# **"THOUGHTS THAT COUNT"**

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#### PSYCHOLOGICAL PROCESSES AND OUTCOMES IN RECOVERY FROM COVID-19







# (NEURO) PSYCHOLOGICAL IMPACTS OF COVID-19

> UK Office of National Statistics COVID-19 Study (May 2022 data):

- > 2.8% of UK population currently experiencing PACS > 120 days.
  - > 57% of those experiencing symptoms for > 1 year.
- > 4.3% to 7.3% of *double-vaccinated* people report 'activity limiting' PACS > 90 days after infection with Omicron BA.I
- $> 3^{rd}$  most commonly reported issue (after fatigue and shortness of breath) is cognitive dysfunction.
- High levels of anxiety, depression and post-traumatic stress disorder amongst those attending (adult) PACS clinics.\*
  - > 74% with a GAD score > 8 (clinical anxiety).
  - > 73% with a PHQ-9 score > 10 (clinical depression).
  - > 15% with trauma symptoms indicative of PTSD.
  - > 14% with suicidal ideation.
  - > Sleep disturbance commonly reported.
  - > Other significant impacts, including relationship difficulties, difficulties coping, impacts on children and dependents.

• \*\* Please note that this is almost entirely data for adults.

 <sup>\*</sup> Data kindly provided by Liverpool Heart & Lung NHS Trust

#Longcovid2022

"I'm never going to get better!"

THINGS THAT CAN IMPACT UPON "Nobody is taking me **PSYCHOLOGICAL RECOVERY** seriously" Actual physical recovery. "It's just like a cold, no big Availability of structured rehabilitative and psychological supports. deal" • Presence of Acute Respiratory Distress (PTSD, additional issues) "My doctor says there's nothing Breathing support, admission to ICU, presence of delirium in ICU. wrong with me" Expectations of recovery (person themselves and others). 'Am I going to Minimisation of impact of COVID-19 have a heart attack?! Am I Pressure to return to work/exercise/family roles quickly developing Catastrophic interpretation of symptoms/fear of triggering further symptoms Self-blame (or stigma from others) at catching COVID-19. "What if I • Psychosocial supports and resources (social, financial, make myself worse?" occupational, previous physical and mental health).

"I'm the one who looks after everyone else, I feel

"I can't go to work and provide for my family, I need to "man up""

"It's my own stupid fault, I should have been more careful"

# WHAT WE THINK CAN HELP

- People are likely to need support with BOTH the process (expectations, adjustment, navigating services) and the outcomes (managing symptoms of distress).
- Clients need access to timely, good quality information and well-informed MDT support to help them (and others) form realistic expectations of their recovery, as well as to help manage their symptoms.
- Experience suggests that positive contact with others who are experiencing similar issues is *extremely* helpful for that process- making coordinated group- and peer-based approaches of particular benefit.
- 'The process' may not be the same for everyone. We need to give particular consideration to support for Māori.
- There is strong research evidence for individual and group-based programmes to help people manage many of the commonly reported issues:
  - Anxiety, depression, sleep, fatigue management, breathlessness, pain, managing cognitive issues, adjustment to injury.
  - Research studies are not specific to PACS (too early) but anecdotally clinics/services overseas report good success with these approaches.
- Given the high number of people experiencing symptoms of trauma/PTSD, suicidal ideation and cognitive difficulties, many clients may need to be supported to access more intensive (neuro) psychological interventions.



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