

# **Kiadis Pharma**

# Price

€10.83

€152m

Pharma & biotech

# Market cap



#### **Share details**

Code	KDS
Shares in issue	14.0m
Net cash (€m) at 30 June <sup>‡</sup> 2016 (excluding Hospira debt)	16.2

#### **Business description**

Kiadis Pharma is a Dutch biotech company focused on cell-based immunotherapies to overcome complications associated with stem-cell transplants in blood diseases. Lead product ATIR101 for leukaemia is undergoing a Phase II trial and will file for EU approval in Q117. ATIR201 is in preclinical stage and has potential for thalassemia. A Phase I/II study will start in H216.

#### Bull

- Conditional EMA approval for ATIR101 in leukaemia possible in Q118.
- Theralux platform could have broader use in HSCT despite underlying cause of disease.
- Sufficient cash to fund operations until early 2018.

### Bear

- Full Phase III study needed in US for ATIR101 approval.
- Cash burn to increase significantly over 2017 and 2018.
- ATIR201 still at the preclinical stage.

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Kiadis Pharma is a client of Edison Investment Research Limited

# A smart approach to stem cell transplantation

Kiadis Pharma is developing T cell-based therapies to address the issues associated with haematopoietic stem cell transplantation (HSCT). The company is leveraging its Theralux technology to develop ATIR101 and ATIR201 as adjunct therapies to HSCT in leukaemia and thalassemia, respectively. On the back of Phase II data, Kiadis is aiming for accelerated filing of ATIR101 with the European Medicines Agency (EMA) in Q117. A Phase III trial will start in H216. ATIR201 will start a Phase I/II trial in H216. Cash at end June 2016 was €23.7m, sufficient to fund operations until early 2018. We value the company at €327.3m or €27.1/share.

### Looking to a fast path to market

Following ATIR101 Phase II data, Kiadis has decided to file for conditional approval with the EMA in Q117, setting a potential approval date in Q118. We believe it is possible that Kiadis may get approval given the precedent set by MolMed, which recently received Conditional Marketing Authorisation (CMA) from the European Commission on data from a small Phase II study that showed a one-year survival rate of 49% vs 37% for historical control. The regulatory pathway is also clear in the US; the Phase III primary endpoint and active comparator arm have been defined after an end of Phase II meeting with the FDA. This Phase III trial is needed for full approval in both the US and EU.

### A smart approach in a rapidly growing market

Kiadis's Theralux platform is a photodynamic system that removes donor cells that are reactive to the host's immune cells and may cause complications after HSCT, thereby providing immunological support post-transplantation without increasing the risk of graft vs host disease (GVHD). This allows for a better response to tumour (graft vs leukaemia effect), reduces opportunistic infections, diminishes treatment-related mortality (TRM) and prolongs survival. The EMA has granted orphan drug designation expansion to ATIR101 for its use in HSCT regardless of the underlying cause of disease, expanding its market potential.

# Valuation: Starting with an rNPV of €327.3m

We estimate a risk-adjusted value of €27.1/share, using a 12.5% discount rate. We assume ATIR101 will be approved in the EU in Q118 and sales will ramp up quickly to combined peak sales of \$501m. We estimate a 70% probability to reach the market in the EU and 50% in the US, where a full Phase III study is needed for approval. Our valuation includes net cash of €7.7m at end of June 2016.

Edison estimates							
Year end	Revenue (€m)	PBT* (€m)	EPS* (€)	DPS (€)	P/E (x)	Yield (%)	
12/14	0.00	(7.21)	(0.07)	0.0	N/A	N/A	
12/15	0.00	(17.35)	(0.14)	0.0	N/A	N/A	
12/16e	0.00	(9.99)	(80.0)	0.0	N/A	N/A	
12/17e	0.00	(13.47)	(0.11)	0.0	N/A	N/A	

Source: Company data, Edison Investment Research



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