Healthcare

Genmab

Price DKK923.50

Bloomberg Reuters 12-month High, Market Cap (DK Ev (BG Estimate Avg. 6m daily vo 3y EPS CAGR	GEN DC GEN.CO 954.0 / 514.0 55,113 51,732 412.9 -12.3%			
	1 M	3 M	6 M 31	1/12/15
Absolute perf.	7.6%	0.7%	49.2%	0.7%
Healthcare	-4.1%	-13.4%	-8.7%	-13.4%
DJ Stoxx 600	-1.6%	-8.9%	-3.8%	-8.9%
YEnd Dec. (DKKm)	2015	2016e	2017e	2018e
Sales	1,133	866.7	906.4	1,306
% change		-23.5%	4.6%	44.1%
EBITDA	554	50.1	2.7	351
EBIT	730.4	50.1	2.7	351.2
% change		-93.1%	-94.5%	
Net income	587.3	85.1	42.7	396.2
% change		-85.5%	-49.8%	
	2015	2016 e	2017e	2018 e
Operating margin	64.5	5.8	0.3	26.9
Net margin	67.4	9.8	4.7	30.3
ROE	21.9	2.4	1.2	9.9
ROCE	-15,400	44.2	11.1	69.2
Gearing	-100.2	-94.6	-89.4	-85.8
(DKK)	2015	2016 e	2017 e	2018e
EPS	9.71	1.41	0.71	6.55
% change	-	-85.5%	-49.8%	
P/E	95.1x	NS	NS	NS
FCF yield (%)	0.3%	0.0%	0.1%	NM
Dividends (DKK)	0.00	0.00	0.00	0.00
Div yield (%)	NM	NM	NM	NM
EV/Sales	45.6x	59.7x	57.2x	39.6x
EV/EBITDA	93.1x	1031.6x	18875.1x	147.1x
EV/EBIT	70.7x	1031.6x	18875.1x	147.1x



Daratumumab's European approval recommended by the CHMP

Fair Value DKK1300 (+41%)

BUY-Top Picks

The CHMP has issued a positive opinion recommending the grant of a conditional market authorization for Darzalex (daratumumab, an anti-CD38) in the EU, and as a treatment for heavily pre-treated patients with multiple myeloma. While largely expected (don't forget that the compound has already been approved by the FDA back in November 2015 for this very same label), we think the market will positively respond to this news as the timing of an European approval is key for respecting/beating the FY 2016 sales guidance that was given for this very compound (USD250-300m). For now, we stick to our sales estimate of EUR219m for this very year... But we have to admit we're quite conservative. We'll see how dara performed in the US during Q1 16, but 1) we're pretty sure the data presented during the 2015 ASH meeting have had a very positive impact on prescriptions (be it for on-label or off-label use); and 2) we can't rule out that CASTOR and POLLUX will further enhance this trend. BUY reiterated with a FV of DKK1,300.

ANALYSIS

- The CHMP (Committee for Medicinal Products for Human Use) has issued a positive opinion recommending the grant of a conditional marketing authorization for Darzalex (daratumumab, an anti-CD38) 1) in the European Union, and 2) as a treatment for heavily pre-treated patients with multiple myeloma. Consequently, we can say the European Commission should give its green light in coming weeks (and we'd say by mi-may at the earliest based).
- While largely expected (don't forget that the compound has already been approved by the FDA back in November 2015), we think the market will positively respond to this news as (i) the timing of an European approval is also key for respecting/beating the FY 2016 sales guidance that was given for this very compound (USD250-300m), especially since the management said they were expecting it by mid-2016; (ii) we assume JNJ will quickly reallocate a part of its marketing effort to daratumumab as Velcade (for which they currently retain the ex-US rights) is set to lose ground because of an increasing competition from new generation of proteasome inhibitors (Amgen's Kyprolis (carfilzomib), Takeda's Ninlaro (ixazomib)) and the advent of generics.
- For now, we stick to our sales estimate of EUR219m for this very year... But we have to admit
 we're quite conservative. We'll see how dara performed in the US during Q1 16, but 1) we're
 pretty sure the data presented during the ASH meeting have had a very positive impact on
 prescriptions (be it for on-label or off-label use); and 2) we can't rule out that CASTOR and POLLUX
 will further enchance this trend.

VALUATION

• BUY reiterated with a FV of DKK1,300.

NEXT CATALYSTS

- Q2 16: European approval of daratumumab as a monotherapy for the treatment of heavily pretreated patients with multiple myeloma
- H1 16: Top-line results from the POLLUX study (Phase III evaluating daratumumab in combination
 with lenalidomide and dexamethasone in R/R patients with myeloma who received at least one
 prior therapy).
- H2 16: Read-across analysis for ofatumumab from the (likely) approval of Roche's ocrelizumab for the treatment of relapsing-remitting multiple sclerosis.

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Stock rating

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NEUTRAL

Opinion recommending not to trade in a stock short-term, neither as a BUYER or a SELLER, due to a specific set of factors. This view is intended to be temporary. It may reflect different situations, but in particular those where a fair value shows no significant potential or where an upcoming binary event constitutes a high-risk that is difficult to quantify. Every subsequent published update on the stock will feature an introduction outlining the key reasons behind the opinion.

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Distribution of stock ratings

BUY ratings 64%

NEUTRAL ratings 29.4%

SELL ratings 6.6%

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