Deltex Medical Group plc

Half year results to 30 June 2023

September 2023

Andy Mears

Chief Executive Officer

Natalie Wettler

Group Finance Director



Real-time oesophageal Doppler haemodynamic monitoring:

improves patient outcomes; increases hospital throughput

Disclaimer

The information contained in this document ("Presentation") and the presentation made to you verbally has been prepared by Deltex Medical Group Plc (the "Company"). Deltex Medical Group Plc is a company quoted on AlM, a market operated by London Stock Exchange plc. This Presentation has not been fully verified and is subject to material updating, revision and further verification and amendment without notice. This Presentation has not been approved by an authorised person in accordance with Section 21 of the Financial Services and Markets Act 2000 (as amended) ("FSMA") and therefore it is being provided for information purposes only.

While the information contained herein has been prepared in good faith, neither the Company nor any of its directors, officers, agents, employees or advisers give, have given or have authority to give, any representations or warranties (express or implied) as to, or in relation to, the accuracy, reliability or completeness of the information in this Presentation, or any revision thereof, or of any other written or oral information made or to be made available to any interested party or its advisers (all such information being referred to as "Information") and liability therefore is expressly disclaimed. Accordingly, neither the Company nor any of its directors, officers, agents, employees or advisers take any responsibility for, or will accept any liability whether direct or indirect, express or implied, contractual, tortious, statutory or otherwise, in respect of, the accuracy or completeness of the Information or for any of the opinions contained herein or for any errors, omissions or misstatements or for any loss, howsoever arising, from the use of this Presentation.

The views of the Company's management/directors and/or its partners set out in this document could ultimately prove to be incorrect. No warranty, express or implied, is given by the presentation of these figures herein and investors should place no reliance on the Company's estimates cited in this document.

This Presentation may contain "forward-looking statements" that involve substantial risks and uncertainties, and actual results and developments may differ materially from those expressed or implied by these statements. These forward-looking statements are statements regarding the Company's intentions, beliefs or current expectations concerning, among other things, the Company's results of operations, performance, financial condition, prospects, growth, strategies and the industry in which the Company operates. By their nature, forward-looking statements involve risks and uncertainties because they relate to events and depend on circumstances that may or may not occur in the future. These forward-looking statements are not guarantees of future performance of the Company and reflect assumptions and subjective judgements by the Company that are difficult to predict, qualify and/or quantify. These forward-looking statements speak only as of the date of this Presentation and the Company does not undertake any obligation to publicly release any revisions to these forward-looking statements to reflect events or circumstances after the date of this Presentation.

This Presentation should not be considered as the giving of investment advice by the Company or any of its directors, officers, agents, employees or advisers. In particular, this Presentation does not constitute or form part of any offer or invitation to subscribe for or purchase any securities and neither this Presentation nor anything contained herein shall form the basis of any contract or commitment whatsoever. No reliance may be placed for any purpose whatsoever on the information or opinions contained in these slides or the Presentation or on the completeness, accuracy or fairness thereof. In particular, any estimates or projections or opinions contained herein necessarily involve significant elements of subjective judgment, analysis and assumptions and each recipient should satisfy itself in relation to such matters.

The distribution of this document in or to persons subject to jurisdictions outside the UK may be restricted by law and persons into whose possession this document comes should inform themselves about, and observe, any such restrictions. Any failure to comply with these restrictions may constitute a violation of the laws of the relevant jurisdiction.

Allenby Capital Limited ("Allenby Capital"), which is authorised and regulated by the Financial Conduct Authority, is acting as the nominated adviser and broker to the Company. Accordingly, the recipients should note that Allenby Capital is neither advising nor treating as a client any other person and will not be responsible to anyone other than the Company for providing the protections afforded to clients of Allenby Capital and nor for providing advice in relation to the matters contained in this Presentation.

H1 2023 – key commercial dynamics

Release of new TrueVue monitor

- Good progress made on new TrueVue monitor during H1 2023 with the device being launched on 10 July into the UK in EU
- Evaluations have now commenced in UK hospitals.
 This process is expected to last six weeks
- Revenues from TrueVue monitor expected to commence in November 2023 from UK and EU customers

Latin American Tender

- Continuing to work on potentially significant Latin America tender order
- Tender was due to start in May 2023, but delayed due to government not releasing funding
- Funding released at end of August and hospitals now starting to select equipment from national tender

Restricted access in UK/US: "new normal"

- Modified our commercial plans, assuming that 'restricted access' to hospitals for DEMG's sale teams is the "new normal"
- Commercial activities have been focused on selling the new TrueVue monitor to existing customers to generate capital pipeline and revenues
- Mitigating effect using targeted digital/remote initiatives: e.g. online training academy that was launched in 2022 now in regular use

Non-invasive Suprasternal device

- Progress on developing the new non-invasive device was hampered by the Company's financial position, but is now moving forward
- The prototype probe is due to be completed in Q4 2023 and can be used with the new TrueVue monitor
- Subject to gaining regulatory approvals, a clinical evaluation in a leading UK hospital has been approved to commence this year

Deltex Medical

Summary P&L information to 30 June 2023

£000 (unaudited)	H1 20	023	H1 2022	
Probe revenues	815	77%	912	79%
Other revenues	244	23%	246	21%
Total revenues	1,059	100%	1,158	100%
Gross profit	728	69%	852	74%
Administrative costs	(642)		(779)	
Sales & distribution costs	(427)		(554)	
R&D and Q&R costs	(116)		(120)	
Total costs	(1,185)		(1,453)	
Adjusted EBITDA*	(361)		(418)	
Loss for the period	(537)		(662)	



Revenues reduced by 9% to £1.1M - offset by cost reductions of 18% to £1.2m



Summary balance sheet information at 30 June 2023

£000	30 June 2023	30 June 2022		30 June 2023	30 June 2022
Property, plant &	237	274	Borrowings	(1,147)	(700)
equipment			Trade & other payables	(1,744)	(1,419)
Intangible assets	3,986	3,419	Current liabilities	(2,891)	(2,119)
Financial assets at amortised cost	159	171	Borrowings	(998)	(1,048)
Non-current assets	4,382	3,864	Trade & other payables	(148)	(203)
Inventories	824	835	Provisions	(67)	(60)
Trade receivables	440	540	Non-current liabilities	(1,213)	(1,311)
	-		Total liabilities	(4,104)	(3,430)
Other current assets	191	206			
Cash	107	611			
Current assets	1,562	2,192			
Total assets	5,944	6,056	Net assets	1,840	2,626

H1 2023 trading performance and fundraise summary

- Having made a solid start to the year, trading performance in Q2 2023 was weaker than expected due to unexpected delays to both the launch of the new monitor and in the award of a national tender in Latin America
- Notwithstanding all loan facilities being fully drawn, cashflows were under severe pressure with the Company's cash position at £0.1 million as at 30 June 2023 creating an immediate need for additional capital
- > Subsequently a fundraise was completed in August 2023, raising new cash for the business of £1.89 million and £350,000 debt converted to equity to strengthen the Company's balance sheet
- Use of funds raised include:
 - headcount reduction equating to cost-savings of c.£1.0 million p.a.
 - focusing on generating revenues from existing single use probes to seek positive "real" EBITDA
 - fund the market launch of the next generation TrueVue monitor
 - fund further development of the non-invasive Suprasternal device
- Restructuring of £1.0 million short-term debt due to Imperialise Limited (controlled by Nigel Keen):
 - £0.35 million converted into equity
 - Repayment of £0.25 million postponed to 30 June 2025
 - Remaining £0.4 million postponed to 31 December 2025

Next generation monitor - TrueVue



- Combines independent measurements of blood flow and blood pressure across each heartbeat in real time
- Immediate assessment of haemodynamic instability during surgery and in ICU patients: sepsis, heart failure etc
- Portable multi-technology device
- New monitors currently in-build at Chichester HQ.
- Evaluations have commenced in UK hospitals to ensure there are no teething issues prior to making first sales into UK and EU
- Will be the platform for the non-invasive device which has significantly broader applications: awake patients - A&E, wards, paramedics



New monitor is expected to help increase activity levels in all territories, with orders in the short term from international distributors within EU.



Developing the new, non-invasive Doppler-based haemodynamic monitoring device is a key part of the Group's future growth and long-term strategy

Next generation monitor – expected regulatory pathway



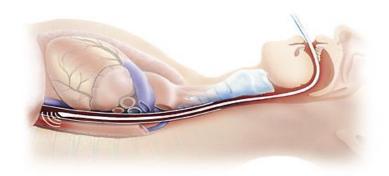


Expected release roadmap – existing distribution

- UK, EU and CE MDD/MDR regulatory approvals
 Released 10 July 2023
- USA FDA approval: 6-9 months from submission
 Expected Target H1 2024
- Latin America: 3-6 months (country dependent)
 Expected Targets H1/H2 2024
- South Korea KFDA approval: 6-9 months from submission
 Expected Target H2 2024

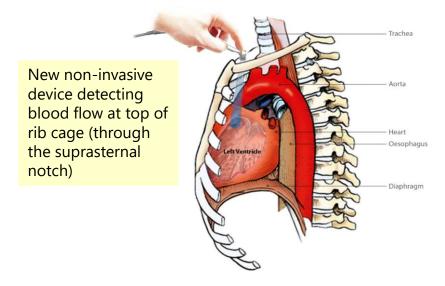


Currently in development - non-invasive Suprasternal device



ODM probe inserted down the oesophagus of a sedated patient (e.g. elective surgery) to generate high-quality data, which is used to provide rapid, optimised treatment

- High margin product, built on substantial published evidence, including 24 RCTs
- Utilises Deltex's proprietary algorithm which has been clinically proven over many years
- Existing user base in over 40 countries worldwide
- Focused on elective surgery and ventilated ICU patients



- Is an on-going R&D activity
- Quick and easy to use provides immediate snapshot of awake patient's haemodynamics
- Developed in collaboration with the UK's National Physical Laboratory
- Introduces awake patient applications (Covid, sepsis, A&E, ward)
- Complementary technology, generating an additional revenue stream

improves patient outcomes; increases hospital throughput

Conclusion and prospects





The Group has refinanced and restructured the Company based on:

- implementing cost savings of c.£1.0m p.a.
- focusing on generating positive monthly EBITDA at high gross margins by the end of the year
- launching the new, next generation TrueVue monitor into UK and EU, which will increase activity levels and revenues, particularly from distributors
- continuing to grow International revenues, especially if successful in winning new business from the national tender in Latin America
- further development of the non-invasive Suprasternal device, and subject to regulatory approval, starting a clinical trial in leading UK hospital



Deltex Medical Group plc

Half year results to 30 June 2023

September 2023

Andy MearsChief Executive Officer

Natalie WettlerGroup Finance Director

