

Thema: ME/CFS: Zahnmedizin unter prekären Umständen für schwer erkrankte Patientengruppe

Autorin: Marlene Hartinger mit Inhalten der Deutschen Gesellschaft für ME/CFS e.V. (DG ME/CFS e.V.)

Literatur

- 1) Deutsche Gesellschaft für ME/CFS e.V. (DG ME/CFS e.V.), <https://www.mecfs.de/>
- 2) Daniell, J. *et al.* The rising cost of Long COVID and ME/CFS in Germany. (2025).
- 3) Falk Hvidberg, M., Brinth, L. S., Olesen, A. V., Petersen, K. D. & Ehlers, L. The health-related quality of life for patients with Myalgic Encephalomyelitis / Chronic Fatigue Syndrome (ME/CFS). *PloS One* 10, e0132421 (2015).
- 4) Institute of Medicine. *Beyond Myalgic Encephalomyelitis/Chronic Fatigue Syndrome: Redefining an Illness*. National Academies Press, Washington (DC), 2015. doi:10.17226/19012.
- 5) Conroy, K., Bhatia, S., Islam, M. & Jason, L. A. Homebound versus Bedridden Status among Those with Myalgic Encephalomyelitis/Chronic Fatigue Syndrome. *Healthcare* 9, 106 (2021).
- 6) Bateman, L. *et al.* Chronic fatigue syndrome and co-morbid and consequent conditions: evidence from a multi-site clinical epidemiology study. *Fatigue Biomed. Health Behav.* 3, 1–15 (2014).
- 7) Angelsen, A. & Schei, T. EMEA survey of ME/CFS patients in Europe: Same disease, different approaches and experiences. (2024).