

Genkyotex

Buy → | Target 2.90 EUR

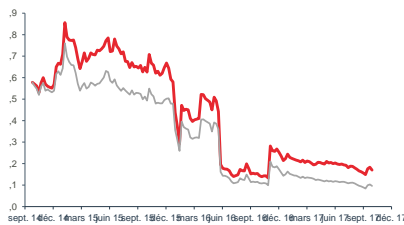
Price (27/10/2017) : 1.70 EUR | Upside : 71 %

Est.chg	2017e	2018e
EPS	ns	ns

Future worldwide specialist in fibrosis

Publication date 30/10/2017 07:51

Writing date 30/10/2017 07:50



Source ; Oddo BHF Securities, Fininfo

Capital

GKTX FP GKTX.PA	
Market Cap (EURm)	132
Enterprise value (EURm)	139
Extrema 12 months	1.30 - 3.25
Free Float (%)	9.6

Performance (%)	1m	3m	12m
Absolute	8.3	-11.9	16.4
Perf. rel. Country Index	6.1	-14.4	1.1
Perf. rel. CAC Small	5.7	-15.3	-12.8

P&L	12/17e	12/18e	12/19e
Sales (EURm)	0.0	0.0	0.0
EBITDA (EURm)	-27.2	-16.2	-38.4
Current EBIT (EURm)	-27.3	-16.2	-38.5
Attr. net profit (EURm)	-27	-16	-39
Adjusted EPS (EUR)	-0.35	-0.21	-0.49
Dividend (EUR)	0.00	0.00	0.00

P/E (x)	ns	ns	ns
P/B (x)	ns	ns	ns
Dividend Yield (%)	0.0	0.0	0.0
FCF yield (%)	ns	ns	ns
EV/Sales (x)	ns	ns	ns
EV/EBITDA (x)	ns	ns	ns
EV/Current EBIT (x)	ns	ns	ns
Gearing (%)	ns	ns	ns
Net Debt/EBITDA(x)	0.3	ns	ns

Next Events

Genkyotex is a listed biotech since 2017 following its reverse take-over of Gentecel. A worldwide specialist in NOX inhibitors it aims to become a leader in fibrotic pathologies such as the very attractive NASH market. We are initiating coverage of the stock on Buy, with a target price of € 2.9, pointing to upside of 70%.

At the leading edge of NOX science

Genkyotex's defining feature is its thorough knowledge of NOX, the enzymes responsible for oxidative stress in humans. The company was founded 11 years ago by three international experts in the field. Today, it is still one of the very few companies with a real capacity to discover these compounds. It is also one of the rare NOX biotech companies to have reached the human testing phase. Finally, it is, to our knowledge, the only NOX company in the world to specialise in fibrotic diseases such as NASH. Genkyotex has already built up a considerable body of encouraging preclinical data. It is preparing to deliver its first clinical data for liver fibrosis in H1 2018.

Fibrosis a potential market worth \$ 6bn in 2022e

In targeting fibrosis, Genkyotex is looking to carve out a position on a fledgling but potentially massive market. The principal indications that could be targeted by the group (NASH and PBC in the liver, diabetic nephropathy, pulmonary fibrosis) represents, according to EvaluatePharma, a potential market of \$ 6bn in 2022 (growing at a pace of 24% per annum). In our view, the advantage of targeting a market that is largely still not addressed is that the competitive field is still relatively open. By way of example, in NASH, only cenicriviroc (Allergan/Novartis) and selonsertib (Gilead), still in development, clearly have mechanistically anti-fibrotic properties. All of the other treatments that are currently in clinical trials (anti-metabolic, anti-inflammatory) would not actually be competing but would rather round out treatment by the future anti-fibrotic products like Genkyotex's GKT831.

Valuation: € 2.9 per share, 70% upside

We value Genkyotex via the sum of its cash position and the indications of its main compound, GKT831. We are excluding, for the time being, the group's second compound, GKT771 (too early) and the agreement on Vaxicase with Serum Institute (a legacy from the former Gentecel which milestones are difficult to model). In the event of clinical success and regulatory approval, GKT831, could, on our estimates, generate peak sales (2029e) of € 230m in PBC (assuming it becomes commercially available in 2021), € 2.7bn in NASH (assuming it becomes commercially available in 2024e) and € 1.9bn in diabetic nephropathy (assuming it becomes commercially available in 2024). We value Genkyotex at € 2.9 per share (€ 0.4 for PBC, € 0.3 for nephropathy, € 2.2 for NASH, -€ 0.1 for cash). The stock offers upside of 70%, on our estimates.

Main risks

Genkyotex is a company with strong potential but still in the fledgling stage. Whilst the rewards could be significant in the event of a blue-sky scenario materialising (potentially x8 out to 2023), the principal risk remains the failure of clinical trials (GKT831 has mainly been tested in the pre-clinical trial stage to date). In the longer term, the company's capacity to effect its transition towards the commercial phase must also be monitored. The stock's main risks are detailed on page 10.

Sebastien Malafosse (Analyst)

+33 (0)1 44 51 88 46
sebastien.malafosse@oddo-bhf.com

Conflict of interests: