

Kritik af den danske Sundhedsstyrelses håndtering af ME i Danmark

I 2023 udgav Cand.med. lektor emeritus Bernard Jeune [en rapport](#) om den sundhedspolitiske og videnskabelige udvikling i og omkring sygdommen ME i Danmark

Per Julin korrekser Sundhedsstyrelsen om forholdene i Sverige

<https://me-foreningen.dk/wp-content/uploads/2021/04/NyRapport-Per-Julin-kopi.pdf>

Kristian Sommerfelt korrekser Sundhedsstyrelsen om forholdene i Norge

<https://me-foreningen.dk/wp-content/uploads/2021/03/SST-NV-.pdf>

Opfordring fra 97 internationale ME-forskere og -læger

<https://me-foreningen.dk/wp-content/uploads/2020/03/Open-Letter-Minister-of-HHealth-100-signatures-02-03-2020-1.pdf>

En lang fortælling om den miskendte sygdom Myalgia Encephalomyelitis (ME)

Bidrag til vores diskussion om ME i Enhedslistens Sundhedspolitisk udvalg

Bernard Jeune. Første udgave i marts 2023, sidste stærkt udvidede udgave i august 2023.

<https://drive.google.com/file/d/1PLQn9QYA75wlzcS47EVYV-9nxWgtavnk/view>

Myalgisk Encephalomyelitis (ME G93.3) Statusrapport august 2016. Oplæg fra ME Foreningen til Dialogmøde om funktionelle lidelser i Sundhedsstyrelsen d. 1. september 2016

Vibeke Vind, Rebecca Hansen, Catrine Schultz Engsig, D. 30/8-2016 udgave 1.0

ME-patienters møde med sundheds- og social-systemer

Do diagnostic criteria for ME matter to patient experience with services and interventions? Key results from an online RDS survey targeting fatigue patients in Norway.

Kielland A, Liu J, Jason LA *Journal of Health Psychology*. 2023;28(13):1189-1203

<https://doi.org/10.1177/13591053231169191>

Tjenesten og Meg Samlerapport fra et prosjekt om ME-sykes møte med den norske velferdsstaten

<https://www.fafo.no/images/pub/2023/969.pdf>

<https://www.fafo.no/zoo-publikasjoner/andre-fafo-utgivelser/tjenesten-og-meg-samlerapport>

[Implications of the quality of the doctor-patient relationship on health in adult ME/CFS patients. A qualitative public health study from a patient perspective]

Auswirkungen der Qualität der Arzt-Patient-Beziehung auf die Gesundheit von erwachsenen ME/CFS-Erkrankten : Eine qualitative Public-Health-Studie aus Patientensicht.

Habermann-Horstmeier L, Horstmeier LM. *MMW Fortschr Med*. 2023 Dec;165(Suppl 5):16-27. German. Doi: 10.1007/s15006-023-2894-z. PMID: 38062324.

<https://pubmed.ncbi.nlm.nih.gov/38062324/>

Identifying the mental health burden in Myalgic Encephalomyelitis/Chronic Fatigue Syndrome (ME/CFS) patients in Switzerland: A pilot study

Rahel Susanne König, Daniel Henry Paris, Marc Sollberger, Rea Tschopp, Heliyon, Volume 10, Issue 5, 2024, e27031, ISSN 2405-8440

<https://doi.org/10.1016/j.heliyon.2024.e27031>

Improving the experiences of people with ME/CFS: interim delivery plan

Department of Health and Social Care, Department for Education, and Department for Work and Pensions Published 9 August 2023, Last updated 4 September 2023

<https://www.gov.uk/government/consultations/improving-the-experiences-of-people-with-mecfs-interim-delivery-plan>

EMEA survey of ME/CFS patients in Europe: Same disease, different approaches and experiences. European MEAlliance (EMEA). Arild Angelsen and Trude Schei. 2024.

<https://www.europeanmealliance.org/documents/emeaeusurvey/EMEAMEsurveyreport2024.pdf>

<https://www.europeanmealliance.org/emea-pan-european-survey-uk.shtml>

Press release: A Shocking Indictment of European Research and Healthcare Policies for Myalgic Encephalomyelitis <https://www.europeanmealliance.org/emea-pan-european-survey-pr-uk.shtml>