



Visualizing Dementia's Impact: Analyzing MRI Scans to Identify Key Brain Regions through Python-Based Image Comparison

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Abstract:

MRI scans of dementia patients were analyzed to identify commonalities and differences in specific areas when compared to non-demented patients. Using Python, average images of the brain scans for both groups were created, and the differences were plotted to visually highlight regions of significant variation. This approach identified critical areas affected by dementia, such as the hippocampus and ventricles, which display distinct patterns of atrophy and enlargement, consistent with previous studies. By generating these comparative images, this analysis aimed to enhance understanding of how dementia structurally impacts the brain, providing a visual representation that can aid in more accurate diagnosis and assessment of the disease's progression. These visual insights contribute to the broader goal of utilizing computational methods to pinpoint early indicators of dementia, offering potential pathways for future AI models to further refine the detection and prediction of this condition.

Introduction:

Alzheimer's disease (AD) is a progressive neurodegenerative disorder that primarily affects the elderly, leading to cognitive decline and memory loss[1]. It is the most common cause of dementia, accounting for an estimated 60-80% of cases worldwide [1]. The diagnosis of Alzheimer's disease often occurs at a later stage of the disease progression when significant brain damage has already occurred, making treatment and intervention less effective [2].

Current diagnostic methods for Alzheimer's disease include clinical evaluations, cognitive testing, neuroimaging, and biomarker analysis from cerebrospinal fluid (CSF) and blood samples [3,4,5]. While these methods are helpful, they each have notable limitations. Clinical evaluations and cognitive testing are inherently subjective, with outcomes often dependent on the clinician's expertise. Furthermore, these methods are less effective at detecting early-stage Alzheimer's when symptoms are mild [3]. Neuroimaging techniques, such as magnetic resonance imaging (MRI) and positron emission tomography (PET) scans, can identify brain changes linked to Alzheimer's, but they are costly, not widely accessible, and may only be helpful to detect late-stage dementia [4]. Biomarker analysis from CSF and blood samples, while promising, can be invasive and is not always conclusive in providing a definitive diagnosis [5].

The early detection of Alzheimer's disease is essential for several reasons. Early intervention has been shown to slow disease progression, improving patients' quality of life [6]. Identifying Alzheimer's in its initial stages also allows patients and their families to plan for the future, seek support, and make informed decisions regarding long-term care [7]. Additionally, the identification of early-stage patients is critical for advancing research and development, as these patients can participate in clinical trials that test new treatments and therapies [8].

The primary objective of this project is to analyze patterns in MRI scans of patients with and without dementia and to understand patterns in patients with dementia.

Literature Review:

Neuroanatomical differences between dementia patients and those without dementia are well-documented in the literature, primarily in relation to brain structure [3]. These structural changes are critical for understanding the progression of dementia and for improving diagnostic accuracy through imaging techniques like MRI. One of the most affected regions is the hippocampus, a key area for memory formation, which is often among the first regions impacted in dementia, particularly in Alzheimer's disease [3]. Studies consistently show significant atrophy in the hippocampus and surrounding medial temporal lobe structures in patients with dementia compared to healthy individuals [1,2]. This atrophy correlates closely with the degree of cognitive impairment, as more advanced stages of dementia are associated with more severe hippocampal shrinkage [3].

Beyond hippocampal atrophy, other brain regions such as the entorhinal cortex, amygdala, and parietal lobes, also exhibit varying degrees of shrinkage in dementia patients. These structural changes contribute directly to hallmark symptoms of dementia, including memory loss, disorientation, and difficulties with spatial awareness and navigation [1]. Additionally, white matter lesions and a general reduction in brain volume have been reported in patients with dementia, further distinguishing them from non-dementia individuals and providing additional markers for disease progression [5].

Understanding these neuroanatomical differences is crucial for interpreting MRI scans within the context of dementia diagnosis. The identification of specific patterns of brain atrophy and other structural changes allows for more precise differentiation between types of dementia, as well as for tracking the progression of the disease over time [12].

In analyzing the MRI images of dementia and non-dementia patients across sagittal, transverse, and coronal views, distinct neuroanatomical differences that highlight the impact of dementia on the brain. The coronal view consistently revealed pronounced atrophy in dementia patients, particularly in the hippocampus and surrounding medial temporal lobes—regions critical for memory formation and cognitive function. This atrophy is coupled by significantly enlarged ventricles, a common occurrence in many forms of dementia that reflects the substantial loss of brain tissue in adjacent areas. The enlargement of the ventricles is most evident in the transverse and coronal views, where the increased size results in wider spaces filled with cerebrospinal fluid, visible as darkened regions around the ventricles. These changes are stark when compared to the brains of non-dementia patients, which show smaller ventricular spaces and well-preserved cortical thickness, indicating healthier brain structures. The visible degeneration and tissue loss underscore how dementia affects critical brain regions and contributes to the cognitive decline characteristic of the disease. These MRI differences provide crucial insights into the neurodegenerative processes underlying dementia, illustrating the profound structural impact of the condition and reinforcing the importance of advanced imaging techniques in early diagnosis, monitoring, and understanding the progression of dementia.

Methods and Results:

MRI and clinical patient data were obtained from the Oasis 1 dataset [12]. Data for subjects 22, 24, 26, 29, 34, 28, 30, 33, 35, and 39 were used. Subjects with a dementia score ≥ 1 were classified as dementia patients, while all others were classified as non-dementia patients. Images of both non-dementia and dementia patients were plotted to compare differences that could indicate affected areas of the brain. The MRI scans were averaged for each view for both non-dementia and dementia patients. Subsequently, the average dementia images were subtracted from the average non-dementia images to create an average image from each category from three different perspectives.

A caveat in this analysis is that the images for dementia and non-dementia patients were not perfectly aligned, leading to conflicting skull outlines, which made the average images blurry and the analysis harder to interpret. It is important to note that raw data were used to create the average images; using processed data would have simplified and improved the accuracy of this process. Specific tools, such as FreeSurfer software, would have enabled better alignment of the images, resulting in a clearer analysis.

Discussion:

The next step of this project would be to develop an Artificial Neural Network (ANN) specifically to detect dementia. This would involve expanding the dataset to include more diverse MRI scans and patient information. An ANN would be created by adding layers, like convolutional layers, which are great for image analysis. Techniques, such as backpropagation, would be used to fine-tune its accuracy. Additionally, including clinical data along with MRI images could help the model make more precise predictions, aiding in earlier and more reliable dementia diagnoses.

To detect dementia, the ANN processes input data such as MRI scans and medical history through multiple layers of neurons. Each layer extracts and transforms features from the data, progressively refining the model's understanding. The final output layer provides a decision or classification based on the transformed features, indicating whether the patient shows signs of dementia.

The primary data used in an ANN project includes MRI scans of patients' brains [12], as well as supplementary medical history information [13]. This includes age, dominant hand, education level, and socioeconomic status. These diverse data sources provide a comprehensive view, enabling the ANN to learn from both visual and non-visual patterns associated with dementia [8]

Effective data preprocessing is crucial for training robust neural networks. Cleaning the data involves removing irrelevant or noisy information, handling missing values, and addressing outliers. Normalizing the data ensures that each feature contributes equally to the model, preventing any single feature from disproportionately influencing the results [10].

Missing data can be handled through techniques such as imputation, where missing values are replaced with estimated ones based on the remaining data. Outliers, which are data points significantly different from others, can be identified and either removed or transformed to minimize their impact on the model [9].

To maintain consistency and improve model performance, all images are resized to a standard dimension of 224 by 224 pixels. This uniform size ensures that the ANN processes each image in the same way, facilitating efficient learning. Normalizing the images also involves scaling pixel values to a standard range, typically between 0 and 1, which helps in stabilizing the training process and speeding up convergence [10].

By resizing the images to 224 by 224 pixels, we established a consistent input format, ensuring that our ANN can effectively learn from the data and accurately detect signs of dementia [10].

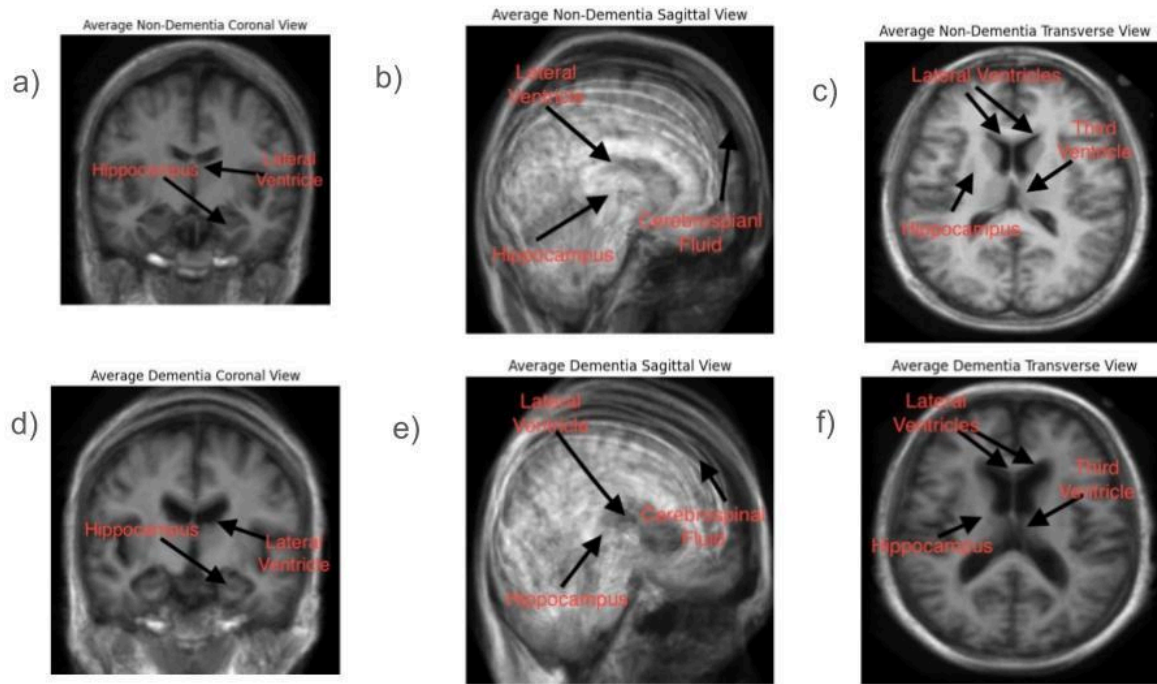


Figure 1: Panels a-f are MRI's scans obtained from Oasis 1 [12]

Works Cited

1. **Alzheimer's Association.** (2023). Alzheimer's Disease Facts and Figures. *Alzheimer's & Dementia*, 19(1), 159-229. Retrieved from [Alzheimer's Association](#).
2. **Jack, C. R., et al.** (2018). NIA-AA Research Framework: Toward a Biological Definition of Alzheimer's Disease. *Alzheimer's & Dementia*, 14(4), 535-562.
3. **McKhann, G. M., et al.** (2011). The Diagnosis of Dementia Due to Alzheimer's Disease: Recommendations from the National Institute on Aging and the Alzheimer's Association Workgroups. *Alzheimer's & Dementia*, 7(3), 263-269.



4. **Mosconi, L., et al.** (2010). Role of PET and MRI Imaging in Predicting Alzheimer's Disease. *Neurobiology of Aging*, 31(1), 17-24.
5. **Blennow, K., et al.** (2015). Cerebrospinal Fluid Biomarkers in Alzheimer's Disease. *Nature Reviews Neurology*, 11(2), 103-118.
6. **Cummings, J., et al.** (2019). Early Intervention in Alzheimer's Disease: Evidence and Implications for Clinical Practice. *Therapeutic Advances in Neurological Disorders*, 12, 1-12.
7. **Brodaty, H., & Donkin, M.** (2009). Family Caregivers of People with Dementia. *Dialogues in Clinical Neuroscience*, 11(2), 217-228.
8. **Hampel, H., et al.** (2018). The Future of Alzheimer's Disease Biomarkers in Clinical Trials. *Journal of Alzheimer's Disease*, 62(3), 1217-1231.
9. **Goodfellow, I., Bengio, Y., & Courville, A.** (2016). Deep Learning. *MIT Press*.
10. **LeCun, Y., Bengio, Y., & Hinton, G.** (2015). Deep Learning. *Nature*, 521(7553), 436-444.
11. **Nair, V., & Hinton, G. E.** (2010). Rectified Linear Units Improve Restricted Boltzmann Machines. *Proceedings of the 27th International Conference on Machine Learning (ICML-10)*, 807-814.
12. **Daniel S. Marcus, Tracy H. Wang, Jamie Parker, John G. Csernansky, John C. Morris, Randy L. Buckner**; Open Access Series of Imaging Studies (OASIS): Cross-sectional MRI Data in Young, Middle Aged, Nondemented, and Demented Older Adults. *J Cogn Neurosci*2007; 19 (9): 1498–1507. doi: <https://doi.org/10.1162/jocn.2007.19.9.1498>



13. **Boysen, J.** (n.d.). *MRI and Alzheimers Magnetic Resonance Imaging Comparisons of Demented and Nondemented Adults*. Retrieved July 2024, from <https://www.kaggle.com/datasets/jboysen/mri-and-alzheimers/data>.

Acknowledgements

Data was provided by Oasis: OASIS-1: Cross-Sectional: Principal Investigators: D. Marcus, R, Buckner, J, Csernansky J. Morris; P50 AG05681, P01 AG03991, P01 AG026276, R01 AG021910, P20 MH071616, U24 RR021382

Daniel S. Marcus, Tracy H. Wang, Jamie Parker, John G. Csernansky, John C. Morris, Randy L. Buckner; Open Access Series of Imaging Studies (OASIS): Cross-sectional MRI Data in Young, Middle Aged, Nondemented, and Demented Older Adults. *J Cogn Neurosci*2007; 19 (9): 1498–1507. doi: <https://doi.org/10.1162/jocn.2007.19.9.1498>