







Investor Presentation

July 2024

Ticker: HVO

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Who we are

A fast-growing specialist contract research organisation and the world leader in testing infectious and respiratory disease products using human challenge clinical trials

Mission

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

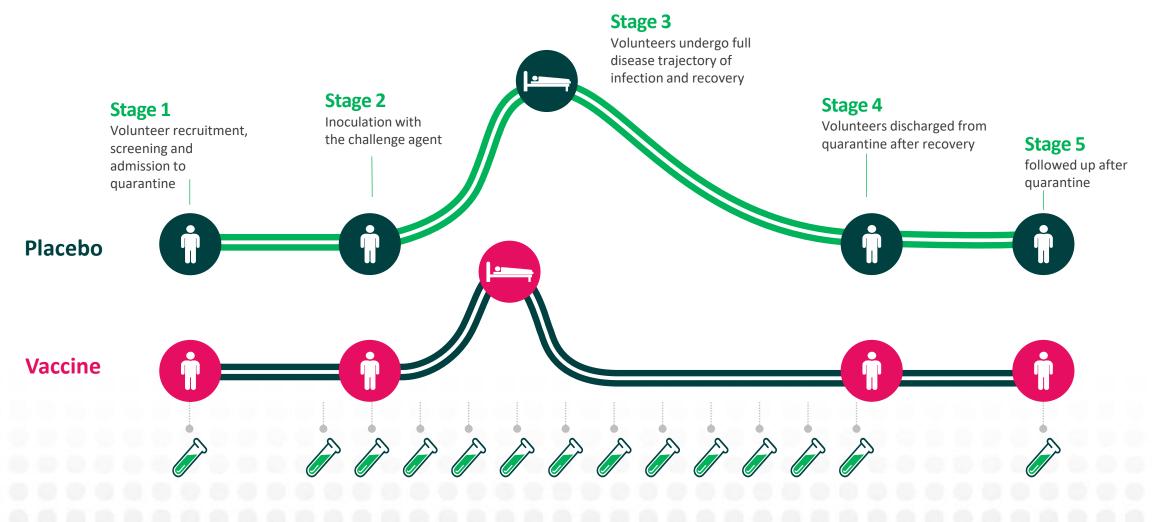
Vision

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

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



- Generates invaluable dosing, safety and stentific 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"

Benefits of Human Challenge **Trials** Regulatory Financial

Clinical D

elopment

Clinical Development

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

Regulatory

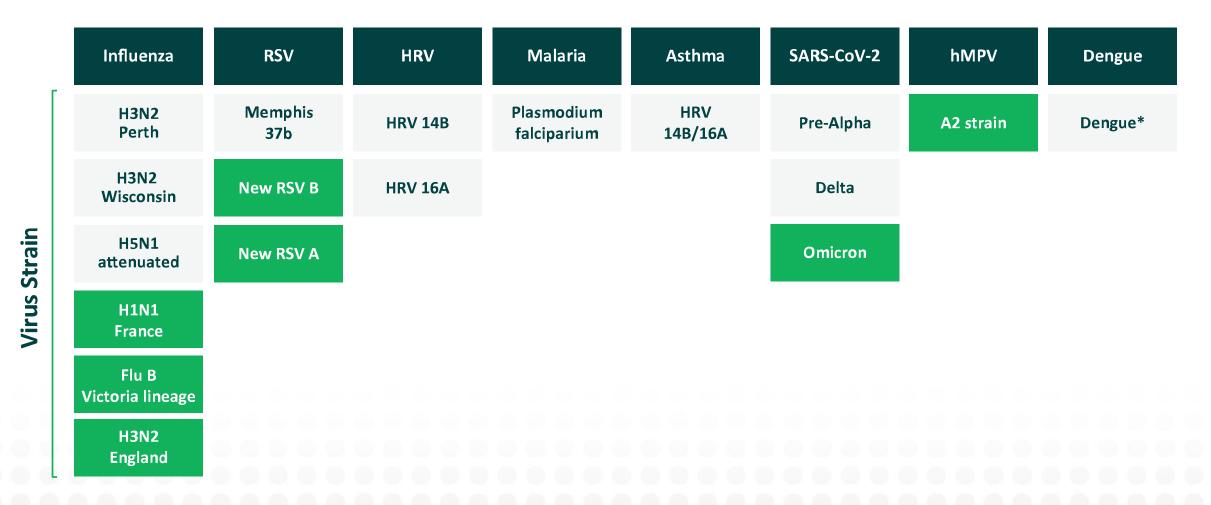
- Potential for Fast Track or Breakthrough designation - accelerating time to market
- Potential approval and Emergency Use Authorisation

World Leading Human Challenge Model Offering



6

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



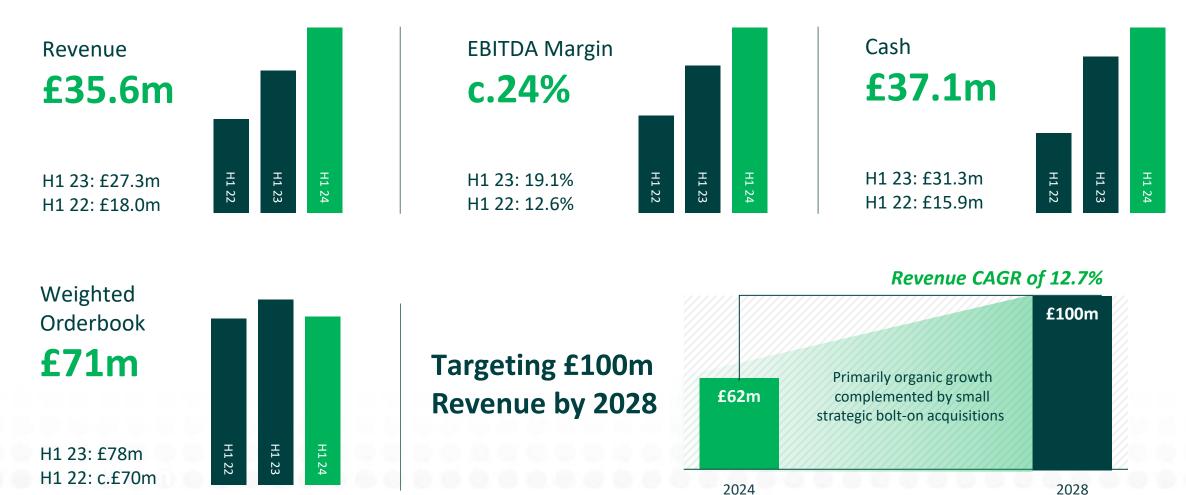
New to hVIVO in the past 2 years

* In development

Record H1 revenue and EBITDA margin



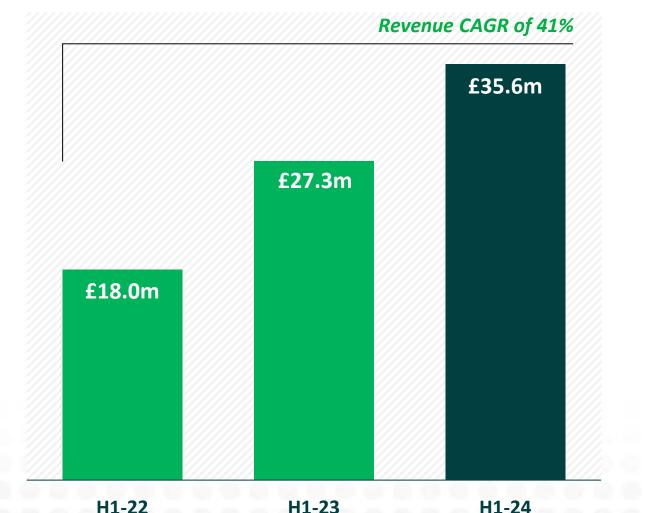
H1 2024 Trading Update – July 2024



Record H1 Revenue Growth

hvivo

H1 2024 Trading Update – July 2024



Revenue • 30% Revenue growth • Acceleration of client projects • Record number of volunteers inoculated • Increase in challenge revenue per volunteer • Six studies in quarantine across five different viruses • Venn Life Sciences underlying revenue up 10%, accounts for 11% of total revenue • £62m full year revenue guidance reaffirmed

An Increasingly Profitable Business



H1 2024 Trading Update – July 2024

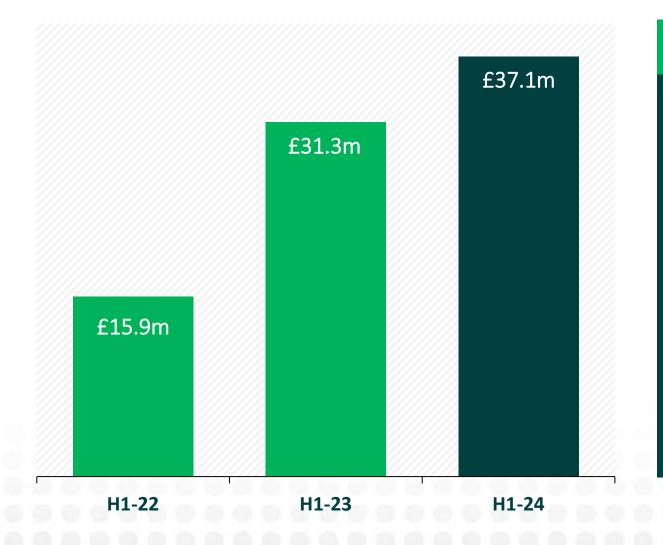


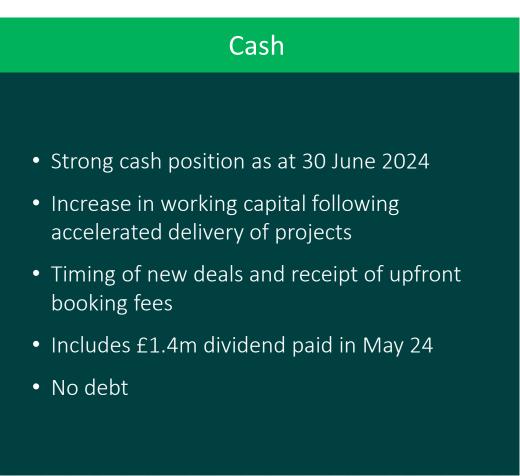
EBITDA Margin

- Significant volume of work leads to margin uplift in H1 24
- Good utilisation of the overlapping facilities
- Significant Flucamp synergies delivering lower volunteer acquisition costs
- Includes manufacturing costs for replacement viruses & facility move costs
- Full year EBITDA margin at the upper end of market expectations

Cash Generative Business

H1 2024 Trading Update – July 2024





hVIVO's Growing & World Leading Services





End-to-End Human Challenge Trial Services



Specialist Analytic Lab Services





Patient Recruitment Services

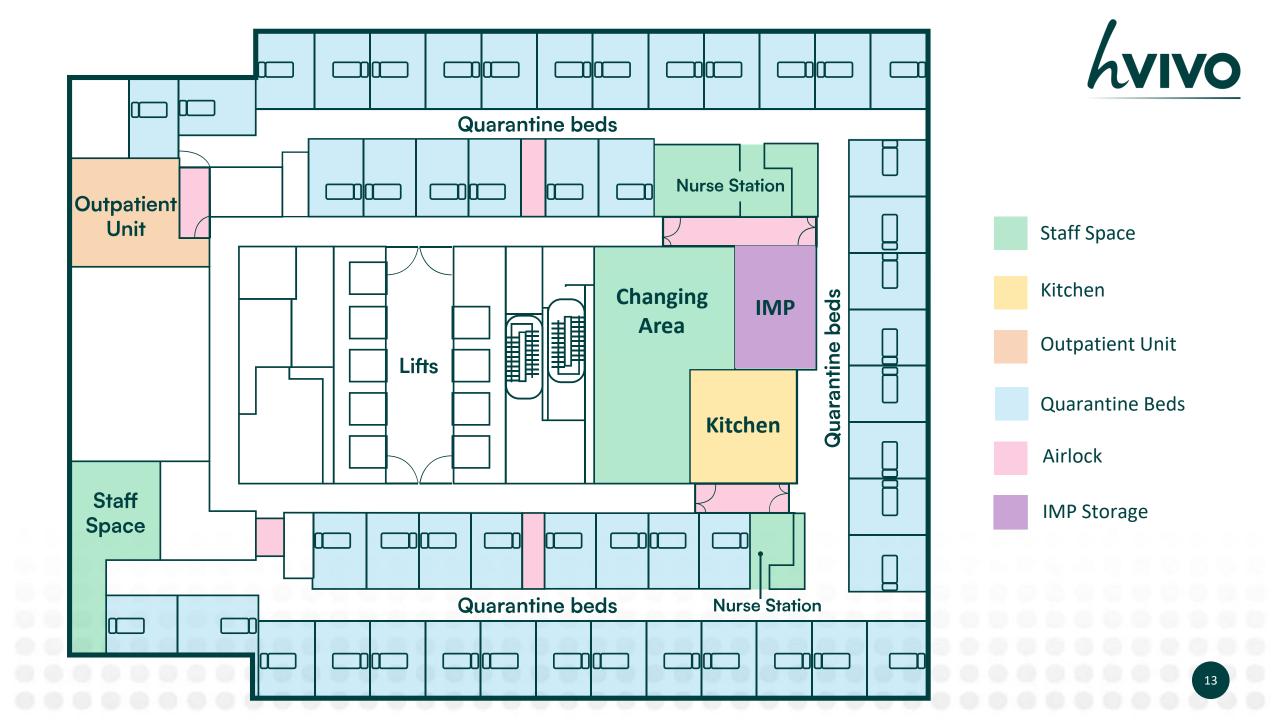
New facility providing the foundation and capacity for future growth

The World's Largest Human Challenge Trial Unit

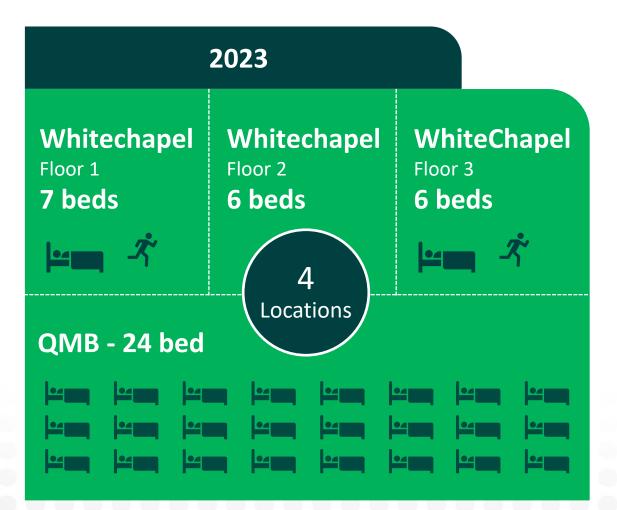


Floor 25, Canary Wharf





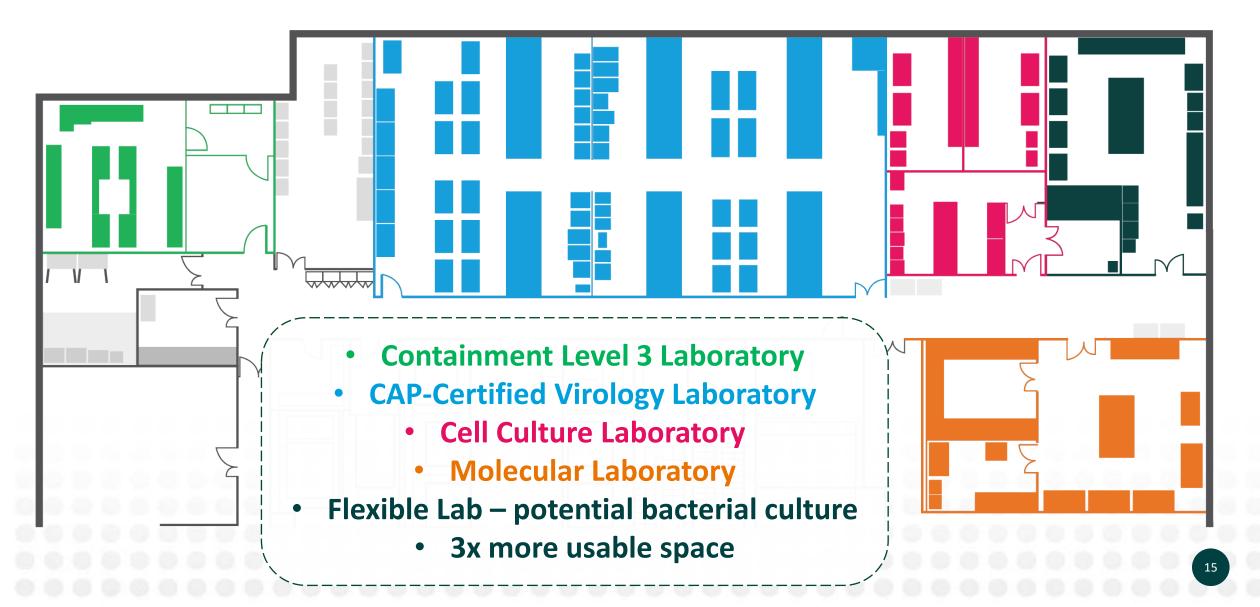
Improved resource utilisation





*h*LAb – Increased Capacity





Case Study: Phase 2b Field Study





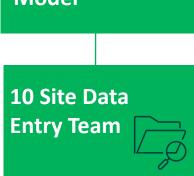
FluCamp

Enrolling up to 1,000 volunteers in 8 weeks



Flexible Resourcing Model





Expansion of Out-Patient Unit to Facilitate Future Field Studies

Increasing Vaccination Capacity from 100 to 200 volunteers per week



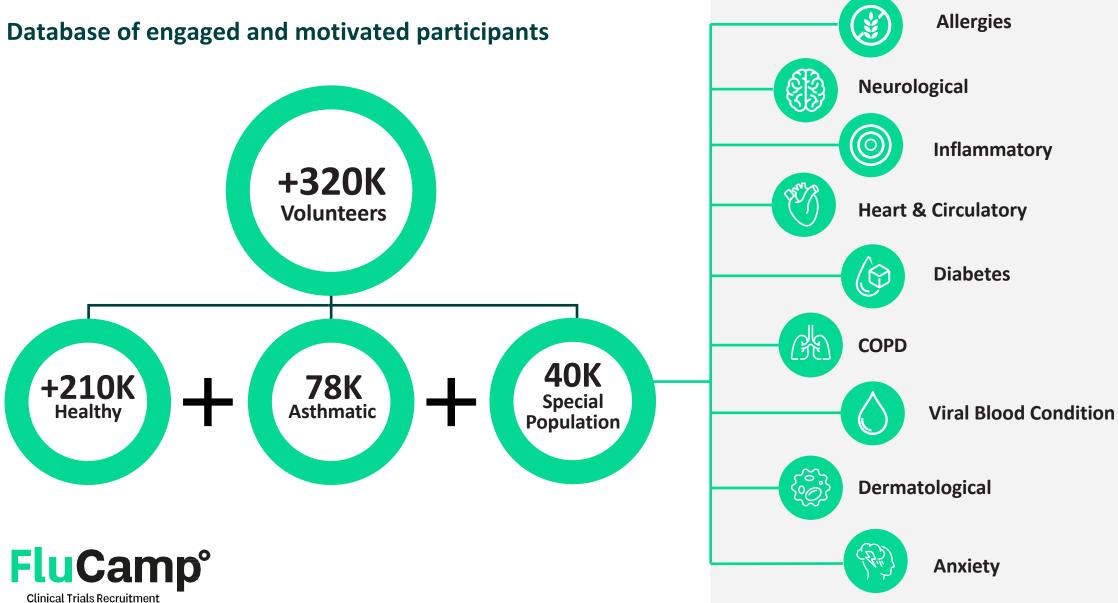




Different Collaboration Scenarios

Full-Scope Phase II studies Acting as a Site in large Phase II-III studies Stand-alone Recruitment Services (subject referral)

FLUCAMP DATABASE



TIERED RECRUITMENT OFFERING



CASE STUDY - RECENT CONTRACT

FluCamp approached by a large Global CRO to support with a difficult to recruit population

Target of 50 referrals

FluCamp°

Clinical Trials Recruitmen

- Essential package database review & outreach
- 98% of participants engaged gave consent
- All 50 referrals completed within 4 weeks
- Discussion on moving to Advance model



Venn Life Sciences Service Offering



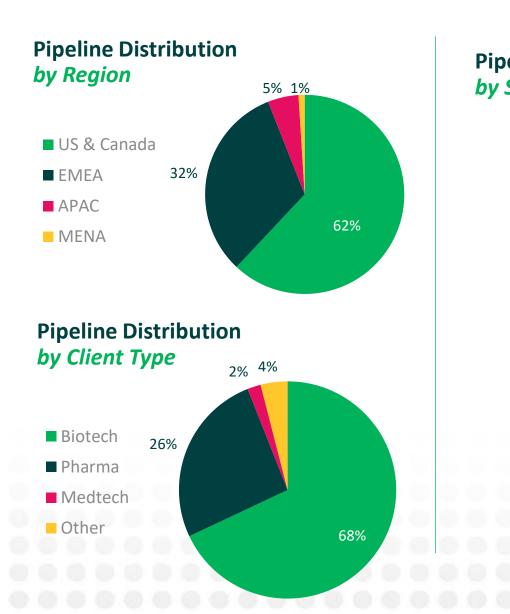
				part of hVIVO		
Discovery/Lead Optimization	Drug Development	NDA, BLA,	MAA	Lifecycle Management		
Gap Analysis, Due diligence						
	Quality (Chemistry, Manufacturing, and Controls)		V	ariations/ Changes		
	Quality development (CMC): process development, analytical development, process development, analytical development, process development, and characterization, specification setting, stability studies, formulation, CDMO sel and management			Process changes provements		
(Q)TPP, DDP	Nonclinical					
	Nonclinical development: proof of concept, ADME, toxicology,Toxicokinetics, safety pharmacology etc					
	Biomarker Safe starting dosePhase IPhase IIPh	nase III	Pos	st-Marketing Studies		
	Clinical development: clinical trial design, PK/PD, M&S, project management (Phase I), medical writing (clinical operations only for phase 1)					
Regulatory Affair	s CTA, IND CTA, IND CTA	A, IND	Mar	keting Authorization		
	Scientific Advice meetings	CTD				

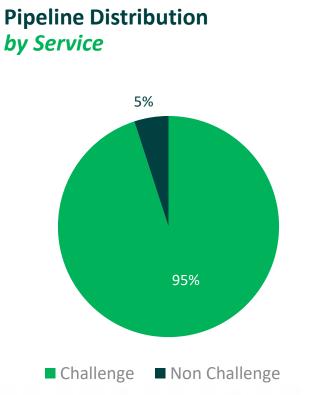
Scientific Advice meetings Pre-IND meeting, EOP1 meeting

CTD authoring

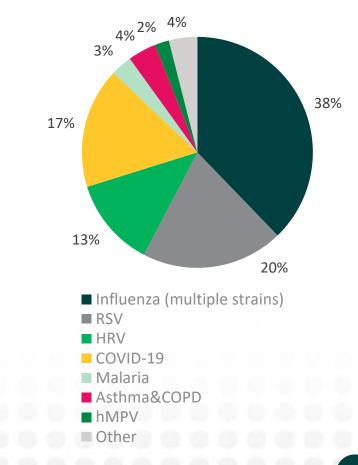
Diverse and Growing Sales Pipeline







Pipeline Distribution by Model



Investment Case



Continued Delivery

- Strong revenue, EBITDA, and sales
- Robust cash position
- Record inoculations
- Improving operational efficiencies and margins

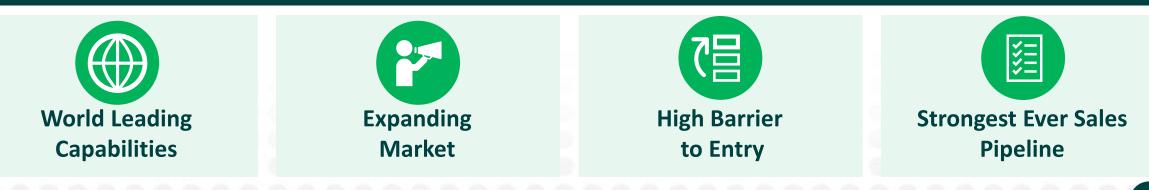
Scaling & Driving Efficiencies

- New 50-bedroom facility
- Increased lab & outpatient capacity
- New challenge models
- New services
- Improved automation

Well Positioned for Future Growth

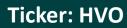
- Guiding £62m revenue 2024 & EBITDA margins at upper end of market expectations
- £71m weighted contracted orderbook
- 100% of 2024 revenue contracted & good visibility into 2025
- Medium-term target of £100m revenue

A long-term sustainable growth model





Appendix





Conducting Trials with Hazard Group 3 Pathogens

Rooms are built to hospital isolation standards with HEPA filters on all air extracts in each isolation room

Air Handling Unit

- Maintains negative pressure in each room
- Between -5 and -15 pascals
- 8-10 air changes per hour
- The failover system

Back up Power

- Building is connected to 2 separate power station via 4 power lines
- Should the power from all 4 lines fail, there are back-up generators on the roof



Previous Site	Previous Site	Canary Wharf	
246 m ²	Usable Lab Space	580m ²	
3 rd Party	CL3 Lab	✓ In House	
-	Automated Sample Dispatch	\checkmark	
3 rd Party	Autoclave Room	✓ In House	
-	Dedicated Storeroom	\checkmark	
-	Additional Flexible Lab Space	\checkmark	









- Brand Awareness
- *h*LAB is a leader in virology clinical trial support
- **Dedicated website**
- Conferences



Scientific Expansion

Large expansion in assay portfolio to include new specialties and Biospecimen Services

- Targeting year-on-year growth of standalone *h*LAB services
- Increased lab capacity
- Increase in subject matter experts

ABWARE Results Count

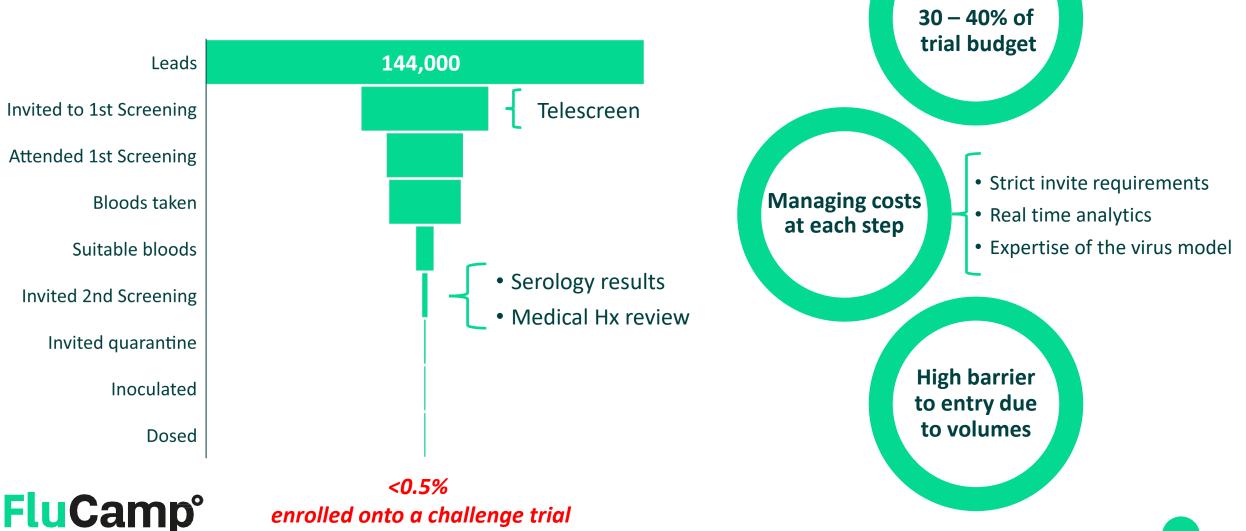
Systems & Technology

- Increased automation: PCR & Cell culture
- LIMS & eSource integration •



Lead to inoculation - 2023

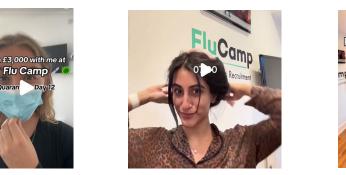
Clinical Trials Recruitment



Recruitment

FLUCAMP AS A SERVICE

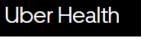
- Loyal volunteer community
- Collaborate on material generation
- 135 User Generated Content





Key partnerships

 Access to 7 million primary care patients





National Institute for Health Research NIHR



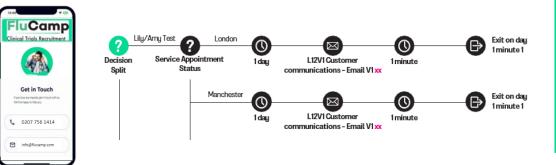












EMIS

FLUCAMP PORTAL Launch 2024!



LIVE DATABASE ACCESSIBLE BY CLIENTS









Scan for Video





Ticker: HVO