

Cognitive Fitness in Aging (COFITAGE), the open-access release of a neuroscience dataset

Antoine Jacquemin^{1,2}, Jiqing Huang^{1,2}, Nikita Belyi^{1,3}, Christian Degueldre¹, François Meyer^{1,4}, Daphne Chylinski^{1,5}, Justinas Narbutas^{1,6}, Maxime Van Egroo^{1,7}, Eric Salmon⁸, Puneet Talwar^{1,9}, Fabienne Collette^{1,3}, Gilles Vandewalle^{1,9}, Christine Bastin^{1,3}, Mohamed Ali Bahri^{1,6,10,*}, Christophe Phillips^{1,2,*}

¹GIGA – CRC Human Imaging, University of Liège, Liège, Belgium

²Montefiore Institute, Department of Electrical Engineering and Computer Science, University of Liège, Liège, Belgium

³Department of Psychology, University of Liège, Liège, Belgium

⁴Memory Clinic, Department of Neurology, University Hospital of Liège, Liège, Belgium

⁵Department of Neuropsychology and Speech Therapy, Hôpital Universitaire de Bruxelles (H.U.B), Brussels, Belgium

⁶Department of Psychology and Neurosciences, Leibniz Research Centre for Working Environment and Human Factors (IfADo) Dortmund, Germany

⁷Faculty of Health, Medicine and Life Sciences, Mental Health and Neuroscience Research Institute, Alzheimer Centre Limburg, Maastricht University, Maastricht, The Netherlands

⁸Department of Clinical Sciences, University of Liège, Liège, Belgium

⁹Department of Biology, Ecology and Evolution, University of Liège, Liège, Belgium

¹⁰Department of Physics, University of Liège, Liège, Belgium

*These authors contributed equally as co-last authors.

Abstract

Understanding early brain aging requires integrating molecular, structural, physiological and cognitive measures within the same individuals. Subtle alterations in amyloid deposition, hippocampal structure, cortical microstructure and sleep-wake regulation may emerge years before clinical decline. However, few open-access datasets combine PET, MRI, sleep phenotyping, genetics and extensive cognitive assessment. The COFITAGE dataset addresses this gap by providing a multimodal, longitudinal resource in healthy late middle-aged adults, enabling investigation of interactions between amyloid and tau burden, brain structure, sleep physiology, cognitive reserve and cognitive trajectories.

The cohort includes 101 participants (33 men, 68 women) aged 50–69 years (mean 59.4 ± 5.3). Phenotypic data collected at baseline and 2-year follow-up include demographics, detailed sleep metrics (e.g., sleep stages, efficiency, awakenings) and a comprehensive neuropsychological battery covering memory, executive functions, attention, processing speed, language, mood and cognitive reserve proxies.

40 Baseline MRI acquisitions comprise anatomical (T1w, T2w), quantitative multi-
41 parameter mapping (MTsat, PD, T1 maps with field maps) and diffusion imag-
42 ing (multi-shell NODDI). PET imaging includes amyloid tracers (Florbetapir or
43 Flutemetamol) and tau-sensitive tracers. All data were anonymized and organized ac-
44 cording to the Brain Imaging Data Structure (BIDS) standard, ensuring reproducibility
45 and interoperability.

46
47 COFITAGE uniquely combines multimodal imaging, sleep and cognition within a
48 narrow age range, allowing investigation of preclinical brain aging. Its scientific value
49 is supported by multiple publications on sleep, Alzheimer’s biomarkers, cognition and
50 cognitive reserve. It constitutes a robust open-access resource for multimodal and
51 longitudinal research on early brain aging.