

Innovating Antibodies, Improving Lives

Investor Presentation
December 2020



Forward Looking Statement

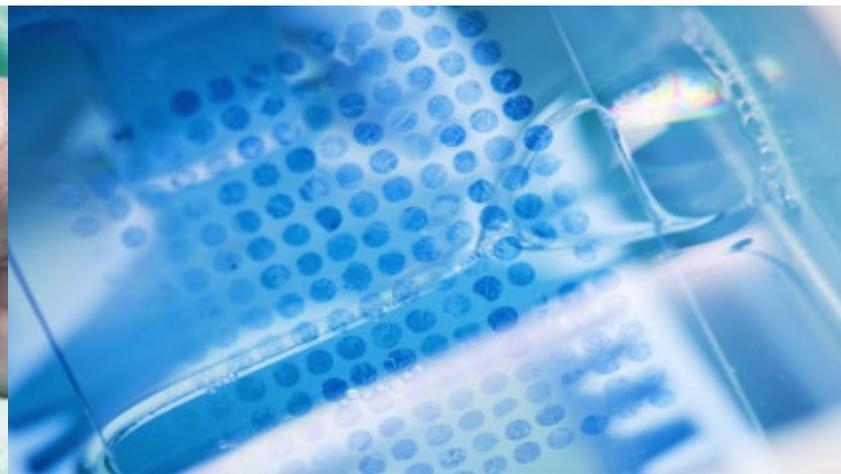
This presentation contains forward looking statements. The words “believe”, “expect”, “anticipate”, “intend” and “plan” and similar expressions identify forward looking statements. All statements other than statements of historical facts included in this presentation, including, without limitation, those 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.

Our Core Purpose, Strategy & Vision Guide Our Work



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

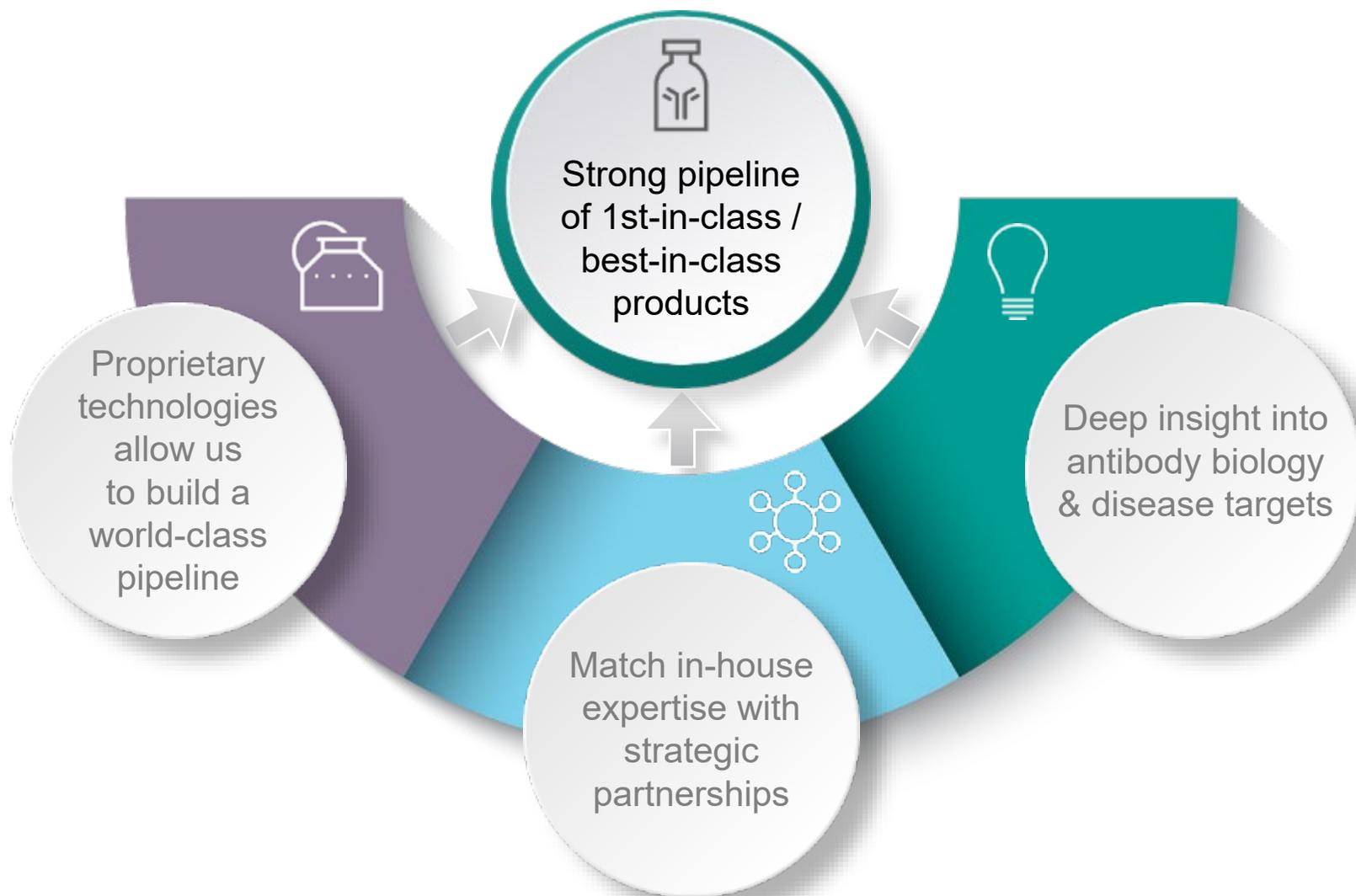


Vision

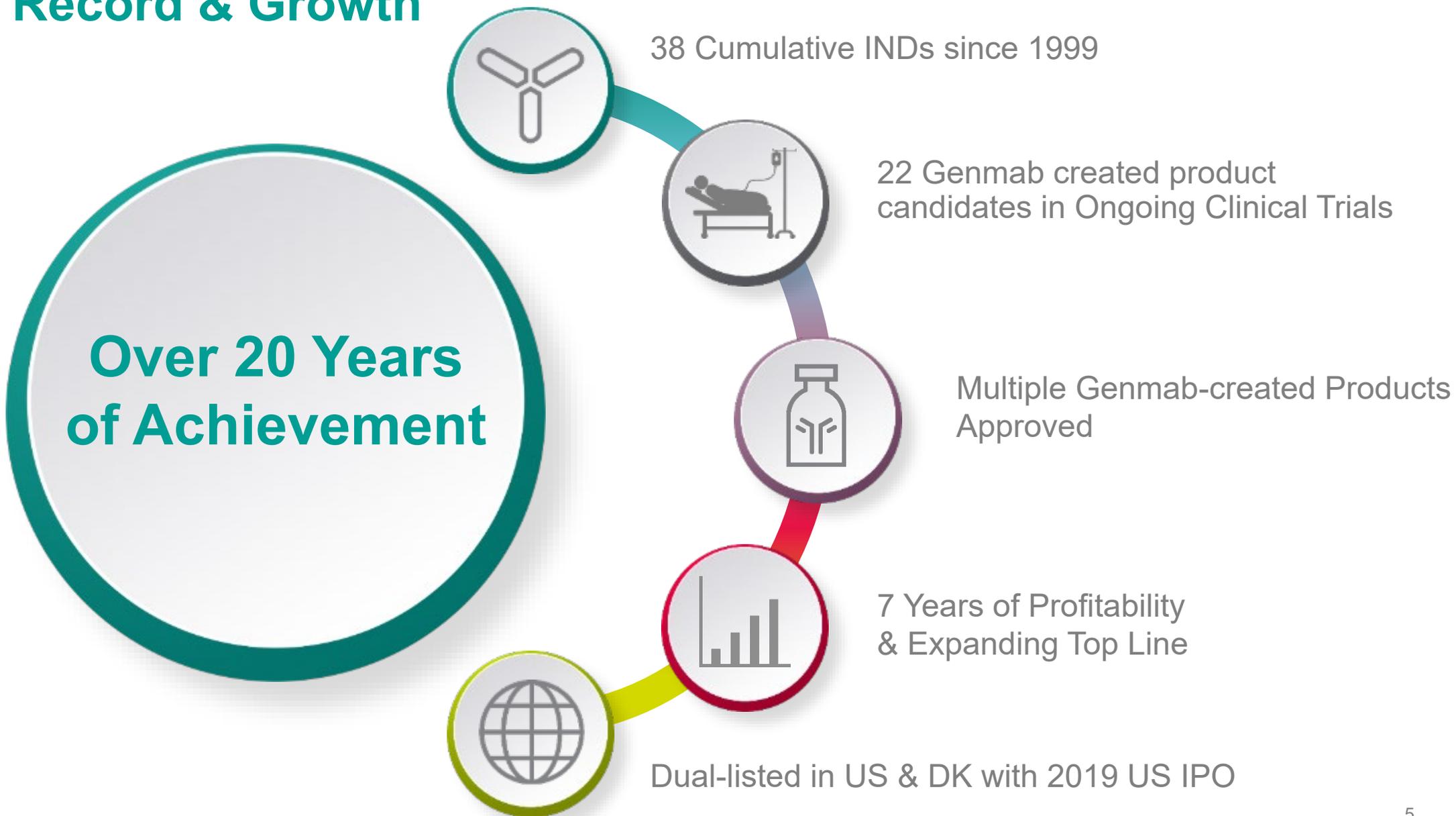
By 2025, our own product has transformed cancer treatment and we have a pipeline of knock-your-socks off antibodies

The Genmab Difference

Innovation Powerhouse Transforming Cancer Treatment & Creating Value



Track Record & Growth



Solid Foundation Built on a Differentiated Pipeline

Potential 1st-in-Class/Best-in-Class

Our Own Clinical Pipeline

- Tisotumab Vedotin⁵
- Epcoritamab (DuoBody[®]-CD3xCD20)⁶
- DuoBody-PD-L1x4-1BB⁷
- DuoBody-CD40x4-1BB⁷
- HexaBody[®]-DR5/DR5
- DuoHexaBody[®]-CD37⁶
- DuoBody-CD3x5T4⁶

R&D Engine

Technologies & Pre-Clinical

- DuoBody technology
- HexaBody technology
- HexElect[®] technology
- DuoHexaBody[®] technology
- Rich Pre-Clinical Pipeline incl. HexaBody-CD38⁸

Solid Financial Base

Approved Partnered Products

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

Programs Built on Genmab's Innovation

Partner-owned Programs in the Clinic

- 15 product candidates in clinical development w/ partners
- Incl. 7 DuoBody products with Janssen, 1 with Novo Nordisk

DARZALEX[®] (daratumumab) & DARZALEX FASPRO[™] (daratumumab and hyaluronidase-fihj): Redefining Treatment of Multiple Myeloma



First-in-class CD38 antibody in development to treat cancer



Collaboration with Janssen: Genmab entitled to tiered royalty of 12-20% of net sales



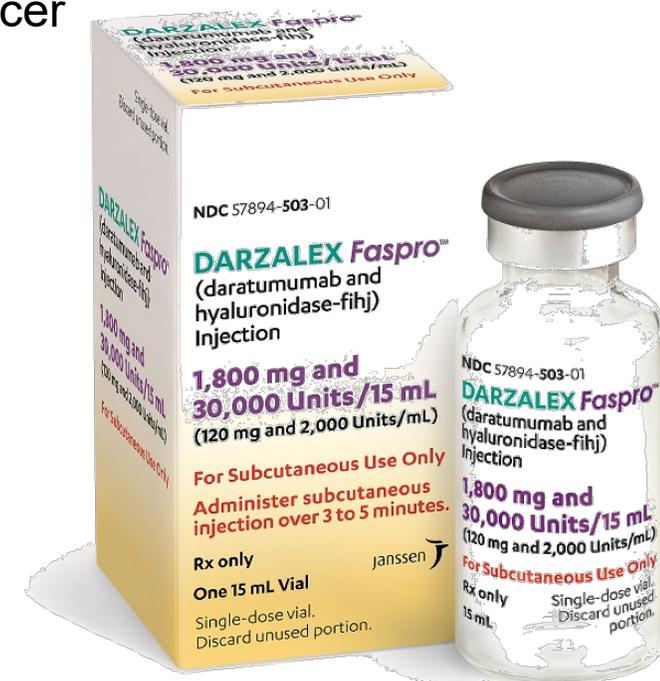
Approved in certain territories for various multiple myeloma (MM) indications¹



DARZALEX FASPRO first and only SubQ CD38 mAb approved in U.S. for treatment of MM



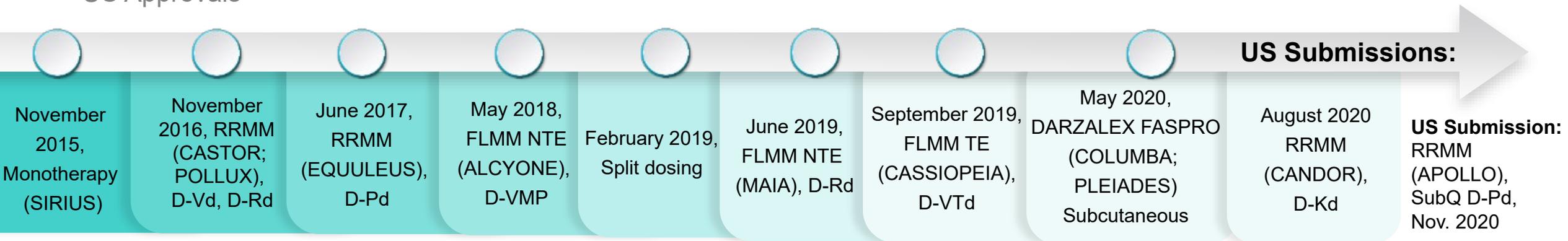
2019 WW net sales by J&J: \$2,998M



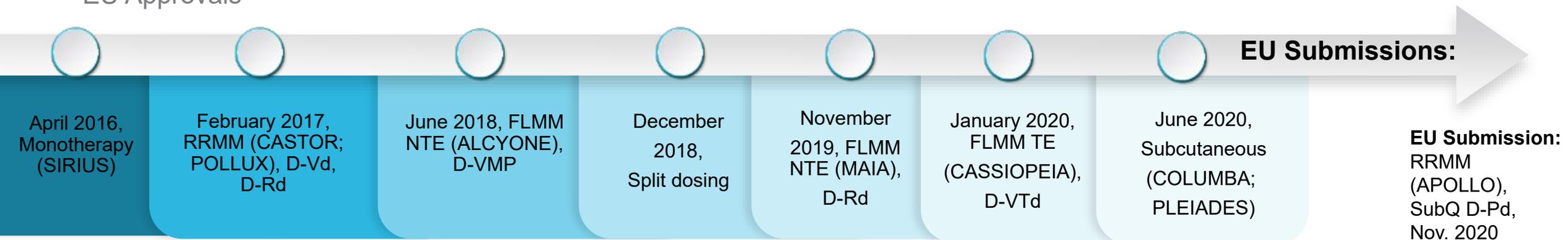
DARZALEX Approvals: US and EU

On Track for Approval Across All Lines of MM Treatment

US Approvals



EU Approvals



Daratumumab

Proving to be the Critical Driver Across Different Combinations & Treatment Lines

Frontline

Relapsed/Refractory

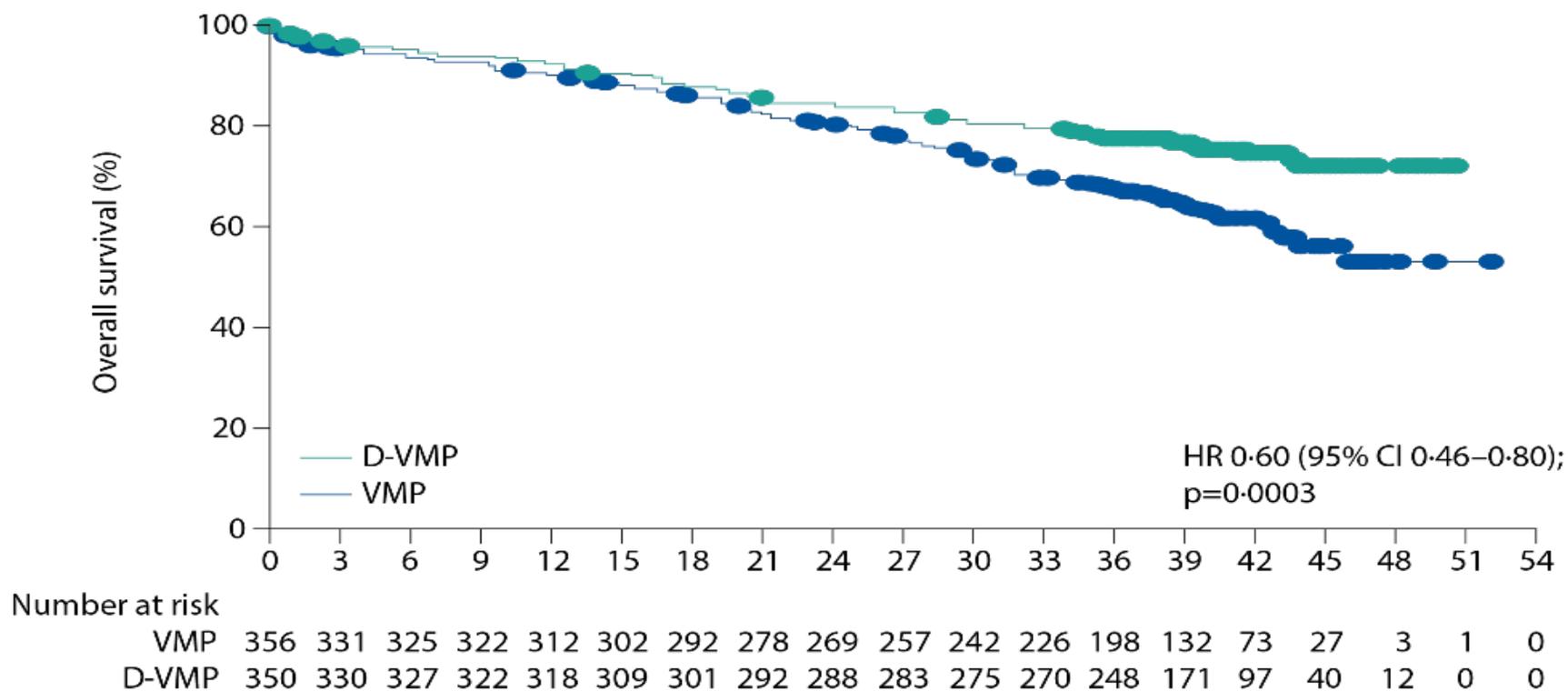
	Transplant Eligible		Transplant Ineligible				
	Ph 3 CASSIOPEIA ^{1,3} (D-VTd vs. VTd)	Ph 2 GRIFFIN ^{1,4} (D-VRd vs VRd)	Ph 3 ALCYONE ^{2,4} (D-VMP vs. VMP)	Ph 3 MAIA ^{2,4} (D-Rd vs. Rd)	Ph 3 POLLUX ^{2,4} (D-Rd vs. Rd)	Ph 3 CASTOR ^{2,4} (D-Vd vs Vd)	Ph 3 CANDOR ^{2,4} (D-Kd vs Kd)
sCR Odds Ratio ¹ or CR ⁺²	1.60	1.57	~2x	~2x	>2x	3x	~3x
MRD-neg rate	1.5x	2.5x	4x	>3x	~5x	>7x	~10x
PFS risk reduction	53% (HR, 0.47)	NA	58% (HR, 0.42)	44% (HR, 0.56)	56% (HR, 0.44)	69% (HR, 0.31)	37% (HR, 0.63)

Ongoing Phase 3: CEPHEUS (D-VRd, NDMM NTE), PERSEUS (D-VRd, NDMM TE)

³Data as per ASCO 2019; ⁴Data as per ASH 2019

Improved Survival for Patients with Multiple Myeloma

Overall Survival Analysis from ALCYONE Trial



Kesimpta® (ofatumumab)

Approved in Relapsing Multiple Sclerosis



Human CD20 Antibody – well validated target



Injection for SubQ use approved for RMS in the US



First B-cell therapy that can be self-administered by patients at home using Sensoready® autoinjector pen



Developed by Novartis: Regulatory submission also made in EU



Genmab 10% royalty payment of net sales

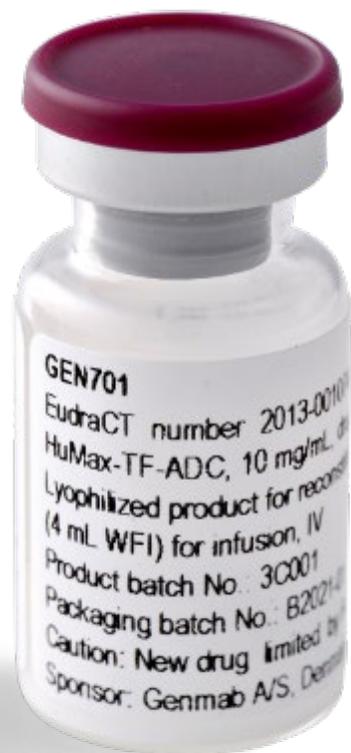


Second Genmab-created product with blockbuster potential



Tisotumab Vedotin

Genmab's Most Advanced Asset with Potential in Solid Tumors



Fully human antibody-drug conjugate (ADC) targeting Tissue Factor (TF) in development to treat solid tumors



License and collaboration agreement with Seattle Genetics 50:50



Very favorable topline results, Phase 2 recurrent or metastatic cervical cancer



Ongoing trials in cervical, ovarian cancer, other solid tumors



Expanding development, additional studies planned

Tisotumab Vedotin in Cervical Cancer

Designed to Address a High Unmet Medical Need

Recurrent or metastatic cervical cancer

- Poor prognosis advanced / recurrent cervical cancer
 - RR standard therapies generally <15%
 - Median OS 6-8 months
- Data ORR & survival after progression on 1L bevacizumab + doublet chemotherapy are limited

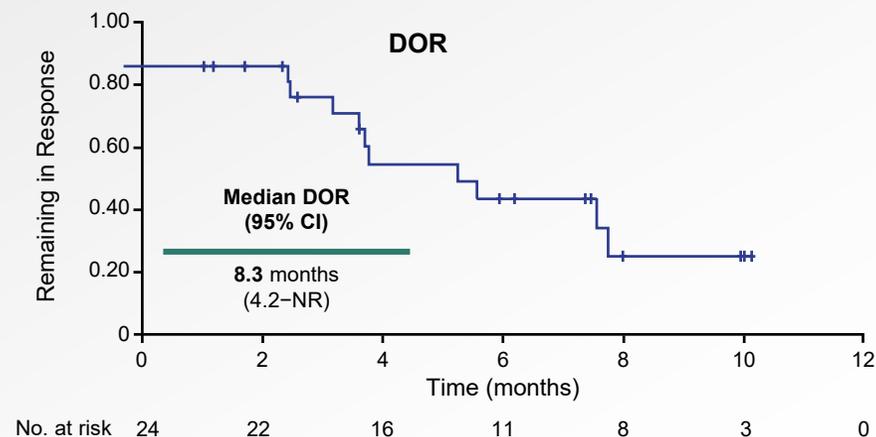
Conclusions*

(previously treated recurrent or metastatic cervical cancer)

- Compelling and durable antitumor activity with manageable and tolerable safety profile
- ORR 24%; CR: 7%
- Median DOR 8.3 mo
- Median PFS (4.2 mo) and OS (12.1 mo) encouraging

Clinically meaningful and durable responses observed*

	N=101
Confirmed ORR (95% CI),^a %	24 (15.9–33.3)
CR, n (%)	7 (7)
PR, n (%)	17 (17)
SD, n (%)	49 (49)
PD, n (%)	24 (24)
Not evaluable, n (%)	4 (4)



Epcoritamab (DuoBody-CD3xCD20)

Potential for Improved Efficacy & Safety in B Cell Malignancies



Potential best-in-class therapeutic



T cell-mediated killing of CD20-expressing cells



SubQ Ph 3 trial in DLBCL announced



50:50 co-development Genmab and AbbVie

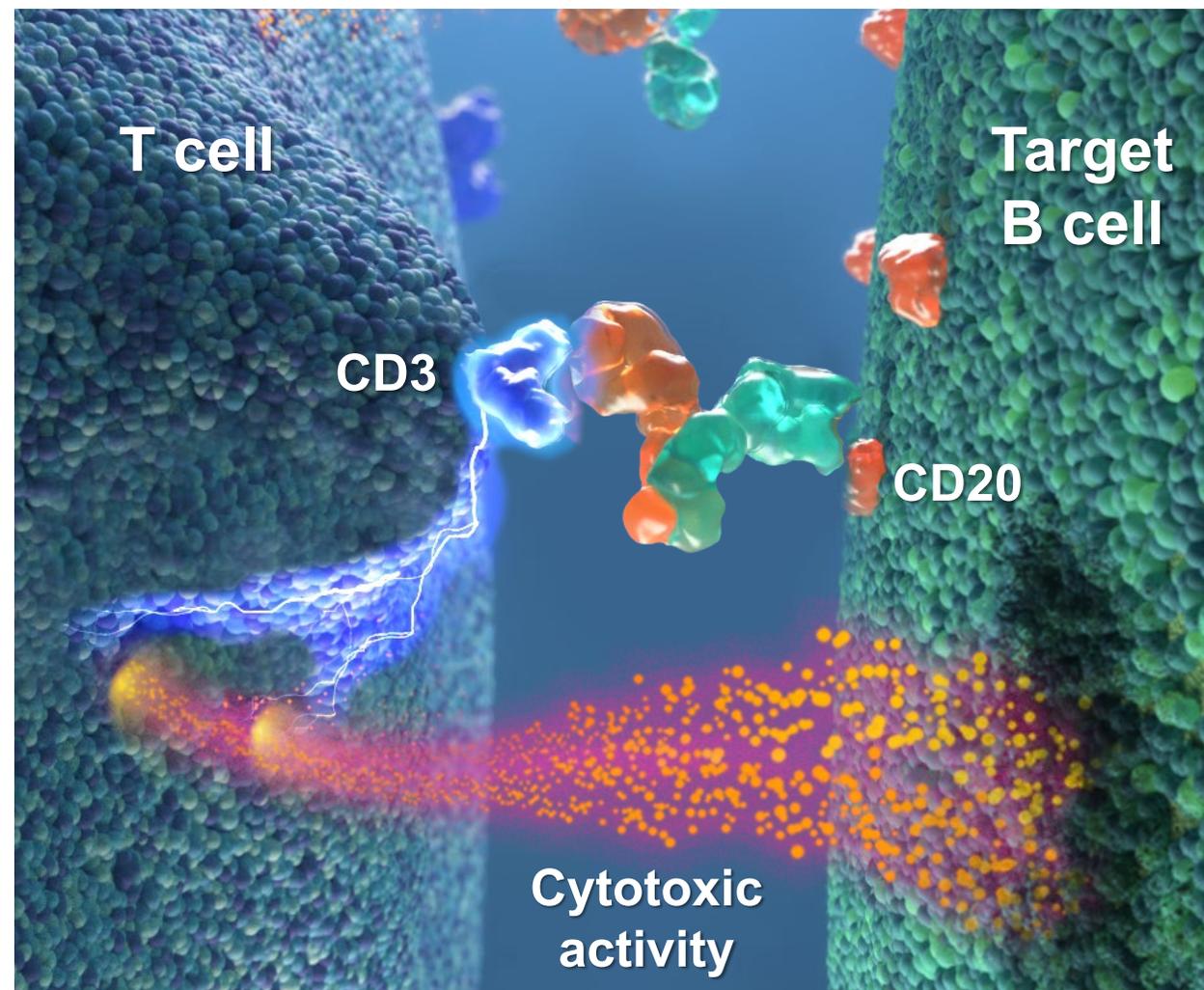
Epcoritamab: Dose Escalation Data Presented at EHA25 Virtual Congress 2020*

Anti-tumor activity

- 86% ORR in FL \geq 0.76mg
- 60% ORR, incl. 3 pts who failed prior CAR-T treatment, in DLBCL/HGBCL \geq 12 mg
- Emerging prelim. data highly encouraging with substantial single-agent efficacy
- Induces rapid and deep responses in heavily pretreated pts with B-NHL across different subtypes

Safety

- No DLTs observed; MTD has not been reached
- No treatment-related deaths
- No discontinuation due to AEs unrelated to disease progression
- No Grade \geq 3 CRS events observed



Dose-escalation data with subcutaneous epcoritamab indicate potential for best-in-class therapy

DuoHexaBody-CD37 (GEN3009)

Building Our Pipeline: First DuoHexaBody in the Clinic



Combination of DuoBody & HexaBody platforms



Novel target for hematologic malignancies



Unique mechanism-of-action



Dose escalation ongoing



50:50 co-development Genmab and AbbVie

DuoBody-CD3x5T4 (GEN1044)

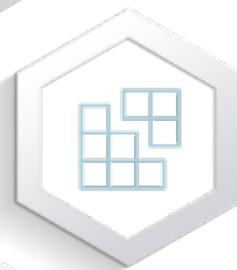
Latest in the Clinic



Based on proprietary DuoBody technology



CD3 bispecific, T cell mediated cytotoxicity of 5T4+ tumor cells



5T4 expressed on multiple solid tumors limited expression in healthy tissue



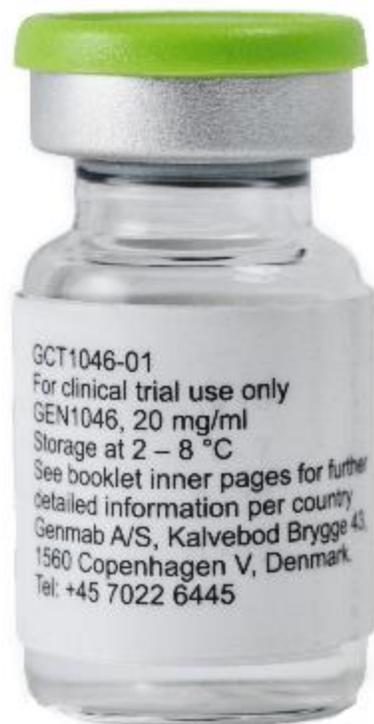
Dose-escalation ongoing



50:50 co-development Genmab and AbbVie

DuoBody-PD-L1x4-1BB (GEN1046)

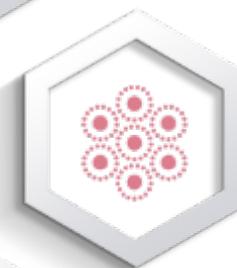
Bispecific Next Generation Checkpoint Immunotherapy



First-in-Class Bispecific antibody targeting PD-L1 & 4-1BB (CD137)



Designed to activate T cells through conditional 4-1BB co-stimulation, while simultaneously blocking the PD1/PD-L1 axis



Combining T cell stimulation with checkpoint blockade



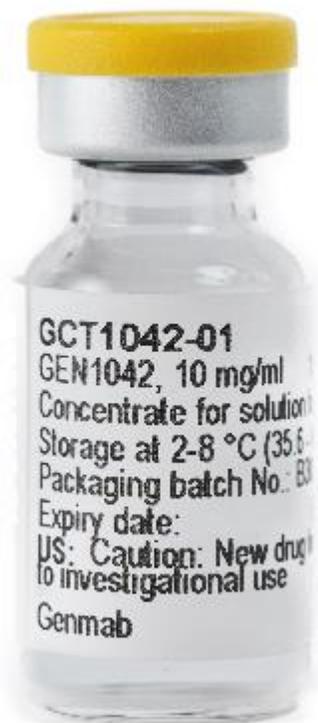
Ph 1/2 study ongoing in solid tumors



50:50 co-development Genmab and BioNTech

DuoBody-CD40x4-1BB (GEN1042)

Bispecific Agonistic Antibody



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



Conditionally activates T cells and APC in presence of CD40-expressing cells



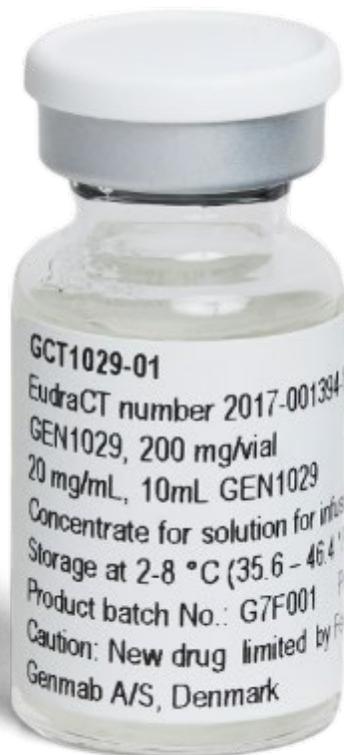
Phase 1/2 study ongoing in solid tumors



50:50 co-development Genmab and BioNTech

HexaBody-DR5/DR5 (GEN1029)

First HexaBody in Clinical Development



Targets 2 distinct DR5 epitopes



HexaBody platform - DR5 clustering & DR5 agonist activity



First 100% Genmab-owned HexaBody product in clinic



Phase 1/2 study ongoing in multiple solid tumors

2020 Guidance: Recurring Revenue Growth and Focused Investments

Income Statement	DKKM	~USDM*	Key Observations
Revenue	9,250 – 9,850	1,423 – 1,515	Summary P&L <ul style="list-style-type: none"> • DARZALEX royalties of ~DKK 4.1bn to ~DKK 4.5bn to drive recurring revenue growth • Nearly 90% of USD 750M upfront from AbbVie collab. recognized immediately • Growth in operating expenses driven by expanding and accelerating our clinical pipeline DARZALEX Sales of USD 3.9bn – USD 4.2bn <ul style="list-style-type: none"> • Significant opportunity for growth in 1L MM market • SubQ DARZALEX approvals in H1 in U.S. & EU • Market share gain in the U.S. and RoW driven by uptake in all lines of treatment • 8 approved indications in U.S., late stage to 1L MM
Operating Expenses	(3,850) – (3,950)	(592) – (608)	
Operating Income	5,350 – 5,950	823 - 915	

Key 2020 Priorities

Building a Strong Differentiated Product Pipeline

Priority	✓	Targeted Milestones
Genmab proprietary* products	✓	» Tisotumab vedotin ¹ - Phase 2 innovaTV 204 safety & efficacy analysis in recurrent/metastatic cervical cancer and engage U.S. FDA for BLA submission subject to trial results
	**	» Tisotumab vedotin - data on other solid tumor types
	***	» Enapotamab vedotin – data to support late stage development
	✓	» Epcoritamab (DuoBody-CD3xCD20) ² Phase 1/2 – decision on recommended Phase 2 dose & initiate expansion cohorts
		» HexaBody-DR5/DR5 Phase 1/2 - advance dose escalation
	✓	» DuoBody-PD-L1x4-1BB ³ Phase 1/2 – initiate expansion cohorts
	✓	» DuoBody-PD-L1x4-1BB initial data in H2 2020
	✓	» File INDs and/or CTAs for 2 new products
Daratumumab ⁴	✓	» U.S. FDA and EMA decision on Phase 3 COLUMBA multiple myeloma SubQ submission
	✓	» sBLA and MAA Submission Phase 3 ANDROMEDA amyloidosis
	✓	» sBLA and MAA submission Phase 3 APOLLO multiple myeloma
Ofatumumab ⁵	✓	» U.S. FDA decision on regulatory dossier submission in multiple sclerosis
Teprotumumab ⁶	✓	» U.S. FDA decision on Phase 3 OPTIC active thyroid eye disease submission

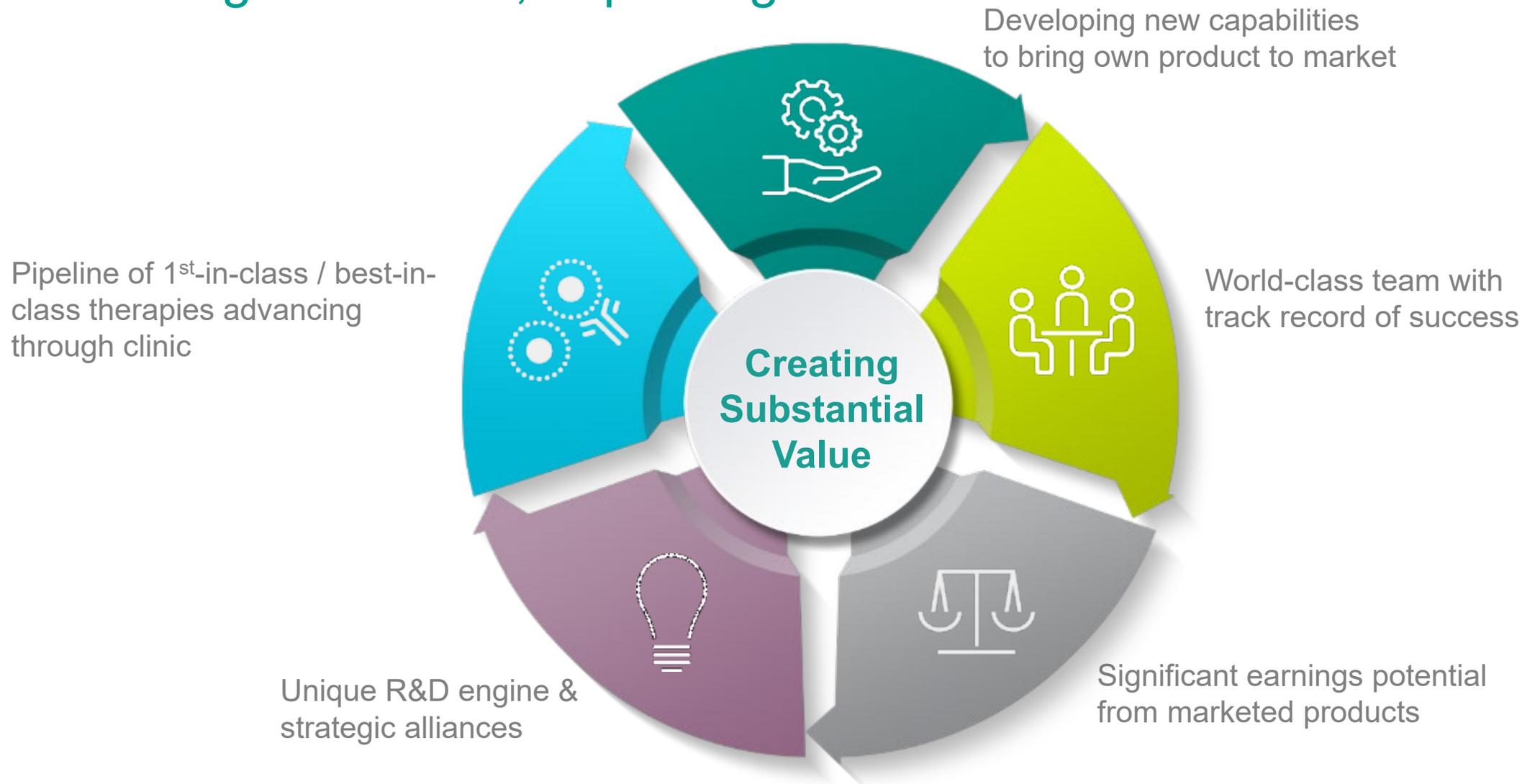
*Certain product candidates in development with partners, as noted.

**Data anticipated in 2021

***Development will not advance

1. 50:50 dev. w/ Seagen; 2. 50:50 dev w/ AbbVie; 3. 50:50 dev. w/ BioNTech; 4. In dev. by Janssen; 5. In dev. by Novartis; 6. In dev. by Horizon Therapeutics

Delivering on Genmab's Promise: Innovating Antibodies, Improving Lives



Innovating Antibodies, Improving Lives

Appendix



A Leading International Biotech With Large Free Float



Genmab & AbbVie: Collaboration Overview

A broad, long-term oncology collaboration with Genmab and AbbVie working together to jointly make all strategy, clinical development and commercialization decisions



50/50 partnership across three clinical next-generation bispecific antibody product candidates (epcoritamab, DuoHexaBody-CD37, DuoBody-CD3x5T4)



Genmab to book epcoritamab sales in the U.S. and Japan; AbbVie to commercialize epcoritamab RoW - Genmab to receive tiered royalties on RoW net sales



Worldwide co-commercialization and profit split of all other programs



Discovery Research Collaboration



Fourth* largest oncology partnership with total potential value ~USD 3.9bn (up-front cash + milestone payments) to Genmab

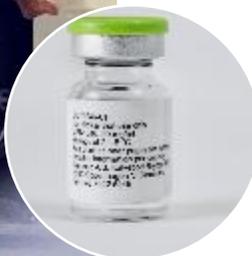
Advancing Pipeline: Delivering on Our Promise & Creating Value

Accelerating Development of Potential “Next Winners”



DuoBody-CD3xCD20 (epcoritamab)

- **Potential best-in-class:** SubQ administration
- Pre-clinical / preliminary clinical data shows encouraging safety & efficacy
- Expeditious and comprehensive clinical development plan
- RP2D decision & expansion cohorts initiation
- 50:50 AbbVie



DuoBody-PD-L1x4-1BB (GEN1046)

- **Potential first-in-class:** Next generation IO
- Unmet medical need
- FiH clinical study: escalation phase is ongoing
- 50:50 BioNTech

Track Record of Success

Advancing Pipeline: Delivering on Our Promise & Creating Value



Delivering on
Genmab's
Promise to
Patients



Bolstering early stage portfolio

- DuoBody-CD40x4-1BB¹; DuoHexaBody-CD37²; DuoBody-CD3x5T4²; HexaBody-CD38³



Adding new technologies



Data sciences



Expanding early stage discovery programs



Enhancing clinical development capabilities

Track Record of Success

Genmab's Commitment to Society

Building a Socially Responsible & Sustainable Company



Anchored in our Core Purpose & Vision

- To improve the lives of patients by creating and developing innovative antibody products
- By 2025 our own product has transformed cancer treatment and we have a pipeline of knock-your-socks-off antibodies



CSR Committee comprised of representatives from variety of functions, chaired by CEO

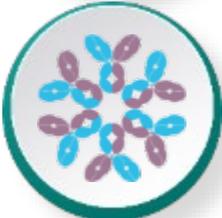
- Ensures that Genmab carries out CSR activities effectively & communicates clearly and openly
- Focus on Environment, Society and Governance reporting



Focus on four main areas

- Employee well-being, including health, safety & development
- Ethics in relation to pre-clinical and clinical studies
- Environment, including waste management & recycling
- Business ethics & transparency

Innovation Powerhouse: Cutting Edge Proprietary Technologies

Technology	Principle	Applications
DuoBody 	Bispecific antibodies	Dual targeting
HexaBody 	Target-mediated enhanced hexamerization	Enhanced potency
DuoHexaBody 	Bispecific antibodies with target-mediated enhanced hexamerization	Dual targeting + enhanced potency
HexElect 	Two co-dependent antibodies with target-mediated enhanced hexamerization	Dual targeting + enhanced potency & selectivity

Innovative Clinical and Pre-Clinical Pipeline

Genmab's Proprietary¹ Products

Product	Target	Developed By	Disease Indications	Most Advanced Development Phase					
				Pre-Clinical	1	1/2	2	3	Approved
Tisotumab vedotin	TF	50:50 Genmab / Seattle Genetics	Cervical cancer	█	█	█	█		
			Ovarian cancer	█	█	█	█		
			Solid tumors	█	█	█	█		
Epcoritamab (DuoBody-CD3xCD20)	CD3, CD20	50:50 Genmab / AbbVie	Hematological malignancies	█	█	█			
DuoBody-PD-L1x4-1BB (GEN1046)	PD-L1, 4-1BB	50:50 Genmab / BioNTech	Solid tumors	█	█	█			
HexaBody-DR5/DR5 (GEN1029)	DR5	Genmab	Solid tumors	█	█				
DuoBody-CD40x4-1BB (GEN1042)	CD40, 4-1BB	50:50 Genmab / BioNTech	Solid tumors	█	█				
DuoHexaBody-CD37 (GEN3009)	CD37	50:50 Genmab / AbbVie	Hematologic malignancies	█	█				
DuoBody-CD3x5T4 (GEN1044)	CD3, 5T4	50:50 Genmab / AbbVie	Solid tumors	█	█				
IND/CTAs in 2020 HexaBody-CD38 (GEN3014) ²		Genmab		█					

Products Created by Genmab*

Including Proposed Label Expansions for Marketed Products

Product	Target	Developed By	Disease Indications	Most Advanced Development Phase					
				Pre-Clinical	1	1/2	2	3	Approved
DARZALEX (daratumumab) & DARZALEX FASPRO (daratumumab and hyaluronidase-fihj) <hr/> Daratumumab	CD38	Janssen (Tiered royalties to Genmab on net global sales)	Multiple myeloma ¹	█	█	█	█	█	█
			AL Amyloidosis	█	█	█	█	█	
			Non-MM blood cancers	█	█	█	█		
Kesimpta (ofatumumab)	CD20	Novartis (Royalties to Genmab on net global sales)	Relapsing multiple sclerosis ¹	█	█	█	█	█	█
Arzerra (ofatumumab)	CD20	Novartis	Chronic lymphocytic leukemia ^{1,2}	█	█	█	█	█	█
TEPEZZA (teprotumumab-trbw) <hr/> Teprotumumab	IGF-1R	Horizon Therapeutics (under sublicense from Roche, royalties to Genmab on net global sales)	Thyroid eye disease ¹	█	█	█	█	█	█
			Diffuse cutaneous systemic sclerosis	█	█				

*Out-licensed products marketed by partner ¹See local country prescribing information for precise indications, ²Not in active development

Partner-owned Products Incorporating Genmab's Innovation*

Product	Target	Developed By	Disease Indications	Most Advanced Development Phase					
				Pre-Clinical	1	1/2	2	3	Approved
Amivantamab (JNJ-61186372)	EGFR, cMet	Janssen	Non-small-cell lung cancer (NSCLC)	█	█	█	█	█	
Teclistamab (JNJ-64007957)	BCMA, CD3	Janssen	Relapsed or refractory MM	█	█	█	█		
PRV-015 (AMG 714)	IL-15	Provention Bio	Celiac disease	█	█	█	█		
Camidanlumab tesirine (ADCT-301)	CD25	ADC Therapeutics	Relapsed /Refractory Hodgkin Lymphoma	█	█	█	█		
			Solid tumors	█	█				
Mim8	FIX(a), FX	Novo Nordisk	Healthy volunteers & hemophilia A	█	█	█			
Talquetamab (JNJ-64407564)	GPRC5D, CD3	Janssen	Relapsed or refractory MM	█	█				
JNJ-63709178	CD123, CD3	Janssen	Acute Myeloid Leukemia (AML)	█	█				
JNJ-63898081	PSMA, CD3	Janssen	Solid tumors	█	█				
JNJ-67571244	CD33, CD3	Janssen	Relapsed or refractory AML or MDS	█	█				
JNJ-70218902	Undisclosed	Janssen	Solid tumors	█	█				
HuMax-IL8	IL8	BMS	Advanced cancers	█	█				
Lu AF82422	alpha-Synuclein	Lundbeck	Parkinson's disease	█	█				

Solid Foundation Built on a Differentiated Pipeline

Tisotumab Vedotin Clinical Program

innovaTV 204

Recurrent or metastatic cervical cancer

- Potentially registrational 102 pts
- Single arm, monotherapy
- 1° endpoint: confirmed ORR
- 2° endpoints: duration of response, PFS, OS

innovaTV 205

Recurrent or metastatic cervical cancer

- In combo or mono
w/ bevacizumab, pembrolizumab, or carboplatin or weekly monotherapy recurrent or stage IVB cervical cancer
- Up to 170 pts
- 1° endpoint: ORR
- 2° endpoints: Safety, duration of response, time to response, PFS, OS

innovaTV 207

Solid tumors

- Basket study
- Up to 250 pts
- Single arm, monotherapy
- 1° endpoint: ORR
- 2° endpoints: Safety, disease control rate, duration of response, time to response, PFS, OS

innovaTV 208

Ovarian cancer

- Ovarian cancer, fallopian tube cancer, peritoneal cancer
- Up to 182 pts, incl 12 pt safety run-in
- Monotherapy
- 2 schedules: q3wk & dose dense
- 1° endpoints: Safety & ORR

Tisotumab Vedotin

Cervical Cancer Market Size

United States³

New Diagnoses	Deaths
12,578	4,115

3rd most common gynecologic cancer in US⁴

Japan⁶

New Diagnoses	Deaths
9,390	3,654

2nd most common gynecologic cancer in Japan⁶

Europe²

New Diagnoses	Deaths
58,373	24,404

3rd most common gynecologic cancer in Europe^{2*}

In developed countries, incidence rates are low (<7.9 per 100,000 women) compared with *developing countries* in sub-Saharan Africa and Central and South America, where incidence is especially high (>30 per 100,000 women)⁵

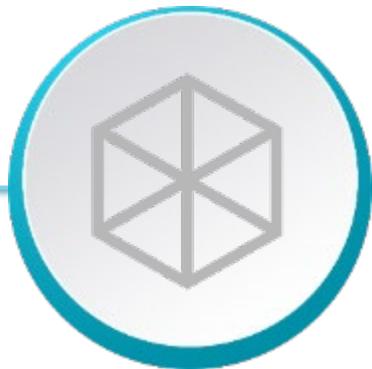
*Europe is defined as the 40 countries in the four United Nations-defined areas of Europe and the European Union (EU-27).

References: 1. American Cancer Society 2. EUCAN (2012) 3. Centers for Disease Control and Prevention. Cervical Cancer Statistics (2017) 4. UpToDate.

5. Ginsburg O et al. *Lancet* 2017 6. HPV Information Centre Japan (2017)

HexaBody-CD38 (GEN3014)

Expanding the Potential of CD38 Antibodies



Incorporates
proprietary
HexaBody
technology



Highly promising
data pre-clinical
models for MM,
lymphoma & AML



Could potentially
add to and broaden
DARZALEX
franchise



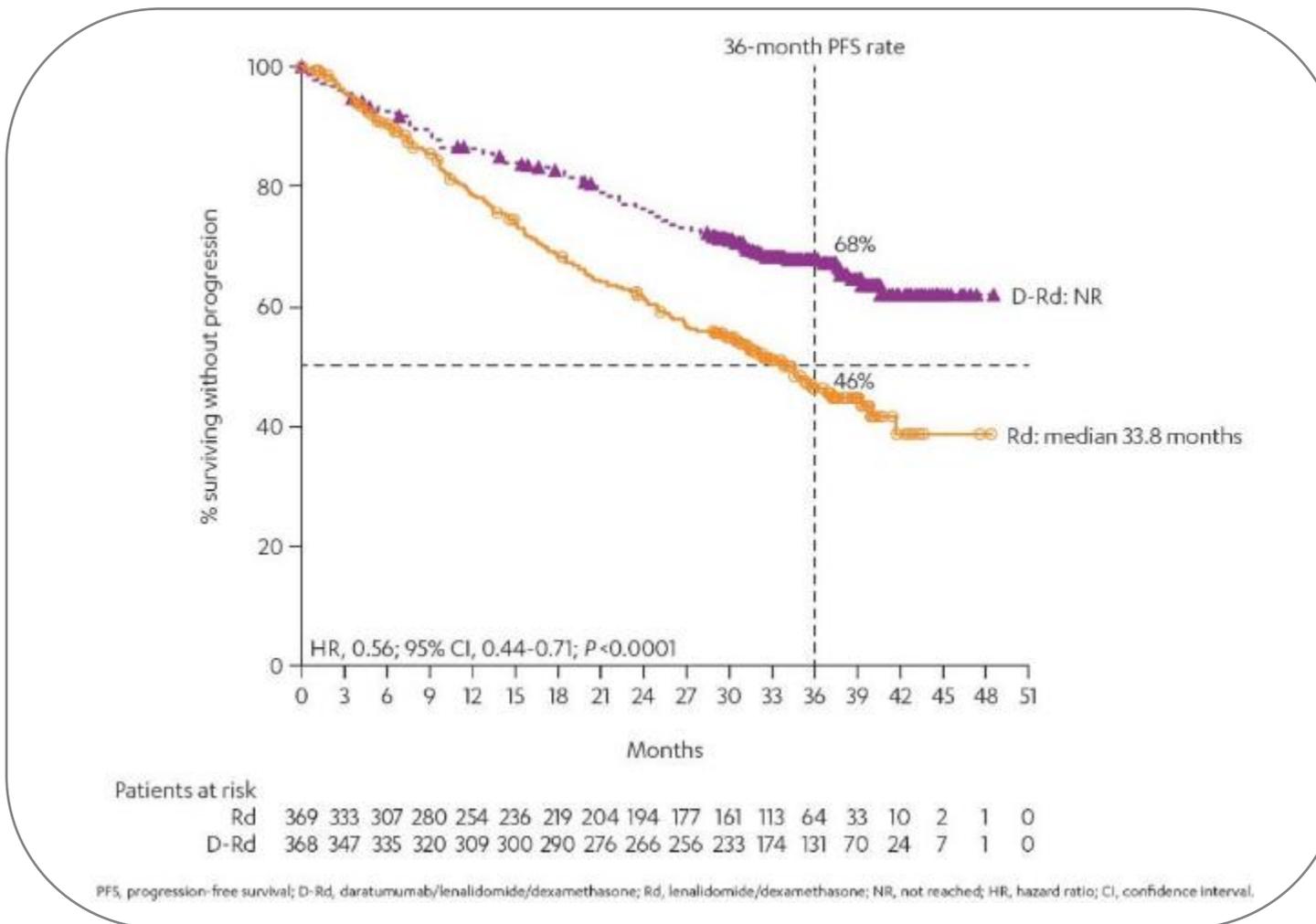
IND/CTA planned
in H2 2020

Covering All Stages of MM and Beyond: Key Ongoing* Industry Sponsored Trials

Disease	Therapy	Development Phase				
		Pre-Clinical	1	1/2	2	3
High Risk Smoldering MM	Subcutaneous		✓ AQUILA			
	Monotherapy		✓ CENTAURUS			
Front line MM (transplant & non-transplant)	Dara + VRd		✓ CEPHEUS			
	Dara + VMP (Asia Pacific)		✓ OCTANS			
	Dara + VRd		✓ PERSEUS			
	Dara + R (maintenance)		AURIGA			
			NINLARO® (Ph II), Venclexta® (Ph II), Selinexor (Ph I/II)			
Relapsed or Refractory MM	Dara + combinations					
	Dara + I.O. (PD1 & PDL1)					
ALL	Dara + SoC chemo		DELPHINUS			

Daratumumab Efficacy in Newly Diagnosed Multiple Myeloma

Updated Phase 3 MAIA Trial (D+Rd, NTE): ASH Dec 2019



- Median PFS not reached in D-Rd arm
- MRD-negativity significantly higher with D-Rd vs. Rd (29% vs 9%; $P < 0.0001$)
- No new safety concerns
- Results continue to support use of D-Rd in 1st line treatment of T1E pts with NDMM

Ongoing Daratumumab Clinical Trials

Janssen Sponsored Phase 3 & 4

Daratumumab Trials Sponsored by Pharma / Biotech

Ct.gov Identifier	Phase	Sponsor	Indication	Therapy
NCT03768960	4	J&J Private Ltd	Relapsed or Refractory MM	Daratumumab (MMY4008)
NCT02252172	3	Janssen	Untreated MM	Daratumumab + Rd (MAIA)
NCT02195479	3	Janssen	Untreated MM	Daratumumab + VMP (ALCYONE)
NCT02541383	3	Janssen	Untreated MM	Daratumumab + VTd (CASSIOPEIA)
NCT02076009	3	Janssen	Relapsed or Refractory MM	Daratumumab + Rd (POLLUX)
NCT02136134	3	Janssen	Relapsed or Refractory MM	Daratumumab + Vd (CASTOR)
NCT03180736	3	Janssen	Relapsed or Refractory MM	Daratumumab + Pom-d (APOLLO)
NCT03201965	3	Janssen	Amyloidosis	Daratumumab + CyBorD (ANDROMEDA)
NCT03217812	3	Janssen	Untreated MM	Daratumumab + VMP (Asia Pacific) (OCTANS)
NCT03234972	3	Janssen	Relapsed or Refractory MM	Daratumumab + Vd vs Vd (LEPUS)
NCT03277105	3	Janssen	Relapsed or Refractory MM	Daratumumab SubQ vs IV (COLUMBA)
NCT03301220	3	Janssen	Smoldering MM	Daratumumab SubQ (AQUILA)
NCT03652064	3	Janssen	Untreated MM	Daratumumab + VRd (CEPHEUS)
NCT03710603	3	Janssen/EMN	Untreated MM	Daratumumab + VRd (PERSEUS)
NCT03901963	3	Janssen	Untreated MM / Maintenance	Daratumumab + R (AURIGA)

Ongoing Daratumumab Clinical Trials

Janssen Sponsored Phase 1 & 2

Daratumumab Trials Sponsored by Pharma / Biotech

Ct.gov Identifier	Phase	Sponsor	Indication	Therapy
NCT03384654	2	Janssen	Relapsed / Refractory ALL / LL	Dara + Vincristine + Prednisone + Doxorubicin (DELPHINUS)
NCT02951819	2	Janssen	Untreated and Relapsed MM	Daratumumab + CyBorD (LYRA)
NCT02874742	2	Janssen	Untreated MM	Daratumumab + VRd (GRIFFIN)
NCT02316106	2	Janssen	Smoldering MM	Monotherapy (CENTAURUS)
NCT02927925	2	Janssen	NKTCL, Nasal Type	Monotherapy (VOLANS)
NCT03412565	2	Janssen	Newly diag. & relapsed / refractory MM	Daratumumab SubQ + Rd, VMP & VRd (PLEIADES)
NCT03871829	2	Janssen	Dara retreatment	Daratumumab SubQ+ Kd vs Kd (LYNX)
NCT03011034	2	Janssen	MDS	Daratumumab (or talacotuzumab) (MDS2002)
NCT01615029	1/2	Janssen	Relapsed and Refractory MM	Daratumumab + Rd (GEN503)
NCT02852837	1	Janssen	Relapsed or Refractory MM	Monotherapy (in China) (MMY1003)
NCT02519452	1	Janssen	Relapsed or Refractory MM	Monotherapy, subcutaneous (PAVO)
NCT02918331	1	Janssen	Untreated MM	Daratumumab + Rd (Japan) (MMY1006)
NCT03242889	1	Janssen	Relapsed or Refractory MM	Daratumumab subq (Japan) (MMY1008)
NCT01998971	1	Janssen	Various MM	Daratumumab + backbone regimens (Vd, VMP, VTd, Pom-d, Kd, KRd) (EQUULEUS)
NCT04108195	1	Janssen	Multiple Myeloma	Daratumuamb + either talquetamab or teclistamab (MMY1002)
NCT04121260	1	Janssen	Multiple Myeloma	Subcutaneous monotherapy (in China) (MMY1010)

Ongoing Daratumumab Clinical Trials

Other Industry Sponsored Trials

Daratumumab Trials Sponsored by Pharma / Biotech

Ct.gov Identifier	Phase	Sponsor	Indication	Therapy
NCT03158688	3	Amgen	Relapsed or Refractory MM	Daratumumab + Kd (CANDOR)
NCT01946477	2	Celgene	Relapsed or Refractory MM	Daratumumab + Pom-d
NCT02807454	2	Celgene	Relapsed and Refractory MM	Daratumumab + Imfinzi (FUSION)
NCT03439293	2	Takeda	Relapsed or Refractory MM	Daratumumab + NINLARO (ixazomib) + Dex
NCT03314181	2	AbbVie	Relapsed or Refractory MM	Daratumumab + Venetoclax + Dex (w/ or w/out bortezomib)
NCT02807558	2	Syros Pharma	AML or MDS	Daratumumab + SY-1425
NCT02773030	1/2	Celgene	Relapsed or Refractory MM	Daratumumab + CC-220 + Dex
NCT02343042	1/2	Karyopharm	Relapsed or Refractory MM	Daratumumab + Selinexor + Dex (STOMP)
NCT03481556	1/2	Oncopeptides AB	Relapsed or Refractory MM	Daratumumab + Melflufen + Dex (ANCHOR)
NCT01592370	1/2	BMS	Relapsed or Refractory MM	Daratumumab + nivolumab
NCT03837509	1/2	Incyte	Relapsed or Refractory MM	Daratumumab + INCB001158
NCT03989414	1/2	Celgene	Various MM	Daratumumab + CC-92480
NCT02431208	1	Roche	Resistant or Refractory MM	Daratumumab + Tecentriq (atezolizumab)
NCT03068351	1	Roche	Resistant or Refractory MM	Daratumumab + RO6870810
NCT04045028	1	Genentech	Relapsed or Refractory MM	Daratumumab + tiragolumab
NCT04136756	1	Nektar Thera.	Salvage for MM	Daratumumab + NKTR-255

