

GENMAB

HexaBody-CD38 overhang gone

JNJ has decided not to opt in for HexaBody-CD38 in multiple myeloma, prompting us to remove HexaBody-CD38 from our SOTP. We note Hexabody-CD38 showed better data than Darzalex in the head-to-head trial (ORR 55% for HexaBody-CD38 versus 52% for Darzalex). With the overhang gone, we expect share-price sentiment to improve and see significant value potential. We reiterate our BUY, but have cut our target price to DKK2,400 (2,500), based on our SOTP NPV.

JNJ has decided not to opt in for HexaBody-CD38. Despite showing better data than Darzalex in the head-to-head trial, JNJ has decided not to opt in for HexaBody-CD38. In the preliminary data (including 84 evaluable patients), HexaBody-CD38 demonstrated an overall response rate of 55% (Darzalex 52%), a very good partial response rate of 29% (Darzalex 17%) and a complete response rate of 7% (Darzalex 2%). Genmab will not pursue further clinical development of HexaBody-CD38.

Removing HexaBody-CD38 from our valuation. Ahead of JNJ's decision, we had included HexaBody-CD38 at a NPV/share of DKK108 in our SOTP, reflecting a 60% probability of opt-in and a 50% probability of approval. However, with JNJ choosing not to opt in, we have removed HexaBody-CD38 from our valuation. We are positive on Genmab's decision not to pursue further clinical development of HexaBody-CD38 and focus on progressing its existing pipeline, and potentially expanding through M&A.

HexaBody-CD38 a clearing event; share-price sentiment set to improve. Despite the negative outcome, we see JNJ's opt-in decision as a clearing event, improving sentiment and renewing investor interest. We focus on Rina-S, with phase II data in 2L+ endometrial cancer due in H1, and we expect it to advance to phase III by end-2025e. We note Tecvayli and Talvey show strong momentum, with JNJ guiding for USD5bn+ peak sales, well above pre-Q4 consensus (USD3,565m and USD2,894m, respectively).

BUY reiterated but target price cut to DKK2,400, based on our SOTP NPV. We remain positive on Genmab's products and pipeline, and expect significant value potential to be realised in the coming weeks. We also expect greater investor interest and focus on the pipeline, given the overhang from JNJ's HexaBody-CD38 decision has been removed.

Year-end Dec	2024	2025e	2026e	2027e
Revenue (USDm)	3,124	3,529	4,213	4,906
EBITDA adj (USDm)	1,000	1,156	1,414	1,594
EBIT adj (USDm)	973	1,133	1,353	1,556
PTP (USDm)	1,330	1,145	1,477	1,669
EPS rep (USD)	17.73	13.82	17.70	19.87
EPS adj (USD)	17.61	13.73	17.58	19.74
Revenue growth (%)	nm	13.0	19.4	16.4
EBITDA growth adj (%)	nm	15.6	22.4	12.7
EPS growth adj (%)	nm	-22.1	28.0	12.3
EBITDA margin adj (%)	32.0	32.7	33.6	32.5
EV/Sales adj (x)	3.29	3.48	2.64	2.02
EV/EBITDA adj (x)	10.3	10.6	7.9	6.2
EV/EBIT adj (x)	10.6	10.8	8.2	6.4
P/E adj (x)	11.8	17.6	13.8	12.3
P/Book (x)	2.50	2.78	2.31	1.95
ROE (%)	nm	16.3	18.5	17.4
ROCE (%)	nm	18.1	19.1	18.6
FCF yield (%)	25.0	5.7	7.9	8.2

Source: Company (historical figures), DNB Markets (estimates)

BUY

TP: DKK2,400

GMAB versus OMXC20 (12m)



Source: FactSet

SUMMARY

Recommendation (prev.)	BUY (BUY)
Share price (DKK)	1,660
Target price (previous) (DKK)	2,400 (2,500)
Upside/downside potential (%)	45
Tickers	GMAB DC

CAPITAL STRUCTURE

No. of shares (m)	64.6
No. of shares fully dil. (m)	65.1
Market cap. (DKKm)	107,255
NIBD adj end-2025e (USDm)	-3,362
Enterprise value adj (USDm)	12,269
Net debt/EBITDA adj (x)	-2.91

Source: Company, DNB Markets (estimates)

Note: Unless otherwise stated, the share prices in this note are the last closing price.

NEXT EVENT

Q1 2025	08/05/2025
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ESTIMATE CHANGES (USDm), (USD)

Year-end Dec	2025e	2026e	2027e
Sales (old)	3,529	4,213	4,902
Sales (new)	3,529	4,213	4,906
Change (%)	0.0	0.0	0.1
EPS adj (old)	13.73	17.58	19.69
EPS adj (new)	13.73	17.58	19.74
Change (%)	0.0	0.0	0.2

Source: DNB Markets, Company consensus

ANALYSTS

Rune Majlund Dahl

rune.dahl@dnb.no

+45 25550122

Oliver Røst Benneballe

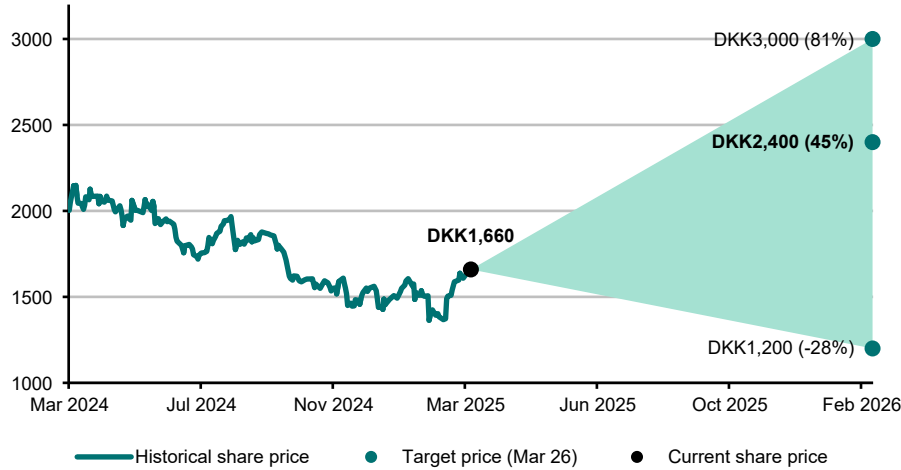
oliver.rost.benneballe@dnb.dk

+45 25550117

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Investment case overview

Share-price performance, DNB Markets' target price, bear- and bull-case scenarios



Source: FactSet, DNB Markets

Downside risks to our investment case

- Lower-than-expected Darzalex sales could have a major effect on the share price.
- Delays in clinical development or the discontinuation of pipeline products could also affect the share price.
- Multiple myeloma (MM) is considered an attractive treatment area, making it a focus area for other companies as well.

Source: DNB Markets

DNB Markets investment case and how we differ from consensus

- We estimate peak Darzalex sales of cUSD16bn (2028), including the subcutaneous version of the drug.
- We see potential in the pipeline and await the publication of further data.
- We still see upside potential to consensus peak sales for Tecvayli and Talvey, which we believe do not yet reflect the potential communicated by partner JNJ.

Source: DNB Markets

Target price methodology

- We continue to base our target price on our SOTP NPV, including 20 years of forecasts, a combined WACC of 9.0%, and 3.0% growth in the terminal period for the pre-clinical pipeline.
- Our bull-case fair value reflects 10% higher Darzalex sales and a de-risked late-stage pipeline (Acasunlimab and Epcoritamab in follicular lymphoma).
- Our bear-case fair value reflects 10% lower Darzalex sales and a pipeline failure (see above).

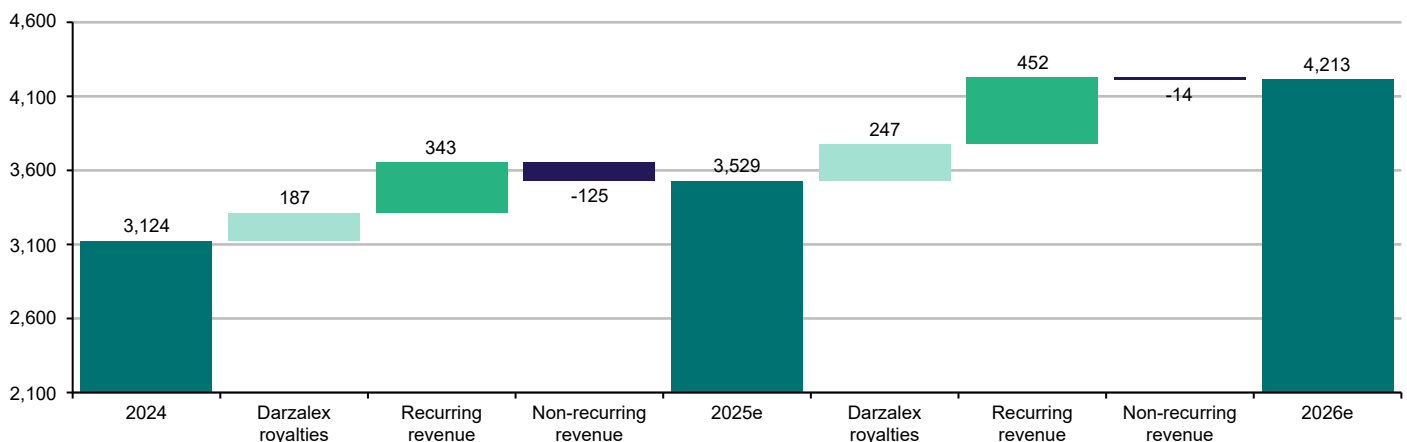
Source: DNB Markets

Upside risks to our investment case

- Higher-than-expected Darzalex sales due to better pricing, treatment duration, or market share gains.
- Positive pipeline news and market penetration/uptake of new products could boost sales and earnings.
- Setbacks in competitors' pipelines could be a positive.
- Effects of the war in Ukraine on the global economy are neutralised and the business climate normalises faster than expected.

Source: DNB Markets

Revenue bridge 2024–2026e (USDm)



Source: Company (historical figures), DNB Markets (estimates)

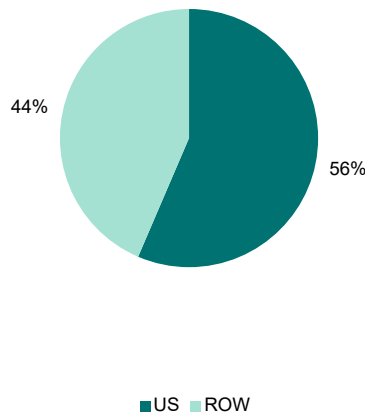
Company overview and SWOT analysis

Company description

- Genmab is a biotechnology company specialising in antibody therapeutics for the treatment of cancer, with four proprietary antibody platforms: DuoBody, HexaBody, DuoHexaBody, and HexElect.
- With its partner, Johnson & Johnson, Genmab has developed Darzalex, which is regarded as the standard of care for multiple myeloma and is Genmab's leading product.

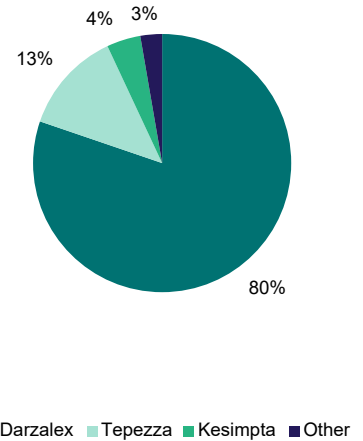
Source: DNB Markets

Darzalex sales by region (2024)



Source: Company

Royalty income by product (2024)



Source: Company

Financial targets

- 2025 revenue of USD3,340m–3,660m.
- 2025 operating profit of USD895m–1,365m.

Source: Company

Key management

- Jan van de Winkel – CEO.
- Anthony Pagano – CFO.

Source: Company

Largest shareholders

- BlackRock – 6.8% (January 2024).

Source: Company

SWOT analysis

Strengths

- **Royalty powerhouse.** Genmab receives significant royalties from its strategic partners (Darzalex c80% of 2024 revenue).
- **Broad market access.** Genmab's strategic collaborations with global pharmaceutical companies enable access to resources and a broader market reach.
- **R&D.** Genmab invests significantly in R&D, focusing on the discovery and development of antibody therapies.

Opportunities

- **Growing oncology market.** Genmab can capitalise on the growing oncology market by developing and commercialising more oncology therapies.
- **Expansion of product portfolio.** Genmab can expand its product portfolio by advancing its pipeline candidates, enhancing the company's market presence.
- **Acquisitions.** Genmab's strong balance sheet enables acquisitions to expand into new disease areas.

Source: DNB Markets

Weaknesses

- **High dependency on partnerships.** Genmab relies heavily on collaborative partnerships to market its products.
- **Competitive pressure.** The industry is competitive and Genmab sees competition from companies with similar expertise and biosimilars.
- **Ethical considerations.** Increased focus on ethical considerations regarding animal testing can affect the public perception and regulatory requirements.

Threats

- **Rising R&D costs.** Rising R&D costs could put pressure on Genmab's profitability.
- **Delays or failures of pipeline candidates.** Delays of clinical trials or failures of pipeline candidates could lead to additional costs and product failures.
- **Regulatory changes.** Changes in regulatory standards could affect the development and approval of Genmab's products, leading to delays or additional costs.

ESG overview

Sustainability assessment

	Positive	Negative
Conclusions	<ul style="list-style-type: none"> ■ We see scope for a positive effect from ESG trends. The group has identified ESG targets, addresses a key health care need with the development of antibodies, and has initiated staff well-being, gender equality, compliance, sustainability, and ethics policies. 	<ul style="list-style-type: none"> ■ With an R&D focus, Genmab's strategy for getting products to market centres on collaborating with biotech and pharma companies, which have ESG policies beyond Genmab's control. ■ With increasing competition for efficacious therapies to treat late-stage disease, there is a potential patient safety-related risk with regard to the drugs in the pipeline.
Actions being taken by company	<ul style="list-style-type: none"> ■ We see scope for a positive effect from ESG trends. The group has identified ESG targets, addresses a key health care need with the development of antibodies, and has initiated staff well-being, gender equality, compliance, sustainability, and ethics policies. 	<ul style="list-style-type: none"> ■ With an R&D focus, Genmab's strategy for getting products to market centres on collaborating with biotech and pharma companies, which have ESG policies beyond Genmab's control. ■ With increasing competition for efficacious therapies to treat late-stage disease, there is a potential patient safety-related risk with regard to the drugs in the pipeline.

Key ESG drivers

Short-term	<ul style="list-style-type: none"> ■ Staff development, knowledge and internal communication initiatives. ■ Controlled waste management of toxic chemicals, and recycling of waste products. ■ 76.8% of Genmab's electricity consumption was from renewables in 2024. 	<ul style="list-style-type: none"> ■ The focus on reducing the environmental effect at Genmab B.V in the Netherlands should be replicated at other sites. ■ Genmab has yet to adopt any environmental indicators. ■ Given the regulatory backdrop, Genmab has not initiated its own human rights policy.
Long-term	<ul style="list-style-type: none"> ■ Genmab has policies in place to limit its environmental effect, but aims to continue to cut waste and switch to more green energy through various initiatives. ■ Genmab has introduced employee satisfaction surveys and strives to ensure it can attract and retain qualified staff. ■ Limited use of natural resources in its research, given the focus on developing compounds. 	<ul style="list-style-type: none"> ■ No structural changes made to reduce the risk of a potential takeover by a pharma company less focused on ESG. ■ The use of hazardous/toxic compounds in research that cannot be replaced by less-toxic alternatives, posing an environmental threat.

Source: DNB Markets

HexaBody-CD38 overhang gone

Despite showing better data than Darzalex in the head-to-head trial, JNJ has decided not to opt-in for HexaBody-CD38. In the preliminary data (including 84 evaluable patients), HexaBody-CD38 demonstrated an overall response rate of 55% (Darzalex 52%), a very good partial response rate of 29% (Darzalex 17%) and a complete response rate of 7% (Darzalex 2%). Genmab will not pursue further clinical development of HexaBody-CD38.

Due to the short-follow up period, secondary efficacy endpoints such as the duration of response, progression-free survival, and overall survival remain immature. We see in the HexaBody-CD38 arm, treatment-emergent events (TEAs) occurring in more than 20% of patients, including neutropenia, infusion-related reactions, anaemia, and thrombocytopenia. We note one death in the HexaBody-CD38 arm and two in the Darzalex arm due to TEAs; however, these were not related to treatment.

Ahead of JNJ's decision, we had included HexaBody-CD38 at an NPV/share of DKK108 in our SOTP, reflecting a 60% probability of opt-in and a 50% probability of approval. However, with JNJ choosing not to opt in, we have removed HexaBody-CD38 from our valuation. We are positive on Genmab's decision not to pursue further clinical development of HexaBody-CD38 and focus on progressing its existing pipeline, and potentially strengthening itself through M&A.

Figure 1: HexaBody-CD38 Efficacy Data

Response Type	HexaBody-CD38 IV arm	Darzalex arm
Overall Response Rate (ORR)	55% (95% CI: 39%, 70%)	52% (95% CI: 36%, 68%)
Very Good Partial Response (VGPR)	29%	17%
Complete Response (CR) or Better	7%	2%

[Source: Genmab]

R&D expectations

We expect phase II data for Rina-S 2L+ in endometrial cancer in H1 and the phase III trial to start by end-2025 (discussions with the FDA already ongoing). Also, we await an update on the phase II data for Acasunlimab 2L+ in small cell lung cancer and phase II data and a decision on the next steps for Rina-S 2L+ in endometrial cancer in 2025.

Figure 2: Pipeline news flow in 2025e

Programme	Indication	Event	Expected timing
Epcoritamab	3L + R/R FL	JP regulatory decision & launch	Q1 2025
Tivdak	2L R/M cervical cancer	EU regulatory decision	2025
Tivdak	2L R/M cervical cancer	JP regulatory decision & launch	2025
Acasunlimab	2L + NSCLC	Phase II data update	2025
Rina-S	2L + endometrial cancer	Phase II data and next steps	H1 2025
DuoBody-CD40x4-1BB	1L HNSCC	Decision on next steps	2025
Epcoritamab	2L DLBCL monotherapy (EPCORE DLBCL-1)	Phase III primary completion	2025

Source: Company (underlying data)

Pipeline overview

Figure 3: Products in development

Priorities	Platform	Product	Target	Disease indications	Development	Partner
Products created by Genmab	UltiMab	Daratumumab	CD38	Outside MM	Phase III	JNJ
	UltiMab	Teprotumumab	IGF-1R	Diffused cutaneous systemic sclerosis	Phase III	Horizon
Genmab's proprietary products	ADC	Tisotumab vedotin	TF	Solid tumours including cervical cancer	Phase II/III	Seagen
	ADC	Rina-S	Fr α	Ovarian cancer and Fr α -expressing tumours	Phase II/III	-
	DuoBody	Epcoritamab	CD3, CD20	Haematological malignancies	Phase I/II/III	AbbVie
	DuoBody	Acasunlimab	PD-L1X4-1BB	Solid tumours and non-small lung cancer	Phase II/III	-
	DuoBody	GEN1042	CD40x4-1BB	Solid tumours	Phase I/II	BioNTech
	HexaBody	GEN3014	CD38	Haematological malignancies	Phase I/II	JNJ ¹
Partner-owned product candidates	DuoBody	Amivantamab	cMET	Advanced or metastatic gastric cancer	Phase II	JNJ
	DuoBody	Mim8	FIXa, FX	Haemophilia A	Phase III	Novo Nordisk
	ADC	Camidanlumab tesirine	CD25	Lymphoma, AML, and solid tumours	Phase II	ADCT
	mAb	PRV-015	IL-15	Celiac disease	Phase II	Provention Bio
	DuoBody	Talquetamab	CD3, GPRC5D	Relapsed or refractory MM	Phase III	JNJ
	mAb	Lu AF82422	α -synuclein	Parkinson's disease	Phase II	Lundbeck
	DuoBody	JNJ-63709178	CD3, CD123	Acute myeloid leukaemia (AML)	Phase I	JNJ
	DuoBody	JNJ-63898081	CD3, PSMA	Solid tumours	Phase I	JNJ
	DuoBody	JNJ-67571244	CD33, CD3	AML or MDS	Phase I	JNJ
	DuoBody	JNJ-70218902	Undisclosed	Solid tumours	Phase I	JNJ
	UltiMab	HuMax-IL8	IL8	Advanced cancers	Phase I	BMS
	All platforms			c20 active pre-clinical programmes ²	Pre-clinical	Various

Source: Company

Notes: ¹ Genmab entered an exclusive worldwide licence and option agreement with JNJ Biotech Inc. to develop and commercialise HexaBody-CD38. ² Pre-clinical pipeline includes partnered products and in-house programmes based on Genmab's proprietary technologies (HuMab, DuoBody, DuoHexaBody, and HexaBody)

2025 guidance

The 2025 guidance is for revenue of USD3,340m–3,660m (we forecast USD3,529m, consensus USD3,594m), including Darzalex royalties of USD2,200m at the midpoint of the guidance (we forecast USD2,207m, consensus USD2,213m), opex of USD2,055m–3,420m (we forecast USD2,203m, consensus USD2,358m), and operating profit of USD895m–1,365m (we forecast USD1,133m, consensus USD1,025m).

Figure 4: 2025 guidance (USDm)

	2025 guidance	DNB Markets forecasts	Consensus*
Revenue	3,340–3,660	3,529	3,594
Royalties	2,785–3,015	2,903	2,879
Darzalex royalties	2,200	2,207	2,213
- Darzalex global sales	12,600–13,400	13,063	1,861
Net product sales/collaboration revenue	415–460	460	520
Milestone/reimbursement revenue	140–185	165	166
Gross profit	3,120–3,420	3,336	3,383
Operating expenses	2,055–2,225	2,203	2,358
Operating profit	895–1,365	1,133	1,025

Source: Company (consensus and guidance), DNB Markets (estimates)

Note: *the company-compiled consensus is reported in DKK, but we have converted it to USD using consensus' USD/DKK7.0 for 2025

Valuation

Our target price is based on a risk-adjusted SOTP NPV for each product, including 20 years of forecasts (as for the other Nordic pharmaceutical companies we cover). Our c9% group WACC is based on the WACCs we have assigned to each product. We have assigned 8.5–9.0% WACCs for approved products and 11% for pipeline products, risk-adjusted for the probability of success. Based on these assumptions, our SOTP NPV is DKK2,400/share (2,500), which is also our target price.

Figure 5: NPV analysis

Product	Indication	Phase	Launch year	Peak sales		WACC	Risk-adj. NPV	NPV/share	% total	
				(USDm)	Probability		(DKKm)	(DKK)		
Darzalex	MM and AL	Marketed	2015–2021	15,730	100%	8.5%	53,017	790	33%	
Tepezza	Thyroid eye disease	Marketed	2020	3,035	100%	8.5%	4,906	75	3%	
Kesimpta	MS	Marketed	2020	5,164	100%	8.5%	13,274	204	8%	
Rybrevant	NSCLC	Marketed/III	2021	4,350	100%/80%	8.5%	8,153	125	5%	
TivDak	Cervical cancer	Marketed	2021	542	100%	8.5%	3,523	54	2%	
Tecvayli	MM	Marketed/I/III	2023	5,758	100%/80%	9.0%	4,993	77	3%	
Talvey	MM	Marketed/I/III	2023	5,637	100%/80%	9.0%	3,850	59	2%	
Epkinly	B-cell lymphoma	Phase I/II	2023	3,249	100%/80%	9.0%	17,566	270	11%	
Total approved products/indications NPV							109,281	1,655	69%	
Acasunlimab	Solid tumours	Phase I/II	2026e	1,925	50%/20%	11.0%	7,231	111	5%	
GEN1042	Solid tumours	Phase I/II	2028e	3,566	20%	11.0%	2,426	37	2%	
Rina-S	Solid tumours	Phase I/II	2027e	2,006	50%	11.0%	7,580	116	5%	
Mim8	Haemophilia A	Phase III	2025e	3,361	100%	11.0%	4,520	69	3%	
Platform/other pipeline	Various					11.0%	7,225	111	5%	
Total pipeline NPV							28,981	445	19%	
Net cash (debt), end-2025e								20,093	309	13%
Warrant programme, end-2025e								(593)	(9)	0%
SOTP							9.1%	157,762	2,400	100%

Source: DNB Markets (estimates)

Forecast changes – P&L

(USDm)	New			Old			Change		
	2025e	2026e	2027e	2025e	2026e	2027e	2025e	2026e	2027e
Revenues	3,529	4,213	4,906	3,529	4,213	4,902	0	0	4
Cost of sales	-192	-326	-690	-192	-326	-690	0	0	0
Gross profit	3,336	3,887	4,217	3,336	3,887	4,213	0	0	4
Operating expenses	-2,203	-2,534	-2,661	-2,203	-2,534	-2,661	0	0	0
EBITDA	1,156	1,414	1,594	1,156	1,414	1,590	0	0	4
EBITDA adj	1,156	1,414	1,594	1,156	1,414	1,590	0	0	4
EBITDA margin (%)	32.7	33.6	32.5	32.7	33.6	32.4	0.0	0.0	0.1
Depreciation	-20	-5	-35	-20	-5	-35	0	0	0
EBITA	1,135	1,409	1,558	1,135	1,409	1,554	0	0	4
Amortisation	-2	-56	-2	-2	-56	-2	0	0	0
EBIT	1,133	1,353	1,556	1,133	1,353	1,552	0	0	4
EBIT adj	1,133	1,353	1,556	1,133	1,353	1,552	0	0	4
Net interest	13	124	113	13	124	113	0	0	0
Net financial items	13	124	113	13	124	113	0	0	0
PBT	1,145	1,477	1,669	1,145	1,477	1,665	0	0	4
Taxes	-252	-325	-367	-252	-325	-366	0	0	-1
Net profit	893	1,152	1,302	893	1,152	1,299	0	0	3
Adjustments to net profit	0	0	0	0	0	0	0	0	0
Net profit adj	893	1,152	1,302	893	1,152	1,299	0	0	3
<i>Per share data (USD)</i>									
EPS	13.82	17.70	19.87	13.82	17.70	19.82	0.00	0.00	0.05
EPS adj	13.73	17.58	19.74	13.73	17.58	19.69	0.00	0.00	0.05
<i>Other key metrics (%)</i>									
Revenue growth	13.0	19.4	16.4	13.0	19.4	16.4	0.0	0.0	0.1
EBIT adj growth	16.5	19.4	15.0	16.5	19.4	14.7	0.0	0.0	0.3
EPS adj growth	-22.1	28.0	12.3	-22.1	28.0	12.0	0.0	0.0	0.3
Avg. number of shares (m)	65	66	66	65	66	66	0	0	0
Capex	-59	-70	-82	-59	-70	-82	0	0	0
OpFCF	1,097	1,344	1,512	1,097	1,344	1,508	0	0	4
Working capital	444	473	427	444	473	427	0	0	0
NIBD adj	-3,362	-4,627	-5,950	-3,362	-4,627	-5,947	0	0	-3

Source: DNB Markets

Annual P&L

(USDm)	2024	2025e	2026e	2027e
Revenues	3,124	3,529	4,213	4,906
Cost of sales	-143	-192	-326	-690
Gross profit	2,981	3,336	3,887	4,217
Operating expenses	-2,008	-2,203	-2,534	-2,661
EBITDA	1,000	1,156	1,414	1,594
Depreciation	-27	-20	-5	-35
EBITA	973	1,135	1,409	1,558
Amortisation	-1	-2	-56	-2
EBIT	973	1,133	1,353	1,556
Net interest	180	13	124	113
FX gains	177	0	0	0
Net financial items	357	13	124	113
PBT	1,330	1,145	1,477	1,669
Taxes	-192	-252	-325	-367
Effective tax rate (%)	14	22	22	22
Net profit	1,138	893	1,152	1,302
Net profit adj	1,138	893	1,152	1,302
Avg. number of shares	65	65	66	66
<i>Per share data (USD)</i>				
EPS	17.73	13.82	17.70	19.87
EPS adj	17.61	13.73	17.58	19.74
<i>Growth and margins (%)</i>				
Revenue growth	nm	13.0	19.4	16.4
EPS adj growth	nm	-22.1	28.0	12.3
Gross margin	95.4	94.5	92.3	85.9
EBITDA margin	32.0	32.7	33.6	32.5
EBITDA adj margin	32.0	32.7	33.6	32.5
Depreciation/revenues	-0.9	-0.6	-0.1	-0.7
EBIT margin	31.1	32.1	32.1	31.7
EBIT adj margin	31.1	32.1	32.1	31.7
PBT margin	42.6	32.5	35.0	34.0
Net profit margin	36.4	25.3	27.3	26.5

Source: Company (historical figures), DNB Markets (estimates)

Cash flow

(USDm)	2024	2025e	2026e	2027e
Net profit	1,138	893	1,152	1,302
Depreciation and amortisation	27	23	61	38
Other non-cash adjustments	3	0	132	0
Change in net working capital	2,222	30	-29	46
Cash flow from operations (CFO)	3,391	947	1,316	1,386
Capital expenditure	-52	-59	-70	-82
Acquisitions/Investments	-3,023	-1,986	-2,209	-2,180
Divestments	1,637	1,788	1,988	1,963
Cash flow from investing (CFI)	-1,438	-257	-291	-299
Free cash flow (FCF)	1,953	689	1,025	1,086
Share issue (repurchase)	-563	-607	0	0
Other	171	19	19	19
Cash flow from financing (CFF)	-392	-588	19	19
Total cash flow (CFO+CFI+CFF)	1,560	102	1,044	1,106
<i>FCFF calculation</i>				
Free cash flow	1,953	689	1,025	1,086
Less: net interest	-180	-13	-124	-113
Less: acquisitions	3,023	1,986	2,209	2,180
Less: divestments	-1,637	-1,788	-1,988	-1,963
Free cash flow to the firm	3,159	875	1,122	1,191
Growth (%)				
CFO	nm	-72.1	39.0	5.3
CFI	nm	82.1	-13.1	-3.0
FCF	nm	-64.7	48.7	6.0
CFF	nm	-49.8	103.2	1.0
FCFF	nm	-72.3	28.2	6.1

Source: Company (historical figures), DNB Markets (estimates)

Balance sheet

(USDm)	2024	2025e	2026e	2027e
Assets	6,648	7,003	8,257	9,610
Trade receivables	980	999	1,111	1,096
Cash and cash equivalents	3,062	3,362	4,627	5,950
Current assets	4,042	4,361	5,738	7,047
Property, plant and equipment	142	180	245	292
Other intangible assets	2,292	2,289	2,234	2,231
Deferred tax assets	132	132	0	0
Non-current financial assets	41	41	41	41
Non-current assets	2,606	2,642	2,519	2,564
Total assets	6,648	7,003	8,257	9,610
Equity and liabilities	6,648	7,003	8,257	9,610
Total equity to the parent	5,326	5,631	6,802	8,124
Total equity	5,326	5,631	6,802	8,124
Trade payables	505	554	638	670
Other payables and accruals	265	265	265	265
Total current liabilities	770	819	903	935
Other non-current liabilities	552	552	552	552
Total non-current liabilities	552	552	552	552
Total liabilities	1,323	1,372	1,455	1,487
Total equity and liabilities	6,648	7,003	8,257	9,610
<i>Key metrics</i>				
Net interest bearing debt	-3,062	-3,362	-4,627	-5,950
Invested capital	617	625	718	719

Source: Company (historical figures), DNB Markets (estimates)

Valuation ratios

(USDm)	2018	2019	2020	2021	2022	2023	2024	2025e	2026e	2027e
<i>Enterprise value</i>										
Share price (USD)	163.46	222.18	407.27	400.74	422.08	319.34	207.73	241.86	241.86	241.86
Number of shares (m)							64.18	64.63	65.08	65.52
Market capitalisation							13,333	15,632	15,739	15,847
Net interest bearing debt							-3,062	-3,362	-4,627	-5,950
Net interest bearing debt adj							-3,062	-3,362	-4,627	-5,950
EV							10,271	12,269	11,112	9,897
EV adj							10,271	12,269	11,112	9,897
<i>Valuation</i>										
EPS							17.73	13.82	17.70	19.87
EPS adj							17.61	13.73	17.58	19.74
P/E							11.7	17.5	13.7	12.2
P/E adj							11.8	17.6	13.8	12.3
P/B							2.50	2.78	2.31	1.95
Average ROE								16.3%	18.5%	17.4%
Earnings yield adj							8.5%	5.7%	7.3%	8.2%
EV/SALES							3.29	3.48	2.64	2.02
EV/SALES adj							3.29	3.48	2.64	2.02
EV/EBITDA							10.3	10.6	7.9	6.2
EV/EBITDA adj							10.3	10.6	7.9	6.2
EV/EBIT							10.6	10.8	8.2	6.4
EV/EBIT adj							10.6	10.8	8.2	6.4
EV/capital employed							1.7	1.9	1.4	1.1
EV/NOPLAT							13.5	13.9	10.5	8.2
EV/OpFCF (taxed)							14.0	14.5	10.6	8.5

Source: Company (historical figures), DNB Markets (estimates)

Key accounting ratios

	2024	2025e	2026e	2027e
<i>Profitability (%)</i>				
ROA		13.1	15.1	14.6
ROCE		18.1	19.1	18.6
ROCE after tax		14.1	14.9	14.5
<i>Return on invested capital (%)</i>				
Net PPE/revenues	4.5	5.1	5.8	5.9
Working capital/revenues	15.2	12.6	11.2	8.7
Revenues/invested capital (pre-GW)	506.6	564.8	586.7	682.8
Pre-tax ROIC (incl. goodwill)		182.9	209.8	216.9
After-tax ROIC (incl. goodwill)		142.3	157.2	168.9
<i>Cash flow ratios (%)</i>				
FCF/revenues	62.5	19.5	24.3	22.1
FCF yield (%)	25.0	5.7	7.9	8.2
CFO/revenues	108.5	26.8	31.2	28.2
CFO/market capitalisation	25.4	6.1	8.4	8.7
CFO/capex	6507.9	1608.3	1872.7	1693.7
CFO/current liabilities	440.2	115.5	145.8	148.3
Cash conversion ratio	171.6	77.2	89.0	83.4
Capex/revenues	1.7	1.7	1.7	1.7
Capex/depreciation	196.1	288.1	1290.4	231.9
OpFCF margin	30.3	31.1	31.9	30.8
<i>Leverage and solvency (x)</i>				
Interest cover	67.18	48.72	62.52	high
EBIT/interest payable	55.85	47.20	56.37	nm
EBITA adj/interest payable	55.88	47.30	58.69	nm
Cash coverage	-5.56	-91.72	-11.44	-14.05
Net debt/EBITDA	-3.06	-2.91	-3.27	-3.73
<i>Cash conversion cycle</i>				
Receivables turnover days	114.5	103.3	96.2	81.6
Credit period	1290.3	1051.6	712.9	354.4

Source: Company (historical figures), DNB Markets (estimates)

Important Information

Company: Genmab
 Coverage by Analyst: Rune Majlund Dahl
 Date: 10/3/2025

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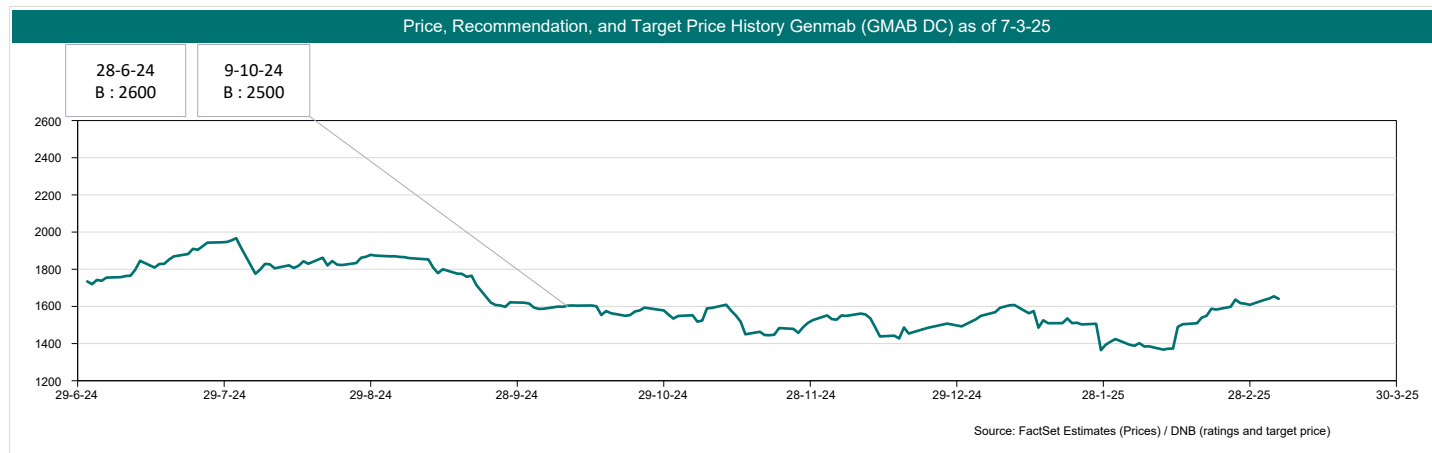
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