#MILLIONSMISSING

is a global campaign for ME health equality

Myalgic Encephalomyelitis (ME), is a devastating, complex, multi-system disease, which affects the immune system, nervous system, endocrine system and metabolism.

WE DEMAND THE FOLLOWING



Parity of research funding:

The government must urgently invest in biomedical research after decades of neglecting people with ME. Funding for research must reflect the severity of ME and its prevalence in the UK.



Suspend GET and CBT treatments



Update Professional Training



Stop Unjustified Child Protection Proceedings

#MillionsMissing

#CanYouSeeMEnow?



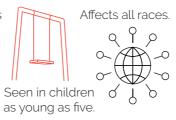
FACTS ABOUT ME

Myalgic encephalomyelitis

WHO IS AT RISK?





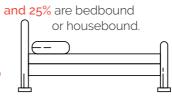




Affects between 15-30 million worldwide

HOW SERIOUS IS THE DISEASE?





Recovery is rare. estimated at just 5%, leaving many patients sick and disabled for the rest of their lives.

RESEARCH



ME receives a tiny percentage of the research funding awarded to other chronic illnesses - 20 times less than MS patients. Over the past few years, funding has decreased further.

TREATMENT



Graded Exercise Therapy (GET) and Cognitive Behavioural Therapy (CBT) are the main treatments recommended in the NHS.

GET is harming people with ME. CBT is ineffective.

CHII DRFN



Estimated 25,000 children with ME in the UK. ME is the leading cause of long-term school sickness absence in the UK. 1 in 5 families of children with ME are subjected to unjustified child protection proceedings.

TRAINING



GPs receive little to no training about ME.

Current continuing professional development courses are outdated and inaccurate