**Update from Linda Coy**

The Secretary of State for Health and Social Care, Sajid David, noted on 19/11/21 that more needs to be done, so will focus now be set on real funding for biomed research for ME, training for all medical professionals and a scheme to report harm?I was thinking about using comments from the Secretary of State for Health and Social Care, Sajid Javid, who stated on Twitter 19/11/21  'We need to do more on ME/Chronic Fatigue Syndrome. There’s been a real lack of research over many years.'

I did email him in September and hope it helped spur the meeting he had with Chris Pontin, Nancy Klimas, Sonya Chowdry and Professors Steven Holgate and Karl Johan Tronstad.

As you know we need funding for biomed research & prevention of harm to people with ME, training for all medical people and a scheme to report harm.  No more money should go to Sir Simon Wessely, Prof Michael Sharp, Trudie Chalder or Prof Peter White who have ruined the lives of millions of ME patients worldwide for decades.

So - a question, lots to cover but need to be concise.

How about:

In view of the recent review of the NICE Guidance for the treatment of ME when will the UK Government condemn that section of the British Psychological Society who have ruined the lives of millions of ME patients worldwide for decades, despite their own Code of Ethics and Conduct focussing on 4 primary principles of respect, competence, responsibility, and integrity.

Focus must be shifted to set up real funding for biomed research for ME, training for all medical professionals and a scheme to report harm.