# BrainParc: Progressive Framework for Lifespan Brain Region Parcellation: Towards Comprehensive Understanding of Development and Aging

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# **ABSTRACT**

A persistent challenge in automatic brain region parcellation from structural MR images in the literature is how to address the issue of generalization and robustness amidst substantial *variations in intensity* caused by innate brain maturation and MR scanner manufacturers. Conventional methods for brain region parcellation are often sensitive to varying image contrast; even within the same individual scanned on different MR manufacturers or ages, performance tends to degrade across different scanners and ages. In this work, to address the issues mentioned above, we present a brain region parcellation (UniParc) framework, the first fully automatic segmentation model designed to handle the changes in intensity contrast across the lifespan of the human brain. Notably, UniSeg adopts a cascaded coarse-to-fine brain region parcellation framework that capitalizes on hierarchical, intensity-agnostic image representations (*i.e.*, edge, boundary, and tissue) to effectively address the inherent intensity variation issues. UniParc is trained on the largest dataset to date, comprising over 20k T1w image samples spanning ages from 0-100 years, and sourced from diverse global populations (including variations in race, age, and disease status). Consequently, UniParc can automatically and accurately segment T1w brain images from a wide array of target domains without requiring fine-turning, accomplishing this task within two minutes. This capability streamlines the analysis of vast brain data, facilitating efficient research and clinical applications. Extensive experimentation on both internal and external testing datasets consistently demonstrates the superior performance and broad applicability of our proposed UniParc framework.

Please note: Abbreviations should be introduced at the first mention in the main text no abbreviations lists. Suggested structure of main text (not enforced) is provided below.

## Introduction

The Introduction section, of referenced text<sup>1</sup> expands on the background of the work (some overlap with the Abstract is acceptable). The introduction should not include subheadings.

## Results

Subsection

Third-level section

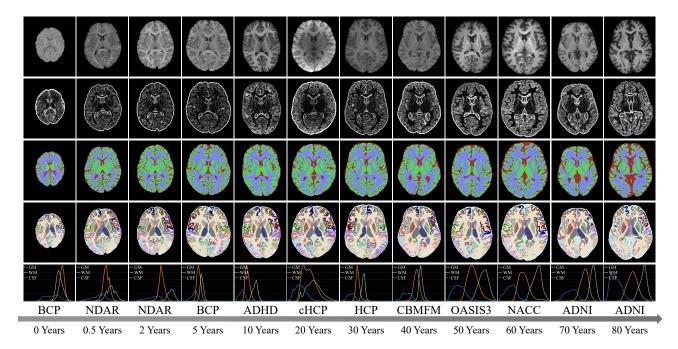
Topical subheadings are allowed.

#### **Discussion**

The Discussion should be succinct and must not contain subheadings.

#### **Methods**

Topical subheadings are allowed. Authors must ensure that their Methods section includes adequate experimental and characterization data necessary for others in the field to reproduce their work.



**Figure 1.** Representative intensity appearance across human brain development periods. From left to right present the brain at 0, 0.5, 2, 5, 10, 20, 30, 40, 50, 60, 70, 80 years old. From top to bottom present intensity data, corresponding edge maps, tissue maps, dk-struct maps, and intensity distributions of three tissues, respectively.

## References

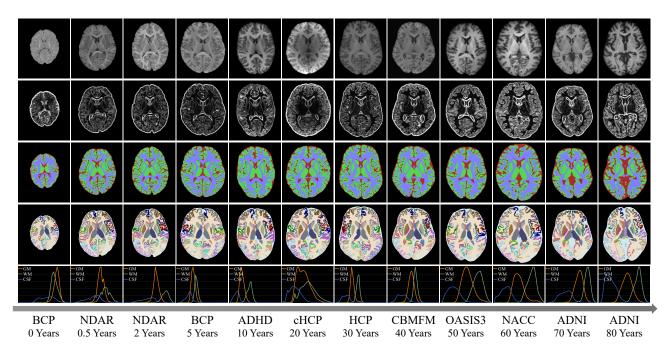
- 1. Figueredo, A. J. & Wolf, P. S. A. Assortative pairing and life history strategy a cross-cultural study. *Hum. Nat.* 20, 317–330, DOI: https://doi.org/10.1007/s12110-009-9068-2 (2009).
- **2.** Hao, Z., AghaKouchak, A., Nakhjiri, N. & Farahmand, A. Global integrated drought monitoring and prediction system (GIDMaPS) data sets. *figshare* http://dx.doi.org/10.6084/m9.figshare.853801 (2014).

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#### **Author contributions statement**

Must include all authors, identified by initials, for example: A.A. conceived the experiment(s), A.A. and B.A. conducted the experiment(s), C.A. and D.A. analysed the results. All authors reviewed the manuscript.



**Figure 2.** Representative intensity appearance across human brain development periods. From left to right present the brain at 0, 0.5, 2, 5, 10, 20, 30, 40, 50, 60, 70, 80 years old. From top to bottom present intensity data, corresponding edge maps, tissue maps, dk-struct maps, and intensity distributions of three tissues, respectively.