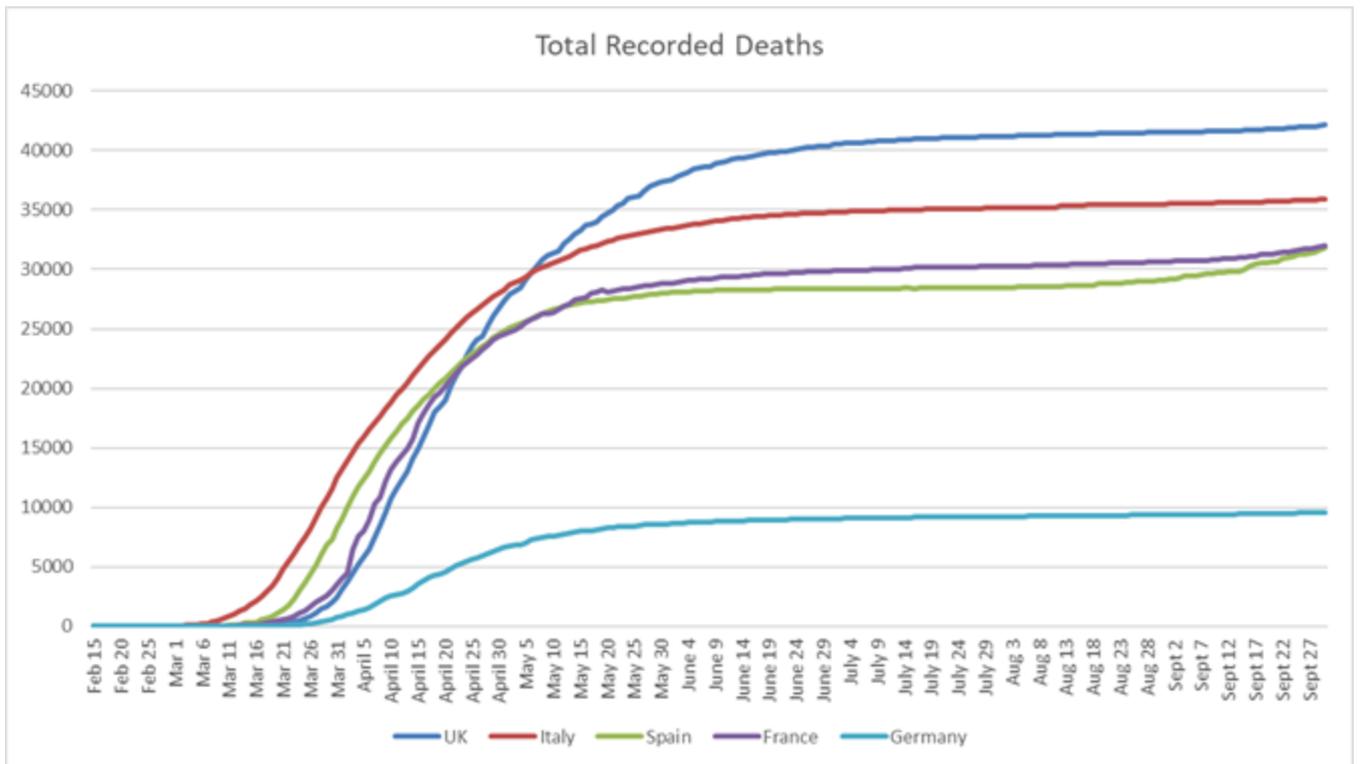
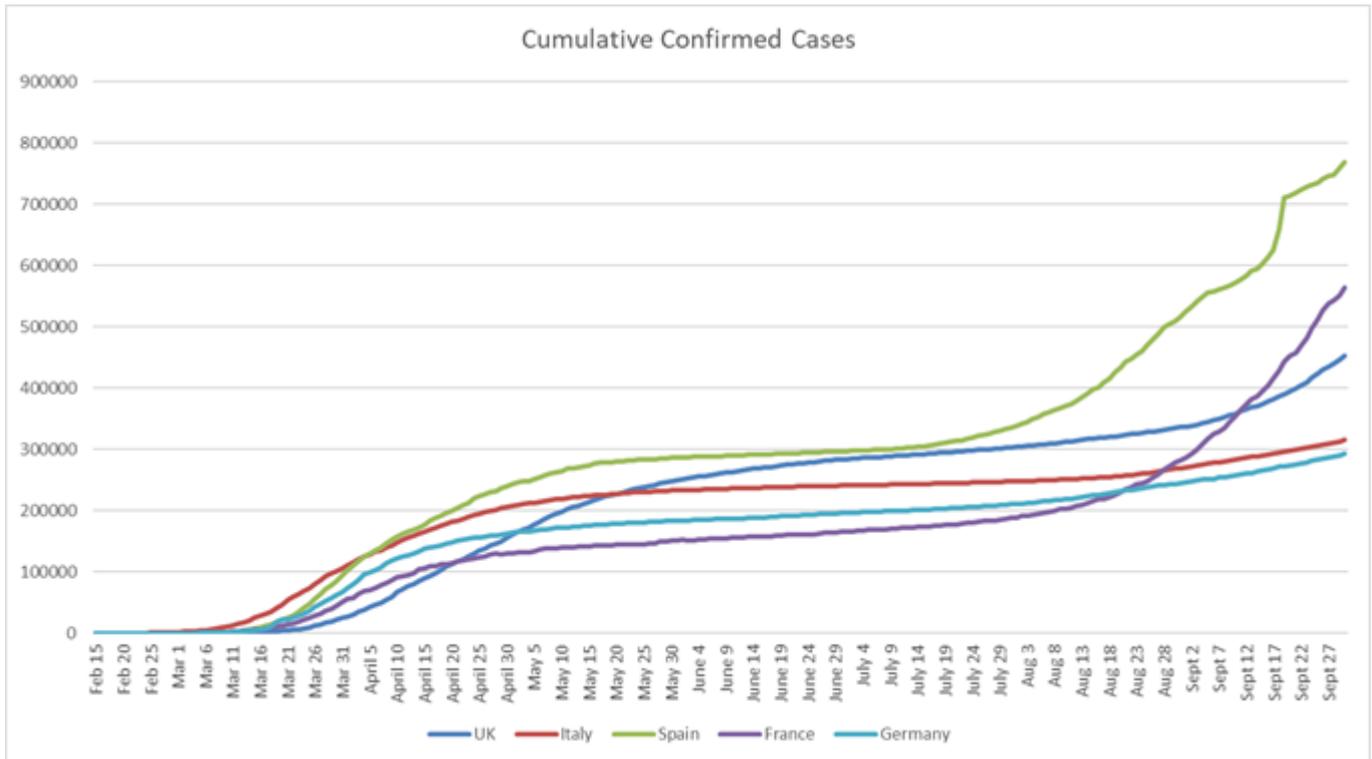
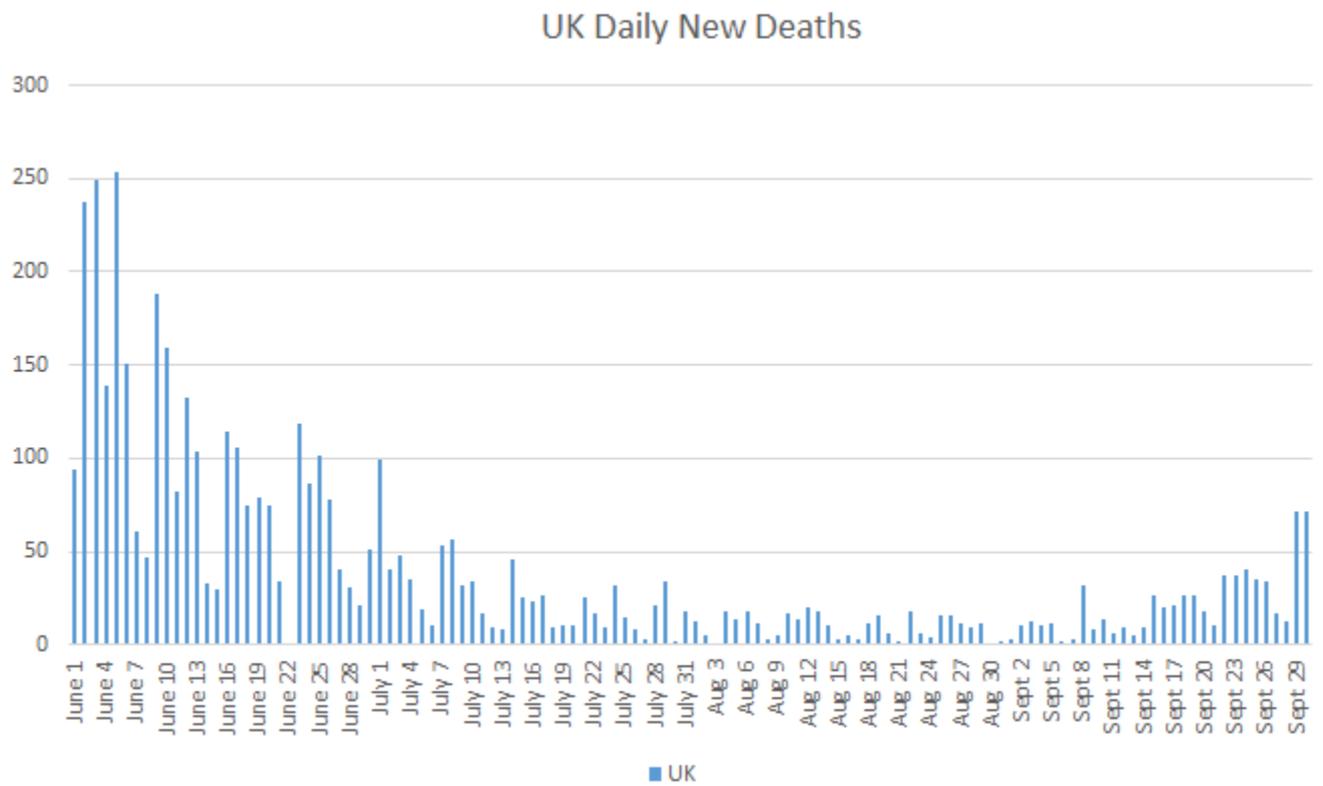
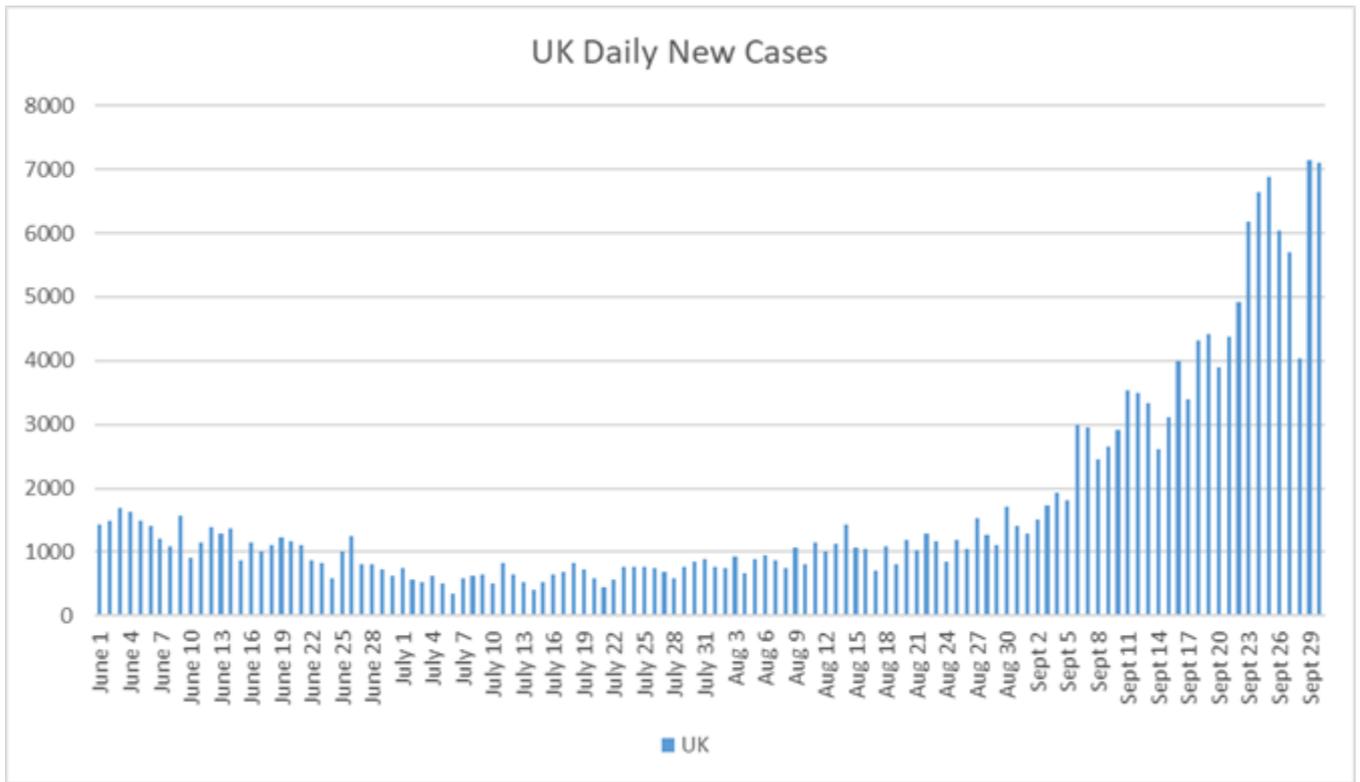


Weekly thoughts on the healthcare sector from finnCap's life sciences analysts: Mark Brewer and Arshad Ahad

Market	Last	-1D	-1M	-3M	-12M	YTD	Health	Last	-1D	-1M	-3M	-12M	YTD
MSCI World Index	1,794	0.2%	-3.2%	6.3%	6.8%	-0.3%	MSCI World Pharma/Biotech	219	0.2%	-0.5%	0.6%	15.2%	2.6%
FTSE All Share	3,303	-0.3%	-1.8%	-3.8%	-19.2%	-21.8%	FTSE All Share Health	12,389	-0.4%	0.5%	-3.2%	-1.3%	-3.8%
AIM All Share	967	0.3%	-0.4%	8.6%	9.9%	0.2%	AIM Health	11,316	1.0%	-2.0%	6.1%	22.8%	9.1%
AIM 100	4,944	0.3%	0.0%	7.9%	9.9%	-0.7%							

- The second wave of COVID-19 cases across Europe continues
- As an example, there are currently only 9 countries to which Britons can travel without either quarantining or having to take a COVID-19 test, and 4 of these are at risk of being taken off the safe travel list (Italy, Greece, Poland and Sweden).
- New social restrictions imposed in the UK have not yet had an effect on infection rates. On September 29, there were 7,143 cases, which is the second highest daily number of cases recorded in the UK since the beginning of this pandemic (although it must be noted that testing rates are much higher now). On September 30, there were a similar number of cases: 7,108.
- Worryingly, COVID-19 deaths have also started to rise again. On both September 29, and September 30, there were 71 deaths, the highest number of deaths since 1 July.
- Sources for the following graphs: Worldometer, finnCap.





Treatment and Testing Update

Avacta*: Launch of SARS-CoV-2 BAMS Research Test

- Avacta's* (AIM:AVCT) bead-assisted mass spectrometry (BAMS) assay has been launched as a research kit by Adeptrix. The assay uses the Affimer reagents specific to the SARS-CoV-2 virus to capture the virus spike protein from the sample for rapid detection by mass spectrometry.
- Up to 1,000 samples per day can be analysed by a single technician
- Avacta continues to work with the UK government's CONDOR programme to clinically evaluate the assay to obtain regulatory approval/CE marking. Avacta will receive a royalty on the research kit sales.
- In terms of Avacta's rapid **antigen test**, the company is aiming to have it validated in Q4 2020. Much of the technical risk has been removed, particularly following recent ELISA antigen data, which showed exquisite sensitivity and specificity.
- Demand for an appropriate rapid SARS-CoV-2 antigen test far outstrips supply
- We introduce a target price of 310p, with a range of 211-796p, based on a SOTP valuation of its various technologies, platforms and profit streams. Modelling the potential impact of a SARS-CoV-2 rapid antigen test is based on the production capacity of its existing partners, with further upside possible as capacity is expanded.

Open Orphan*: Potential COVID-19 challenge contract within 1-2 weeks

- This week Open Orphan* (AIM:ORPH) announced a £4.3m challenge study contract and announced its interim results. In H1 2020, it laid the foundations for transformation and the company has stated that a potential COVID-19 challenge study contract with the UK government may be signed within 1-2 weeks.
- COVID-19 challenge study contracts could generate revenues of £7-£10m each
- Open Orphan is close to having its hVIVO quarantine clinic booked until December 2021 with traditional challenge study contracts, and thus COVID-19 challenge studies would take place in an alternative quarantine facility, which the company is in advanced negotiations for
- We believe Open Orphan has laid the groundwork for a transformational H2, given the contracts it has won this year, its strong forward order book and advanced negotiations for COVID-19 challenge model contracts. We reiterate our target price of 19p.

Genedrive*: SARS-CoV-2 Kit approved in South Africa

- genedrive's* (AIM:GDR) SARS-CoV-2 PCR test kit has received regulatory approval in South Africa
- The kit can now be distributed and sold within South Africa, and the test will be supplied in the country by Sysmex
- This is the first approval outside of the CE marking and provides further validation of genedrive's test, and may provide a catalyst for other approvals in Africa.
- genedrive is awaiting approval in various African countries, India and the US.

- genedrive's test is particularly well-suited to Africa and India, as its freeze-dried formulation eliminates the need for cold storage, and its 'one-step' PCR bead format eliminates the need for time-consuming and error-prone preparation.

Novacyt: Launch of COVID-19 antibody test and UK government contract

- Novacyt (AIM: NCYT) has launched a CE-Mark approved serology 96-well plate ELISA test for the detection of IgG antibodies to SARS-CoV-2 derived from plasma and serum samples.
- The test complements Novacyt's existing COVID-19 product portfolio, including its PCR test to detect active infections.
- The antibody test was validated in a study where 1,673 patient samples were evaluated and the test demonstrated 100% sensitivity and 99.4% specificity, in patients that were tested 14 days after testing positive for COVID-19 by a PCR test.
- Novacyt has capacity to deliver more than 3m antibody tests per month initially, with potential to increase this depending on demand.
- While some experts have questioned the usefulness of antibody tests, due to the fact that we do not yet know if antibodies to COVID-19 can prevent reinfection, the test may aid in increasing the understanding of the disease through screening of populations for infection rates.
- Novacyt also announced this week that it will supply 300 PCR machines and test kits to the UK government for £150m, with the potential for a further £100m of supply.

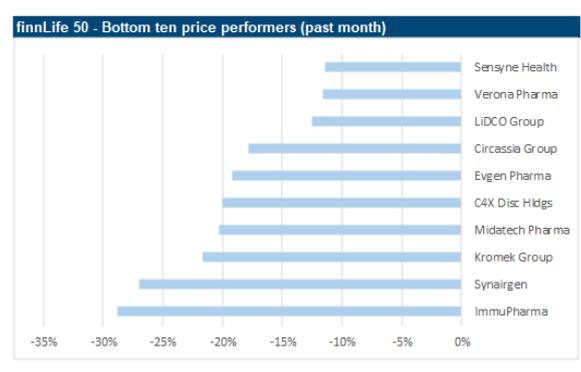
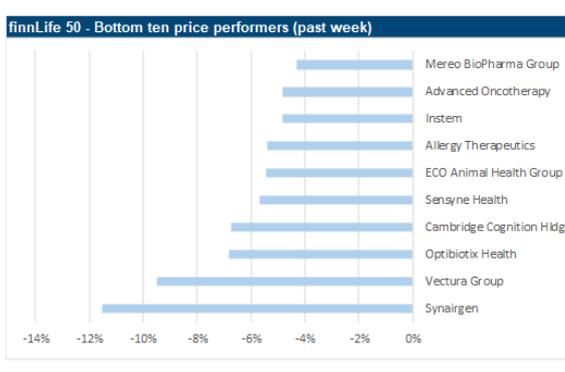
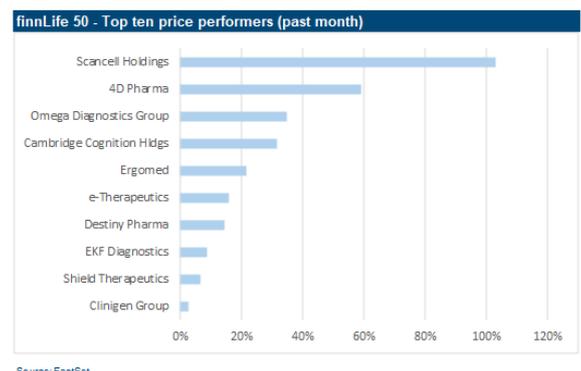
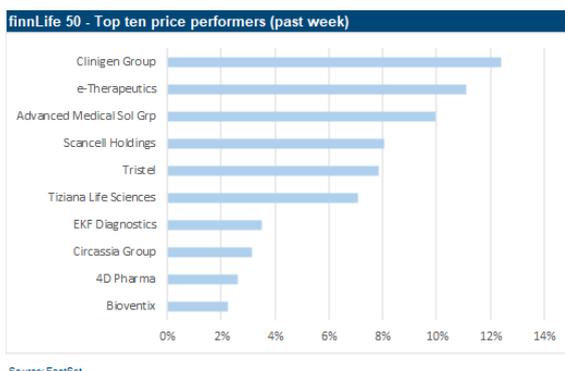
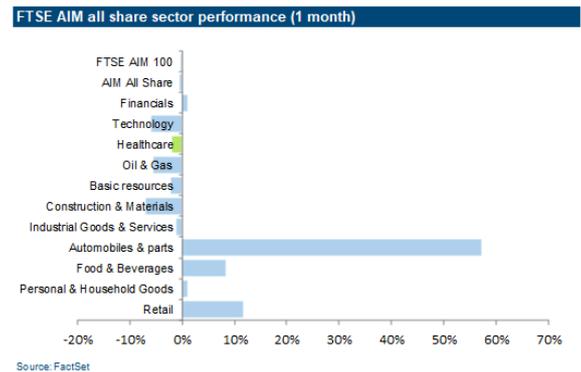
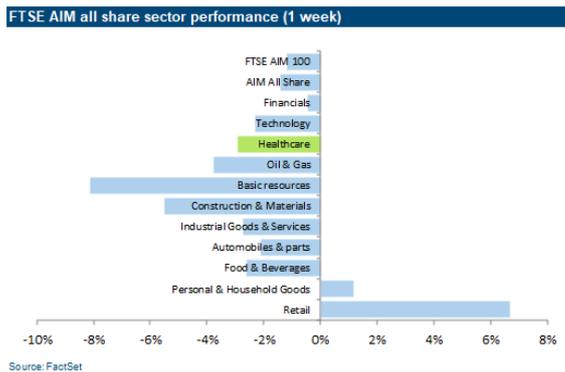
Synairgen*: Fuller Phase II COVID-19 trial data analysis and managed access program

- A fuller analysis of Synairgen's* (AIM:SNG) Phase II trial of inhaled interferon (SNG001) in hospitalised COVID-19 patients confirms our earlier optimism at the time of its first headline disclosure in July. Synairgen has a potentially significant asset to treat hospitalised COVID-19 patients.
- A Managed Access Program (MAP) has been launched, with Clinigen to provide SNG001 to hospitalised COVID-19 patients in the UK and Europe and is a significant step towards first commercial revenues before regulatory approval.
- Managed Access Programs allow doctors and patients access to investigational drugs outside of the clinical trial setting and prior to regulatory approval.
- Synairgen is in discussions with regulatory agencies to establish the route to approval of SNG001 as a treatment for COVID-19. It is also investing in supply chain activities to ensure that drug and aerosol delivery system availability can meet potential demand, pending approval, with the aim of being able to supply c.100,000 treatment courses per month in 2021.

Abbott's small, cheap and fast antigen test gets big orders

- In August, Abbott announced its BinaxNOW COVID-19 Ag Card rapid antigen test, a cheap point of care test the size of a credit card promising quick results. Administered by a health-care provider using a nose swab, the card delivers a result within 15 minutes and costs \$5, much like a pregnancy test. Abbott claims the test has 97.1% sensitivity and 98.5% specificity, albeit in very infectious (high viral load) patients

- The US federal government has previously agreed to purchase \$760m worth of tests, and on 28 September it described the plan to distribute the 150m tests from that contract within the next few weeks.
- The WHO also announced a contract for 120m tests, split between Abbott and SD Biosensor of South Korea. The tests will be distributed to low and middle-income countries over the next 6 months. However, the UK government stated it has no plans to purchase the test.
- While Abbott says it can increase production to 50m tests a month in October, this only scratches the surface in terms of global need and demand for such tests.



Research reports and comments in the past two weeks

Company	Date	Title
InnovaDerma*	30-Sep	FY 2020 finals – growth drivers beyond COVID
Open Orphan*	30-Sep	H1 2020 – Laying the foundations for transformation
Synairgen*	29-Sep	Interims and fuller Phase II data analysis

ANGLE*	28-Sep	FDA submission – a significant valuation milestone
Avacta*	28-Sep	Interim results – new forecast, new target price
Open Orphan*	28-Sep	£4.3m contract win with top ten vaccine company
Byotrol*	28-Sep	FY 2020 results – positive outlook drives upgrades
Allergy Therapeutics	28-Sep	FY 2020 – record pre-R&D EBIT
ANGLE*	23-Sep	Peer reviewed publication, using Parsortix in MBC
Cambridge Cognition*	22-Sep	Strong interims with 57% 2021 revenue visibility
Destiny Pharma*	17-Sep	Interims – on track for Q1 2021 Phase 2b readout

Upcoming roadshows and events

Company Name	Ticker	Event Type	Roadshow/Event Date
InnovaDerma Plc	IDP	Preliminary Results Roadshow	30th September & 1st October – Conference Calls
Open Orphan Plc	ORPH	Interim Results Roadshow	6th October – Group conference call at 11.30am
Tristel plc	TSTL	Preliminary Results Roadshow	19th & 20th October – Conference Calls
Bioventix plc	BVXP	Preliminary Results Roadshow	19th – Group Conference Call at 12pm 20th & 21st October – Conference Calls

[An archive of previous Health Matters can be found here.](#)

[Download our latest Q3 Quarterly Report, on the theme of Cell and Gene Therapy here.](#)

To UNSUBSCRIBE, please simply reply to that effect.

*Denotes corporate client of finnCap. This research cannot be classified as objective under finnCap research policy.

^ This company and finnCap have agreed that finnCap will produce and disseminate research and finnCap may receive remuneration in return for this service. This research cannot be classified as objective under the finnCap research policy.

A marketing communication under FCA Rules, this document has not been prepared in accordance with legal requirements designed to promote the independence of investment research and is not subject to any prohibition on dealing ahead of the dissemination of investment research.

This research cannot be classified as objective under finnCap Ltd. research policy. Visit www.finncap.com

MAR Research disclosures can be found at <http://www.finncap.com/disclosures>

The recommendation system used for this research is as follows. We expect the indicated target price relative to the FT All Share Index to be achieved within 12 months of the date of this publication. A 'Hold' indicates expected performance relative to this index of +/-10%, a 'Buy' indicates expected outperformance >10% and a 'Sell' indicates expected underperformance of >10%.

Approved and issued by finnCap Ltd. for publication only to UK persons who are authorised persons under the Financial Services and Markets Act 2000 and to Professional customers. Retail customers who receive this document should ignore it. finnCap Ltd. uses reasonable efforts to obtain information from sources which it believes to be reliable, but it makes no representation that the information or opinions contained in this document are accurate, reliable or complete. Such information and opinions are provided for the information of finnCap Ltd.'s clients only and are subject to change without notice. finnCap Ltd.'s salespeople, traders and other representatives may provide oral or written market commentary or trading strategies to our clients that reflect opinions contrary to or inconsistent with the opinions expressed herein. This document should not be copied or otherwise reproduced. finnCap Ltd. and any company or individual connected with it may have a position or holding in any investment mentioned in this document or a related investment. finnCap Ltd. may have been a

manager of a public offering of securities of this company within the last 12 months, or have received compensation for investment banking services from this company within the past 12 months, or expect to receive or may intend to seek compensation for investment banking services from this company within the next three months. Nothing in this document should be construed as an offer or solicitation to acquire or dispose of any investment or to engage in any other transaction. finnCap Ltd. authorised and regulated by the Financial Conduct Authority, London E14 5HS, and is a member of the London Stock Exchange.