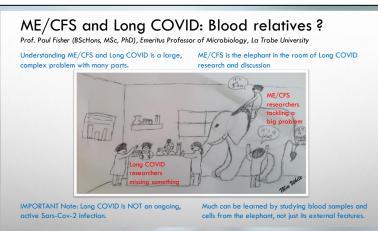
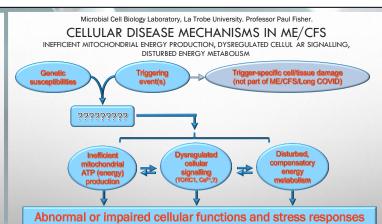


## ME/CFS and Long COVID: blood relatives?

Emeritus Professor Paul Fisher, La Trobe University

ME/CFS research has been neglected and is the elephant in the room of Long COVID and post-infection disease research





So far, ME/CFS and Long COVID share the same clinical presentation and underlying biology

## Comparison of symptoms in ME/CFS and Long COVID (modified from Komoroff and Lipkin, 2023)

Post-exertional malaise Impaired reasoning Secondary depression Decreased smell and taste Myalgia/arthralgia Chemical sensitivities

- Excludes known long term damage/scarring from acute infection or triggering event.
   Decreased smell and taste in Long COVID have now been attributed to SARS-CoV-2 causing damage/loss of cells that support sensory neurons in the nacial macos, so not part of the Long COVID
   S. Every individual experiences a different constellation of symptoms
   Presemb less common

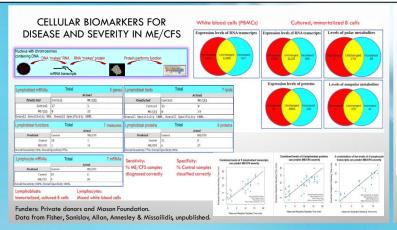
## Underlying biological abnormalities in ME/CFS and Long COVID (Data from Komoroff and Lipkin, 2023 and other sources)

Abnormality	ME/CFS (With number of abnormal subcategories)	Long COVID (With number of abnormal subcategories)
Neurological/neuromuscular abnormalities	✓ 23/24 (1 not tested)	✓ 20/24 (1 -ve, 3 not tested)
Immunological abnormalities	✓ 18/18	✓ 11/18 (1 -ve, 6 not tested)
Viral reactivation (EBV, HHV6,HHV7,CMV)	✓	✓
Gut microbiome dysbiosis	✓	✓
Metabolic and cell signalling abnormalities (energy metabolism, hypometabolic state, redox imbalance, kynurenine pathway, cell stress signalling, Ca <sup>2</sup> signalling)	<b>√</b> 15/15	✓ 10/15 (5 not tested)
Cardiopulmonary/vascular abnormalities	<b>√</b> 14/14	✓ 6/14 (8 not tested)

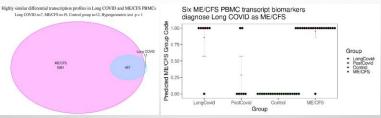
- In most subcategories there were a minority of dissenting studies for one or both illnesses

  From a total of 73 subcategories of biological abnormality, only 2 not present in both illnesses (each tested in a
  single Long COVID study), 22 not yet tested in Long COVID and 1 not yet tested in ME/CFS.

Small panels from thousands of blood markers of ME/CFS predict illness severity and diagnose ME/CFS and Long COVID



ME/CFS AND LONG COVID MOLECULAR SIGNATURE MAY BE IDENTICAL - ME/CFS BIOMARKERS WORK FOR LONG COVID.



Note: Number of RNAs with significantly abnormal levels was lower in Long COVID because small number of Long COVID samples (r=7) limited the statistical power of the test. Data from Fisher, Sanislay, Allan, Missailidis, Gras & Annesley, unpublish

The AusME Registry and Biobank (La Trobe Uni, Emerge Australia et al, Mason Foundation funds) provides an easier path into Australian ME/CFS/Long COVID research - 13 active research projects from 5 states/territories, 3 blood processing sites, partnering 9 current funding applications, 1520 registrants, 126 blood samples.....and growing.

All that is needed to hasten progress in diagnosis, understanding and treatment of ME/CFS (and Long COVID) is a fair share of biomedical research funds.