A new study led by the University of Portsmouth has identified that one of the major factors of age-related brain deterioration is the loss of a substance called myelin.

Myelin acts like the protective and insulating plastic casing around the electrical wires of the brain—called axons. Myelin is essential for superfast communication between nerve cells that lie behind the supercomputer power of the human brain.
The image depicts myelin (Cyan), OPCs (Magenta) and cell nuclei in blue in the Corpus Callosum of the brain. The Corpus callosum is one of the biggest tracts of white matter and it connects the left and right hemisphere of the brain together. It is also one of the white matter tracts affected by aging. Credit: Dr Andrea Rivera

"A key feature of the ageing brain is the progressive loss of white matter and myelin, but the reasons behind these processes are largely unknown. The brain cells that produce myelin—called oligodendrocytes—need to be replaced throughout life by stem cells called oligodendrocyte precursors. If this fails, then there is a loss of myelin and white matter, resulting in devastating effects on brain function and cognitive decline. An exciting new finding of our study is that we have uncovered one of the reasons that this process is slowed down in the aging brain."

Dr. Rivera, lead author of the study while he was in University of Portsmouth and who is now a Fellow at the University of Padua, explained: "By comparing the genome of a young mouse brain to that of a senile mouse, we identified which processes are affected by ageing. These very sophisticated analysis allowed us to unravel the reasons why the replenishment of oligodendrocytes and the myelin they produce is reduced in the aging brain.

"We identified GPR17, the gene associated to these specific precursors, as the most affected gene in the ageing brain and that the loss of GPR17 is associated to a reduced ability of these precursors to actively work to replace the lost myelin."

The work is still very much ongoing and has paved the way for new studies on how to induce the 'rejuvenation' of oligodendrocyte precursor cells to efficiently replenish lost white matter.

Dr. Azim of the University of Dusseldorf said: "This approach is promising for targeting myelin loss in the aging brain and demyelination diseases,"
including Multiple Sclerosis, Alzheimer's disease and neuropsychiatric disorders. Indeed, we have only touched the tip of the iceberg and future investigation from our research groups aim to bring our findings into human translational settings.”

Dr. Rivera performed the key experiments published in this study while at the University of Portsmouth and he has been awarded the prestigious MSCA Seal of Excellence @UniPD Fellowship to translate these findings and investigate this further in the human brain, in collaboration with Professors Raffele De Caro, Andrea Porzionato and Veronica Macchi at the Institute of Human Anatomy of the University of Padua.

The study was funded by grants from the BBSRC and MRC to Professor Butt, together with the UK and Italian MS Societies (to Professors Butt and Abbracchio, respectively), and the Swiss National Funds Fellowship and German Research Council (Dr. Azim). Dr. Andrea Rivera was supported by an Anatomical Society Ph.D. Studentship (with Professor Butt), and the MSCA Seal of Excellence @UniPD (Dr. Rivera).

Dr. Emma Gray, Assistant Director of Research at the MS Society, said: “MS can be relentless and painful, and there are sadly still no treatments to stop disability progression. We can see a future where no one has to worry about MS getting worse but, for that to happen, we need to find ways to repair damaged myelin. This research sheds light on why cells that drive myelin repair become less efficient as we age, and we’re really proud to have helped fund it. By improving our understanding of ageing brain stem cells, it gives us a new target to help slow the progression of MS, and could have important implications for future treatment.”
