Rapid improvement in Alzheimer’s disease symptoms following fecal microbiota transplantation: a case report

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Abstract
Alzheimer’s disease (AD), the most common form of dementia, is a leading cause of death and a major cause of morbidity in older people. The disease is characterized by progressive memory loss, cognitive impairment, and the cerebral accumulation of amyloid-β peptide. Given the health and economic impacts of AD, treatments that target the underlying etiology of AD or modify the course of the disease are of significant interest. The gut microbiome has been increasingly implicated in the pathogenesis of several neurological diseases, including multiple sclerosis and Parkinson’s disease. Furthermore, emerging evidence has demonstrated that there are alterations in gut microbiome composition in patients with AD, suggesting involvement of the microbiome–gut–brain axis. We present symptom improvement in a patient with AD following fecal microbiota transplantation for a *Clostridioides difficile* infection.

Keywords
Alzheimer’s disease, fecal microbiota transplantation, gastrointestinal microbiome, microbiota, neuroinflammation, *Clostridioides difficile*

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Introduction
Alzheimer’s disease (AD) is a devastating neurodegenerative disease characterized by a deterioration in memory and other cognitive domains and the cerebral accumulation of amyloid-β peptide. Advancing age is the...
most significant risk factor for the disease, with incidence doubling every 5 years after the age of 65.\textsuperscript{1} Approximately 46.8 million people worldwide currently have AD or a related dementia.\textsuperscript{2} However, this number is expected to increase exponentially in the coming years, largely as a result of the ageing population.\textsuperscript{5} The associated economic burden of AD is substantial; in 2015, the total estimated worldwide cost of dementia was US$818 billion, which is projected to rise to US$2.0 trillion by 2030 because of the increased prevalence of AD.\textsuperscript{2} However, despite decades of research, the etiology of AD remains unknown and there are currently no preventative or disease-modifying treatments.

A growing body of experimental and clinical data implicates the gut microbiome in the pathogenesis of several neurological conditions, including autism spectrum disorder (ASD),\textsuperscript{3} Parkinson’s disease (PD),\textsuperscript{4} and multiple sclerosis (MS).\textsuperscript{5,6} More recently, alterations in gut microbiome composition have been observed in patients with AD,\textsuperscript{7} suggesting a potential role for the microbiome in AD pathogenesis. This hypothesis has been supported by animal models;\textsuperscript{8,9} for example, germ-free amyloid-\(\beta\) precursor protein transgenic mice have markedly less cerebral amyloid-\(\beta\) pathology compared with control mice with intestinal bacteria.\textsuperscript{8} Certain bacteria within the gut microbiota are also capable of secreting large amounts of amyloids and lipopolysaccharides,\textsuperscript{10} a hallmark feature of AD.

Fecal microbiota transplantation (FMT) is the infusion of fecal material from a healthy donor into the gastrointestinal tract of an individual with disease. This procedure represents the most powerful means of modulating the gut microbiome. The therapy has risen to prominence in the last decade following several outbreaks of severe \textit{Clostridioides difficile} infection (CDI) in North America and Europe, which were caused in part by the emergence of the hypervirulent NAP1/B1/027 strain. In these difficult-to-treat cases as well as in subsequent randomized, controlled trials, FMT has consistently achieved cure rates >90\%.\textsuperscript{11} Serendipitous improvements following FMT have since been reported in a number of extra-intestinal conditions, including ASD,\textsuperscript{3} MS,\textsuperscript{12,13} and myoclonus dystonia.\textsuperscript{14} To our knowledge, ours is the first report of a patient who experienced rapid improvement in their AD symptoms following FMT for recurrent CDI.

**Case report**

An 82-year-old man presented for opinion and management of recurrent CDI following hospitalization for methicillin-resistant \textit{Staphylococcus aureus} pneumonia. The patient had previously failed several courses of antibiotics for CDI, including vancomycin, vancomycin with metronidazole, fidaxomicin, and bezlotoxumab, with relapse confirmed via symptom recurrence and positive stool test.

At the time of presentation, the patient was under the care of his primary care physician and his neurologist for the treatment of AD, following a gradual 5-year decline in memory and cognition. The patient was taking memantine (28 mg once daily) and donepezil (23 mg once daily). The patient’s dementia symptoms included confusion, memory loss, depression, and flattened affect. On his most recent Mini-Mental State Examination (MMSE) administered by the neurologist, the patient scored 20, indicating mild cognitive impairment. This result reflected the gastroenterologist’s findings and was within the expected range for patients with AD. The patient’s wife reported that he no longer appeared to enjoy socializing, and required considerable assistance with basic tasks such as food preparation, bathing, and taking his medication. Neuropsychiatric testing revealed significant impairments in the areas of
memory and semantic language abilities, nonverbal learning, and divided attention and response inhibition. His scores in these areas were within the first through fifth percentiles for function.

Following a detailed discussion regarding the potential risks and benefits associated with the procedure, the patient underwent a single 300 mL FMT infusion (per the Borody method) using stool from the patient’s 85-year-old wife as a donor. The patient’s wife was intellectually acute, with normal affect and stable mood. Following the procedure, the patient’s CDI symptoms resolved, and repeat stool testing 2 months later was negative.

At the follow-up visit 2 months post-FMT, the patient’s wife reported improvements in the patient’s mental acuity and affect. The MMSE was re-administered by the gastroenterologist (and subsequently by the neurologist) and the patient scored 26, indicating normal cognition. Four months post-FMT, the patient reported continued improvement in memory, with no progression in symptoms. The patient now remembered his daughter’s birthday, which he had not been able to recall previously, and was able to correct the physician’s recollections of his symptoms. Six months post-FMT, the patient reported a marked improvement in mood, was more interactive, and showed more expressive affect. Readministration of the MMSE revealed that the patient’s score had further increased to 29.

Because no study was conducted, no ethics approval or consent was required. Verbal consent was obtained from the patient for the publication of the report; however, this was merely a courtesy because all data have been de-identified.

**Discussion**

To our knowledge, this is the first report of a case of rapid reversal of AD symptoms in a patient following FMT for recurrent CDI. Improvements in AD symptoms occurred as early as 2 months post-FMT and continued to the 6-month follow-up visit (the date of the last follow-up), with no noted reversion of symptoms. The resolution of symptoms occurred in a stepwise manner: first, there was increased mental acuity and improved affect observed at 2 months post-FMT, and this was followed by marked improvements in memory and mood by 4 and 6 months post-FMT, which were accompanied by more expressive affect. These findings paralleled improvements in the patient’s MMSE scores, which increased to within the range of normal cognition by 2 months post-FMT. Eradication of CDI was also confirmed at the 2-month follow-up visit, via the resolution of symptoms and a negative stool test.

The central role of the gut microbiome in neurological dysfunction has been increasingly recognized, and alterations in gut microbiome composition have consistently been reported in ASD, PD, and MS. Rapid and dramatic improvements have also been observed in several of these conditions following manipulation of the gut microbiome, including with FMT, which further supports a causal association. In an open-label study in 18 children with ASD, 2 weeks of vancomycin treatment followed by daily FMT infusions for 7 to 8 weeks resulted in significant improvements in core ASD symptoms, which persisted at the 2-year follow-up. Furthermore, in a case series of three patients with MS and underlying gastrointestinal symptoms, 5 to 10 days of FMT infusions resulted in a profound reversal of major neurological symptoms, and led to longstanding disease remission. Similarly, a 61-year old female with secondary progressive MS achieved disease stability for over 10 years following FMT for recurrent CDI. In addition, in a patient with myoclonus dystonia, significant
improvement in tremor and functional deficits were reported following the manipulation of gut microbiota with vancomycin, rifaximin, and metronidazole for chronic diarrhoea.\textsuperscript{14} In another study, a 71-year old male with longstanding constipation and PD experienced a rapid and dramatic reduction in PD symptoms, including an absence of persistent tremors, glabellar tap reflex, and cogwheel rigidity, when his constipation was treated with vancomycin, metronidazole, and colchicine.\textsuperscript{16}

Gut bacteria may contribute to the pathogenesis of AD via a number of mechanisms. Microbe-mediated bidirectional communication pathways exist between the gut and the central nervous system, and many of the neurotransmitters that regulate mood and cognition in the brain, such as gamma-aminobutyric acid (GABA) and dopamine, are also synthesized and catabolized by gut bacteria. GABA dysfunction has been shown to play a role in AD,\textsuperscript{17} and abnormally high GABA concentrations have recently been identified in reactive astrocytes of post-mortem AD patients.\textsuperscript{18} Molecular mimicry is a well-documented strategy that is commonly employed by pathogens to gain a competitive advantage over their host. It occurs when similarities between microbial proteins and host peptides result in the cross-activation of autoreactive T or B cells and a loss of self-tolerance.\textsuperscript{19} Rheumatic fever after \textit{Streptococcus pyogenes} infection is a classic example of this; antigen cross-reactivity between streptococcal M protein and cardiac myosin results in the targeting of myocardial tissue.\textsuperscript{19} However, a number of other well-documented molecular mimicry events exist, and a recent database study catalogued 261 validated host–microbial mimicry interactions.\textsuperscript{20} Although the focus to date has been on the role of molecular mimicry as a cause of autoimmune diseases, this mechanism may also be exploited in the development of AD. The progressive accumulation of \(\beta\)-amyloid protein plaques between neurons represents a hallmark of AD. However, the initiating factor responsible for the development and propagation of these prion-like proteins is unknown. Amyloids are secreted by a wide range of pathogenic and non-pathogenic bacteria, and play an important role in cell adhesion and biofilm formation.\textsuperscript{10} Similarities between the tertiary structure of microbe-derived amyloids and \(\beta\)-amyloid in AD may trigger antigen cross-reactivity, thus potentially initiating the formation of AD. Interestingly, both microbe-derived amyloids (such as curli) and AD-related \(\beta\)-amyloid are recognized by the same Toll-like receptor (TLR)2/TLR1 complex.\textsuperscript{21} Finally, dysbiosis itself may contribute to the development of AD. A dysbiosis-induced increase in gut permeability may potentially lead to persistent systemic inflammation, disruption of the blood–brain barrier, and ultimately neurodegeneration.

Neuroinflammation plays a central role in the pathogenesis of AD,\textsuperscript{22,23} and is characterized by elevated levels of pro-inflammatory cytokines,\textsuperscript{23} nuclear factor \(\kappa\)B signalling,\textsuperscript{24} and aggregation of activated microglia in damaged areas.\textsuperscript{25} The influence of the gut microbiome on neuroinflammation in AD remains poorly understood, as do the exact mechanisms of action of FMT in this disease. However, the potent immunomodulatory effects of FMT are well recognized. FMT has been demonstrated to ameliorate chronic intestinal colitis in both humans and animal models via the downregulation of pro-inflammatory cytokines, promotion of anti-inflammatory responses, and inhibition of nuclear factor \(\kappa\)B activity.\textsuperscript{26–28} In addition, similarly beneficial effects have since been observed in several other conditions with an inflammatory component.\textsuperscript{13,29,30} Thus, FMT may have a positive effect on cognitive function in AD via alterations in the levels of circulating cytokines.
Alternatively, by restoring the previously impaired intestinal barrier function, FMT may prevent the translocation of neuroactive compounds and metabolites within the central nervous system that regulate mood and cognition and contribute to inflammation. However, further studies are urgently required to elucidate the exact mechanisms by which FMT may ameliorate symptoms in AD.

To our knowledge, this is the first documented case of a rapid reversal of AD symptoms in a patient following FMT for recurrent CDI. Although this is only a preliminary report, the remarkable resolution of AD symptoms following FMT for CDI is instructive and adds to the evidence that suggests a causal association between the gut microbiome and neurological dysfunction. Given the probable role of the gut microbiome in the pathogenesis of AD, modulation of the microbiome represents a promising avenue of treatment. A randomized, double-blind, placebo-controlled trial is currently underway to evaluate the efficacy of oral FMT in AD, the results of which are eagerly anticipated.

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Sabine Hazan has a pecuniary interest in Ventura Clinical Trials and ProgenaBiome.

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