	R/R AML; MDS					-	
JNJ-70218902	Undisclosed	Janssen	Solid tumors						
HuMax-IL8	IL8	BMS	Adv. cancers						
Lu AF82422	alpha-synucleir	n Lundbeck	Parkinson's						
~20 active pre-clinical progra	•	oprietary progran aBody and Hexal							

Products & Technologies

Tisotumab vedotin

- · ADC in development to treat solid tumors
- Very favorable data, Ph 2 cervical cancer
- Ph 2 studies in ovarian & other solid tumors
- 50:50 agreement w/ Seagen

Enapotamab vedotin

- · ADC in development to treat solid tumors
- Ph 1/2 trial for multiple solid tumors

HexaBody-DR5/DR5 (GEN1029)

- · Proprietary antibody therapeutic
- · Phase 1/2 trial in solid tumors

Epcoritamab (DuoBody-CD3xCD20)

- · Bispecific antibody created w/ DuoBody tech
- Phase 1/2 trial in B-cell malignancies
- 50:50 collaboration w/ AbbVie

DuoBody-PD-L1x4-1BB (GEN1046)

- Bispecific antibody created w/ DuoBody tech
- Phase 1/2 trial in solid tumors
- 50:50 agreement w/ BioNTech

DuoBody-CD40x4-1BB (GEN1042)

- · Bispecific antibody created w/ DuoBody tech
- Phase 1/2 trial in solid tumors
- 50:50 agreement w/ BioNTech

DuoHexaBody-CD37 (GEN3009)

- · Based on combo DuoBody & HexaBody
- · FiH trial commenced
- 50:50 collaboration w/ AbbVie

DuoBody-CD3x5T4 (GEN1044)

- · Bispecific antibody created w/ DuoBody tech
- · FiH trial commenced
- 50:50 collaboration w/ AbbVie

Kesimpta (ofatumumab)

- SubQ formulation approved in US for RMS¹⁴
 Combines dual targeting with enhanced
- · In development by Novartis

Founded: 1999

Exchanges: CSE / Nasdag

Symbol: GMAB

Shares outstanding: 65,498,346 Market Cap as of Nov. 4, 2020:

DKK 154B (~USD 24,064M)

DARZALEX (daratumumab) / DARZALEX FASPRO (daratumumab & hyaluronidase-fihj)

- · First-in-class CD38 antibody
- Approved in certain territories for various multiple myeloma (MM) indications¹⁴
- First & only SC CD38 antibody approved
- In development by Janssen
- 2019 net sales by Janssen: \$2,998M, DKK 3,132M royalties to Genmab

TEPEZZA (teprotumumab)

- Dev. & manufactured by Horizon Therapeutics
- Approved in US for thyroid eye disease¹⁴
- Created under collaboration with Roche
- Under terms of agreement with Roche, Genmab to receive mid-single digit royalties on sales

DuoBody Platform

- · Bispecific antibody technology platform
- Potential in cancer, autoimmune, infectious, cardiovascular, central nervous system diseases & hemophilia
- · Multiple commercial & research collaborations

HexaBody Platform

- Enhanced potency antibody technology platform
- Broadly applicable technology that builds on natural antibody biology

DuoHexaBody Platform

- Antibody technology that combines
- DuoBody & HexaBody platforms
- Creates bispecific antibodies with targetmediated enhanced potency

HexElect Platform

- Antibody technology platform inspired by HexaBody platform
- Combines dual targeting with enhanced selectivity & potency

Executive Management:

Jan G. J. van de Winkel, Ph.D. President & CEO Anthony Pagano, EVP & CFO Judith Klimovsky M.D., EVP & CDO Anthony Mancini, EVP & COO

¹¹Certain product candidates in development with partners, as noted

¹²Genmab is developing HexaBody-CD38 in an exclusive worldwide license and option agreement with Janssen Biotech, Inc.

¹³Out-licensed products marketed by partner

¹⁴See local country prescribing information for precise indications

¹⁵Also approved as Arzerra for certain CLL indications, not in active clinical development

¹⁶Products under development by a third-party incorporating Genmab's technology and innovation.