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Company Comment	Healthcare	Sweden	18 September 2023
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Catalyst receives new FDA clearance

FDA has approved OssDsign Catalyst for use in interbody cages in spinal surgery. This will allow surgeons to use Catalyst on-label in the interbody space. Today a lot of the synthetic bone graft is believed to be used off-label which represents a risk to the hospital and surgeons. We view this news as very positive as it will most likely increase the number of hospital approvals which should lead to more users and larger volumes. Our mid-point of SEK 12 is under review.

Key figures				
(SEKm)	2022	2023E	2024E	2025E
Revenues (m)	57	98	136	189
Adj. EBIT	(90)	(84)	(62)	(13)
PTP (m)	(100)	(84)	(62)	(13)
EPS	(1.74)	(1.17)	(0.87)	(0.19)
EPS (adjusted)	(1.74)	(1.17)	(0.87)	(0.19)
DPS	0.00	0.00	0.00	0.00
Revenue growth (%)	79.6	72.2	38.7	39.3
EPS growth (%)	n.a.	n.a.	n.a.	n.a.
Operating margin (%)	(158.8)	(85.7)	(45.8)	(7.0)
ROCE (%)	(34.9)	(41.9)	(36.7)	(7.6)
Net Debt/EBITDA (x)	1.4	0.2	(1.1)	(45.0)
PER (adjusted)	(3.3)	(4.2)	(5.7)	(26.8)
Dividend yield (%)	0.0	0.0	0.0	0.0
Free Cash Flow Yield (%)	(26.8)	(23.5)	(18.9)	(6.0)
P/BV (x)	1.8	2.5	4.4	5.3
EV/EBIT (x)	(3.4)	(4.0)	(6.6)	(33.2)
EV/Sales (x)	5.34	3.43	3.04	2.32

Source: SEB

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