

August 12, 2021

EMERGING COMPANY
SPECULATIVE BUY (no change)

Stock code:	ANP AU
Price:	A\$0.185
12-month target price:	A\$0.44
Previous target price:	A\$0.44
Up/downside to target price:	137.8%
Dividend yield:	0.00%
12-month TSR*:	120.3%
Market cap:	A\$106m
Average daily turnover:	A\$0.33m
Index inclusion:	N/A

* Total stock return – Up/downside to target price + 12-month forward dividend yield.

Price performance

(%)	1M	3M	12M	3Y
Absolute	0.0	21.2	90.5	769.6
Rel ASX/S&P200	-4.0	14.7	66.7	748.8



Source: Bloomberg

Financial summary

	Jun-20A	Jun-21F	Jun-22F	Jun-23F
Revenue (A\$m)	3.6	6.3	2.8	0.0
EBITDA Norm (A\$m)	-6.2	-8.0	-22.8	-8.0
Net Profit (A\$m)	-5.2	-7.8	-22.7	-7.7
EPS Norm (A\$)	-0.01	-0.01	-0.03	-0.01
EPS Growth Norm (%)	-1846.4%	27.7%	125.3%	-70.6%
P/E Norm (x)	NA	NA	NA	NA
DPS (A\$)	0.00	0.00	0.00	0.00
Dividend Yield (%)	0.0%	0.0%	0.0%	0.0%
EV/EBITDA (x)	-16.5	-12.6	-4.0	-12.6
Gearing (Net Debt/EBITDA)	0.66	0.68	0.64	0.67

Source: Company data, Morgans estimates

Related research
[ANP \(SPEC BUY - TP A\\$0.441\) - 01 Jul 2021](#)
[ANP \(SPEC BUY - TP A\\$0.38\) - 28 Apr 2021](#)
Iain WILKIE

(61) 7 3334 4521

iain.wilkie@morgans.com.au

Scott POWER

Analyst(s) own shares in the following stocks mentioned in this report:

– Antisense Therapeutics

Antisense Therapeutics

The chicken and the egg

- ANP has received feedback from the FDA that its Fast-Track Designation (FTD) request will require resubmission post the lifting of its partial hold.
- While the announcement delays the potential issuance of a valuable short-term catalyst, it does not change any long-term timelines to the program and may in fact prompt ANP to engage and address the clinical hold sooner, in our view.
- Receipt of this designation, while beneficial to the long-term value of a drug asset is meaningless unless you have the approval to commence the trial, so a push to address the hold and achieve acceptance from the FDA on these higher doses is necessary and will allow ANP to progress its US trials to align with ANP's global regulatory plans.
- While the delay of the FTD may create short-term weakness as traders move to more immediate-term catalysts, we view any weakness as an opportunity to continue to add to holdings.
- We remain positive in our view of the stock. No changes to our base case valuation and price target of A\$0.44. Speculative Buy recommendation retained.

Event: Fast-track submission requires partial hold to be lifted

- ANP has received feedback from the FDA on its FTD request which notes the requirement to submit its revised clinical and toxicology protocols as part of its complete response letter (CRL) to lift the partial hold on the ATL-1102 drug for use above 25mg/week.
- The technicality arose as its submission includes trial protocols with higher and longer administration levels beyond the limits of current restrictions.
- With the hold limiting dosage to US patients to 25mg/week paired with the current thinking on trial design to be split into 3 groups including a higher dose arm (placebo / 25mg / >25mg), ANP were always going to be required to address this issue prior to commencing the studies in the US.
- Overall, no impact to the timing of the trial commencement or completion although it does push out a price catalyst which investors would view favourably if received.

No changes to timing for overall program

- Whether ANP received an FTD this month or 12 months from now doesn't really matter as the receipts are used post clinical trial completion and are designed to facilitate and expedite the review of a new drug application and not shorten the clinical trial itself.
- We would expect an update on timing in the coming months although anticipate timing on this to line up with the lodgment of its nine month toxicology protocols which we expect by the end of CY21.

Forecast and valuation update:

- No changes to forecasts or valuation.

Investment view: using short-term weakness as another entry point

- We continue to be buyers in this name. Any weakness would present an opportunity to continue to add to holdings.
- While considerable trial risk and hurdles remain, we view ANP as one of the best risk/reward plays in the healthcare space given the data produced to date and heading into a catalyst rich 12-24 month period.

Risks:

- Prolonged delays in trial recruitment and commencement.
- Failure of DMD in Ph2b program.
- Funding requirements.

Antisense Therapeutics

as at August 12, 2021

Rating	SPECULATIVE BUY	Price (A\$):	0.185
Market cap (A\$m):	106	12-month target price (A\$):	0.44
Shares outstanding (m):	574.0	Up/downside to target price (%):	137.8
Free float (%):	100.0	Dividend yield (%):	0.00

Company description

Antisense Therapeutics Limited, a biopharmaceutical company, engages in the research and development of novel antisense pharmaceuticals in Australia. Its product pipeline comprises ATL1102, an antisense inhibitor of CD49d that has completed Phase IIa for the treatment of multiple sclerosis, Duchennes Muscular Dystrophy, acromegaly, asthma, and other inflammatory indications. The company's product pipeline also includes ATL1103, a second generation antisense drug designed to block growth hormone receptor expression thereby reducing levels of the hormone insulin-like growth factor-I in the blood, as well as to treat diseases associated with excessive growth hormone action that has completed Phase II clinical trial.

Market considerations for ATL1102	Near-term milestones (CY21)
<div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <p>ANTI-INFLAMMATORY Anti-inflammatory Therapeutics Market^a is expected to garner US\$106.1 billion by 2020 (Allied Market Research) <small>^aMS, Arthritis, Psoriasis, Respiratory, IBD</small></p> </div> <div style="width: 45%;"> <p>CORTICOSTEROIDS The global steroid market is forecast to attain the value of US\$17 Billion by the end of 2025 (QV Research)</p> </div> </div> <div style="margin-top: 10px;"> <p>DMD THERAPIES The global DMD drug market is expected to reach over US\$4 Billion by 2023 (Grand View Research)</p> </div>	<ul style="list-style-type: none"> * ATL1102 drug product manufacture - ACHIEVED * ATL1102 in new indications - IN PROGRESS * US FDA DMD progress - IN PROGRESS * ATL1102 DMD Ph2b EU trial prep and initiation - IN PROGRESS

SOURCE: ANTISENSE THERAPEUTICS

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Product pipeline	ATL1102 mechanism of action																												
<table border="1"> <thead> <tr> <th>PRODUCT</th> <th>INDICATION</th> <th>RESEARCH</th> <th>PRECLINICAL</th> <th>PHASE I</th> <th>PHASE II</th> <th>PHASE III</th> </tr> </thead> <tbody> <tr> <td>ATL1103 s.c. injection</td> <td>Acromegaly</td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> </tr> <tr> <td>ATL1102 s.c. injection</td> <td>Multiple Sclerosis</td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> </tr> <tr> <td>ATL1102 s.c. injection</td> <td>DMD</td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> <td style="background-color: red;"></td> </tr> </tbody> </table>	PRODUCT	INDICATION	RESEARCH	PRECLINICAL	PHASE I	PHASE II	PHASE III	ATL1103 s.c. injection	Acromegaly						ATL1102 s.c. injection	Multiple Sclerosis						ATL1102 s.c. injection	DMD						<p>Mechanism of translation inhibition by ASOs</p> <ul style="list-style-type: none"> • ASO binds near start codon • ASO sterically blocks translation initiation machinery or ribosome
PRODUCT	INDICATION	RESEARCH	PRECLINICAL	PHASE I	PHASE II	PHASE III																							
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SOURCE: ANTISENSE THERAPEUTICS

SOURCE: CureFFI.org

MARKET DATA	#	Key Drivers
Population of target market ('000s)	48.0	Licensing deal value for late stage assets
Regulatory approval weight	25.0%	Potential for early commercialisation
Non-ambulant population	50.0%	Key risks:
Number of Cases Forecast for Year 1 ('000s)	6.0	Timing / execution risks
Annual Population Growth	0.70%	Trial risks
Peak Market Penetration	50.0%	Alternative therapies
Revenue Per Unit (\$US)	\$ 150,000	COVID-19 related impact
Market Ramp Time to Peak Penetration (Years)	5	Funding risk
Hold peak	10	
Life cycle of drug	20	
Royalty Rate	20.0%	

SOURCE: MORGANS

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Figure 1: Financial summary

Income statement	2019A	2020A	2021F	2022F	2023F	Closing price (A\$)	0.185	Price target (A\$)	0.44			
Milestone payments	0.0	0.0	0.0	0.0	0.0	Valuation metrics						
Royalty	0.0	0.0	0.0	0.0	0.0	Methodology -DCF-PER Comp		Target Price	\$0.44			
R&D rebate	3.0	3.6	6.3	2.9	0.0	DCF valuation inputs						
Total revenue	3.0	3.6	6.3	2.9	0.0	Rf	3.50%					
EBITDA	0.0	-6.2	-8.0	-22.8	-8.0	Rm-Rf	6.00%					
Associate income	0.0	0.0	0.0	0.0	0.0	Beta	1.53					
Depreciation	0.0	0.0	0.0	0.0	0.0	CAPM (Rf+Beta(Rm-Rf))	12.7%					
EBITA	0.0	-6.2	-8.0	-22.8	-8.0	E/EV*Ke+D/EV*Kd(1-t)		NPV cash flow (A\$m)	249.7			
Amortisation/impairment	0.0	0.0	0.0	0.0	0.0	Equity (E/EV)	100.0%	Minority interest (A\$m)	0.0			
EBIT	0.0	-6.2	-8.0	-22.8	-8.0	Debt (D/EV)	0.0%	Net debt (A\$m)	-5.5			
EBIT(incl associate profit)	0.0	-6.2	-8.0	-22.8	-8.0	Interest rate	5.00%	Investments (A\$m)	0.0			
Net interest expense/FX	0.0	0.3	1.0	-0.2	-0.1	Tax rate (t)	30.0%	Equity market value (A\$m)	255.2			
Pre-tax profit	0.3	-5.2	-8.2	-22.9	-8.3	WACC	12.7%	Diluted no. of shares (m)	574.0			
Income tax expense	0.0	0.0	0.0	0.0	0.0			DCF valuation	\$0.44			
After-tax profit	0.3	-5.2	-8.2	-22.9	-8.3							
Minority interests	0.0	0.0	0.0	0.0	0.0	Multiples						
NPAT	0.3	-5.2	-8.2	-22.9	-8.3	Enterprise value (A\$m)	103.3	2019A	102.1	2021F	2022F	2023F
Significant items	0.0	0.0	0.0	0.0	0.0	EV/Sales (x)	n.a.	n.a.	n.a.	n.a.	n.a.	n.a.
NPAT post abnormals	0.3	-5.2	-8.2	-22.9	-8.3	EV/EBITDA (x)	n.a.	-16.5	-12.6	-4.0	-12.6	
						EV/EBIT (x)	-108648.8	-16.5	-12.6	-4.0	-12.5	
						PE (pre-goodwill) (x)	298.0	-17.5	-13.6	-6.5	-19.0	
						PEG (pre-goodwill) (x)	1.2	0.0	0.0	0.1	0.1	
Cash flow statement	2019A	2020A	2021F	2022F	2023F	At target price	2019A	2020A	2021F	2022F	2023F	
EBITDA	0.0	-6.2	-8.0	-22.8	-8.0	EV/EBITDA (x)	-108648.8	-16.5	-12.6	-4.0	-12.5	
Other cash items	0.0	0.0	0.0	0.0	0.0	PE (pre-goodwill) (x)	716.2	-42.1	-32.7	-15.6	-45.7	
Net interest (pd)/rec	-0.3	-1.0	0.2	0.1	0.3							
Taxes paid	0.0	0.0	0.0	0.0	0.0	Per share data						
Change in working capital	-2.7	3.3	0.9	0.0	-1.5	No. shares	420.1	488.8	574.0	796.0	796.0	
Cash flow from ops (1)	-2.9	-4.0	-6.9	-22.7	-9.2	EPS (cps)	0.1	-1.1	-1.4	-2.9	-1.0	
Capex (2)	0.0	-0.1	0.0	0.0	0.0	EPS (normalised) (c)	0.1	-1.1	-1.4	-2.9	-1.0	
Disposals/(acquisitions)	0.0	0.0	0.0	0.0	0.0	Dividend per share (c)	0.0	0.0	0.0	0.0	0.0	
Other investing cash flow	0.0	0.0	0.0	0.0	0.0	Dividend payout ratio (%)	0.0%	0.0%	0.0%	0.0%	0.0%	
Cash flow from invest (3)	0.0	-0.1	0.0	0.0	0.0	Dividend yield (%)	0.0%	0.0%	0.0%	0.0%	0.0%	
Incr/(decr) in equity	1.5	5.2	8.3	30.0	0.0	Growth ratios						
Incr/(decr) in debt	0.0	0.0	0.0	0.0	0.0	Sales growth	n.a.	n.a.	n.a.	n.a.	n.a.	
Ordinary dividend paid	0.0	0.0	0.0	0.0	0.0	Operating cost growth	n.a.	n.a.	29.0%	185.3%	-64.8%	
Preferred dividends (4)	0.0	0.0	0.0	0.0	0.0	EBITDA growth	n.a.	n.a.	-29.1%	-185.1%	64.8%	
Other financing cash flow	0.0	0.0	0.0	0.0	0.0	EBITA growth	n.a.	n.a.	n.a.	n.a.	n.a.	
Cash flow from fin (5)	1.5	5.2	8.3	30.0	0.0	EBIT growth	n.a.	n.a.	n.a.	n.a.	n.a.	
Forex and disc ops (6)	0.0	0.0	0.0	0.0	0.0	NPAT growth	n.a.	n.a.	n.a.	n.a.	n.a.	
Incr/(decr) cash (1+3+5+6)	-1.4	1.1	1.4	7.3	-9.2	Pre-goodwill NPAT growth	n.a.	n.a.	n.a.	n.a.	n.a.	
Equity FCF (1+2+4)	-2.9	-4.1	-6.9	-22.7	-9.2	Pre-goodwill EPS growth	n.a.	n.a.	n.a.	n.a.	n.a.	
						Normalised EPS growth	n.a.	n.a.	n.a.	n.a.	n.a.	
Balance sheet	2019A	2020A	2021F	2022F	2023F	Operating performance	2019A	2020A	2021F	2022F	2023F	
Cash & deposits	2.9	4.1	5.5	14.6	5.4	Asset turnover (%)	0.0	0.0	0.0	0.0	0.0	
Trade debtors	0.6	0.7	1.0	0.5	0.0	EBITDA margin (%)	n.a.	n.a.	n.a.	n.a.	n.a.	
Inventory	0.0	0.0	0.0	0.0	0.0	EBIT margin (%)	n.a.	n.a.	n.a.	n.a.	n.a.	
Investments	0.0	0.0	0.0	0.0	0.0	Net profit margin (%)	n.a.	n.a.	n.a.	n.a.	n.a.	
Goodwill	0.0	0.0	0.0	0.0	0.0	Return on net assets (%)	0.0	-136.0	-158.5	-184.5	-174.1	
Other intangible assets	0.0	0.0	0.0	0.0	0.0	Net debt (A\$m)	-2.9	-4.1	-5.5	-14.6	-5.4	
Fixed assets	0.0	0.1	0.1	0.1	0.1	Net debt/equity (%)	-103.3	-89.1	-108.4	-118.1	-116.8	
Other assets	0.0	0.0	0.0	0.0	0.0	Net interest/EBIT cover (x)	0.0	23.7	7.7	-123.9	-73.5	
Total assets	3.7	5.4	7.1	15.7	6.0	Internal liquidity	2019A	2020A	2021F	2022F	2023F	
Short-term borrowings	0.0	0.0	0.0	0.0	0.0	Current ratio (x)	3.3	5.1	2.6	4.4	3.9	
Trade payables	0.6	0.3	1.6	2.8	0.9	Receivables turnover (x)	0.0	0.0	0.0	0.0	0.0	
Long-term borrowings	0.0	0.0	0.0	0.0	0.0	Payables turnover (x)	0.0	14.7	8.6	10.4	4.4	
Provisions	0.3	0.5	0.5	0.5	0.5							
Other liabilities	0.0	0.0	0.0	0.0	0.0							
Total liabilities	0.9	0.8	2.1	3.3	1.4							
Share capital	63.9	69.1	77.4	107.4	107.4							
Other reserves	0.0	2.4	2.4	2.4	2.4							
Retained earnings	-61.1	-67.0	-74.8	-97.5	-105.3							
Other equity	0.0	0.0	0.0	0.0	0.0							
Total equity	2.8	4.6	5.0	12.4	4.6							
Minority interest	0.0	0.0	0.0	0.0	0.0							
Total shareholders' equity	2.8	4.6	5.0	12.4	4.6							
Total liabilities & SE	3.7	5.4	7.1	15.7	6.0							

Source: Morgans estimates, company data

Queensland		New South Wales		Victoria		Western Australia	
Brisbane	+61 7 3334 4888	Sydney	+61 2 9043 7900	Melbourne	+61 3 9947 4111	West Perth	+61 8 6160 8700
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Regulatory disclosures

Analyst owns shares in the following mentioned company(ies): Antisense Therapeutics

Morgans Corporate Limited was Joint Lead Manager to the Placement and Share Purchase Plan of shares in Antisense Therapeutics Limited in November 2020 and received fees in this regard.

Recommendation structure

For a full explanation of the recommendation structure, refer to our website at morgans.com.au/research_disclaimer

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Research independence statement

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