Innovating Antibodies, Improving Lives

Investor Presentation
August 2018

Genmab

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 does not undertake any obligation to update or revise forward looking statements in this presentation nor to confirm such statements to reflect subsequent events or circumstances after the date made or in relation to actual results, unless required by law.

Genmab At-A-Glance

Core Purpose, Strategy & Vision

Core Purpose

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

Our Strategy

- Turn science into medicine
- Build a profitable & successful biotech
- Focus on Core Competence

Vision

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

Genmab At-A-Glance
Solid Foundation

DARZALEX®
Arzerra®

2 marketed products generating royalty income

Tisotumab vedotin
HuMax®-AXL-ADC
HexaBody-DR5/DR5
DuoBody-CD3xCD20

4 exciting proprietary clinical programs

DuoBody® Platform
HexaBody® Tech.

2 proprietary next generation technologies for robust pre-clinical pipeline

Solid financial base

Aim to own at least 50% of product rights
Allows for building capabilities to market own product in future

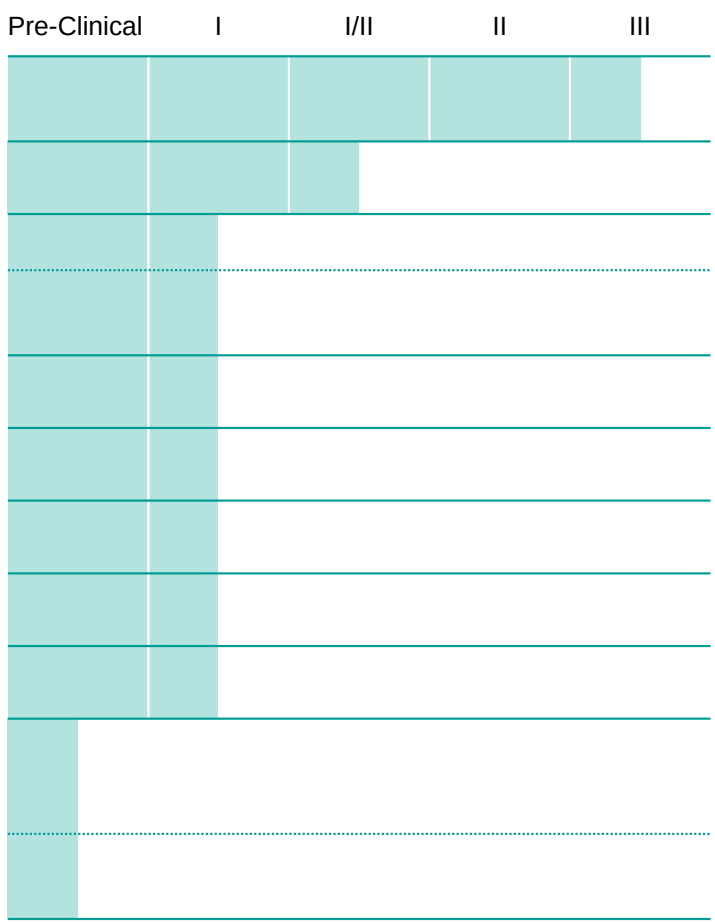
Innovative Clinical & Pre-clinical Pipeline

Development for Marketed & Genmab Proprietary Products

Product	Disease Indications	Development Phase
		Pre-Clinical / I / I/II / II / III
Daratumumab BTD (2 - MM) Target: CD38 Partner: Janssen	Multiple myeloma (MM)	
	Amyloidosis	
	Non-MM blood cancers	
Ofatumumab (OMB157) BTD (CLL) Target: CD20 Partner: Novartis	Relapsing multiple sclerosis (RMS) (SubQ)	
Tisotumab vedotin Target: TF Partner: Seattle Genetics	Cervical cancer	
	Solid tumors	
HuMax-AXL-ADC Target: AXL	Solid tumors	
HexaBody-DR5/DR5 Target: DR5	Solid tumors	
DuoBody-CD3xCD20 Targets:CD3, CD20	Hematological malignancies	

Innovative Clinical & Pre-clinical Pipeline
Additional Shots on Goal

Product	Disease Indications	Development Phase
Teprotumumab (RV001) BTD Target: IGF-1R, Partner: Horizon Pharma	Graves' orbitopathy	
HuMax-IL8 Target: IL8, Partner: BMS	Advanced cancers	
Camidanlumab tesirine (ADCT-301) Target: CD25, Partner: ADCT	Lymphoma	
	Acute myeloid leukemia (AML) or acute lymphoblastic leukemia (ALL)	
JNJ-61186372 Targets: EGFR, cMet, Partner: Janssen	Non-small-cell lung cancer (NSCLC)	
JNJ-63709178* Targets: CD3, CD123, Partner: Janssen	Acute Myeloid Leukemia (AML)	
JNJ-64007957 Targets: BCMA, CD3, Partner: Janssen	Relapsed or refractory MM	
JNJ-64407564 Targets: CD3, GPRC5D, Partner: Janssen	Relapsed or refractory MM	
Lu AF82422 Target: alfa-Synuclein, Partner: Lundbeck	Parkinson's disease	
~20 Active Pre-clinical programs incl. DuoBody CD40x4-1BB	Proprietary programs: HuMab, HuMab-ADC, DuoBody, DuoBody-ADC & HexaBody	
	Partnered programs: HuMab, DuoBody & HexaBody	



Aim 4 INDs in 4 Years

*As per clinicaltrials.gov, trial currently on hold due to Grade 3 event.

Cutting Edge Capabilities

Additional Value Created by Technologies

DuoBody

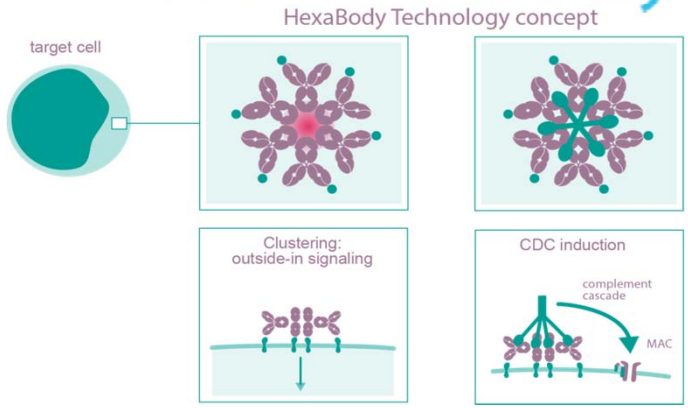
DuoBody Platform

- Efficient & versatile bispecific Ab platform
- Applicable to any antibody from any platform
- Regular IgG format
- Large scale production validated
- No developability liabilities
- Robotized bispecific library generation
- Multiple ongoing collab. incl. with Novo Nordisk, Gilead & Janssen

HexaBody Technology

- Robust effector function enhanced Ab
- Enables antibodies to readily form clusters of 6 (hexamers)
- Induces & enhances target cell killing after binding (CDC and apoptosis)
- Creates innovative products in cancer & infectious diseases
- Multiple ongoing research collaborations

HexaBody



Daratumumab (Marketed as DARZALEX®)
Approved in US, EU & Japan

First-in-class antibody targeting CD38 – 2 FDA BTDs

Marketed as monotherapy in US & EU for double refractory MM

Approved in US, EU & Japan in combo. w/ Revlimid® & dex or Velcade® & dex for relapsed / refractory MM

Approved in the US in combo. w/ Velcade®, melphalan & prednisone for newly diagnosed MM pts ineligible for ASCT & in combo. w/ Pomalyst® & dex for pts w/ MM who have received at least 2 prior therapies

Industry sponsored clinical studies ongoing in MM, NKT-cell lymphoma, MDS, and amyloidosis

Blockbuster status – growing royalty income
- Royalty rate: 12% - 20%

Collaboration w/ Janssen Biotech
- Up to $1bn total in dev., reg. & sales milestones, Janssen responsible for all costs assoc. w/ dev. & commercialization

See local country prescribing information for precise indications

Covering All Stages of MM: Key Ongoing Trials

Disease Stage	Therapy	No. Pts*	Development Phase
			Pre-Clinical I I/II II III
High Risk Smoldering	Subcutaneous	360	AQUILA
	Monotherapy	126	✓ CENTAURUS
Front line (transplant & non-transplant)	Dara + VMP	706	✓ ALCYONE
	Dara + VMP (Asia Pacific)	210	
	Dara + Rd	745	✓ MAIA
	Dara + VTd	1,080	✓ CASSIOPEIA
	Dara + RVd	224	✓ GRIFFIN
Relapsed or Refractory	Dara + Vd (China)	210	
	Dara + Kd	466	✓ CANDOR
	Dara + Pom + d	302	APOLLO
	Subcutaneous vs IV	480	COLUMBA
	Dara + combinations	>400	NINLARO® (Ph II), Venclexta™ (Ph II), Selinexor (Ph I/II)
	Dara + I.O. (PD1 & PDL1)	>700	Keytruda® (Ph II), Opdivo® (Ph I/II), Tecentriq® (Ph I)

V = Velcade® , MP = melphalan-prednisone , T = thalidomide d= dexamethasone, R = Revlilmid®, K = Kyprolis®, Pom = Pomalyst®

✓ Fully recruited *Number of patients are as per linicaltrials.gov, include full trial recruitment, not just dara arms.

Maintenance integrated into some study protocols

Daratumumab Development

Beyond Multiple Myeloma

Amyloidosis

- Ph III D (SC) + cyclo., bortezomib & dex. (CyBorD)

MDS

- Ph II mono.

ALL

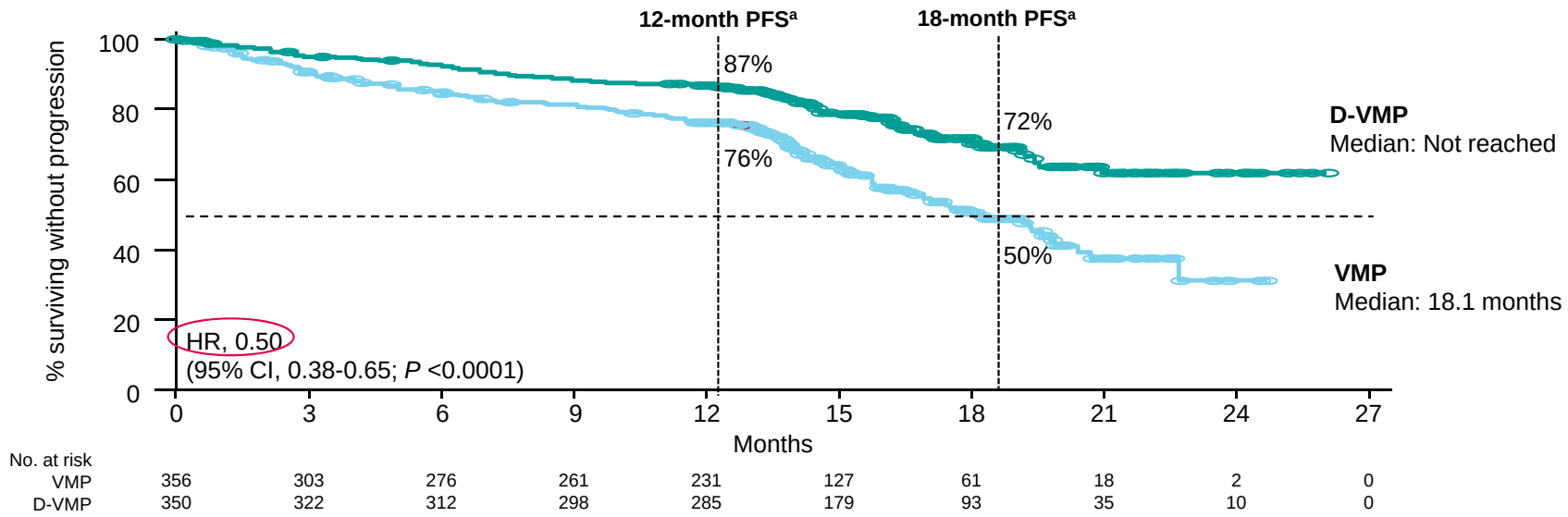
- Ph II D + standard of care chemo.

NKTCL (nasal type)

- Ph II mono.

Front Line Multiple Myeloma: ALCYONE

Ph III Newly Diagnosed Multiple Myeloma



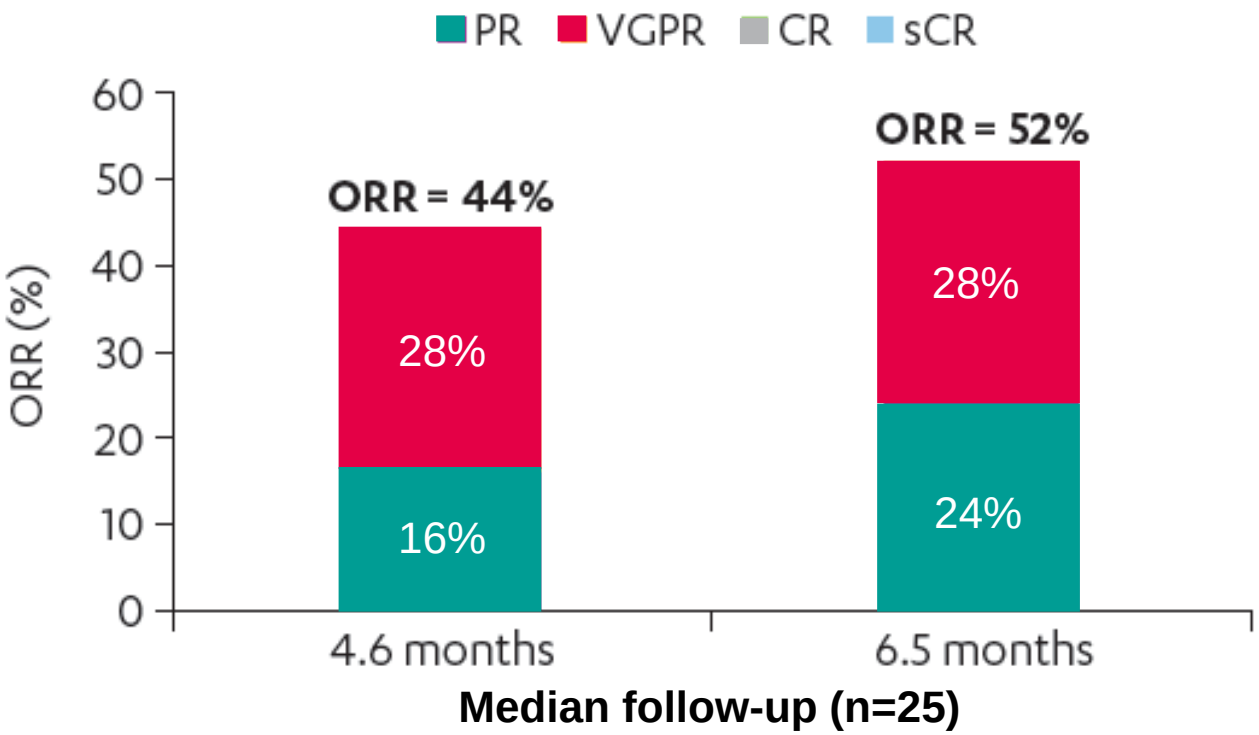
In D-VMP arm:

- 50% reduction risk of disease progression or death in patients receiving D-VMP
- Median PFS not reached
- **>3-fold higher MRD-negative rate**

Data Presented at ASH – Atlanta, December 2017 / Basis of FDA Approval (May 2018) & Positive CHMP Opinion (July 2018)

Subcutaneous Daratumumab

PAVO Study in Relapsed or Refractory MM: ORRs in Part 2 (Dara SC 1,800 mg)



ORR, overall response rate; DARA, daratumumab; SC, subcutaneous; PR, partial response; VGPR; very good partial response; CR, complete response; sCR, stringent complete response

Presented at ASCO – Chicago, June 2018

Faster Infusion time

- Dosing in 3-5 min.
  - Ph III study underway
- First IV infusion: 7 hrs

Well tolerated

- IRRs w/ dara SC: 16%
- IRRs w/ dara IV: 45% - 56%

- High clinical response rates that improved w/ longer follow-up observed
- Median PFS not reached after median follow-up of 6.5 mo

Ofatumumab (Arzerra®)

Human antibody targeting CD20

Two Phase III studies in relapsing MS ongoing

MS Advantages: Dosing
- Better disease management, subcutaneous dosing

MS Advantages: Attributes
- Potential for low immunogenicity, manageable safety profile

Marketed in various territories for certain CLL indications*
In non-US markets, Novartis intends to transition from commercial to compassionate use programs

Collaboration with Novartis
- Cash flow positive for Genmab

*See local country prescribing information for precise indications

Clinical Projects: Tisotumab vedotin

Phase II for Cervical Cancer

Fully human antibody-drug conjugate (ADC)

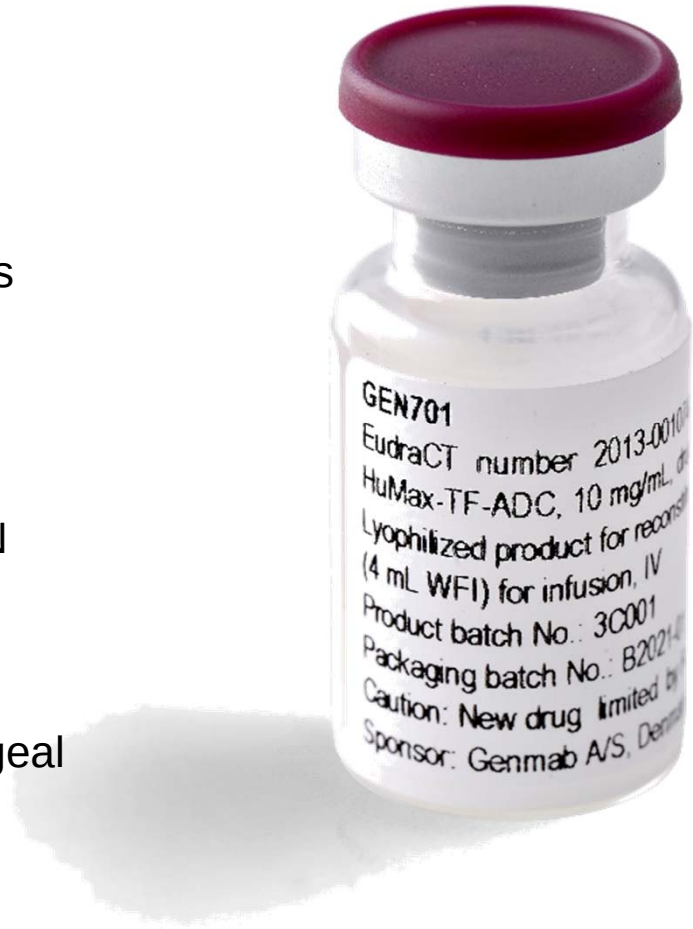
Targets Tissue Factor (TF)
- Therapeutic potential in broad range of solid tumors

Ph II study in cervical cancer
- Potential registrational pathway

Ph II study in colorectal, NSCLC, pancreatic, SCCHN

Studies ongoing in solid tumors
- Indications incl. gynecologic (ovarian, cervical, and endometrial) cancers, prostate, bladder, & esophageal cancers, NSCLC & SCCHN

50:50 Co-development with Seattle Genetics



Clinical Projects: HuMax-AXL-ADC

Efficacy in *in vivo* Tumor Model

Human ADC

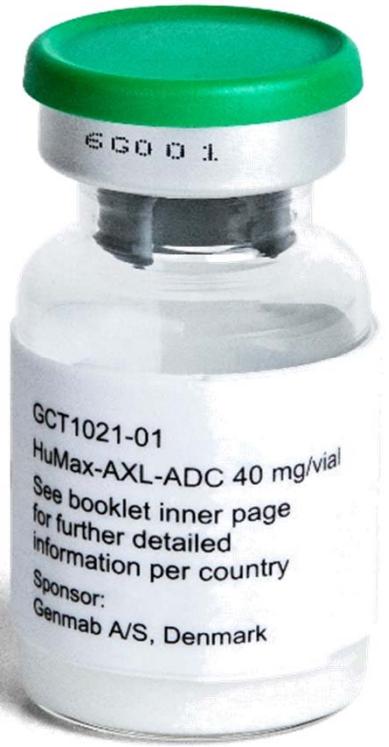
Targets tumor-associated AXL

Therapeutic potential in solid tumors

First-in-human Phase I/II study

- Indications incl. gynecologic (ovarian, cervical, & endometrial) cancers, thyroid cancer, NSCLC, melanoma and sarcoma
- Expansion cohorts initiated in 2018 (NSCLC, melanoma, sarcoma)

ADC technology licensed from Seattle Genetics



Clinical Projects: HexaBody-DR5/DR5

Potential in Solid Tumors

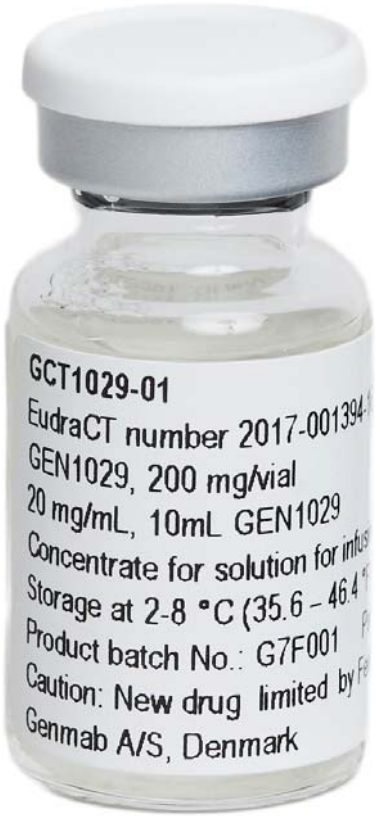
Proprietary HexaBody technology

Targets DR5

Phase I/II study initiated in Q2 2018

Potential in solid cancers

Colorectal, NSCLC, triple neg. breast cancer, renal cell cancer, gastric cancer, pancreatic cancer & urothelial cancer



Clinical Projects: DuoBody-CD3xCD20

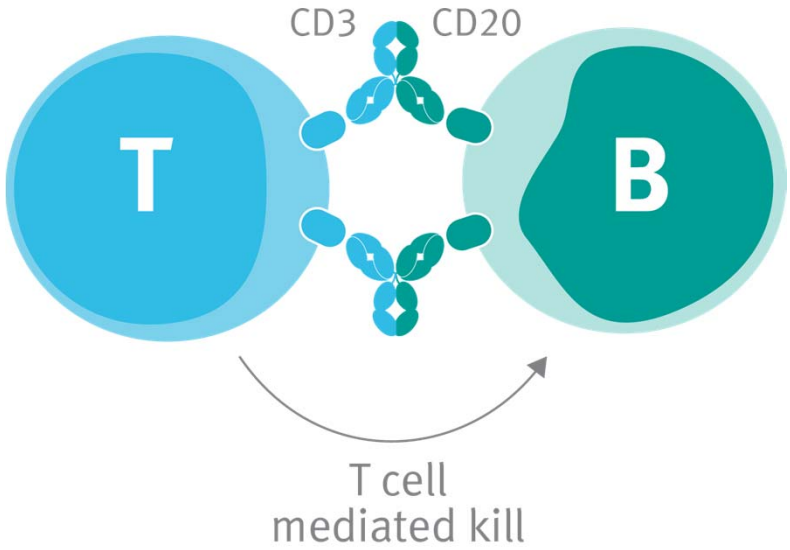
Phase I/II Study Planned

Proprietary DuoBody Technology

Simultaneous binding to CD20 on B cells and CD3 on T cells

Phase I/II study initiated in Q3 2018

Potential in B-cell malignancies



Well-Capitalized Biotech – 2018 Guidance

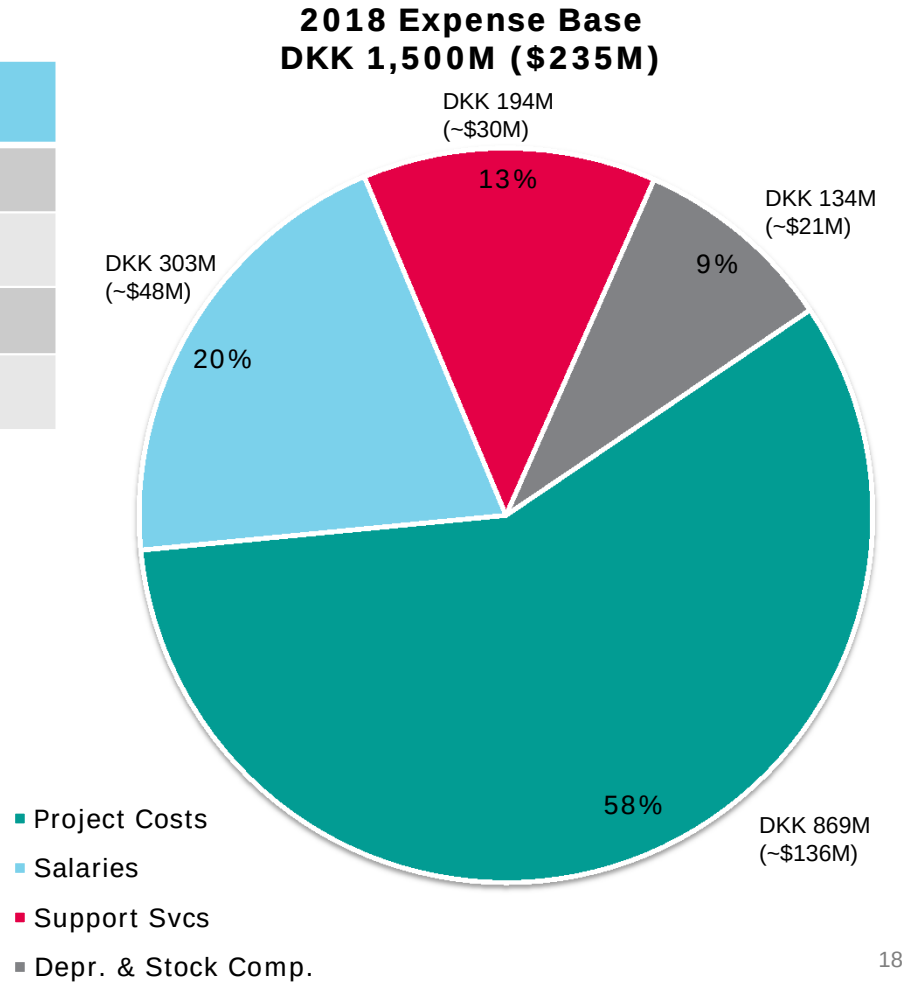
Income Statement	DKKM	~USDM*
Revenue	2,700 – 3,100	422 - 485
Operating expenses	(1,400) – (1,600)	(219) – (250)
Operating income	1,300 – 1,500	203 - 235
*USD 1.00 = DKK 6.3958		

2018 Guidance – August 8, 2018

DARZALEX sales
- 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 1,500
- 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	X	» 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 ofatumumab value	✓	» Complete recruitment Phase III subcutaneous ofatumumab relapsing MS studies
Maximize tisotumab vedotin progress	✓	» Start two Phase II studies in 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

Creating Value for Patients & Shareholders

Building on 3 central pillars: Focus, Innovation & Execution

- 2 marketed products
- 4 proprietary early stage clin. programs
- 2 proprietary technologies
- Robust pre-clinical pipeline
- World-class antibody & R&D expertise
- Strategic collaborations
- Building commercial expertise
- Solid financials
- Proven track record

Innovating Antibodies, Improving Lives

Appendix

Genmab

Publicly Listed Company with Large Free Float

Large cap, listed on Nasdaq Copenhagen, Denmark & ADR in US

Rest of shares held across world incl.
- USA
- UK
- DK
- NL

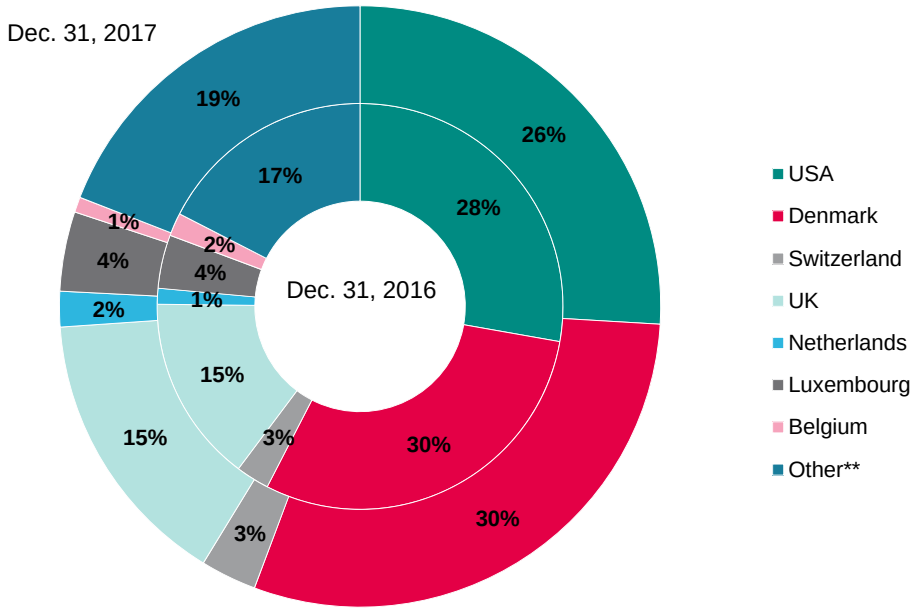
Approx. Market Cap
- DKK 65 bn
- USD 10 bn

Approx. shares outstanding: 61.5M

Warrants outstanding: 1.3M (2%)

Approx. diluted shares: 63M

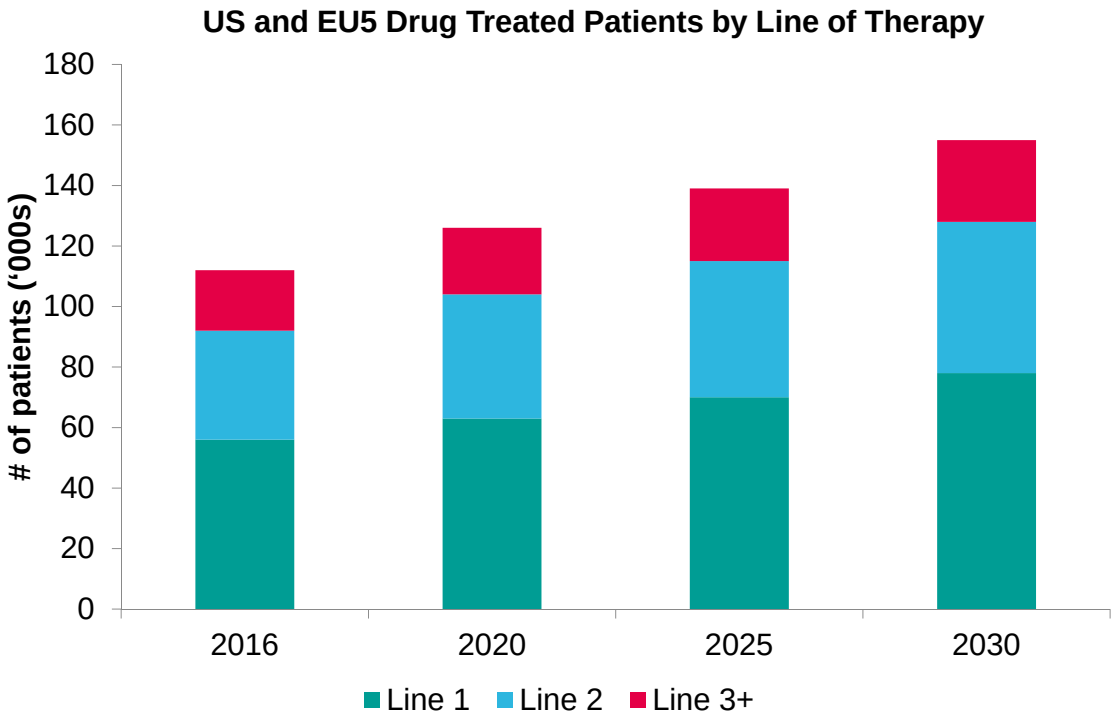
Geographical Shareholder Distribution*
December 31, 2017



* Based on figures from the internal shareholder register per December 31, 2017
** "Other" includes shares held in other countries and shares not held in nominee accounts, including OTC traded shares

Market Opportunity in MM

- Current projections assume a larger frontline patient population and greater rate of growth over time
- As a disease of the elderly, MM prevalence is expected to rise in line with the growing elderly population
- Incidence is expected to increase in Europe in line with the growing elderly population
- Mortality has significantly decreased due to effectiveness of newer treatments
  - Average lifespan of a patient diagnosed with MM is 7-8 years



Source: Kantar Health, 2015 US and EU5

DARZALEX® (daratumumab) Sales Potential

$1,242M

Net sales
Full Year 2017

$2 – 2.3B

Genmab projected 2018 sales

$9B

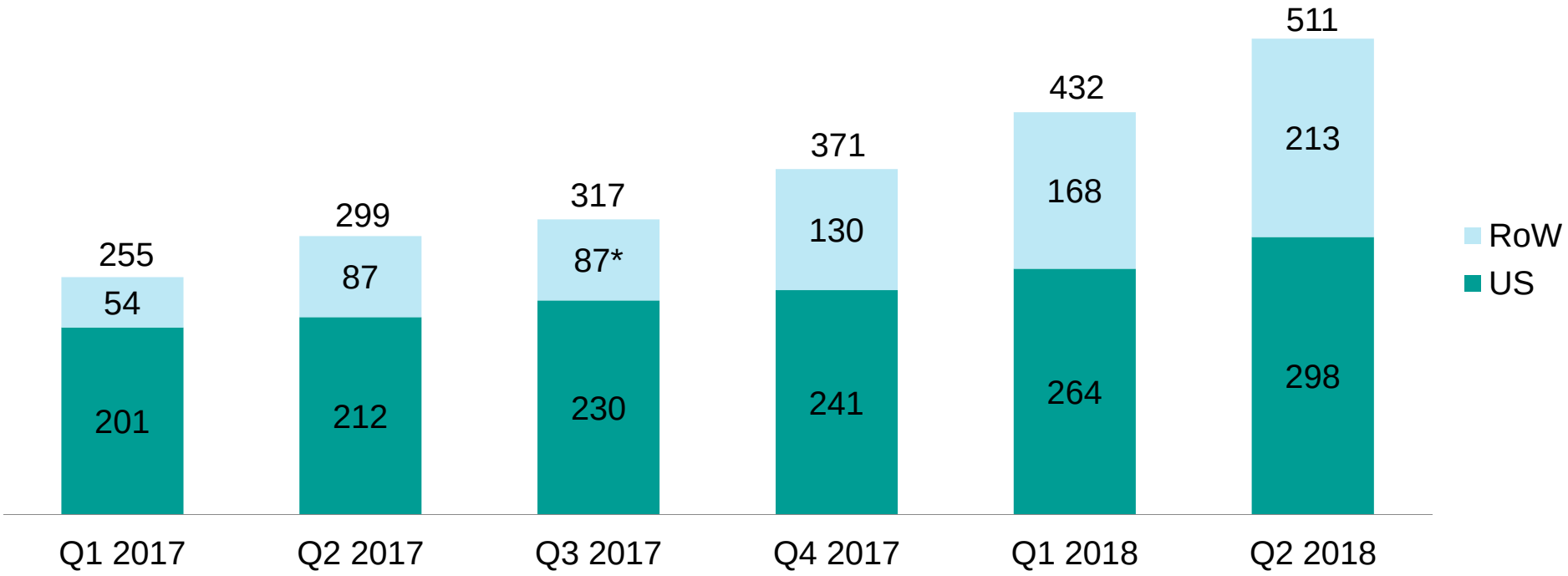
Average analyst*
projected peak MM sales

Potential upside:
smoldering disease, other blood cancers, rheumatoid arthritis

*Average sales projections of analysts covering Genmab as of June 2018

DARZALEX Quarterly Sales

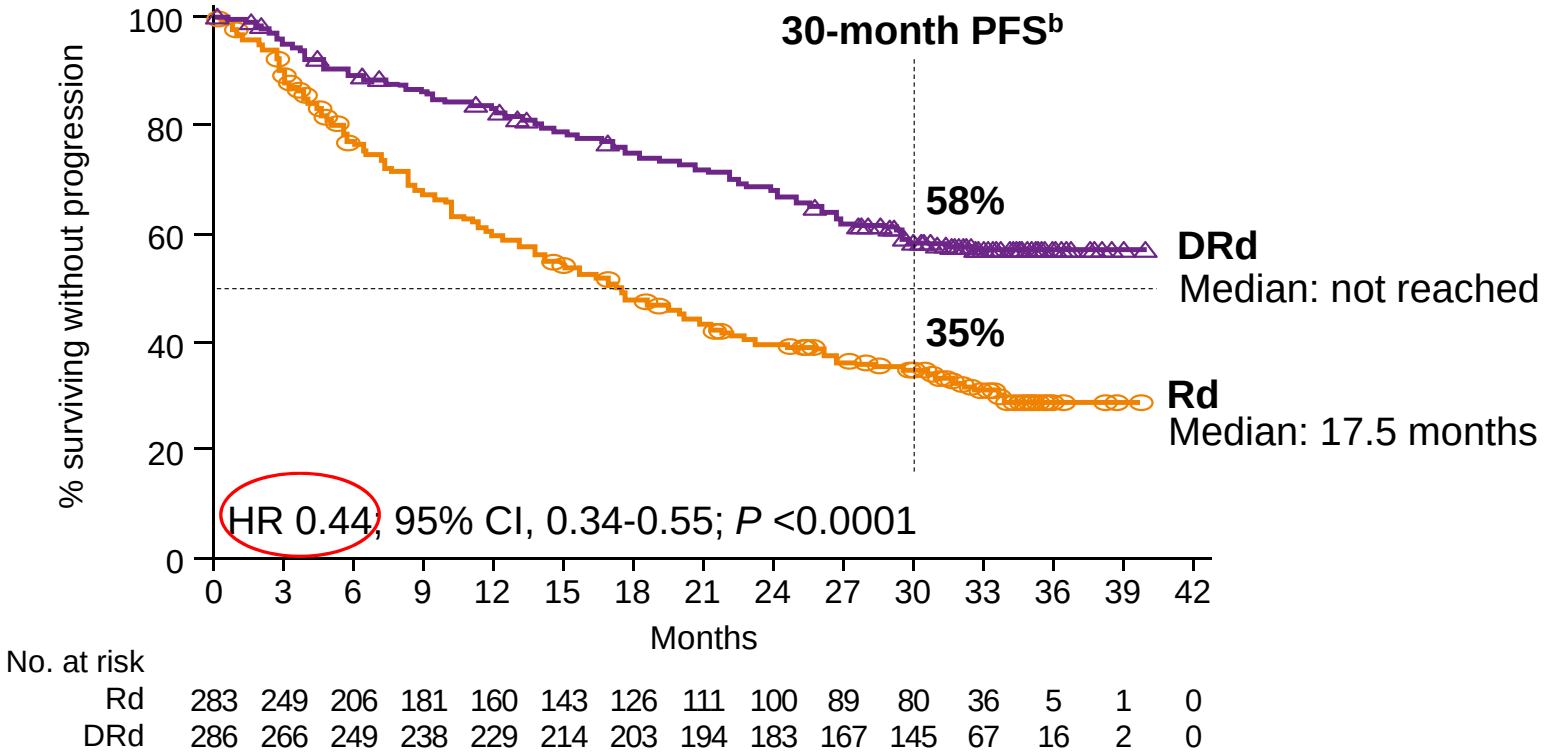
Q1 2017 – Q2 2018, USD M



*RoW sales negatively impacted by one time adjustment of $20M related to retroactive reimbursement matters in Germany and France.

Updated Efficacy: POLLUX

Presented ASH 2017



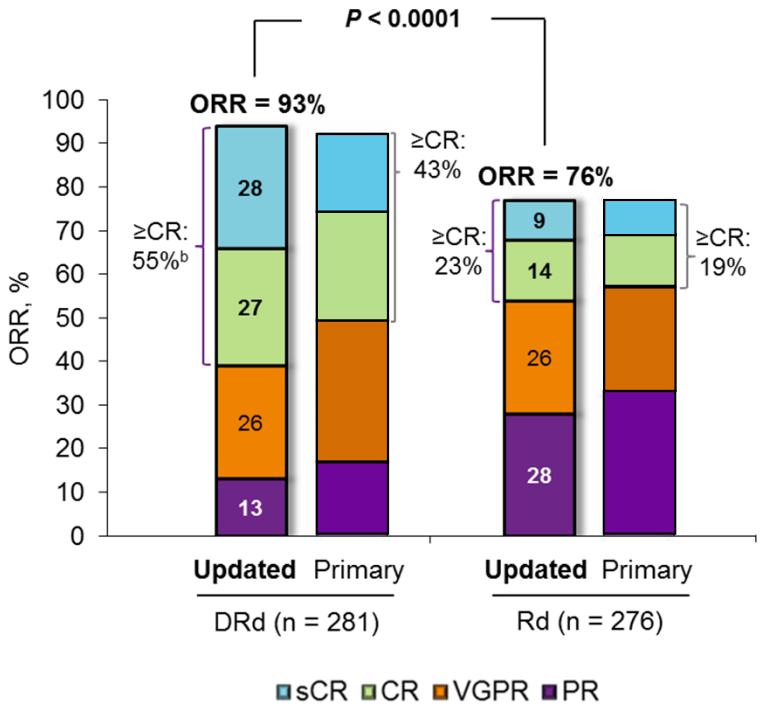
56% reduction in risk of progression/death for DRd versus Rd

HR, hazard ratio; CI, confidence interval.
[a]Exploratory analyses based on clinical cut-off date of October 23, 2017.
[b]Kaplan-Meier estimate.

Updated Efficacy: POLLUX

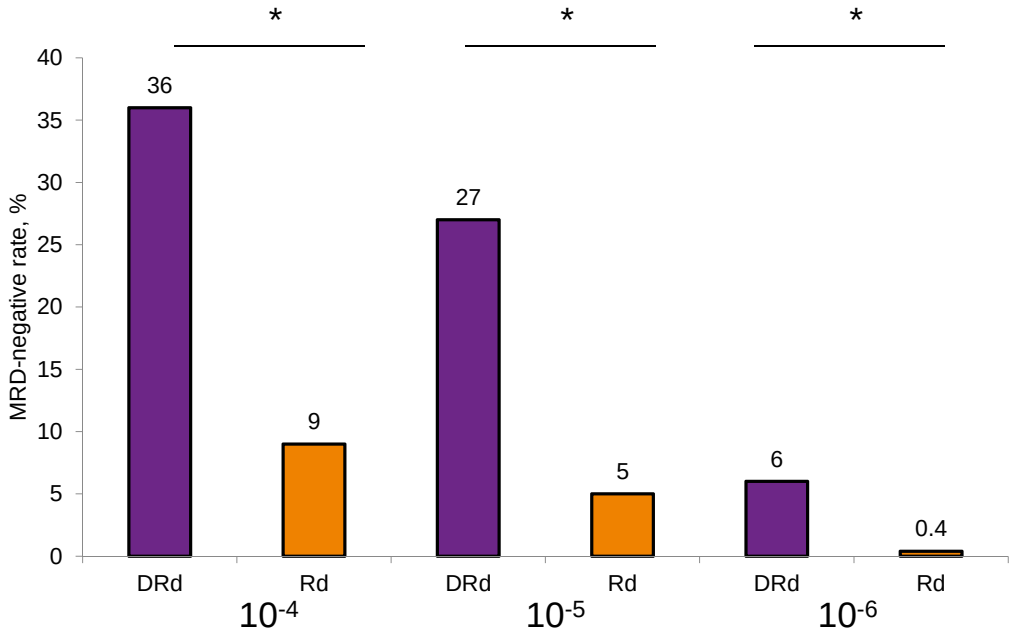
Presented ASH 2017

ORR



MRD-negative Rates

*P <0.0001



MRD assessed using clonoSEQ® assay V2.0

- **Responses continued to deepen in the DRd group**
- **Significantly higher (>3-fold) MRD-negative rates for DRd versus Rd**

sCR, stringent complete response; PR, partial response.

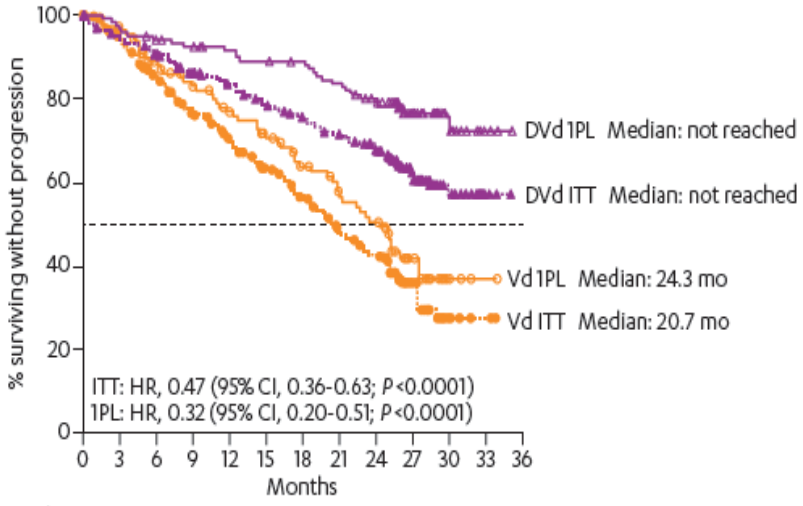
Primary analysis reported in Dimopoulos MA, et al. *N Engl J Med.* 2016;375(14):1319-1331.

[a]Exploratory analyses based on clinical cutoff date of October 23, 2017; [b]*P* <0.0001 for DRd versus Rd.

Updated Efficacy: CASTOR

Presented ASH 2017

PFS

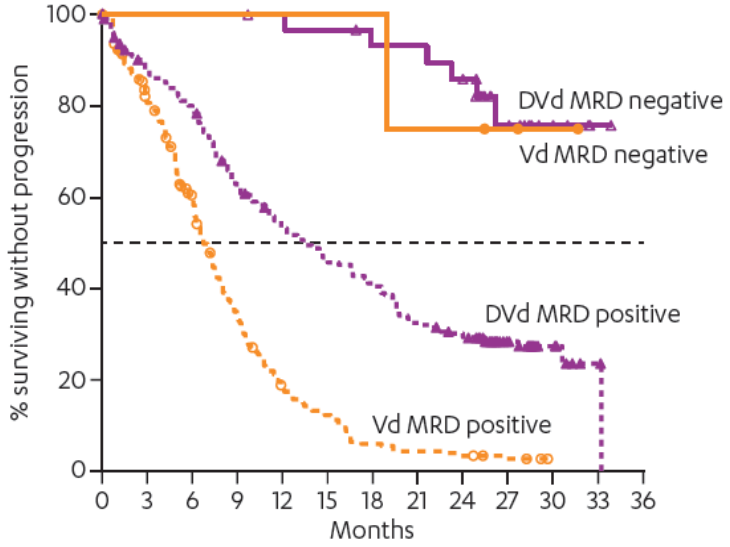


No. at risk

Vd ITT	247	214	188	162	140	119	101	81	70	31	5	2	0
DVd ITT	251	229	218	201	191	178	167	155	144	66	25	4	0
Vd 1PL	113	105	92	82	73	65	55	46	41	18	3	1	0
DVd 1PL	122	115	111	107	104	101	99	92	85	42	17	3	0

PFS2, progression-free survival on subsequent line of therapy; ITT, intent-to-treat; 1PL, 1 prior line of therapy; DVd, daratumumab/bortezomib/dexamethasone; Vd, bortezomib/dexamethasone.

MRD-negative Rates

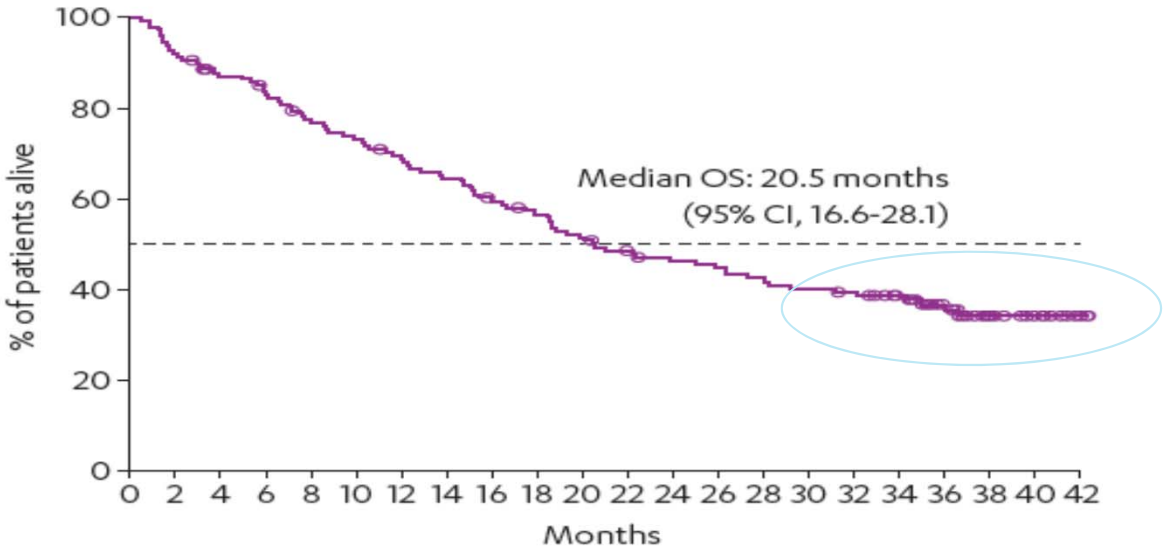


No. at risk

Vd MRD negative	4	4	4	4	4	4	4	3	3	2	1	0	0
DVd MRD negative	30	30	30	30	29	28	26	26	24	12	6	1	0
Vd MRD positive	243	178	125	70	35	23	11	8	6	3	0	0	0
DVd MRD positive	221	185	168	131	109	95	83	66	59	28	13	2	0

Updated Efficacy: Monotherapy

Dara monotherapy in RR MM→ tail effect



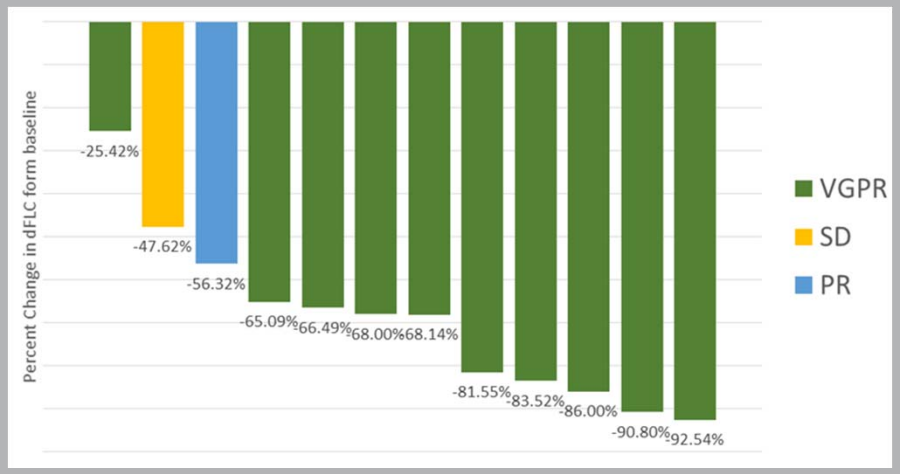
Overall survival (OS): combined analysis of GEN501 Part 2 and SIRIUS data.

Usmani, et al. ASH 2017. Poster #3107

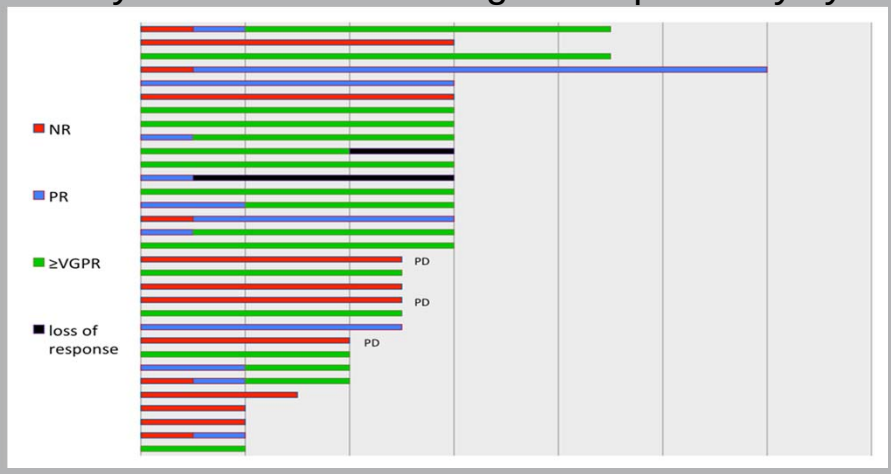
Daratumumab in AL Amyloidosis

Presented at ASH Annual Meeting, Dec. 2017

Ph II Daratumumab in relapsed AL amyloidosis*: % reduction in dFLC after 1 infusion



Ph II Daratumumab in previously treated systemic AL amyloidosis**: Hematological response by cycles



Light chain (AL) amyloidosis

- Occurs when amyloid proteins form deposits that damage tissues and organs
- Most frequently affects kidneys, heart, nervous system, liver & digestive tract
- Currently no cure

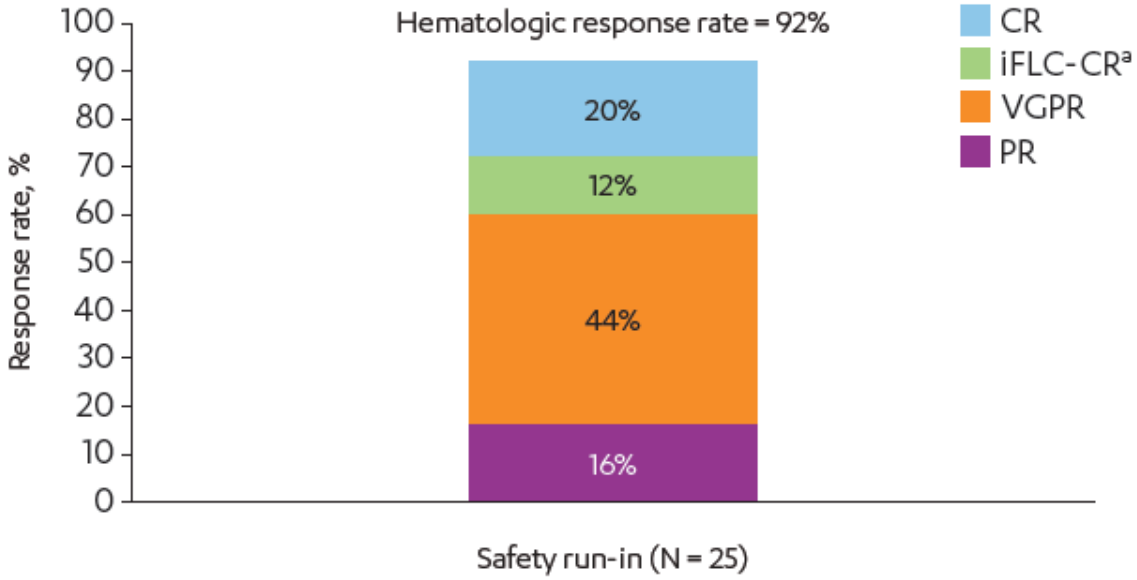
*Safety and Tolerability of Daratumumab in Patients with Relapsed Light Chain (AL) Amyloidosis: Preliminary Results of a Phase II Study, Sanchorawala V. et al
**A Prospective Phase II of Daratumumab in Previously Treated Systemic Light Chain (AL) Amyloidosis, Roussel M. et al

Daratumumab in AL Amyloidosis con't

Subcutaneous daratumumab plus cyclophosphamide, bortezomib and dexamethasone in patients with newly diagnosed amyloid light chain amyloidosis

Summary of overall best hematologic response based on IACC

Preliminary Efficacy: Except for 2 patients, all remaining patients demonstrated hematologic responses based on IACC Guidelines



IACC, International Amyloidosis Consensus Criteria; CR, complete response; LLN, lower limit of normal; iFLC, involved free light chain; VGPR, very good partial response; PR, partial response.
[a]Patients with negative serum and urine immunofixation and normalization of involved FLC level; if uninvolved FLC level is below LLN and FLC ratio is abnormal or normal, patient will be assigned to iFLC-CR (involved FLC CR) response category.

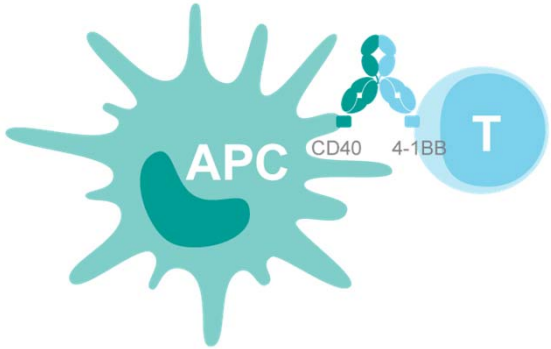
Presented at ASCO Annual Meeting, June 2018: Safety Run-in Results of ANDROMEDA

DuoBody-CD40x4-1BB

Immunomodulation: targeting two checkpoint activators

Bispecific antibody targeting CD40 and 4-1BB (CD137)

- Trans-activating bispecific targeting two checkpoint activators
- Simultaneously activates antigen-presenting cell (APC) and enhances T cell activation
  - Co-engagement of CD40 (APCs) and 4-1BB (T cells) in immune response against tumor
  - Conditional activation and expansion of previously activated cytotoxic $CD8^+$ T cells
  - Inert Fc backbone
- For treatment of solid cancers
- 2018 IND/CTA candidate
- 50/50 Co-development Genmab and BioNTech



Ongoing Daratumumab Clinical Trials

Janssen Sponsored Phase II & III

Daratumumab Trials Sponsored by Pharma / Biotech

Ct.gov Identifier	Phase	Sponsor	Indication	Therapy
NCT02252172	III	Janssen	Untreated MM	Daratumumab + Rd (MAIA)
NCT02195479	III	Janssen	Untreated MM	Daratumumab + VMP (ALCYONE)
NCT02541383	III	Janssen	Untreated MM	Daratumumab + VTd (CASSIOPEIA)
NCT02076009	III	Janssen	Relapsed or Refractory MM	Daratumumab + Rd (POLLUX)
NCT02136134	III	Janssen	Relapsed or Refractory MM	Daratumumab + Vd (CASTOR)
NCT03180736	III	Janssen	Relapsed or Refractory MM	Daratumumab + Pom-d (APOLLO)
NCT03201965	III	Janssen	Amyloidosis	Daratumumab + CyBorD (ANDROMEDA)
NCT03217812	III	Janssen	Untreated MM	Daratumumab + VMP (Asia Pacific)
NCT03234972	III	Janssen	Relapsed or Refractory MM	Daratumumab + Vd vs Vd (China)
NCT03277105	III	Janssen	Relapsed or Refractory MM	Daratumumab SC vs IV (COLUMBA)
NCT03301220	III	Janssen	Smoldering MM	Daratumumab SC (AQUILA)
NCT03384654	II	Janssen	Relapsed / Refractory ALL / LL	Dara + Vincristine + Prednisone + Doxorubicin (ALL2005)
NCT02951819	II	Janssen	Untreated and Relapsed MM	Daratumumab + CyBorD (LYRA)
NCT02874742	II	Janssen	Untreated MM	Daratumumab + RVd (GRIFFIN)
NCT02316106	II	Janssen	Smoldering MM	Monotherapy (CENTAURUS)
NCT02927925	II	Janssen	NKTCL, Nasal Type	Monotherapy (NKT2001)
NCT03011034	II	Janssen	Myelodysplastic Syndromes	Daratumumab or Talacotuzumab (MDS2002)
NCT03412565	II	Janssen	Newly diagnosed & relapsed / refractory MM	Daratumumab SC + Rd, VMP & VRd (MMY2040)

Ongoing Daratumumab Clinical Trials

Janssen Sponsored Phase I & I/II

Daratumumab Trials Sponsored by Pharma / Biotech

Ct.gov Identifier	Phase	Sponsor	Indication	Therapy
NCT01615029	I/II	Janssen	Relapsed and Refractory MM	Daratumumab + Rd (GEN503)
NCT02852837	I	Janssen	Relapsed or Refractory MM	Monotherapy (in China) (MMY1003)
NCT02519452	I	Janssen	Relapsed or Refractory MM	Monotherapy, subcutaneous (PAVO)
NCT02918331	I	Janssen	Untreated MM	Daratumumab + Rd (Japan) (MMY1006)
NCT03242889	I	Janssen	Relapsed or Refractory MM	Daratumumab subq (Japan) (MMY1008)
NCT01998971	I	Janssen	Various MM	Daratumumab + backbone regimens (Vd, VMP, VTd, Pom-d, Kd, KRd) (EQUULEUS)
NCT03320707	I	Janssen	Healthy volunteers	Daratumumab vs placebo (EDI1001)

Ongoing Daratumumab Clinical Trials

Other Industry Sponsored Trials

Daratumumab Trials Sponsored by Pharma / Biotech

Ct.gov Identifier	Phase	Sponsor	Indication	Therapy
NCT03158688	III	Amgen	Relapsed or Refractory MM	Daratumumab + Kd (CANDOR)
NCT01946477	II	Celgene	Relapsed or Refractory MM	Daratumumab + Pom-d
NCT02807454	II	Celgene	Relapsed and Refractory MM	Daratumumab + Imfinzi (FUSION)
NCT03221634	II	Merck	Relapsed or Refractory MM	Daratumumab + Keytruda
NCT03314181	II	AbbVie	Relapsed or Refractory MM	Daratumumab + Venetoclax + dex w/wout bort
NCT02807558	II	Syros	AML & MDS	Daratumumab + SY-1425
NCT03439293	II	Takeda	Relapsed or Refractory MM	Daratumumab + NINLARO (ixazomib) + Dex
NCT02343042	I/II	Karyopharm	Relapsed or Refractory MM	Daratumumab + Selinexor + Dex
NCT03481556	I/II	Oncopeptides AB	Relapsed or Refractory MM	Daratumumab + Melflufen + Dex
NCT01592370	I/II	BMS	Relapsed or Refractory MM	Daratumumab + nivolumab
NCT02431208	I	Roche	Resistant or Refractory MM	Daratumumab + Tecentriq (atezolizumab)
NCT03068351	I	Roche	Resistant or Refractory MM	Daratumumab + RO6870810

Ongoing Daratumumab Clinical Trials

Investigator Sponsored Study (ISS): MM

Investigator Sponsored Studies (ISS) of Daratumumab

Ct.gov Identifier	Phase	Sponsor	Indication	Therapy
NCT02944565	II	ISS	MM	Daratumumab accelerated infusion
NCT02977494	II	ISS	R/R MM & Severe Renal Impairment	Daratumumab + Vd
NCT02626481	II	ISS	Resistant or Refractory MM	Daratumumab + dexamethasone
NCT03004287	II	ISS	Newly diagnosed MM	KTD-Dara-PACE / Dara-KD / Dara-RD
NCT03012880	II	ISS	Newly diagnosed MM	Daratumumab+ Ixazomib, Len & Dex
NCT03143036	II	ISS	RRMM	Daratumumab + thalidomide + Dex
NCT03184194	II	ISS	RRMM	Daratumumab + nivolumab w/ or w/out Len & Dex
NCT03188172	II	ISS	Newly diagnosed MM	Daratumumab + VRd
NCT03215524	II	ISS	RRMM	Daratumumab + Dex, Cy, Pom
NCT03224507	II	ISS	Deep remission in MM	Daratumumab + KRd
NCT03290950	II	ISS	Newly Diagnosed MM	Daratumumab + KRd
NCT03289299	II	ISS	Smoldering MM	Daratumumab + carfilzomib, lenalidomide & dexamethasone
NCT03346135	II	ISS	MM	Dara as maintenance after ASCT

Ongoing Daratumumab Clinical Trials

Investigator Sponsored Study (ISS): MM, con't

Investigator Sponsored Studies (ISS) of Daratumumab

Ct.gov Identifier	Phase	Sponsor	Indication	Therapy
NCT03450057	II	ISS	RRMM w/ renal impairment	Daratumumab
NCT03475628	II	ISS	Effects on bone disease in RRMM	Daratumumab
NCT03477539	II	ISS	MM	Daratumumab, ASCT, lenalidomide
NCT03490344	II	ISS	MM	Daratumumab, lenalidomide short course
NCT03500445	II	ISS	Newly diagnosed MM	Daratumumab, carfilzomib, lenalidomide, low dose Dex
NCT03556332	II	ISS	RRMM	Daratumumab, carfilzomib, lenalidomide, dex
NCT03589222	II	ISS	RRMM	Daratumumab, selinexor, bortezomib, dexamethasone
NCT03590652	II	ISS	RRMM	Daratumumab, ixazomib, pomalidomide, dexamethasone
NCT03606577	II	ISS	Newly diagnosed MM	Daratumumab, carfilzomib, lenalidomide, dex
NCT03622775	II	ISS	Relapsed MM	Daratumumab
NCT03236428	I	ISS	Smoldering MM	Daratumumab
NCT02955810	I	ISS	Untreated MM	Daratumumab + CyBorD
NCT03311828	I	ISS	Relapsed MM	Daratumumab + positron emission tomography
NCT02751255	I/II	ISS	RRMM	Daratumumab + All-trans retinoic acid
NCT01665794	I/II	ISS	RRMM	Daratumumab + K, Pom, dex

Ongoing Daratumumab Clinical Trials

ISS: Other Indications

Investigator Sponsored Studies (ISS) of Daratumumab

Ct.gov Identifier	Phase	Sponsor	Indication	Therapy
NCT02816476	II	ISS	Amyloidosis	Monotherapy
NCT03067571	II	ISS	AML or MDS	Monotherapy
NCT03095118	II	ISS	Membranoproliferative Glomerulonephritis	Monotherapy
NCT03187262	II	ISS	Waldenstrom macroglobulinemia	Monotherapy
NCT03473730	II	ISS	Metastatic Renal Cell Carcinoma (MRCC) or Muscle Invasive Bladder	Monotherapy
NCT02841033	I/II	ISS	Amyloidosis	Monotherapy
NCT03537599	I/II	ISS	AML	Daratumumab + donor lymphocyte infusion
NCT03177460	I	ISS	High-risk localized prostate cancer	Monotherapy with prostatectomy
NCT03432741	I	ISS	RR NHL, Hodgkin lymphoma or Stage IV breast cancer	Intralesional injection
NCT03283917	I	ISS	Amyloidosis	Daratumumab, ixazomib & dexamethasone
NCT03447808	I	ISS	CLL	Daratumumab & ibrutinib
NCT03591744	I	ISS	Plasma cell leukemia	Daratumumab + bortezomib, dexamethasone, lenalidomide, pegylated liposomal doxorubicin hydrochloride

Dex = dexamethasone
Pom = Pomalyst (pomalidomide)
Rd = Revlimid (lenalidomide) + dexamethasone

Pom-d = Pomalyst (pomalidomide) + dexamethasone
CyBorD = Cyclophosphamide, bortezomib, dexamethasone
KRd = Kyprolis (carfilzomib) + Revlimid (lenalidomide) + dexamethasone

VTd = Velcade (bortezomib) + thalidomidde + dexamethasone
VMP = Velcade (bortezomib) + melphalan-prednisone
Kd = Kyprolis (carfilzomib) + dexamethasone

Vd = Velcade (bortezomib) + dexamethasone

As per clinicaltrials.gov, July 2018

Income Statement: Six Months Ended June 30

	2018	2017		2018	2017
	DKK millions		Change	USD millions *	
Darzalex Royalties	695	454	241	109	71
Darzalex Milestones	-	489	(489)	-	76
Other Revenue	496	81	415	78	13
Total Revenue	1,191	1,024	167	187	160
R&D Costs	(632)	(372)	(260)	(99)	(58)
G&A Expenses	(100)	(70)	(30)	(16)	(11)
Operating Expenses	(732)	(442)	(290)	(115)	(69)
Operating Result	459	582	(123)	72	91
Net Financial Items	132	(171)	303	21	(27)
Tax	(132)	(88)	(44)	(21)	(14)
Net Result	459	323	136	72	50

* USD 1.00 = DKK 6.3958 (Danish Central Bank spot rate on June 30, 2018)

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