

Covid-19 and Vaccination Programme

Acute Severe COVID

Professor Nick Hart, Clinical Director for Respiratory, Sleep and Critical Care

Mild to Moderate COVID, Long COVID and Rehabilitation

Professor Nick Hart and Alex Curtis, Deputy Clinical Lead Physiotherapist for Rehabilitation

Vaccine Hesitancy

Dr Ali Hashtroudi, Clinical Director, Occupational Health and Honorary Senior Lecturer



A Year in COVID 'Speed Trumps Perfection'

Mike Ryan, Executive Director WHO

Nicholas Hart Professor of Respiratory & Critical Care Medicine Clinical Director Respiratory, Sleep and Critical Care Cardiorespiratory Critical Care Clinical Group St Thomas' Hospital Director of Research Delivery Guys & St Thomas' NHS Foundation Trust

Thanks to ...

- Critical care team
- Anaesthetic team
- Evelina PICU team
- Critical care surge team
- Acute medicine team
- Turning team
- IT team
- Vascular surgical team
- ENT team
- Surge education and training team
- Infection prevention and control team
- Data team
- Wellbeing team
- Pharmacy team
- Physiotherapy team
- Occupational therapy team
- SALT team
- Dietetic team
- Cleaning team
- Engineering team
- Estates team
- Catering team





- Report the presentations of mild, moderate, severe and very severe COVID
- Report the nature of the disease over two pandemic waves at GSTT
- Report our GSTT outcomes
- Report the acute treatments we are now using based on UK Urgent Public Health COVID-19 Studies
- Report the telephone clinic that was set up to support post-acute discharge
- Prevention is better than cure





451 days after the first patient was admitted under HCID Team on Hillyers ward at St Thomas' Hospital

425 days after the first patient was admitted to HCID ECMO Critical Care Unit at St Thomas' Hospital EW6...



It is understood the woman will be treated at London's St Thomas' Hospital

Ninth coronavirus case found in UK

10 mins ago | UK

A woman who flew into London from China a few days ago is being treated for coronavirus, bringing the total number of UK cases to nine.

Chief medical officer Chris Whitty said the woman had been transferred to a specialist NHS centre at Guy's and St Thomas' in central London.





Evidenced Based Clinical Care

GSTT COVID Clinical Reference Group





Assessment tool for adults with confirmed or suspected COVID-19



Clinical features triggering hospital review or admission:

History

- Breathless at rest or minimal exertion e.g. walking to the bathroom, walking up the stairs, recovery time >2 mins post-exertion.
- Review/admit all from the high-risk group (box 1) with any features of infection. Have a lower threshold for reviewing patients with a moderate risk (box 2) compared to those with no risk.
- Worsening clinical condition >5 days after symptom onset if risk factors (boxes 1-3).
- · Fever ongoing for >72h.
- Delirium in older adults (aged ≥65 years).
- If discharged, does not have a sufficient home support network or is unable to adequately self-isolate from a clinically extremely vulnerable individual (box 1) in the household.

Examination

- · Unable to speak in full sentences, use of accessory muscles of breathing.
- Dry mucous membranes, clinically 'dry'.
- Heart rate >110/min.
- Respiratory rate >20/min.

Observations and investigations

- SpO2 ≤94% on room air.
- SpO2 ≤92% or ≥3% fall on 1 minute sit-to-stand test.
- Urea >7 mmol/L and greater caution if >14 mmol/L; confirmed or suspected acute kidney injury (AKI) stage 2 or 3 (NICE guideline).
- Albumin <35g/L
- Widespread infiltrates on chest X-ray.
- N.B. The list above is not exhaustive. Other findings or tests, typically requiring hospital attendance e.g. troponin, may also determine it in-patient care is indicated.

If the patient is suitable for discharge from the ED or a hospital ward:

- Advise the patient to isolate at home until non-infectious: 10 days from symptom onset for non-hospitalised individuals; 14 days from initial positive test if discharged from a hospital ward, including two consecutive days without fever (and no medication to reduce fever).
- If any risk factors are present (boxes 1-3), refer by EPR for a daily phone review from day 7 to day 14 following onset of symptoms (discontinues after two consecutive days of improvement).
- Advise patient to contact their GP or NHS 111 if their condition deteriorates.

Patient risk groups

HIGH-RISK ('clinically extremely vulnerable') includes.

- Solid organ transplant recipients
- Chemotherapy or immunotherapy for cancer
- Radical radiotherapy for lung cancer
- Blood or bone marrow cancer at any stage of treatment
- · Bone marrow or stem cell transplant within the last six months
- · Severe lung conditions e.g. cystic fibrosis, severe asthma, severe COPD
- Diseases that increase the risk of infection e.g. SCID, sickle cell disease etc.
- HIV: CD4 <50 or opportunistic infection in the last 6 months (BHIVA advice)
- Immunosuppression therapies in the last three months e.g. >40mg prednisolone per day for >1 week or >20mg per day for >14 days; methotrexate >25mg per week; azathioprine >3.0mg/kg/day; 6-mercaptopurine >1.5mg/kg/day
- Chronic kidney disease (CKD) stage 5 (eGFR < 15 mls/min) or on dialysis
- Pregnancy with significant heart disease, congenital or acquired.

MODERATE RISK ('clinically vulnerable') includes:

- ≥70 years of age
- ≥65 years of age with <u>clinical frailty scale</u> of ≥5
- Very obese (BMI ≥40)
- Diabetes (type 1 or 2)
- · Chronic, non-severe, respiratory conditions e.g. asthma, COPD etc.
- · Chronic heart disease e.g. heart failure
- Chronic kidney disease (CKD) stage 3 or 4 (eGFR 15-60mls/min)
- Chronic liver disease
- HIV: CD4 <200, detectable viral load or not on ART (BHIVA advice)
- Chronic and/or progressive neurological disorders e.g. Parkinson's disease, cerebral
 palsy, multiple sclerosis, motor neurone disease
- . Drugs that affect the immune system e.g. low-dose steroids (see also box 1).
- Pregnancy

3. OTHER RISKS include:

- ≥55 years of age
- Obese (BMI ≥30)
- Learning difficulty
- BAME background

Developed by Guy's and St Thomas' NHS Foundation Trust





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COVID | Big 5



Five interventions to help treat suspected or confirmed COVID-19



GSTT COVID CLINICAL REFERENCE GROUP

ACUTE COVID DISCHARGE PATHWAY







Acute Post COVID-19 Discharge Telephone Assessment

Initial assessment

- I understand that you have recently been discharged from hospital with a diagnosis COVID-19
- I am ringing to assess your progress and ensure that you are improving
- Can I ask a few questions please?

Symptoms

- Are you feeling worse, better or the same?
- Are you breathless sitting down in a chair?
- Are you breathless on walking from one room to another in your home?*
- When did you last have a temperature/fever?
- Are you able to drink plenty of fluids?

Specific medical conditions:

Diabetes

- Do you check your blood sugars at home?
- Have you sugar levels been high?

Severe respiratory disease including COPD and ILD

- Have you been issued an oxygen saturation probe?
- What are you oxygen levels and heart rate at rest?

Social support, medications and food provision

Do you live alone?

showing

- Is someone at home or a carer able to support you at home while you are recovering?
- Do you have enough supply of your regular medications and food?

*For patients with long term breathlessness e.g. COPD, heart failure

CLINICAL FEATURES

Worsening breathlessness

Breathless on minimal exertion e.g. walking to the bathroom, walking up

the stairs

Fever > 72 hours

Breathless at rest

Worsening breathlessness after day 5 from onset of symptoms

New onset confusion

Unable to speak in full sentences

Uncontrolled blood glucose levels

Central chest pain lasting >15 minutes (not generalised body aches)

Action From Clinical Assessment

If 1 AMBER, refer to HOT clinic next day If 2 AMBER, refer for urgent F2F review on same day If 1 RED, refer for urgent F2F review on same day

You should always reassess with the same questions as the first assessment

Discharge from acute post COVID-19 telephone assessments if all the following criteria below are met:

- Symptom onset > 10 days
- Last fever >48 hours
- Improved breathlessness for two consecutive telephone reviews
- Improved symptoms and "feeling better"
- Diabetes: stable blood glucose levels





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THANK YOU FOR LISTENING

Prevention is better than cure







LONG COVID

Using a holistic approach to patient care

Nicholas Hart Professor of Respiratory & Critical Care Medicine Clinical Director Respiratory, Sleep and Critical Care Cardiorespiratory Critical Care Clinical Group St Thomas' Hospital Director of Research Delivery Guys & St Thomas' NHS Foundation Trust

- Report the presentations of mild and moderate COVID
- Report the categories of COVID
- Report the common symptoms of Long COVID
- Report the pathway for Long COVID patient referral across NHSL
- Report the current clinical and research activity
- Report of rehabilitation interventions
- Prevention is better than cure





In the UK SARS-CoV-2 has affected over four million people SARS-CoV-2 has resulted in 450 000 hospital admissions SARS-CoV-2 has a 30% in-hospital mortality

LONG COVID AFFECTS 5-10%





COVID CATEGORIES

- ACUTE COVID
- POST ACUTE COVID (Up to 12 weeks)
- LONG COVID (>12 weeks)
- *SELEQUAE OF ACUTE COVID (>12 weeks)

*respiratory, cardiac, renal, neurological and endocrine





Symptoms

Most commonly reported symptoms

Fatigue

Muscle ache

Muscle weakness

Joint pain

Lethargy

Impaired sleep quality

Breathlessness

Short-term memory loss

Slower thinking





POST HOSPITAL COVID RECOVERY STUDY

PHOSP: 92.8 % of patients had at least one persistent symptom with a median of 9 symptoms; <u>only 3/10 patients fully recovered by 5 months</u>.

PHOSP Cluster 1: very severe mental and physical health impairment with most obese, co-morbid (Respiratory and neuropsychiatric co-morbidities). Higher Deprivation.

PHOSP Cluster 2: severe mental and physical health impairment with most obese, comorbid ((Respiratory and neuropsychiatric co-morbidities). Lower Deprivation.

PHOSP Cluster 3: moderate mental and physical health impairment with pronounced cognitive impairment with male predominance, older, overweight, less co-morbid. Higher Deprivation.

PHOSP Cluster 4: mild mental and physical health impairment with male predominance, overweight, less co-morbid. Lower Deprivation.

Physical, cognitive and mental health impacts of COVID-19 following hospitalisation – a multi-centre prospective cohort study Evans R et al on behalf of PHOSP investigators; under review





GSTT APPROACH TO LONG COVID

- Combined service with Kings College Hospital
- Bespoke SEL referral form from GP to Hospital
- GP to undertake F2F consultation with physiological tests, blood tests, ECG and CXR as standard
- It is important that all patients are reviewed in the right clinic
- GSTT has developed a therapy led-medically supported
- Clinic runs 3 times a week with a medical 'wash-up' once per week
- Clinic is holistic, needs-based and person-centred





THANK YOU FOR LISTENING

Prevention is better than cure







Long COVID and Rehabilitation Assessment, Treatments & Outcome Measures

Alexandra Curtis & Jacky Jones, Physiotherapy

GSTT Post COVID Assessment Pathways







Post COVID MDT assessment clinic

- Therapy led by occupational therapy and physiotherapy
- Psychology support
- Medical input from respiratory teams
- Patients are seen at the point of medical stability, >12 weeks after acute COVID. Persistent signs and symptoms consistent with post-COVID syndrome, not explained by an alternative diagnosis
- Face to face appointments rather than remote consultations. Aiming for inclusion for those without access to technology. Enhance the therapeutic effect by being seen and listened to in person
- Clinic launched Friday 30th April





Patient reported Outcome Measures (PROMS)

GSTT IMPARTs

Numerical Rating Scales for breathlessness

Numerical Rating Scales for cough

Numerical Rating Scales for fatigue

Numerical Rating Scales for pain

Numerical Rating Scales for sleep

Modified Medical Research Council Dyspnoea Score

Generalised Anxiety Disorder (GAD7)

Patient Health Questionnaire (PHQ9)

Trauma Screening Questionnaire (TSQ)

Smoking assessment

Post COVID Functional Status Scale

IMPARTS Cognition Question

Vocational Status

Multi-dimensional Assessment of Fatigue (MAF) EQ5D-5L

Nijmegen breathless questionnaire

WASA Work and social adjustment scale

- Completed prior to attending clinic
- Wide ranging measures due to the constellation of symptoms we expect to see





During the clinic visit

- Comprehensive specialist MDT holistic assessment will be based on the PROMS but more importantly the clinical interview
- Other assessments based on emerging data, could include but not limited to:
 - 1-min sit-to-stand test, looking for evidence of desaturation on exertion
 - Breathing Pattern Assessment, other centres report approx. 23% of patients do appear to have a breathing pattern disorder and this seems to be linked to post exertional malaise.
 - Lying and standing blood pressure, autonomic issues and feelings of dizziness are also prominent
 - Cognitive assessment

Aim of the clinic is advice, guidance and sign post onwards





On going evaluation and monitoring

- Health inequalities- we need to make the clinic as accessible as possible and will monitor who comes through the clinic against local demographic data
- Psychological effect of COVID and long COVID. Can the identified pathways can meet this need
- Identifying gaps in existing services for effective rehabilitation





Personalised Intervention

Mild (45%)

Moderate (17%) Severe (21%), V Severe (17%)

Persistent symptom burden (PHOSP-COVID Collaborative Group 2021)

Self-management

- Your COVID Recovery web based resource (Phase I)
- Healthcare professional supported self-management
- Your COVID Recovery web based resource (Phase II)
- MDT intervention
- Rehabilitation programme







Moderate

		Programme	Research Abo	ut Feedback Co	ntact Q
COVID Recovery	What is COVID-19?	Managing The Effects	Your Wellbeing	Your Road To Recovery	NHS

Supporting your recovery after COVID-19

As you find yourself recovering from COVID-19 you may still be coming to terms with the impact the virus has had on both your body and mind.

These changes should get better over time, some may take longer than others, but there are things you can do to help.

Your COVID Recovery helps you to understand what has happened and what you might expect as part of your recovery.















- Current understanding is limited
- Caution is suggested
- Low level initially and cut back if getting muscle ache, increased breathlessness, severe fatigue (Greenhalgh 2020)
- However likely to be clusters of presentations that respond better to a graded exercise approach





Pulmonary rehabilitation approach to COVID recovery



(Spielmanns 2021)





Fatigue reporting in a COPD population



Figure 1. Prevalence of patients with normal, mild and severe fatigue before and after PR, and change in fatigue level following PR. Data in the figure is presented as number of subjects (%). Abbreviations: PR, pulmonary rehabilitation.

(Van Herck et al 2019)





Local GSTT pilot data.

Aim

 To explore if a 6 week face-face rehabilitation course of exercise and education improves symptoms and function of patients post COVID-19

Results

10 patients completed the course







Physical outcomes







Conclusions

- We anticipate a high demand for the MDT assessment clinic
- More data is required to inform effective rehabilitation interventions
- Likely a stratified approach
- We look forward to further understanding the patients lived experiences and refining this clinic to support their needs







Health Seminar: Covid-19 and Vaccination Programme

Dr Ali Hashtroudi

Clinical Director of Occupational Health and Safety

04 May 2021

Natural immunity

• Antigen = part entering the body vs. Antibody = body reaction







How vaccine helps

Part of virus used as Antigen









The basis of all current vaccines







Different vaccines: how the Ag is introduced

Type of COVID vaccine	Example
Live	only Phase 1 COVI-VAC (India)
Inactivated	Covaxin (India), CoronaVac (sinoVac, China) and SinoPharm (China)) Valneva (France)
Protein-based subunit	Novavax & EpiVacCorona (India) Medicago (Canada- quadrivalent VLP)
mRNA	Pfizer BioNTech BNT162b2 mRNA Moderna mRNA-1273
Viral vectors	ChAdOx1 nCoV-19 (AZD1222) Janssen (Ad26.cov2.s) Sputnik V (rAd26_ rAd5)





The COVID vaccine cannot cause infection.

The COVID vaccine does not change the genetic material.





Research

- Pfizer: 44000, age 12 and above, 95% protective
- AstraZeneca: 24000, age 18 and above, 70%
- Moderna: 31000, age 18 and above, 94%
- Different countries
- Not difference in efficacy between age group, gender, ethnicity, medical conditions
- Not difference in safety between age group, gender, ethnicity, medical conditions







Protection against severe disease

- Protection against catching the disease
- Protection against passing the disease
- Laboratory based measurement





Schedule

- Two doses, 4 to 12 weeks apart
- The efficacy is the same





Who?

- JCVI rules
- Age (now >42)
- Vulnerability (clinically extremely vulnerable, at risk)
- Responsibility (work, carers)







How

• National booking system

https://www.nhs.uk/conditions/coro navirus-covid-19/coronavirusvaccination/book-coronavirusvaccination/

- Pharmacies
- Vaccine centres
- Hospital hubs
- PCN







Why not?

- Side effects
- Limited research, speed
- Pregnancy
- Non-clinical
- Myth, fake news







Side effects

- Like all vaccines, can cause side effects
- Not everybody
- <u>Most side effects are mild or moderate and go away</u> within a few days
- Paracetamol
- 48 million people vaccinated in the UK, >200000 by GSTT
- Exceptionally rare to have lasting / concerning side effect





Side effects - Pfizer

- Very common (>10% of people): pain at injection site, tiredness, headache, muscle pain, chills, joint pain, fever
- Common (<10% of people): injection site swelling or redness, nausea
- Uncommon (<1% of people): enlarged lymph nodes, feeling unwell
- Rare side effects (<0.1% of people): temporary one sided facial drooping





Side effects – AstraZeneca

- Very common (>10% of people): tenderness, pain, warmth, itching or bruising where the injection is given, generally feeling unwell, feeling tired, chills or feeling feverish, headache, nausea, joint pain or muscle ache
- Common (<10% of people): swelling, redness or a lump at the injection site, fever, vomiting or diarrhoea, flu-like symptoms (high temperature, sore throat, runny nose, cough and chills)
- Uncommon (<1% of people): feeling dizzy, decreased appetite, abdominal pain, enlarged lymph nodes, excessive sweating, itchy skin or rash
- Rare side effects (<0.1% of people): severe allergic reaction (anaphylaxis)





Allergy and anaphylaxis

- Pfizer and Moderna
- History of immediate onset-unexplained anaphylaxis or anaphylaxis to multiple classes of drugs or an unexplained anaphylaxis
- PEG
- History of reaction to food, venom, one medicine need NOT to be excluded from Pfizer or Moderna





Blood clot

- AstraZeneca
- Blood clot <u>AND</u> low platelet
- Extremely rare, 5 in a million
- Risk of COVID vs. risk of side effect (< 30 not offered AZ)
- History (including after the first dose) of blood clot <u>NOT</u> more at risk

So previous clot e.g. DVT, PE, stroke, heart attack, family history of these NOT a contraindication

Low platelet without blood clot is <u>NOT</u> a contraindication





Blood clot, symptoms

Seek urgent medical advice if experience any of the following symptoms more than 4 days and within 28 days of coronavirus vaccination:

- New onset of severe headache, which is getting worse and does not respond to simple painkillers
- An unusual headache which seems worse when lying down or bending over, or may be accompanied by blurred vision, nausea and vomiting, difficulty with speech, weakness, drowsiness or seizures
- New unexplained pinprick bruising or bleeding
- Shortness of breath, chest pain, leg swelling or persistent abdominal pain





Pregnancy (FAQ)

- No known risk associated with giving non-live vaccines, such as the COVID-19 vaccine
- Vaccines cannot cause infection in either the mother or foetus
- Available data do not indicate any safety concern but insufficient evidence to recommend routine use of COVID-19 vaccines during pregnancy
- Pfizer / Moderna are preferred choice, AZ can be done too
- Should be considered in pregnancy if:
 - Risk of exposure to COVID is high e.g. working in health and social care
 - Underlying conditions
- <u>RCOG</u>





Breast feeding (FAQ)

- No known risk associated with giving non-live vaccines such as COVID-19 vaccines
- Vaccination may be offered to breastfeeding women
- No safety data
- <u>RCOG</u>







- There is no evidence that the COVID-19 vaccine or any other vaccines can cause impotence or affect ability to conceive, fertility
- <u>The British Fertility Society and Association of</u> <u>Reproductive and Clinical Scientists</u> document





Speed of developing the vaccines

- Massive coordinated international effort
- Biggest trials we have seen
- Thousands of people volunteered to participate
- Scientists and trial doctors worked all hours and at speed
- Regulatory agencies worked extra hours and extra speed
- Manufacturers produced at same time as testing was occurring
- Good experience in developing vaccines quickly e.g. Ebola





Religious / believes

- No porcine product
- No animal product or egg
- No problem having during fast time





The best and quickest way out of the pandemic is by getting the vaccine

Thank you



