

# **Allium Medical Solutions**

Wirion approved by FDA; new stents in Europe

Allium has announced FDA approval of its subsidiary Gardia's Wirion, which has become the only embolic protection system approved for all atherectomy procedures. Allium has reported FY17 results, with revenues up 5% y-o-y to NIS7.7m, and announced its first order in Mexico for NIS300k and approval of some of its stents in Russia. However, registration of the remaining stents and IBI Medical (EndoFast soft tissue fixation) in Mexico and Russia and stents in China has been delayed; we believe approval is possible in 2018. Mexico, Russia and China are the bulk of the distribution deals (NIS132m) and delays are the major reason for the slight decline in our valuation to NIS1.64/share (NIS1.68/share previously).

Year end	Revenue (NISm)	PBT*	EPS*	DPS	P/E	Yield
ena	(MISIM)	(NISm)	(NIS)	(NIS)	(x)	(%)
12/16	7.4	(22.0)	(0.49)	0.0	N/A	N/A
12/17	7.7	(21.4)	(0.37)	0.0	N/A	N/A
12/18e	14.0	(13.7)	(0.19)	0.0	N/A	N/A
12/19e	21.0	(9.6)	(0.13)	0.0	N/A	N/A

Note: \*Normalised, excluding amortisation of acquired intangibles and exceptionals.

## FDA approves Wirion; chances of transaction higher

Gardia Medical's Wirion system has received FDA approval for leg artery catheterisation on the back of strong clinical data from the WISE-LE trial, which met its primary and secondary endpoints at interim analysis. Wirion captured a significant amount of debris in all clinical cases and across all atherectomy devices and is now the only embolic protection system cleared by the FDA for all atherectomy procedures. Allium is looking for a strategic transaction, preferably M&A; we believe the FDA approval could make Gardia more attractive to potential partners. We model Wirion as part of Allium's overall business and valuation and project revenue of NIS2.4m after full launch in 2018, rising to NIS8.6m in 2020 as we adjust our price forecast based on the most recent FY17 data.

## Mexico, Russia and China: Approvals and delays

Allium has received the first order for its urological stent products in Mexico for NIS300k, which was paid upfront in Q417. We now expect EndoFast (soft tissue fixation) and the remaining stents to be approved in Mexico and Russia during H118 vs YE17 before. Allium has also received approval in Russia for its stent products and EndoFast could be approved in 2018, as it could in China after it sent a full package to the Chinese FDA in Q417. We reduce our revenue forecast to c NIS14m in FY18 (from NIS16.6m) to reflect these delays, partially offset by a new deal worth NIS6.3m signed in 15 countries in Central and Eastern Europe.

## Launch of new stents; Allevetix starts clinical trial

Allium has launched two new CE-marked products in Europe: the Triangular Prostate Stent (TPS) Plus which is a new approach to treat benign prostate hyperplasia; and Ureteral Stent (URS) 200, which self-expands to 200mm length and 9mm diameter. Allium has started a clinical trial with the Allevetix indwelling gastroduodenal sleeve in 10 patients with obesity and diabetes for three months; Allium expects to complete the study by the YE and start a pivotal trial in 2019.

### Valuation: DCF of c NIS117m or NIS1.64/share

Our DCF valuation of Allium is NIS1.64/share (vs NIS1.68/share) as a result of updating the launch delays in Mexico, Russia and China and net cash.

FY17 results; business update

Medical devices

### 26 April 2018

Price*	NIS1.31
Market cap	NIS94m
*Priced at 20 April 2018	
Net cash (NISm) at end 2017	22.95
Shares in issue	71.4m
Free float	60%
Code	ALMD
Primary exchange	TASE
Secondary exchange	N/A

#### Share price performance

1.5



#### **Business description**

Allium Medical Solutions is a company focused on developing and marketing minimally invasive devices in various areas: cardiovascular, metabolic, genitourinary and gastrointestinal. The company has three selling product lines: Allium Stents, IBI (EndoFast) and Gardia Medical. Allium markets its products mainly through distribution agreements.

#### **Next events**

Potential strategic agreements for Gardia	2018
Regulatory approval in additional markets for Allium and IBI	2018

#### **Analysts**

Juan Pedro Serrate +44 (0)20 3681 2534 Jonas Peciulis +44 (0)20 3077 5728

Q418

healthcare@edisongroup.com

Complete Allevetix first-in-human trial

Edison profile page



## FY17 results overview and valuation update

Allium reported sales of NIS7.7m in FY17, up 5% vs NIS7.4m in FY16, but lower than our forecast of NIS9.7m. Allium has received the first order for its ureteral stent products in Mexico for NIS300k, fully paid in Q417. We now expect EndoFast (soft tissue fixation) and the remaining stents to be approved in Mexico and Russia during 2018 vs YE17 previously. The distribution deals are worth a total of NIS74m over five years. Separately, Allium has received approval in Russia for its stent products. Next is approval of EndoFast, potentially in 2018 (deal is NIS48m over five years). Approval in China could be possible in 2018; the company sent a full package to the CFDA in Q417 (deal is NIS58m over eight years). Additionally, Allium has been strengthening the distribution network and has replaced the distributor of stents in the Czech Republic and Slovakia and EndoFast with a new deal in 15 Central and Eastern European countries. The deal involves a minimum purchase of NIS6.3m over five years. Therefore, we reduce our near-term revenue forecast to reflect these delays, partially offset by the new commercial agreement to c NIS14m in FY18 (from NIS16.6m) and c NIS21m in FY19 (from NIS24.9m).

We expect R&D expenses to decrease in 2018 (NIS10m in FY18 vs NIS5m previously) due to public grants from the Israeli government. R&D expenses in 2018 are mainly associated with the continued clinical development of Allevetix and the preclinical study with TruLeaf. From 2019 onwards we expect a decrease in R&D expenses as the company completes clinical development of Allevetix. General and administrative expenses were NIS8.5m in FY17 vs NIS8.1m in FY16. This is above our estimate of NIS7.3m and is mainly related to non-cash, stock-based compensation.

EBITDA loss in FY17 was NIS20.8m vs NIS20.4m in FY16. This is slightly higher than our FY17 estimate of NIS19.5m. We forecast an EBITDA loss of NIS13.2m in FY18, from a loss of NIS6.8m previously. Net loss was NIS22.7m vs our forecast of a NIS21.9m loss. We now forecast a net loss of NIS15.2m in FY18 (vs NIS8.8m before). As a consequence of our revised forecast, we now expect Allium to reach EBITDA break-even in 2020, one year later than before.

NIS000s	FY16	FY17			FY18e			
	Act.	Est.	Act.	Change (%)	Old	New	Change (%)	
Revenue	7,353	9,735	7,703	-21%	16,612	13,972	-16%	
R&D expenses	(13,494)	(14,000)	(13,914)	-1%	(5,000)	(10,000)	100%	
S&M expenses	(2,895)	(2,531)	(2,458)	-3%	(3,322)	(3,493)	5%	
G&A expenses	(8,129)	(7,250)	(8,527)	18%	(7,000)	(7,000)	0%	
Operating profit/loss	(22,633)	(21,548)	(22,842)	6%	(8,624)	(15,100)	75%	
Profit/loss before tax	(23,917)	(21,909)	(22,679)	4%	(8,770)	(15,236)	74%	
Profit/loss after tax	(23,917)	(21,909)	(22,679)	4%	(8,770)	(15,236)	74%	
EPS (NIS)	(0.53)	(0.38)	(0.39)	3%	(0.12)	(0.21)	75%	

During 2017 Allium consumed NIS18.9m in cash vs NIS17.5m in FY16. This is lower than our FY17 estimate of NIS20.1m. Over the year, Allium raised total net proceeds of NIS19.1m. We estimate that net cash, cash equivalents and short-term deposits of c NIS23m provide runway until 2019 when we project a cash shortfall that for illustrative purposes we cover with long-term debt of NIS20m. Cash will be spent on the first-in-man study with Allevetix's gastroduodenal sleeve (recently started) and the ongoing animal study with TruLeaf.

## Valuation updated to NIS1.64/share

We were expecting approval of all stent products and EndoFast in Mexico in Q417, in Russia in H118 and approval of stents in China in Q118. For modelling purposes, we have lowered our



forecasts to reflect approval and launch in those countries in H218 instead of the initial dates. We have also updated the new deal in Central and Eastern Europe and distribute NIS6.3m over 2018 to 2023 as the company's distributor has already started selling.

As a result of these changes, our revised DCF valuation of Allium moves to NIS1.64 per share, or NIS117m, from NIS119m or NIS1.68/share, which includes net cash of c NIS23m at end 2017.

Exhibit 2: Allium summary DCF valuation							
	Previous valuat	tion	Current valuation				
	\$000s	NIS000s	\$000s	NIS000s			
PV of explicit FCF forecast (2018-26e)	5,131*	18,577	4,733	16,319			
Terminal value (2% TGR)	55,732	201,783	49,995	172,396			
PV of Terminal value	19,308	69,905	19,485	67,190			
Value attributed to Allevetix	2,680	9,703	3,071	10,588			
Total NPV	27,119	98,186	27,288	94,098			
Add net cash (end-2017)	5,723**	20,722	6,656	22,953			
Implied equity value	32,842	118,907	33,945	117,051			
Number of shares (m)	70,501	70,501	71,42	71,423			
Per basic share	\$0.47	NIS1.68	\$0.48	NIS1.64			

Source: Edison Investment Research. Note: US dollar values are based on the spot exchange rate. \*PV of explicit FCF forecast is 2017-26e. \*\*Net cash as per our estimate of 2017.



	NIS'000 2015	2016	2017	2018e	2019
Year end 31 December	IFRS	IFRS	IFRS	IFRS	IFR
PROFIT & LOSS					
Revenue	5,178	7,353	7,703	13,972	20,99
Cost of Sales	(4,421)	(5,171)	(5,687)	(8,579)	(10,520
Gross Profit	757	2,182	2,016	5,392	10,47
EBITDA	(16,333)	(20,377)	(20,826)	(13,174)	(9,014
Operating Profit (before amort. and except.)	(16,759)	(20,759)	(21,219)	(13,549)	(9,347
Intangible Amortisation	(1,705)	(1,579)	(1,623)	(1,551)	(1,378
Exceptionals	(720)	(295)	0	0	(
Operating Profit	(19,184)	(22,632)	(22,842)	(15,100)	(10,725
Net Interest	(1,748)	(1,283)	(163)	(135)	(202
Exceptionals	0	0	0	0	
Other	0	0	0	0	(
Profit Before Tax (norm)	(18,507)	(22,042)	(21,382)	(13,684)	(9,550
Profit Before Tax (IFRS)	(20,932)	(23,916)	(22,679)	(15,236)	(10,928
Tax	0	0	0	0	(
Profit After Tax (norm)	(18,507)	(22,042)	(21,382)	(13,684)	(9,550
Profit After Tax (IFRS)	(20,932)	(23,916)	(22,679)	(15,236)	(10,928
Average Number of Shares Outstanding (m)	28.53	44.97	58.19	71.42	71.42
EPS - normalised (NIS)	(0.65)	(0.49)	(0.37)	(0.19)	(0.13
EPS - IFRS (NIS)	(0.73)	(0.53)	(0.39)	(0.21)	(0.15
Dividend per share (NIS)	0.00	0.00	0.00	0.00	0.00
Gross Margin (%)	15%	30%	26%	39%	50%
EBITDA Margin (%)	N/A	N/A	N/A	N/A	N/A
Operating Margin (before GW and except.) (%)	N/A	N/A	N/A	N/A	N/A
BALANCE SHEET					
Fixed Assets	25,612	23,616	22,244	20,417	18,806
Intangible Assets	24,059	22,465	20,916	19,364	17,986
Tangible Assets	1,472	1,025	1,059	784	550
Restricted cash	81	126	269	269	269
Current Assets	31,342	28,606	29,609	15,815	26,627
Stocks	2,277	2,516	2,661	2,278	2,534
Debtors	889	1,253	1,491	1,531	2,013
Cash, equivalents and short term deposits	27,053	23,203	22,953	9,501	19,576
Other	1,123	1,634	2,504	2,504	2,504
Current Liabilities	(5,620)	(12,660)	(11,962)	(11,798)	(12,147
Creditors	(1,524)	(1,890)	(1,987)	(1,823)	(2,172
Accruals	(1,895)	(936)	(185)	(185)	(185
Other short term liabilities	(2,201)	(4,124)	(4,373)	(4,373)	(4,373
Long Term Liabilities	(6,207)	(1,368)	(1,134)	(913)	(20,692
Long term borrowings	Ó	Ó	0	Ó	(20,000
Other long term liabilities	(6,207)	(1,368)	(1,134)	(913)	(692
Net Assets	45,127	38,194	38,757	23,521	12,593
CASH FLOW					
Operating Cash Flow	(15,874)	(17,258)	(18,418)	(13,131)	(9,605
Net Interest	(13,674)	(17,230)	(10,410)	(13,131)	(9,003
Tax	0	0	0	0	
Capex	(164)	(220)	(501)	(100)	(100
Acquisitions/disposals	(104)	(220)	(4,005)	(100)	(100
• •		12.056	,		
Financing	31,992	13,956 0	19,125 0	0	(
Dividends Other					
Other	(1,841)	(328)	(456)	(221)	(221
Net Cash Flow	14,113	(3,850)	(4,255)	(13,452)	(9,926
Opening net debt/(cash)	(12,940)	(27,053)	(23,203)	(22,953)	(9,501
HP finance leases initiated	0	0	0	0	
Other	0	0	4,005	0	10
Closing net debt/(cash)	(27,053)	(23,203)	(22,953)	(9,501)	42



Edison is an investment research and advisory company, with offices in North America, Europe, the Middle East and AsiaPac. The heart of Edison is our world-renowned equity research platform and deep multi-sector expertise. At Edison Investment Research, our research is widely read by international investors, advisers and stakeholders. Edison Advisors leverages our core research platform to provide differentiated services including investor relations and strategic consulting. Edison is authorised and regulated by the Financial Conduct Authority. Edison Investment Research (NZ) Limited (Edison NZ) is the New Zealand subsidiary of Edison. Edison NZ is registered on the New Zealand Financial Service Providers Register (FSP number 247505) and is registered to provide wholes ale and/or generic financial adviser services only. Edison Investment Research Inc. (Edison US) is the US subsidiary of Edison and is regulated by the Securities and Exchange Commission. Edison Investment Research Pty Limited (Edison Aus) [46085869] is the Australian subsidiary of Edison. Edison Germany is a branch entity of Edison Investment Research Limited [4794244], <a href="https://www.edisongroup.com">www.edisongroup.com</a>

#### EDISON ISRAEL DISCLAIMER

Disclosure regarding the scheme to enhance the awareness of investors to public companies in the technology and biomed sectors that are listed on the Tel Aviv Stock Exchange and participate in the scheme (hereinafter respectively "the Scheme", "TASE", "Participant" and/or "Participants"). Edison Investment Research (Israel) Ltd, the Israeli subsidiary of Edison Investment Research Ltd (hereinafter respectively "Edison Israel" and "Edison"), has nettered into an agreement with the TASE for the purpose of providing research analysis (hereinafter "the Agreement"), regarding the Participants and according to the Scheme (hereinafter "the Analysis" or "Analyses"). The Analysis will be distributed and published on the TASE website (Maya), Israel Security Authority (hereinafter "the ISA") website (Magna), and through various other distribution channels. The Analysis for each participant will be published at least four times a year, after publication of quarterly or annual financial reports, and shall be updated as necessary after publication of an immediate report with respect to the occurrence of a material event regarding a Participant. As set forth in the Agreement, Edison Israel is entitled to see for providing its investment research services. The fees shall be paid by the Participants directly to the TASE, and TASE shall pay the fees directly to Edison. Subject to the terms and principals of the Agreement, the Annual fees that Edison Israel shall be entitled to for each Participant shall be in the range of \$35,000-50,000. As set forth in the Agreement and subject to its terms, the Analyses shall include a description of the Participant participant

#### DISCLAIMER

Copyright 2018 Edison Investment Research Limited. All rights reserved. This report has been commissioned by Allium Medical Solutions and prepared and issued by Edison for publication of publicity available sources that are believed to be reliable, however we do not guarantee the accuracy or completeness of this report Opinions contained in this report represent those of the research department of Edison at the time of publication. The securities described in the Investment Research may not be eligible for sale in all jurisdictions or to certain categories of investors his research is issued in Australia by Edison Investment Research Pty Ltd (Corporate Authorised Representative (1252501) of Myonlineadvisers Pty Ltd (AFSL: 427484)) and any access to it, is intended only for "wholesale clients" within the meaning of the Corporations At 2001 of Australia. The Investment Research is distributed in the United States by Edison US to major US institutional investors only. Edison US is registered as an investment adviser with the Securities and Exchange Commission. Edison US relies upon the "publishers' exclusion" from the definition of investment adviser under Section 202(a)(11) of the Investment Advisers Act of 1940 and corresponding state securities laws. As such, Edison does not offer or provide personalised advice. We publish information about companies in which we believe our readers may be interested and this information provided by us should not be construed by any subscriber or prospective subscriber as Edison's solicitation to effect, or attempt to effect, any transaction in a security. The research in this document is intended for New Zelaand resident professional financial advisers or brokers; join abhabitual investors who are "wholesale clients" for the purpose of the Financial Advisers Act 2008 (FAA) (as described in sections 5(c) (1)(a), (b) and (c) of the FAA). This is not a solicitation or inducement to buy, sell, subscribe, or underwrite any securities mentioned or in the topic of this document. This