





H1 25 Trading Update

July 2025



AIM: HVO

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H1 25 Trading Update



H1 25 Key Highlights

- Revenue of £24.2m
- EBITDA margin (pre-exc.) c.12%
- Cash of £23.3m
- Weighted orderbook of £40m
- Strong sales pipeline, longer sales cycles
- Good progress with new revenue streams
- CRS & Cryostore acquisitions completed
- Integrations on track
- FY25 revenue guidance £47m, low-single digit EBITDA loss
- Preferred candidate for independent Non-Executive Chair

HCT

- hVIVO challenge services
- hVIVO laboratory services
- Venn Life Sciences

Consultancy

Venn Life Sciences

hLAB

- hVIVO laboratory services
- Cryostore

Clinical Services

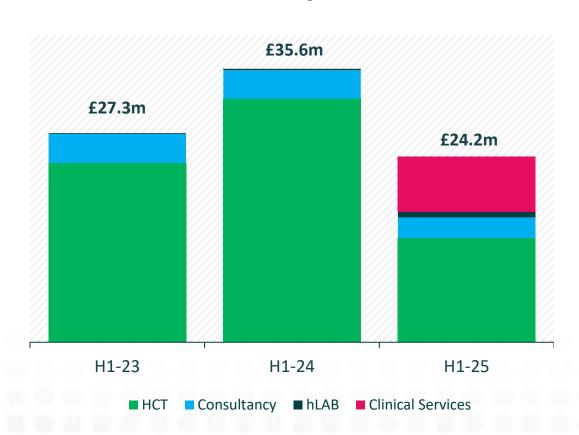
- hVIVO site services
- hVIVO recruitment services
- CRS
- Venn Life Sciences

Full-Service European Contract Research Organisation

Revenue



Revenue by Service

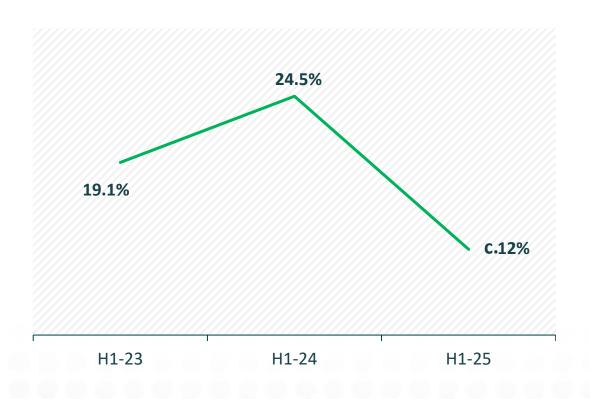


- Delivering on diversification strategy
- hLAB & Cryostore revenue growing
- Clinical services growing, CRS & hVIVO
- Acquisitions delivered £5.5m:
 - CRS £5.2m & Cryostore £0.3m since acquisition date
- Broader therapeutic areas & customer base
- Cancellation / postponement fees higher than normal
- Consultancy & HCT services lower
- Revenue guidance of £47 million for FY25

EBITDA Margin (pre-exceptional items)



EBITDA Margin

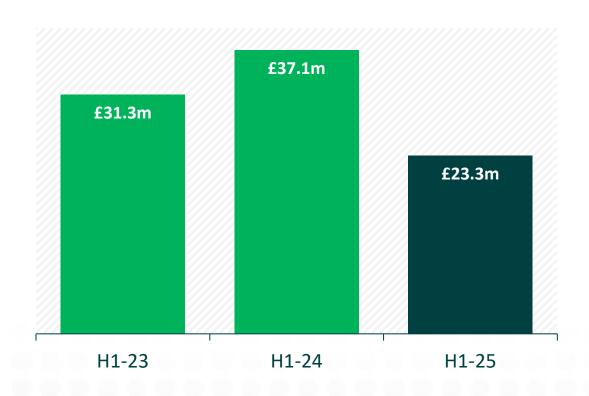


- EBITDA benefitting from the positive impact of operational efficiencies, cost management & cancellation fees
- Headcount:
 - 24% lower vs H1 24
 - Flexibility of using temporary staff
 - Efficiencies driven by consolidation of facilities
- Investments continue in automation to drive efficiencies
- Excludes acquisition & restructuring costs of ~£1.4m
- FY25 EBITDA expected to be low-single digit loss, an improvement on previous guidance

Cash



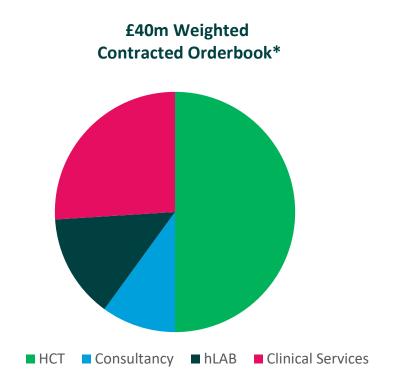
Cash



- Healthy cash balance no debt
- Cash balance impacted by:
 - Acquisitions account for ~ £14m (incl. exceptional items & related working capital movement)
 - Reduction in deferred revenue/ advance receipts due to cancellations
- Tight cost control while driving further diversification

Orderbook & Pipeline – 30 June 2025

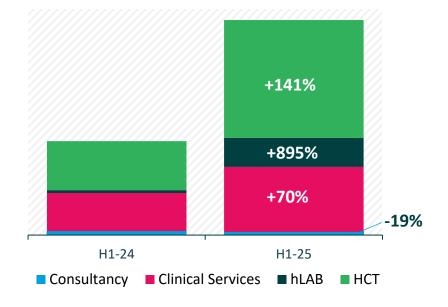






- Weighted orderbook does not include ILiAD HCT*
- Total value of proposals in H1 25 double vs H1 24
- Significant increase in hLAB interest

Proposals by Value H1 24 v H1 25 by Service (incl. CRS & Cryostore)



- Good growth in HCT & clinical services
- Good interest in new models hMPV & Flu B
- Some pipeline projects represent largest ever value HCTs
- 60% growth CRS opps, cross-selling opps CRS & Cryostore

Integration Update



Financial

- Revised pricing models
- Move to hVIVO financial system
- Staff synergies identified
- £1.1m annualised cost savings identified

CRS EXPERTS. EARLY PHASE. Cryostore

Operational

- Operational teams aligned
- Enhanced IT security
- Improved KPI monitoring & management
- Process standardisation

Business Development

- New hLAB & Cryostore BD person
- Marketing & conference attendance
- New cross-selling opportunities realised
- £2.1m Venn opportunities in CRS sales pipeline

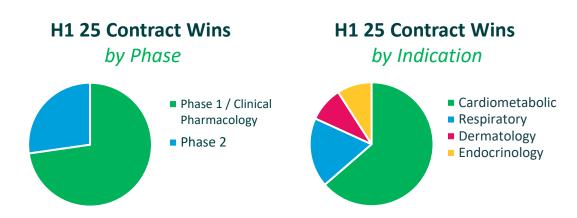
Outlook

- CRS FY25 adjusted EBITDA loss in line with expectations
- CRS expected to be earnings accretive 2026
- Cryostore earnings accretive

CRS – At a Glance



Business Progress



- Average clinical trial contract value c.€1m
- 60% increase in proposals by value H1 24 vs H1 25
- First cross-selling contracts signed with Venn Breda & Paris
- High level client meetings
- Re-established long-standing relationships with German clients

Example Indication – Obesity

- 64% of CRS contract wins H1 24 in cardiometabolic diseases
- 120 ongoing Phase 1-2 clinical trials globally¹
- 2024 record year for obesity trials globally 20% CAGR (2019-2024)²
- Over 300 GLP-1R drugs in an active stage of development³
- \$125Bn GLP-1R expected sales by 2033³
- Indication expansion beyond obesity⁴

Key Team Members



Dr Thomas Forst Chief Medical Officer



Till Mieskes
Managing Director

Enhances hVIVO's early-stage clinical capabilities

Market Trends



Headwinds

- Changes at Health & Human Services (HHS)
- FDA timelines unpredictable
- Drug pricing reforms & tariffs
- Vaccine Policy
- Depressed biotech funding
- Broader industry impacted

Opportunities

- UK Life Sciences Sector Plan
- Regulatory changes UK & Germany
- Diversification in location of trials
- Diversification of services
- Broader therapeutic expertise larger markets
- Broader client base & cross-selling opportunities
- HCT reduced data review, faster, cheaper
- Antiviral R&D
- Vaccine R&D necessary public health risks

The Board is confident that the issues affecting the sector are transitory rather than fundamental

Outlook





Outlook

- Revenue guidance of £47 million for FY25
- Low-single digit EBITDA loss pre-exceptionals, an improvement on previous guidance
- Increase in multi-service contracts
- CRS on track to be earnings accretive in 2026
- Expect to achieve growth in 2026 and beyond

Strong Fundamentals

Strong cash position Unique early-phase clinical provider

Market Headwinds

US market volatility (esp. in vaccines) Depressed biotech funding

Diversification of Revenues

New organic services delivering CRS & Cryostore integration on track

Strong Sales Pipeline

Growing number of RFPs Large HCT opportunities in advanced discussions New disease indications in significant markets

Debt free & well funded to execute on strategy of building a sustainable & diversified business





A Full-Service CRO



Expanding our European footprint, expertise, and recruitment potential

CMC Consulting	Preclinical Consulting	Phase I Consulting	Phase I	Phase Ib & Ila HCT	Phase II & III Site Services		Data Mgmt.	Regulatory
Specialist virology & Immunology laboratory services								
Biobank Services								
Case Study	Participant Recruitment – European Footprint e Study							
Master Protocol	Hu	Phase I Trial Sub-Protocol	CRS EXPERTS. EARLY PHASE.		e 1 Database EDC, DM*)	Venn Life Sciences	Sta Data can be a	ats analysed and
		nan Challenge Trial Sub-Protocol	hvivo		T Database EDC, DM)	Venn Life Sciences	reported as s and/or as a con	single studies
		Field Trial Sub-Protocol	CRS EXPERTS. EARLY PHASE.		Trial Database EDC, DM)	Venn Life Sciences	Venn Life	e Sciences

Strategic Acquisition of CRS Mannheim & Kiel

Long-term track record as early-phase specialist

Expanding hVIVO's Site Services

Phase I-II

SAD/MAD

Proof of Concept

BE/BA, QTc, DDI

Expanding hVIVO's Therapeutic Expertise

Cardiometabolic

Dermatology

Renal / Hepatic Impairment

Immunology /
Inflammation

Expanded European Footprint

94 Beds Mannheim

26 Beds Kiel

37,000 + Subject Pool

100+ Specialists & Experts

A full-service offering supported by Venn

Cross-selling opportunities

Multi-site capability

Strengthening hVIVO's Early Clinical Development Offering

45+

Years of experience

120

Beds (78 long-term)

+1,850

Trials completed

4

Top 10 global pharma clients

12

Clients 2024

EUR19.9m

Revenue FY24

15

Cryostore Acquisition - Strengthening hLAB Offering

A specialist provider of biological and clinical materials storage

GMP & GDP compliant

HTA license

Home officecontrolled drugs licence

GMO approved CL-3 approved







Earnings enhancing, highly stable & recurring revenue stream

1999

Established

32

Freezers

c.2,800 sqft

Scope for future expansion

37 *Clients 2024*

c.9 years

Avg client tenure

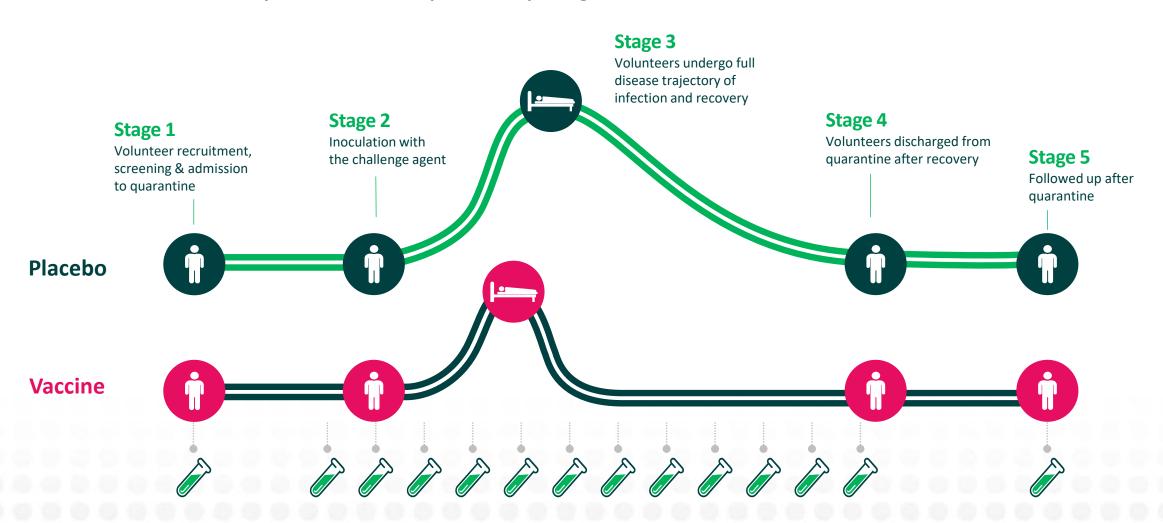
£0.9m

Revenue FY24

What is a Human Challenge Trial?



A clinical trial where healthy volunteers are exposed to a pathogen to test the effectiveness of vaccine and treatments



Benefits of Human Challenge Trials





Scientific

- Generates invaluable dosing, safety and efficacy data
- Helps optimise for larger field trials
- De-risks Phase III programs

Financial

- Significant valuation uplift for Biotech sponsor
- Quick, cost-effective data in a tight funding environment
- Allows products to "Succeed fast" or "Fail Fast"

Clinica/D sientific **Benefits of** Human Challenge **Trials**



- Requires fewer subjects
- Significant time savings
- No seasonal dependence

Regulatory

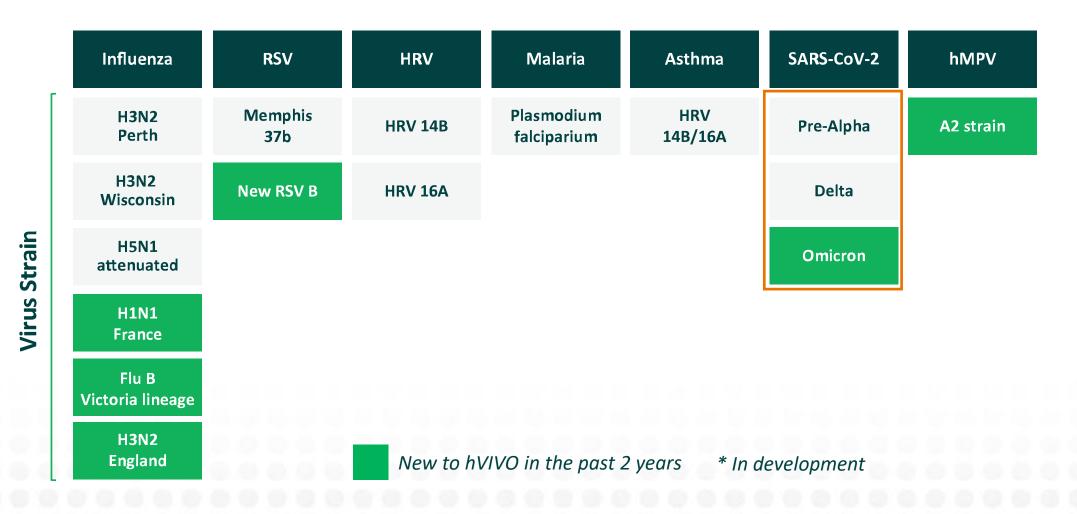
- Potential for Fast Track or Breakthrough designation - accelerating time to market
- Potential approval and Emergency Use Authorisation



hVIVO's Expanding Challenge Agent Portfolio

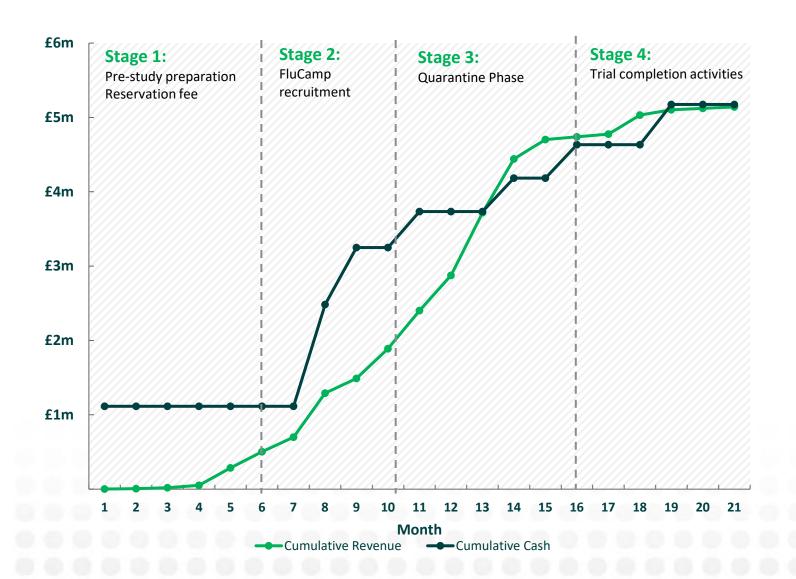


10 challenge agents manufactured in the past three years – investing in sustainable growth



Challenge Trial Revenue Recognition Profile

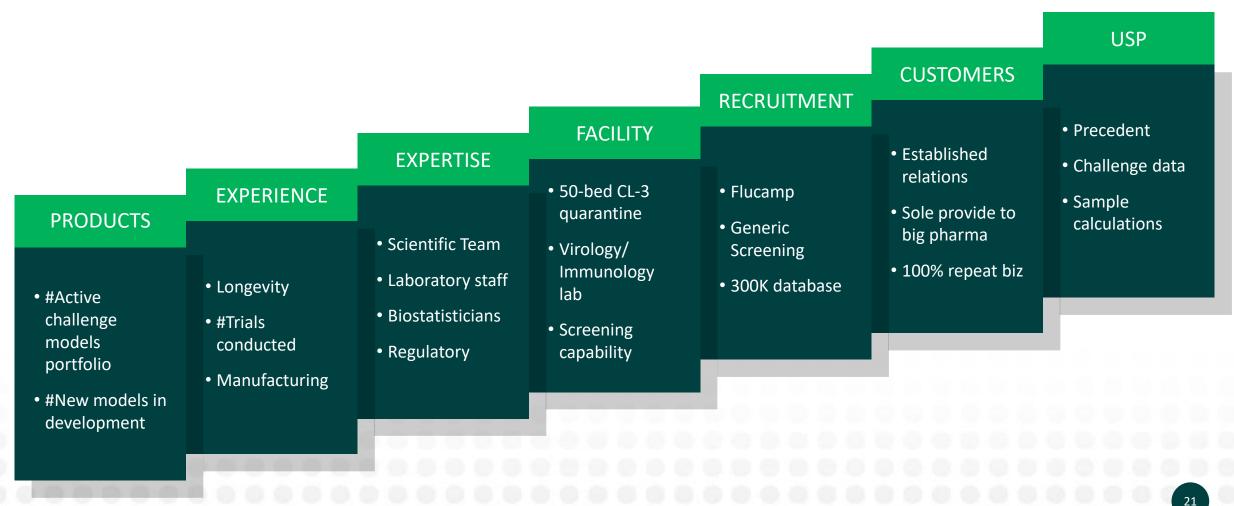




- hVIVO receives an upfront, nonrefundable booking of c.10-20% of total trial value to reserve quarantine space
- This mitigates against the risk of cancellation or client delay
- Majority of revenue recognition relates to the recruitment and quarantine phase of the trial

HCT Services: Significant Barriers to Entry





The World's Largest Human Challenge Unit





CRS: Long History & Recognised Quality





1977

Prof. Dr. Lücker GmbH

Institut für klinische Pharmakologie Bobenheim



1992

Pharm PlanNet

Contract Research



2006

CRS Clinical Research Services

Kiel, Mannheim, Mönchengladbach Member of LTS group Established as a merger of 3 Phase I CROs



2013

CRS Clinical Research Services

Berlin, Wuppertal Strategic Partnership -Take over of BAYER RESEARCH



2017

Management buy-out

Acquisition of LTS shares by APLEONEX



2025

Acquired by hVIVO

01

FDA Inspected & Passed

1991 | 1996 | 2002 | 2008 | 2009 2010 | 2011 | 2014 | 2024



GCP Inspected & Passed

2003 | 2018 (system audit by local & federal authorities)



ANVISA Inspected & Passed

- > 200+ audits by clients since 2006
- 2012 | 2016









CRS Experience (5 Years)



First-in-Human

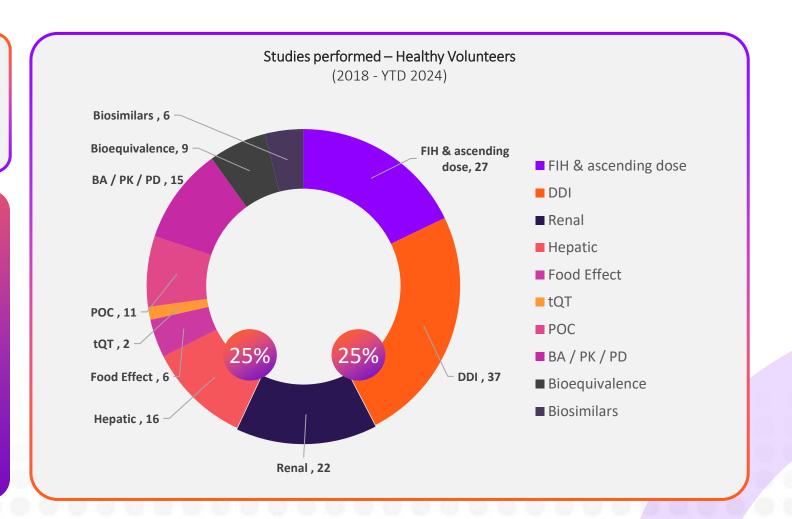
Top 5 CRO in Europe for FIH SAD/MAD, #1 in the DACH region.

Clinical Pharmacology during Later Clinical Development

Largest European CRO for impairment studies with renal and hepatic patients and

strong reputation for subsequent DDI studies (6-10 studies per year)

other pharmacokinetic studies (FE, tQT, special populations)





Venn Life Sciences Service Offering



Discovery/Lead **Drug Development** Lifecycle Management NDA, BLA, MAA **Optimization** Gap Analysis, Due diligence Quality (Chemistry, Manufacturing, and Controls) Variations/ Changes Quality development (CMC): process development, analytical development, product Quality: Process changes characterization, specification setting, stability studies, formulation, CDMO selection and improvements and management (Q)TPP, DDP **Nonclinical** Nonclinical development: proof of concept, ADME, toxicology, Toxicokinetics, safety pharmacology etc... Biomarker Safe Phase I Phase III **Post-Marketing Studies** Phase II starting dose Clinical development: clinical trial design, PK/PD, M&S, project management (Phase I), medical writing (clinical operations only for phase I) CTA, IND CTA, IND CTA, IND Marketing Authorization **Regulatory Affairs** Scientific Advice meetings CTD Pre-IND meeting, EOP1 meeting authoring

Focus on ESG

- Sustainability is integral to our corporate ethos & operational framework
- ESG Group reports to the Audit & Risk Committee
- We play a pivotal role in expediting the development of vital medicines through our full-service offering







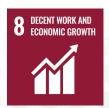






We strive to align with the 17 United Nations Sustainable Development Goals, prioritising specific goal that hold greater relevance to our business operations:











ESG Highlights



ISO 14001 accreditation achieved at Canary Wharf site 2024



Expanded facilities & services support the development of a wider range of medicines



Energy & carbon reporting, waste reduction & Electronic document management



Strong focus on ethical and compliant business practices



Empowering staff to give back to the community through charitable donations & volunteering policies



Staff well-being and development – flexible working, training & development programme



Collaborative culture and ESG focus broadening to new subsidiaries

hVIVO's State-of-the-Art Facilities

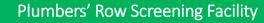


Canary Wharf Quarantine Unit



hLAB Virology & Immunology Laboratories













FluCamp



















Biobank



Watch the walk-through tour of Canary Wharf here

