15th November 2016

Healthcare

ERYTech

Price EUR17.47

Bloomberg Reuters 12-month High Market Cap (EU Avg. 6m daily vo	ERYP FP ERYP.PA 29.0 / 16.1 139 15.30			
	1 M	3 M	6 M 3	1/12/15
Absolute perf.	6.1%	-13.1%	-25.3%	-31.8%
Healthcare	-2.7%	-7.9%	-2.8%	-13.3%
DJ Stoxx 600	-0.5%	-2.3%	1.1%	-7.5%
	2014	2015e	2016e	2017e
P/E	NS	NS	NS	NS
Div yield (%)	NM	NM	NM	NM

Marketing approval further delayed in Europe. FV revised down

Fair Value EUR30 vs. EUR47 (+72%)

BUY

ANALYSIS

- Erytech yesterday announced they it has decided to withdraw the MAA (marketing authorisation application) for GRASPA in Europe. Efficacy and safety-wise, the clinical package has not been called into question. However, the company believes that it would need more time than the procedures allow to address all of the requests from the CHMP (1/ additional data comparability between the old form of asparaginase that was included in the red blood cells in the last Phase III vs the newer one; 2/ development of a new immunogenicity test; 3/ Q&A around the pharmacodynamics). So in very concrete terms, we estimate the marketing approval could be delayed by c.2 years (given that management said it intends to resubmit the MAA by mid-2017).
- Obviously the stock will react quite negatively this morning; and the very next catalyst is quite a
 hard one to play (see our previous comments for further details). That said, we remain positive on
 the fundamentals behind GRASPA in ALL and its rationale in AML. And we believe the latter should
 be seen as a buffer in the very near term (given that it amounts to c.EUR5 per share in our SOP).

VALUATION

• We revise down our FV from EUR47 to EUR30 as 1/ we have increased our average WACC from 12.5% to 15.0%, and 2/ delayed our first sales assumption by nearly two years.

NEXT CATALYSTS

- Q1 2016: Phase II results for ERY-ASP/GRASPA for the treatment of pancreatic cancer.
- H2 2016: Phase II results for GRASPA in acute myeloid leukaemia.

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Mickael Chane Du, mchanedu@bryangarnier.com

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London	Paris	New York	Munich	New Delhi	
Beaufort House	26 Avenue des Champs Elysées	750 Lexington Avenue	Widenmayerstrasse 29	The Imperial Hotel Janpath New Delhi 110 001 Tel +91 11 4132 6062 +91 98 1111 5119	
15 St. Botolph Street	75008 Paris	New York, NY 10022	80538 Munich		
London EC3A 7BB	Tel: +33 (0) 1 56 68 75 00	Tel: +1 (0) 212 337 7000	Germany		
Tel: +44 (0) 207 332 2500	Fax: +33 (0) 1 56 68 75 01	Fax: +1 (0) 212 337 7002	+49 89 2422 62 11	Fax +91 11 2621 9062	
Fax: +44 (0) 207 332 2559	Regulated by the	FINRA and SIPC member		Geneva	
Authorised and regulated by the	Financial Conduct Authority (FCA) and the			rue de Grenus 7	
Financial Conduct Authority (FCA)	Autorité de Contrôle prudential et de			CP 2113	
	resolution (ACPR)			Genève 1, CH 1211 Tel +4122 731 3263	
				Fax+4122731 3243	

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