

Celyad

Allogeneic trial first dose given

The allogeneic version of NRK CAR T-cell therapy, CYAD-101, is underway with the first patient dosed. The study mirrors the current colorectal SHRINK trial in combining NKR CAR cell therapy with FOLFOX chemotherapy. This gives Celyad the lead in a potential high-value massmarket solid cancer where allogeneic therapy is likely to be essential. The indicative value is €1,090m (€89 per share) pending further data.

Year end	Revenue (€m)	PBT* (€m)	EPS* (€)	DPS (€)	P/E (x)	Yield (%)
12/16	8.52	(20.00)	(2.09)	0.0	N/A	N/A
12/17	3.54	(26.80)	(2.79)	0.0	N/A	N/A
12/18e	0.00	(27.25)	(2.43)	0.0	N/A	N/A
12/19e	0.00	(28.50)	(2.38)	0.0	N/A	N/A

Note: *PBT and EPS are normalised, excluding amortisation of acquired intangibles, exceptional items and share-based payments.

Leading position in solid cancers and allogeneic cells

Celyad has a dominant position in CAR therapy for solid cancers (see T-cell cancer therapies). Its NKR CAR T-cell therapy targets 'stressed' cancer cells, for example, after chemotherapy like FOLFOX. As an allogenic approach, CYAD-101 is a more affordable, rapid-response potential therapy for many thousands of patients. Of allogeneic competitors in the US, only Cellectis (UCART123 in AML) and Precision Biosciences (PBCAR0191) are in trials but for haematological cancers. Allogene in solid cancers is in preclinical development; UCART19 targets ALL. However, there are many allogeneic trials in China. Celyad has regained its allogeneic therapy rights in key Asian counties from ONO as ONO failed to exercise its option.

AlloSHRINK

Celyad is already evaluating autologous (customised) CYAD-01 with FOLFOX in the open-label SHRINK trial. The alloSHRINK (NCT03692429) study mirrors the SHRINK design. Initial results from three SHRINK patients were disclosed in November 2018. Of these, one had a pathological complete response (CR) and the other two had partial pathological responses. However, cell numbers after infusion were low and no Cy-Flu preconditioning was used. AlloSHRINK is starting its dose escalation phase to test safety and is focusing on any signs of graft vs host disease. A later expansion phase will test the objective response rate. One important secondary endpoint is host vs graft (HvG) response especially after the second and third doses, as HvG could eliminate CYAD-101 cells or block any cell expansion, stopping any therapeutic benefit. The 36-patient trial could compete in late 2021. Celyad holds key patents in allogeneic therapy (licensed by Novartis).

Valuation: Unchanged at €1,090m pending more data

We assign a nominal value of €100m to the allogenic opportunity; this will be revised as response and safety data emerge. AlloSHRINK open-label data is likely to be reported as the trial progresses. Allogeneic products may come to dominate the valuation. The overall indicative value remains €1,090m, €89 per share. More CYAD-01 data is due on 3 December at ASH.

AlloSHRINK underway

Pharma & biotech

3 December 2018

Price	€24.30
Market cap	€290m
	\$1.18/€
Cash (€m) at 30 September 201	8 55.9
Shares in issue	11.94m
Free float (Edison estimate)	67%
Code	CYAD
Drimany avalance	Comment December

Primary exchange **Euronext Brussels** Secondary exchange NASDAQ

Share price performance



Business description

Celyad is developing an innovative natural killer receptor (NKR) CAR T-cell immune-oncology platform. Celyad has a leading position in CAR for AML and solid tumours and is exploring the use of NKR CAR with chemotherapy. It holds a key granted patent in allogeneic CAR technology.

Next		-1-
next	ever	IIS

ASH data	3 December 2018
FY18 results	Q119

Analysts

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Edison profile page

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	€000s 2016	2017	2018e	2019
Year end 31 December	IFRS	IFRS	IFRS	IFR
PROFIT & LOSS				
Revenue	8,523	3,540	0	
Cost of Sales	(53)	(515)	(500)	(50)
Gross Profit	8,470	3,025	(500)	(500
EBITDA	(21,246)	(22,317)	(26,500)	(27,750
Operating Profit (before amort, and except).	(22,006)	(23,283)	(27,500)	(28,750
Intangible Amortisation	(756)	(748)	(750)	(750
Other income and charges	0	(26,273)	0	
Share-based payments	(2,847)	(2,569)	(2,600)	(2,600
Operating Profit	(25,609)	(52,873)	(30,850)	(32,100
Net Interest	1,997	(3,521)	250	25
Profit Before Tax (norm)	(20,009)	(26,804)	(27,250)	(28,500
Profit Before Tax (FRS 3)	(23,612)	(56,394)	(30,600)	(31,850
Tax	6	1	Ó	•
Profit After Tax (norm)	(20,003)	(26,803)	(27,250)	(28,500
Profit After Tax (FRS 3)	(23,606)	(56,393)	(30,600)	(31,850
Average Number of Shares Outstanding (m)	9.3	9.6	11.2	12.
EPS - normalised (c)	(209)	(279)	(243)	(238
EPS - (IFRS) (c)	(253)	(586)	(243)	(267
Dividend per share (c)	0.0	0.0	0.0	0.
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Gross Margin (%)	N/A	N/A	N/A	N/A
EBITDA Margin (%)	N/A	N/A	N/A	N/.
Operating Margin (before GW and except) (%)	N/A	N/A	N/A	N/A
BALANCE SHEET				
Fixed Assets	53,440	41,232	40,492	39,75
Intangible Assets	49,566	36,508	35,768	35,02
Tangible Assets	3,563	3,290	3,290	3,29
Investments	311	1,434	1,434	1,43
Current Assets	85,366	36,393	50,659	22,14
Stocks	0	0	0	,
Debtors	1,359	233	233	23
Cash (cash plus deposits)	82,587	33,905	48,171	19,66
Other	1,420	2,255	2,255	2,25
Current Liabilities	(11,275)	(7,944)	(7,944)	(7,944
Creditors	(9,960)	(7,509)	(7,509)	(7,509
Deferred revenue	(4,533)	0	0	(.,
Walloon loans and bank loan	(1,315)	(435)	(435)	(435
Long Term Liabilities	(36,646)	(22,146)	(22,146)	(22,146
Loans (non-current) Bank and Walloon	(7,866)	(1,870)	(1,870)	(1,870
Other long term liabilities	(28,780)	(20,276)	(20,276)	(20,276
Net Assets	90,885	47,535	61,061	31,81
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CASH FLOW	(20,005)	(46.007)	(OC E14)	(07.474
Operating Cash Flow	(26,695)	(46,027)	(26,514)	(27,471
Net Interest	1,997	(3,521)	264	(29
Tax	0 (4.700)	0 (0.50)	(4.040)	
Capex	(1,782)	(858)	(1,010)	(1,010
Acquisitions/disposals	(1,561)	0	U 46.440	
Financing	0	625	46,140	
Dividends Others	0	0	(4.044)	
Other	3,109	1,099	(4,614)	(00.54)
Net Cash Flow	(24,932)	(48,682)	14,266	(28,510
Opening net debt/(cash)	(96,131)	(73,406)	(31,600)	(45,866
HP finance leases initiated	0	0	0	
Loan and finance movements	2,207	6,876	(0)	(47.050
Closing net debt/(cash)	(73,406)	(31,600)	(45,866)	(17,356



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