





Interim Results

September 2025



AIM: HVO

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Full-Service Early Phase European CRO & World Leader in Human Challenge Trials



Cryostore

Mission

Delivering today's healthcare by empowering tomorrow's innovation.

Vision

To transform global healthcare by revolutionising the drug development process through scientific ingenuity.

100+

Years Service

5

Clinical Sites

1.9k+

Trials Completed

6

Key Areas of Expertise

10k+

Trial Participants

7

Top 10 Global Pharma Clients



Venn Life Sciences

A Full-Service Early Phase CRO



Service Lines

HCT

- hVIVO challenge services
- hVIVO laboratory
- Venn Life Sciences medical writing & biometry

Clinical Services

- CRS Phase I/II CRO services
- hVIVO & CRS clinical site
- FluCamp recruitment
- Venn Life Sciences medical writing & biometry

hLAB

- hVIVO standalone laboratory services
- Cryostore biobank & storage services

Consultancy

 Standalone Venn Life Sciences consulting services

Key Areas of Expertise











Renal/Hepatic Impairment



Supporting Clients Across the Drug Development Pathway

CMC Consulting

Preclinical Consulting

Phase I-II **CRO Services**

Human **Challenge Trials** Phase II & III **Site Services**

Marketing Authorisation

Specialist virology & immunology laboratory services

Consulting services

Biobank & storage services

Participant recruitment

H1 25 Interim Results



Key Financial Highlights

£24.2m

Revenue

H1 24: £35.6m

12.5%

EBITDA Margin

H1 24: 24.5%

0.29p

Adjusted Basic EPS

H1 24: 0.81p

£3.0m

EBITDA (pre-exceptionals)

H1 24: £8.7m

£23.3m

Cash

H1 24: £37.1m

£40m

Orderbook

H1 24: £71m

Future Financial Guidance

£47m

FY25 Revenue Forecast

Low-Single Digit

FY25 EBITDA Margin Forecast

Returning to Growth in 2026

Operational Highlights

- Synergistic acquisitions of two Clinical Research Units from CRS, and Cryostore
- Integration near completion & sales synergies being realised
- £5.5m CRS contracts signed, majority expected to be recognised 2025
- Strong progress with Clinical Services & hLAB services completed delivery of 817 participant Phase II influenza trial
- £3.2m hLAB contract for multi-site Phase II field trial
- Letter of Intent for world's first pivotal Phase III HCT
- Bacterial lab fit-out ahead of future bacterial HCTs & hLAB contracts.

Post-Period End Highlights

- c.£2m & c.£5m new awards for Clinical Services & hLAB respectively
- Phase III clinical site study awarded
- World's only contemporary-strain hMPV human challenge model now available for HCT
- Appointment of Shaun Chilton as independent Non-Executive Chair





Stephen Pinkerton

Chief Financial Officer

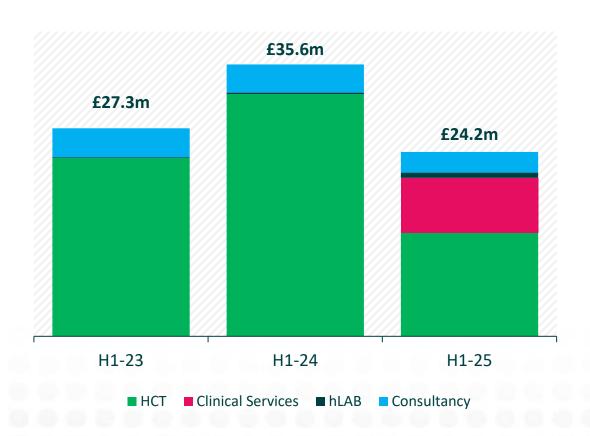


Macroeconomic & sector-specific headwinds impacted H1 25 Expect to return to growth in 2026 and beyond

Revenue

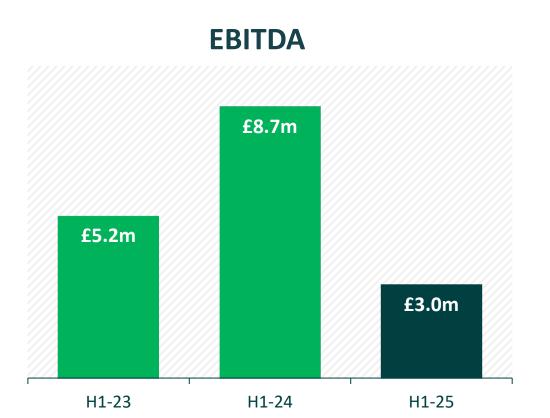


Revenue by Service



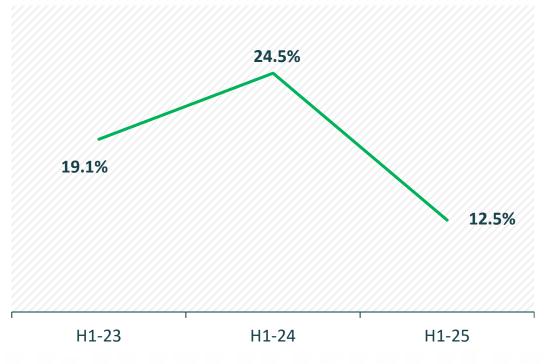
- Delivering on diversification strategy
- hLAB & Clinical Services, including acquisitions, accounted for £7.9m versus minimal revenues in H1 24
- Acquisitions delivered £5.5m:
 - CRS £5.2m & Cryostore £0.3m since acquisition date
- Broader therapeutic areas & customer base
- Cancellation fees higher than normal
- Consultancy & HCT services lower
- Revenue guidance of £47 million for FY25





- Excludes exceptional costs of £1.4m ie acquisition & restructuring
- Acquisitions contributed an expected EBITDA loss of £0.5m
- Benefitted from the positive impact of operational efficiencies, cost management & cancellation fees

EBITDA Margin

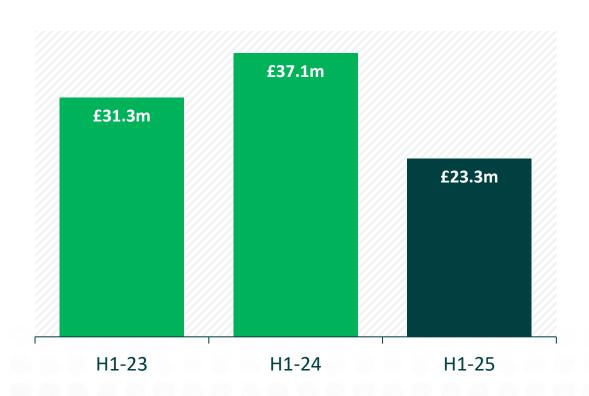


- Headcount 24% lower vs H1 24, flexibility of using temporary staff, efficiencies driven by consolidation of facilities
- Investments continue in automation to drive efficiencies
- FY25 EBITDA expected to be low-single digit loss

Healthy Cash Balance - No Debt



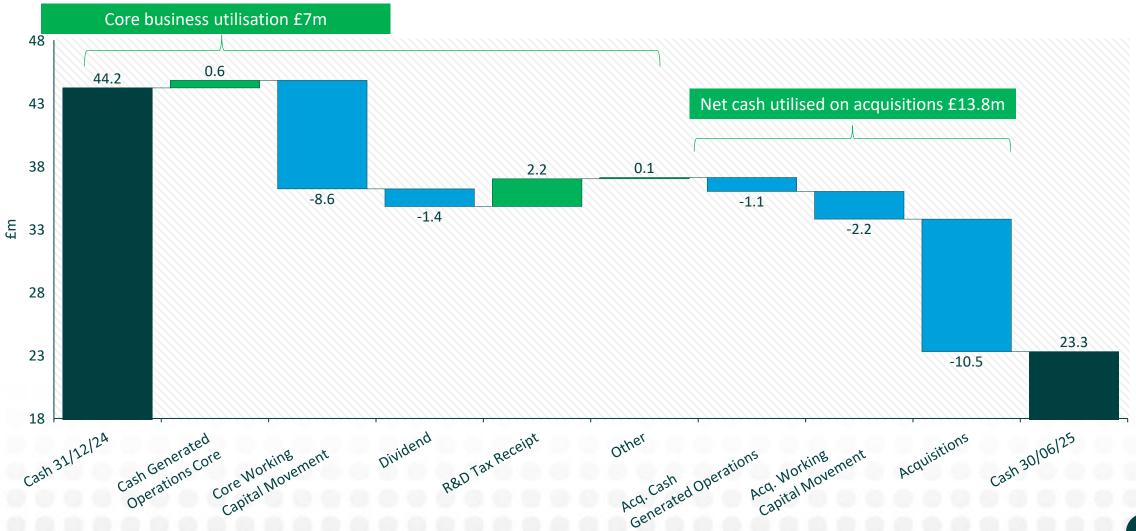
Cash



- Cash balance impacted by:
 - Acquisitions account for net £13.8m (operating loss & related working capital movement)
 - Reduction in deferred revenue/advance receipts due to cancellations & lower volume of HCT contracts signed
- Tight cost control while driving further diversification
- In the absence of material HCT contracts in 2025 expect cash balance to decline further through H2 25
- Sufficient cash to continue to invest & grow the business
- Expect to become cash generative again when HCT business returns to normal activity levels

Cash Utilisation Analysis









Dr Yamin 'Mo' Khan

Chief Executive Officer



A resilient and well-diversified organisation with strong fundamentals, diversified revenue streams, and a healthy sales pipeline

Market Trends



- Changes at Health & Human Services (HHS)
- Changes in US regulatory bodies
- FDA timelines unpredictable
- Drug pricing reforms & tariffs
- Vaccine policy
- Depressed biotech funding
- Broader industry impacted
- Reduced childhood vaccine uptake

Market Dynamics

Opportunities

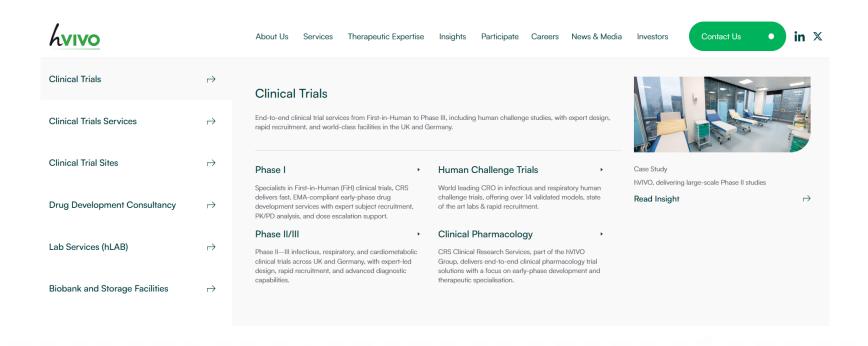
- UK Life Sciences Sector Plan
- Regulatory changes UK & Germany
- Oxford Vaccine Group five-year project
- Diversification in location of trials
- Diversification of services
- Broader therapeutic expertise larger markets
- Broader client base & cross-selling opportunities
- HCT reduced data review,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

Headwinds

Realisation of Synergies – Combined Service Offering

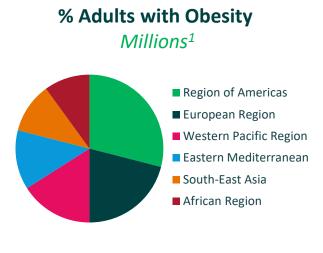


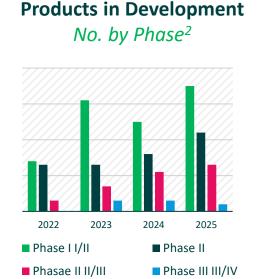
- Integration nearing completion
- Cost rationalization: >£1m annualised cost savings
- Cross-selling being realised
- £2.1m Venn opportunities in CRS sales pipeline
- Establishment of four key hVIVO service lines
- Cryostore earnings accretive
- CRS expected to become earnings accretive 2026

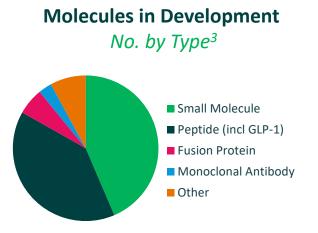


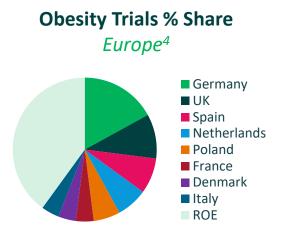
CRS Acquisition Unlocks Evolving Obesity Market











CRS – A Trusted Partner in Obesity Trials



Dr Thomas Forst Chief Medical Officer

Key Opinion Leader in Endocrinology

- \$64B by 2034⁵ global obesity clinical trials market⁴
- 33 cardiometabolic trials completed by CRS since 2019
- c.60% CRS contract wins H1 25 cardiometabolic
- c.150k overweight/obese participants in FluCamp & CRS database

hLAB – At a Glance



Leading Virology & Immunology Laboratory Services

580m²
Laboratories

Bacterial

Lab
operational

9x

Growth in proposals

Bolstered by Cryostore Acquisition

32
Sample storage
freezers

~260m²
Scope for future expansion

~9 years

Avg client tenure, recurring revenue

Key Team Members





Maria Menikou hLAB BD Director

Business Progress

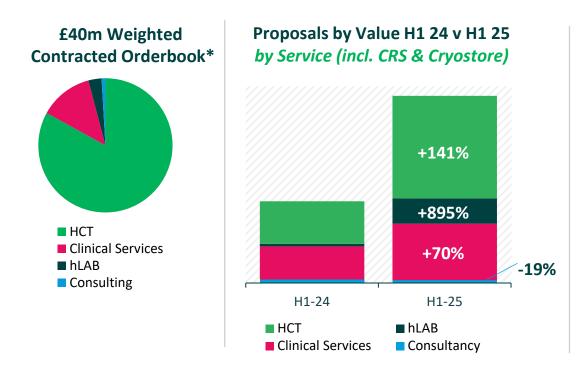
- Revenues grew strongly in H1 25
- Strong sales in H2 25: c.£5 million new awards
- 895% increase in proposals by value H1 24 vs H1 25
- Strong uptick in orderbook
 - o Phase IIb influenza trial signed in H1 25 (£3.2 million)

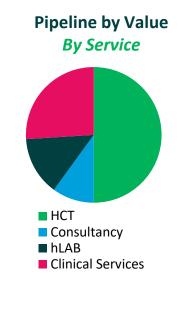
Phase III hLAB Contract

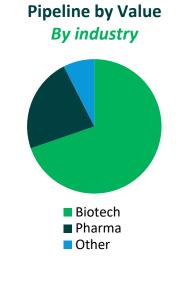
- Clinical Trial Kits & virology analysis for all subjects recruited in the international, multi-site trial
- Expected to commence Q4 25
- Majority of revenue expected to be recognised in 2026
- Expected to meaningfully support ongoing BD efforts

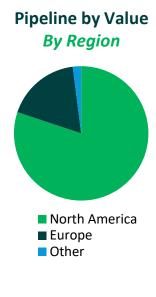
Orderbook & Pipeline – 30 June 2025











- Highly diversified orderbook, does not include ILiAD HCT*
- Total value of proposals in H1 25 exceeds FY24
- Good interest in new models hMPV & Flu B
- Some pipeline projects represent largest ever value HCTs

- Significant increase in hLAB interest
- Good growth in Clinical Services improved conversion rate of proposals to contracts for CRS vs 2024
- Cross-selling opportunities being realised

Outlook



- Revenue guidance of £47 million for FY25
- Low-single digit EBITDA loss pre-exceptionals, improvement on previous guidance
- Increase in multi-service contracts
- CRS on track to become earnings accretive in 2026
- Diversification delivering results, expected to reduce volatility over time
- Expect high-single digit revenue growth in 2026 on the back of anticipated growth in newly diversified services & moving towards a normalisation of HCT activity

Strong Fundamentals

- · Unique early-phase clinical provider
- Trusted partner & experienced team

Diversification of Revenues

- New organic services delivering
- CRS & Cryostore integration on track

Market Headwinds

- US market volatility (esp. in vaccines) impacting HCT
- · Opportunities from headwinds

Strong Sales Pipeline

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

Returning to growth in 2026 and beyond





Experienced Board of Directors





Shaun Chilton
Independent NonExecutive Chair









Dr Yamin 'Mo' Khan CEO







Stepen Pinkerton CFO



Deloitte.

Euromoney Institutional Investor PLC



Elaine Sullivan Senior Independent NED









Prof Brendan Buckley NED



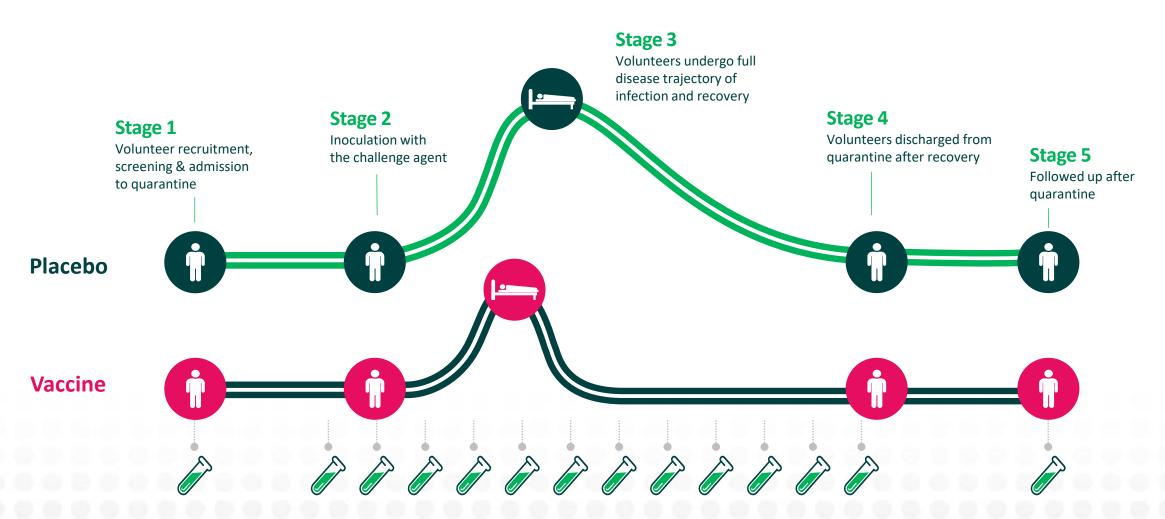




What is a Human Challenge Trial?



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



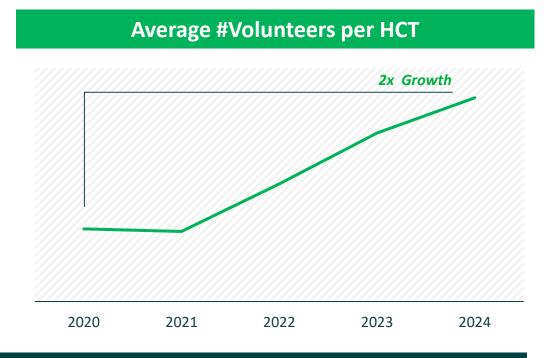
Human Challenge Trials



Benefits

- Generates valuable dosing, safety and efficacy data
- Helps optimise & derisk larger field trials
- Requires fewer subjects, significant time savings
- No seasonal dependence

- Significant valuation uplift for biotech
- Quick, cost-effective data in a tight funding environment
- Potential for Fast Track / Breakthrough designation accelerating time to market
- Potential approval & Emergency Use Authorisation



Key Growth Drivers



Larger trial sizes



13 characterised models



Bivalent / multivalent



Mucosal



Antivirals



hMPV



CL-3 capability



Bacterial laboratory



Transmission studies



Market awareness

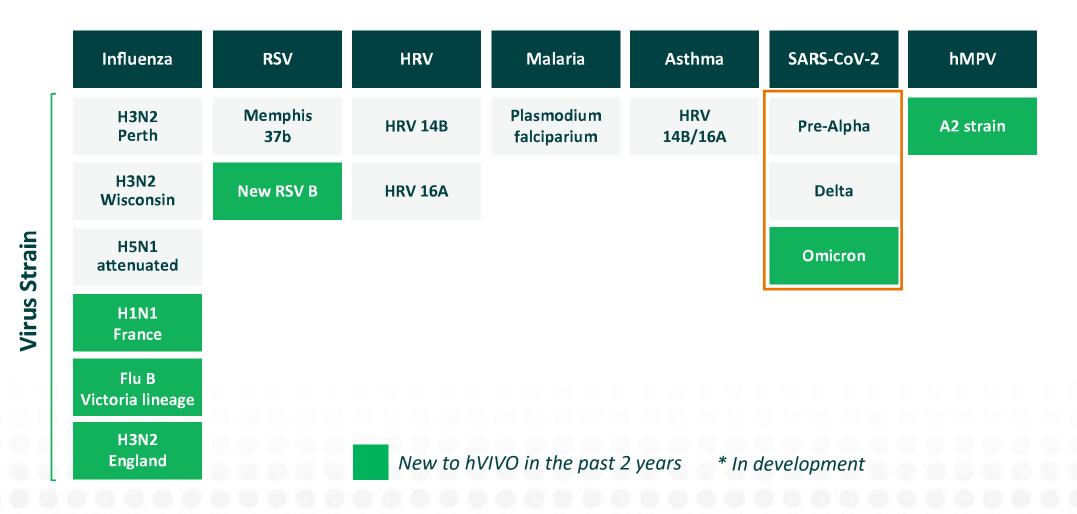


Phase III

hVIVO's Expanding Challenge Agent Portfolio

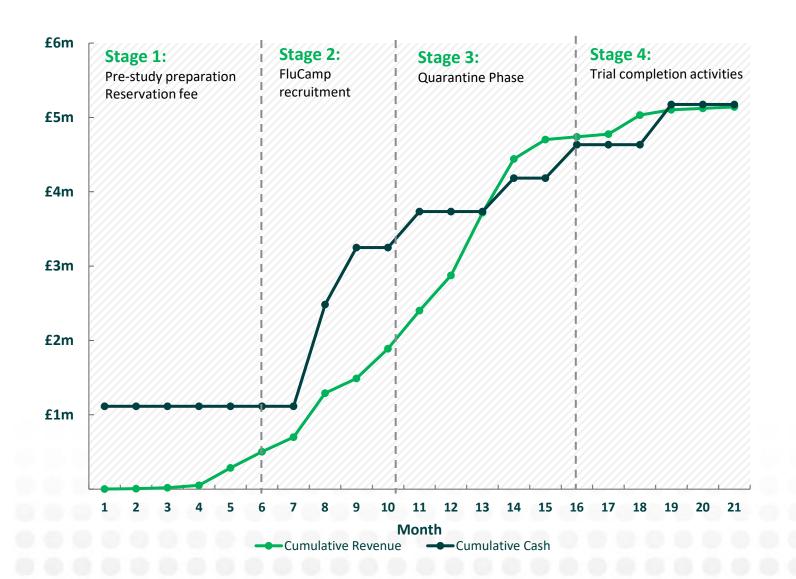


10 challenge agents manufactured in the past three years – investing in sustainable growth



Challenge Trial Revenue Recognition Profile

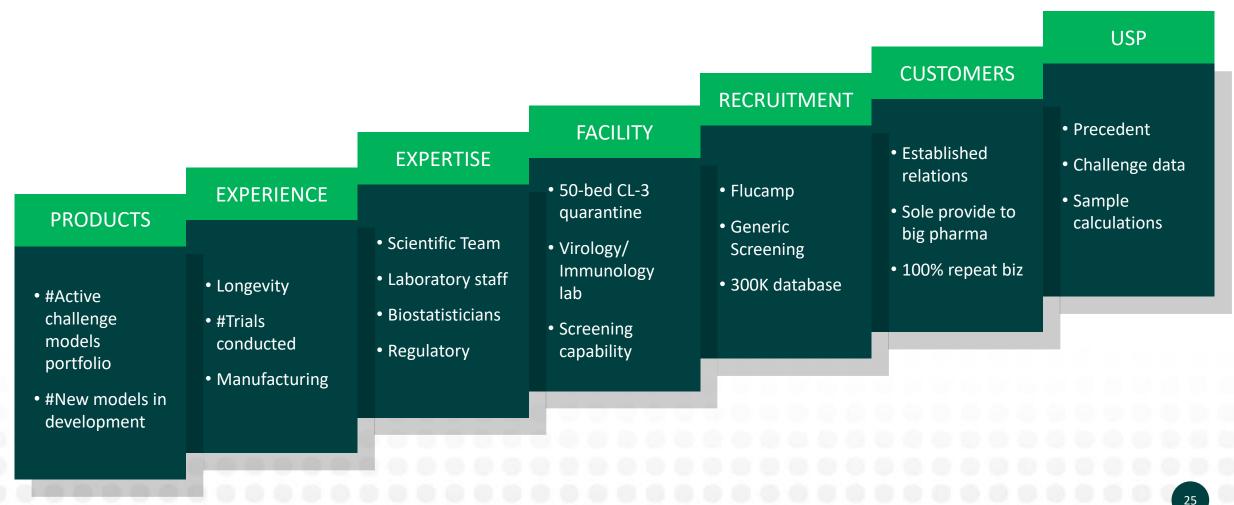




- hVIVO receives an upfront, nonrefundable booking of c.10-20% of total trial value to reserve quarantine space
- This mitigates against the risk of cancellation or client delay
- Majority of revenue recognition relates to the recruitment and quarantine phase of the trial

HCT Services: Significant Barriers to Entry





The World's Largest Human Challenge Unit





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

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

28

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



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.

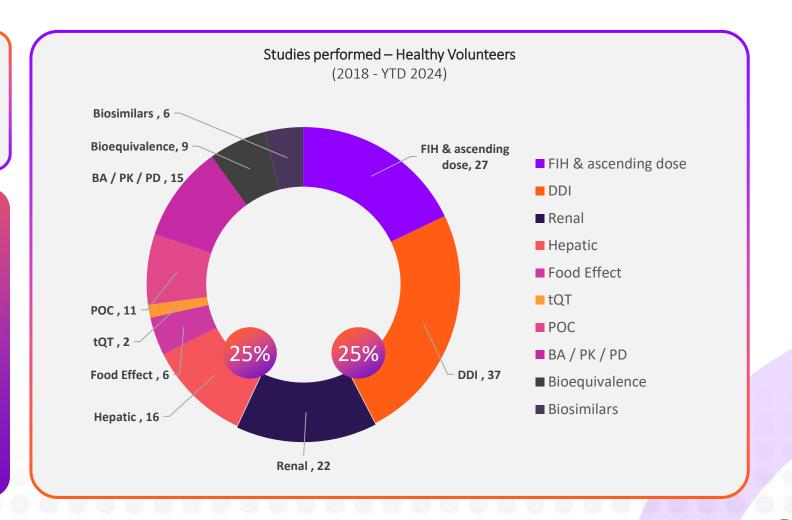
Clinical Pharmacology during Later Clinical Development

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

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

or

other pharmacokinetic studies (FE, tQT, special populations)





Venn Life Sciences Service Offering

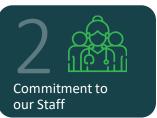


Discovery/Lead **Drug Development** Lifecycle Management NDA, BLA, MAA **Optimization** Gap Analysis, Due diligence Quality (Chemistry, Manufacturing, and Controls) Variations/ Changes Quality development (CMC): process development, analytical development, product Quality: Process changes characterization, 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







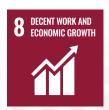






We strive to align with the 17 United Nations Sustainable Development Goals, prioritising specific goal that hold greater relevance to our business operations:











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













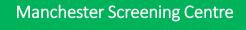












Biobank









