



Research Update

BlinkLab Ltd (ASX:BB1)

Pilot study confirms path to commerciality

Price: \$0.92 | Valuation: \$1.90 | Implied Return: 106% | 28 January, 2026

Recent Developments

Blinklab's pilot study was unblinded in October 2025 and gave us a first proper look at how well the Dx1 software functions in diagnosing autism in children.

- **Pilot Study Results:** The study involved 485 children and achieved 83.7% sensitivity and 84.7% specificity. The results places BB1 as one of the most advanced digital-diagnostic companies targeting autism. The pilot study focused on children who were either in the process of receiving, or had recently undergone, neurodevelopmental assessment at an autism center in the USA. The children ranged from 2-11 years old with the mean age being 5.98. 42.9% were female and 57.1% male. Care was taken to mimic recruitment to mirror real-world population usage of BB1's device. The pilot study was designed to be highly challenging to stress tests.
- **FDA give benchmark:** The FDA have given BB1 a benchmark for the upcoming registrational study with a minimum performance threshold of at least 65% specificity and at least 65% sensitivity. The pilot study results were well above the expectations set by the FDA.
- **Technology peer reviewed:** BB1's technology has been independently validated in a peer reviewed paper and published in Autism Research, a leading journal, giving strong validation to BB1's core technology. The paper focuses on the technology, data acquisition method and the underlying neurobehavioral measures. A follow-up paper in preparation will detail BB1's machine learning models.
- **Pivotal Study to begin:** The first patient is poised to be tested in the current quarter and the trial set to conclude by Q3 '26. The total number of medical centres now onboarded is 8 with the latest being the Rush University Medical Centre in Chicago. FDA clearance is expected in Q1 2027.
- **European ADHD Study results due:** The European ADHD trial is nearing data readout with a dataset from approx. 300 participants. A total of 5 sites participated under the coordination of the Mental Care Group in the Netherlands. The dataset will underpin the upcoming USA based pivotal study for Dx2.
- **European study could lead to commercialisation:** On the back of the European ADHD study BB1 can begin proceedings to apply for CE/MDR clearance (European Union Medical Device Regulation) which opens the path for early revenue in Europe.

Valuation

We have lifted our **valuation to \$1.90** from \$1.30, as the risk from the pilot study is removed. The ASD result gives further credence to the upcoming ADHD study and commercialization potential in a market which is 5 times larger than the ASD market. Previously, we projected ADHD research to begin in later years, but an accelerated timeline could positively impact probability estimates and revenue projections. We anticipate revenue from autism to begin in earnest in FY28 and revenue from ADHD in FY29.

Company Data

Recommendation: Speculative Buy
Shares on Issue: 125.9m
Market Capitalisation: \$115.8m
Enterprise Value: \$111.8m

Board Structure

Dr Anton Uvarov: Executive Director
Brian Leedman: Non-Exec Chairman
Dr Richard Hopkins: Non-Exec Director
Dr Henk-Jan Boele: Managing Director

Major Shareholders

Yulia Uvarova: 7.44%
Cason Holding BV: 5.41%
Inacea Holding BV: 4.61%
Bello Holding BV: 4.61%
Total Top 20: 46.7%

Chart



Source: ASX

BB1 is on the verge of commercialising a smartphone app which aids in diagnosing ASD, ADHD, and other neurodevelopmental conditions.

What do the pilot results really mean

It's worthwhile reminding what sensitivity and specificity mean and how it applies to ASD cases. Sensitivity refers to the test's ability to correctly identify individuals with ASD, while specificity relates to its ability to correctly identify those without ASD. Since diagnosis can at times be subjective and two expert clinicians can, at times, disagree, specificity and sensitivity are not likely to ever be 100%. Thus, a digital biomarker trained on that label can't be perfect either.

BB1's pilot study has shown specificity and sensitivity in line or better than other diagnostic and screening tools and if replicated in the pivotal study will enable Blinklab to begin path to commercialization.

<u>Tool</u>	<u>Sensitivity</u>	<u>Specificity</u>	
Dx 1	83.7	84.7	Screening
ADOS	80-90	70-85	gold standard diagnostic
ADOS-2	89-92	81-85	gold standard diagnostic
CARS DSM-IV	86	79	Diagnostic
CARS-2	90	75	Diagnostic
SRS	85-90	75-90	Screening

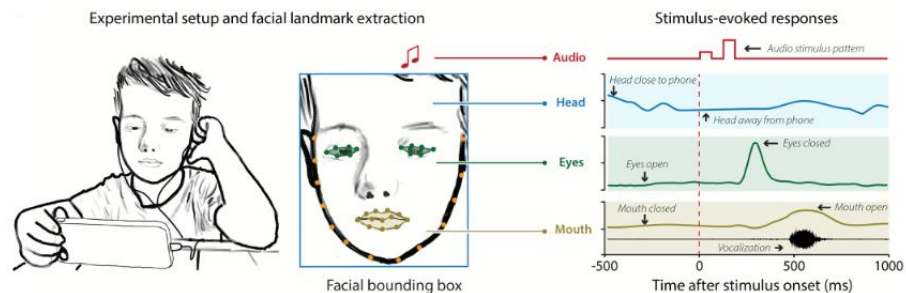
BB1's Dx1 compared to other screening/diagnostic tools. Source: Lodge/BB1

Path to commercialisation

While commercialization hinges on FDA clearance, we look forward to what it may incorporate. Initially, BB1 would target the autism centres which participated in the trials with the value proposition being a 30 minute evaluation test as compared to a 3-8 hour clinical evaluation, thus saving time, money and cuts down on long autism waiting lists. We envisage cost per Dx1 test to be around US\$250, potentially as a subscription model for each centre or enterprise licensing. We believe other target customers will be pediatricians, GP's, psychologists, telehealth platforms, and large health systems seeking a scalable screening tool. Being a smartphone-based screening tool there is potential to rapidly scale.

The cost per test compares favourably to other tests, below showing the typical cost for ADOS-2 (Autism Diagnostic Observation Schedule) which is a standardized assessment tool through structured observations and interactions. Below figures in US dollars.

Infographic of Dx1 in action.
Source: Neurobehavioral Assessment of Sensorimotor Function in Autism Using Smartphone Technology, published in Autism Research, 2026





- Private Practice: \$800-\$2,500 for comprehensive evaluation including ADOS-2.
- Hospital-Based Clinics: \$1,000-\$3,000 depending on additional assessments included.
- University Training Clinics: \$400-\$1,000 with supervised graduate students.
- Community Mental health Centers: \$200-\$800 with sliding scale options available.

Blinklab expects the pivotal study to completed by end of this year with FDA clearance expected in the first quarter of 2027. Should FDA clearance be attained we expect a commercialisation strategy to begin very rapidly.

A US based pivotal study for Dx 2 on ADHD patients is poised to begin early 2027 with onboarding of clinical trial sites in the 3rd quarter 2026. Submission to the FDA is expected late 2027 with a response for Dx 2 in Q1 2028.

Pivotal Study BB1 Dx1 - ASD	Timeline
Finish onboarding of US clinical sites	4Q 2025
Start pivotal study (first patient)	1Q 2026
Complete Pivotal Study	3Q 2026
Submit to FDA	4Q 2026
FDA response on 510(k)	1Q 2027

Pivotal Study BB1 Dx2 - ADHD	Timeline
Finish onboarding of US clinical sites	3Q 2026
Start Dx2 pivotal study (first patient)	1Q 2027
Complete Pivotal Study	3Q 2027
Submit to FDA	4Q 2027
FDA response on 510(k)	1Q 2028

Commercialisation risk and Mitigation

<u>Risk</u>	<u>Impact</u>	<u>Mitigation</u>
Pivotal trial underperformance	Delays FDA clearance	Strong pilot data reduces risk; multi-site design improves robustness
Slow payer adoption	Slower revenue ramp	Build economic evidence; target private pay initially
Competition from other digital diagnostics	Market share pressure	BlinkLab’s involuntary-response biomarkers are differentiated
Regulatory changes	Compliance burden	Early FDA alignment reduces uncertainty

Company Overview

BlinkLab (ASX:BB1) is developing a smartphone-based application to facilitate the diagnosis of neurodevelopmental conditions such as Autism Spectrum Disorder (ASD) and Attention Deficit Hyperactivity Disorder (ADHD) in children. The FDA registrational study is currently underway with the first patient is poised to be tested in the current quarter and is designed as a multi-site, prospective, double-blinded, within-subject comparison.

Year ending June	2024A	2025A	2026E	2027E
NPAT	(1.8)	(5.7)	(5.6)	(5.9)
EPS_{adj} (¢)	(0.0)	(0.1)	(0.1)	(0.1)
EPS growth		0.0%	0.0%	0.0%
P/E ratio	N/A	N/A	N/A	N/A
Enterprise Value (m)	21	15	20	17
EV/Sales (x)	562.19 x	0.00 x	0.00 x	0.00 x
EV / EBIT (x)	-11.9 x	-2.6 x	-3.5 x	-2.9 x
EV / EBITDA (x)	-11.9 x	-2.6 x	-3.5 x	-2.9 x
DPS (\$)	0.00	0.00	0.00	0.00
Dividend Yield	0.0%	0.0%	0.0%	0.0%
Payout Ratio	0.0%	0.0%	0.0%	0.0%
Franking	N/A	N/A	N/A	N/A
FCFPS (¢)	(0.02)	(0.06)	(0.06)	(0.06)
P/FCFPS	(13.28)	(4.17)	(4.24)	(4.03)

Year ending June	2024A	2025A	2026E	2027E
Operating revenue	0.0	0.4	0.6	0.3
EBITDA	(1.8)	(5.7)	(5.6)	(5.9)
D&A	0.0	0.0	0.0	0.0
EBIT	(1.8)	(5.7)	(5.6)	(5.9)
Net interest income	0.0	0.0	0.0	0.0
NPBT	(1.8)	(5.7)	(5.6)	(5.9)
Tax Expense (benefit)	0.0	0.0	0.0	0.0
NPAT	(1.8)	(5.7)	(5.6)	(5.9)
Significant Items	0.0	0.0	0.0	0.0
NPAT	(1.8)	(5.7)	(5.6)	(5.9)
EBITDA Margin	N/A	N/A	N/A	N/A
EBIT Margin	N/A	N/A	N/A	N/A
NPAT Margin	N/A	N/A	N/A	N/A

Year ending June	2024A	2025A	2026E	2027E
Bank Balance	3.0	8.7	4.1	6.4
Receivables	0.3	0.4	0.3	0.3
Inventories	0.0	0.0	0.0	0.0
Other	3.00	0.00	0.00	0.00
Current assets	6.31	9.10	4.41	6.74
Net PPE	0.0	0.0	0.0	0.0
Intangibles	0.35	0.42	0.42	0.42
Right of use assets	0.16	0.10	0.10	0.10
Non-current assets	0.56	0.56	0.56	0.56
Total assets	6.87	9.67	4.97	7.30
Payables	0.21	0.56	0.56	0.56
Lease liabilities	0.17	0.13	0.13	0.13
Employee benefits	0.00	0.00	0.00	0.00
Total liabilities	0.38	0.70	0.70	0.70
NET ASSETS	6.49	8.96	4.27	6.60

Year ending June	2024A	2025A	2026E	2027E
Net Debt	(3)	(9)	(4)	(6)
NTA	6.49	8.96	4.27	6.60
Price / NTA (x)	0.037 x	0.027 x	0.056 x	0.036 x
Return on assets	-26.1%	-59.1%	-113.0%	-81.0%
Return on equity	-27.6%	-63.6%	-43.6%	-69.1%

Year ending June	2024A	2025A	2026E	2027E
Sales Revenue	0.04	0.38	0.65	0.29
Other	-	-	-	-
Group Revenue	0.04	0.38	0.65	0.29

Year ending June	2024A	2025A	2026E	2027E
Discounted Cash Flow		WACC		15.00%
		Discount Period		6 years
		DCF/Price Target		\$ 1.90

Rating

BUY – anticipated stock return is greater than 10%

SELL – anticipated stock return is less than -10%

HOLD - anticipated stock return is between -10% and +10%

SPECULATIVE BUY – high risk stock with price likely to fluctuate by 50% or more and anticipated return is greater than 10%

Disclaimer

In accordance with section 949A of the Corporations Act 2001, any recipient of the information contained in this document should note that information is general advice in respect of a financial product and not personal advice. Accordingly the recipient should note that: (a) the advice has been prepared without taking into account the recipient's objectives, financial situations or needs; and (b) because of that, the recipient should, before acting on the advice consider the appropriateness of the advice, having regard to the recipient's objectives, financial situation and needs.

Although Lodge Partners Pty Ltd ("Lodge") consider the advice and information contained in the document is accurate and reliable, Lodge has not independently verified information contained in the document which is derived from publicly available sources. Lodge assumes no responsibility for updating any advice or recommendation contained in this document or for correcting any error or admission which may become apparent after the document has been issued. Lodge does not give any warranty as to the accuracy, reliability or completeness of advice or information which is contained in this document. Except in so far as liability under any statute cannot be excluded, Lodge, its employees and consultants do not accept any liability (whether arising in contract, in tort or negligence or otherwise) for any error or omission in this document or for any resulting loss or damage (whether direct, indirect, consequential or otherwise) suffered by the recipient of this document or any other person.

General Securities Advice Warning

This report is intended to provide general securities advice. In preparing this advice, Lodge did not take into account the investment objectives, the financial situation and particular needs of any particular person. Before making an investment decision on the basis of this advice, you need to consider, with or without the assistance of a securities adviser, whether the advice is appropriate in light of your particular investment needs, objectives and financial circumstances.

Disclosure of Interests

Lodge directors, consultants and advisers currently hold less than 1% of total shares in the Company and may buy or sell the shares from time to time. Lodge has earned and will continue to earn broking commissions by acting for individual clients that are buying or selling their shares in company.

The Company currently is, or in the past 12 months was, a client of Lodge Partner's affiliated company and authorised representative Lodge Corporate Pty Ltd. During this period, Lodge Corporate provided corporate services to the company. In the past 12 months, Lodge Corporate have received compensation for corporate services from the company. Part of this was commissioned research which includes this research note. Lodge Corporate intends to seek or expect to receive compensation for Investment Banking services from the Company in the next twelve months.

Analyst Verification

I verify that I, Sven Restel, have prepared this research report accurately and that any financial forecasts and recommendations that are expressed are solely my own personal opinions.

Contact Lodge Partners

Lodge Partners Pty Ltd

ABN: 25 053 432 769 AFSL: 246271

Melbourne office:

Level 40, 80 Collins Street, Melbourne VIC 3000

T +61 3 9200 7000

lodgepartners.com.au

Sydney office:

Suite 14.04, Level 14, 109 Pitt Street

Sydney, NSW, 2000

T +61 2 9225 4433