Boundary Based Movement Correction of Functional MR Data Using a Genetic Algorithm

Guo jun Bao, Jagath C. Ra japakse

Sc hool of Applied Science, Nany ang Technological University , Singapore 639798

Abstract. This paper describes a novel image registration method for mov ement correction of fMR time-series. It is important to align the fMR images in the time-series before time-dependent analyses. This registration method aligns the boundaries of brains extracted from the functionalimages. It uses a genetic algorithm to minimize the distance function obtained from the chamfer distance transform. The global search nature of genetic algorithm makes this method robust to the presence of localminima.

1. Introduction

Movement correction of brain images obtained at different time instances of an fMR head time-series of scans is essential for the analyses of time-dependent changes. Despite the restraints to inhibit head movement, subjects could still show sligh t displacement in the scanner. If these motion artifacts are not corrected in the time-series analysis, the detected activities may be artifactual and important activities may be missed.

Although many techniques ha v ebeen proposed for mov ement correction [2, 3, 4, 6, 11], most of these techniques use local hill-climbing optimization approaches. In general, mov ement correction of fMR time-series is a multidimensional optimization problem which may contain local minima. Therefore, these approaches are not appropriate. In this paper, we propose a nov el method which uses boundary based genetic algorithm for mov ement correction of fMR data. We register a series of fMR images to a reference MR image. It conquers the local minimum problem using the global search nature of genetic algorithm.

Consider two images $f: \Omega_f - \to Q_f \subset \mathbf{R}$ and $g: \Omega_g - \to Q_g \subset \mathbf{R}$ where for and g are the domains of images f and g , respectively . Registration of α images f and g is the process of performing spatial transformation on the images so that the voxel positions of images correspond to the same points in the imaging space. Without losing generality , let us consider one image, say g, as the reference image and that the spatial transformation is performed on the other image f to match g. Let S denote the spatial transformation and

 f' denote the image f after transformation. We are in terestedin registering the boundaries of a prominent object. Let ${\cal C}_{f'}$ and ${\cal C}_f$ be the boundaries of the object in f' and f respectively. When $\mathbf{p} \in \Omega_f$ and $\mathbf{q} \in \Omega_g$, one can write

$$
\mathcal{C}_{f'}(\mathbf{q}) = \mathcal{C}_f(\mathcal{S}(\mathbf{p}, \boldsymbol{\alpha}))
$$

where $\mathbf{q} = \mathcal{S}(\mathbf{p}, \alpha)$, and α indicates the set of parameters for spatial transformation.

We assume that the spatial transformation S is an affine transformation. That is when $\mathbf{p} \in \Omega_f$, $\mathcal{S}(\mathbf{p}, \alpha) = \mathbf{M} \mathbf{p}$ where the linear transformation matrix M is a combined matrix of the translation matrix T, cen ter translation matrix α ; rotation matrices α ; α , α , and α , and α be written α , and α and α be written be as $M = TR_xR_yR_zSC$ which contains nine parameters [9].

2. Boundary Based Distance Transform

In our approach, image registration is to minimize a *distance function* that measures mismatch of corresponding boundary positions of the object in f' and g. Let \mathcal{C}_q be the boundary of the object in image $g, \mathcal{D}(\cdot, \cdot)$ be the distance function betw een the boundaries of w o images andg["] be the reference distance image obtained from distance transform of g . Then

$$
\mathcal{D}(\mathcal{C}_{f'}, \mathcal{C}_g) = \sum_{\mathbf{p'} \in \mathcal{C}_{f'}} g''(\mathbf{q''})
$$

where **p**' is a v oxel inf', q'' is the equivalent voxel of **p**' in g'' and $g''(q'')$ is the v alue of that particular voxel. The perfect alignment is achieved when D is minimized.

Sey eral distance transform techniques exist to obtain the reference distance image, and the chamfer distance transform [1] is the most popular one because of its speed and simplicity. For image g with boundary \mathcal{C}_q , chamfer distance transform converts g into a binary image g' consisting of boundary elements and non-boundary elements, and then converts g' into a c hamfer distance imagegⁿ where each element has a value that approximates the distance to the nearest boundary element $q \in \mathcal{C}_g$.

3. Genetic Algorithm

Local hill climbing algorithms are not suitable for optimization of distance function due to local minima. Though some stochastic global search procedures like simulated annealing [7] are available, they are time-consuming and not strongly reliable. In this section, we introduce genetic algorithms (GA) [5, 8, 10] for optimization of the distance function in our study.

3.1. Genetic Optimization

GA is a class of global optimization techniques that model some natural phenomena, namely genetic inheritance and Darwin's strife for natural survival. It attempts to maximize a parametric function $\mathcal{E}(\alpha)$ referred to as the *evaluation* function where $\alpha = {\alpha_1, \alpha_2 ... \alpha_n}$ is the set of parameters and n is the number of parameters. GA finds the optimal set α^* of parameters such that

$$
\boldsymbol{\alpha}^* = \arg\max_{\boldsymbol{\alpha}} \mathcal{E}(\boldsymbol{\alpha})
$$

In our application, the evaluation function is

$$
\mathcal{E}(\boldsymbol{\alpha}) = -\mathcal{D}(\mathcal{C}_{f'}, \mathcal{C}_q) + A
$$

where constant A is added to make $\mathcal{E}(\alpha)$ positive for all parameter sets. The algorithm evolves efficiently in the whole parameter space in a probabilistic manner to realize the global optimal parameters.

To formulate GA, all parameters $\alpha_i \in \alpha$ are represented with binary strings called genes. A chromosome is formed by concatenating genes representing dierent parameters of the evaluation function. If μ are the gene representing parameter α_i , the chromosome c can be written as c = $\alpha_{1,1,2}, \ldots, \alpha_n$ $population \, \mathcal{P}$ consists of a set of chromosomes representing the same parameter set

$$
\mathcal{P} = \{\mathbf{c}_1, \mathbf{c}_2 \dots \mathbf{c}_K\}
$$

where K denotes the population size.

GA performs multidirectional search by maintaining a population of chromosomes encoding potential solutions. Optimization begins by relating the distance function to an evaluation function, and defining a population of chromosomes representing parameters of the evaluation function. The population is randomly initialized and iterativ ely updated b y performing genetic operations until the ev aluation functionis maximized. Genetic operations namely, selection, crossover, mutation, and exchange are performed to update the population in each generation. This can be formulated as follows:

> Randomly initialize P of size K While not end $P = \text{selection}(P)$ $P = \text{crossoverer}(P)$ $P = \text{mutation}(P)$ $P = \text{exchange}(P)$ Repeat

3.2. Genetic Operations

A population ev olves from eac h generation b y incorporating and exc hanging new information in every direction of the multidimensional parameter space.

In ev ery generation, GA tries to get rid of inferior members and encourages reproduction and exchange betw een superior members in an optimal way towards the optimum solution.

3.2.1.Selection

Selection is performed based on the fitness of chromosomes calculated from the ev aluation function. The fitness of a chromosome indicates the closeness of the chromosome to **optimal** solution. Those chromosomes with higher fitness values have a better chance and more copies to appear in the next generation. Before the selection process, a simulated biased roulette wheel with slots sized according to chromosome fitness is constructed. The selection process is then carried out by spinning the roulette wheel K times. Each time, w eselect a single chromosome for a new population. The population size is thus held constant from one generation to the next.

3.2.2.Crossov er

Crossov er operation occurs among the chromosomes selected from the current population with a probability of p_c . This gives us an expected number $p_c \cdot K$ of chromosomes undergoing crossov er operation. The selected chromosomes are paired randomly and the crossov er operation takes place in each pair where the bit patterns beyond a randomly selected bit position of the two chromosomes are in terchanged.

Mutation introduces new information to the current population at bit lev el. E is only bit of each chromosome in a population has a population has a population μ pm for mutation μ b y simply reversing its value. The expected n umber of mutated bits is given by $p_m \cdot N \cdot K$, where N is the length of a chromosome. Mutation should be used sparingly as in nature because it is a random search operator; otherwise the algorithm will become little more than a random search.

3.2.4. Exchange **3.2.4. Exchange 3.2.4. Exchange 3.2.4.** Exchange 3.4. Exchange 3.

Exchange occurs betw een pairs of dromosomes in a subset of the population where each chromosome was selected with a probability of personal μ and μ personal current μ population. The genes at a randomly selected position of a randomly selected pair of chromosomes from the subset are interchanged. p_e κ n umber of chromosomes are expected to undergo exchange operation. The exchange operator forces the population to increase its div ersity and thus prev ents premature conv ergence during optimization process.

4. Motion Correction

We demonstrate our techniques for analysis of a fMRI time-series using images obtained in a visual experiment. All images were acquired on 3T Bruker Medspec 30/100 system at the Max-Planck-Institute of Cognitive Neuroscience, Leipzig, Germany .

4.1. Visual Experiment

while a subject was performing the experiment, $2D/T_2$ -weighted images, each with 64 scans, were acquired using a gradient-echo FLASH sequence. An 8Hz alternating checkerboard pattern with a central fixation point w asprojected on a LCD system and subjects were ask ed to fixate on the point during stimulations. F our successive ON and OFF stimuli were presented each at a rate of 5.162 s/sample. The stimulations were repeated for eight cycles. Extensive experiments show it as \sim 32, 30 iterations for GA, pm = 0:005, pm = 0:005, pm = 0:005, pm = 0:005, pm = 0:005 and per satisfactory per satisfactory results.

The chamfer distances before and after movement correction of the image series are shown in Figure 1(a). The convergence of the algorithm is illustrated by showing a particular slice in Figure 1(b).

Figure 1: (a) Chamfer distances betw een boundaries of fMR slices and a reference image obtained before and after registration (b) Illustration of conv ergence of chamfer distance betw een boundaries of an fMR slice and a reference image.

$4.2.$ **Conclusion**

The experiment results show signicant improv ement in the chamfer distance of the image slices by applying our novel algorithm. The genetic algorithm also shows rapid conv ergence of the distance function. Because only the boundary voxels are involved in computation, our technique is more efficient than those

v olume based techniques where all brain voxels are inv olved in computation. Due to the global search nature of genetic algorithm, this nov el tec hnique does not hav e local minimum problem. We tried gradient descent technique and genetic algorithm show ed superior performance.As a conclusion, our boundary based genetic algorithm gives great efficiency and effectiveness in movement correction of functional MR images.

References

- [1] G. Borgefors. Distance transformations in arbitrary dimensions. Computer Vision, Graphics, and Image Processing, $27:321-345$, 1984.
- [2] D. L. Collins, P . Neelin,T. M. P eters, and A.C. Evans. Automatic 3D intersub ject registration of MR volumetric data in standardized taliarach space. J. Computer Assisted Tomography, $18(2):192-205$, 1994.
- [3] R. Cox. Algorithms for image registration, motion detection and motion correction. $fMRI12Day$, pages 25-42, 1996.
- [4] K. J. Friston, J. Ashburner, C. D. Frith, J. B. Poline, J. D. Heather, and R. S. J. F rac k o wiak. Spatial registration and normalization of images. Human Brain Mapping, $2:165{-}189, 1995$.
- [5] L. J. Holland. A daptation in Natural and Artificial Systems. The University of Michigon Press, 1994.
- [6] B. Kim, J. L. Boes, P. H. Bland, T. L. Chenevert, and C. R. Meyer. Motion correction in fMRI via registration of individual slices into an anatomical volume. Magnetic Resonance in Medicine, 41:964-972, 1999.
- [7] S. Kirkpatrick. Optimization by simulated annealing. *Science*, 220, 1983.
- [8] Z. Michalewicz. Genetic A lgorithm+ Data Structures = Evolution Programs. Springer, Springer-Verlag Berlin Heidelberg New York, 1996.
- [9] J. C. Ra japakse and G. J. Bao. Functional MR image registration using a genetic algorithm. In Pr oceedings of ICONIP'9, 9 volume 3, pages $922-927$, Nov. 1999.
- [10] L. H. Staib and X. Lei. In termodality 3D medical image registration with global search. In Pr oceedings of the IEEE Workshop on Biometical Image A nalysis pages 225-234. IEEE Computer Society, 1994.
- [11] R. P .Woods, S. R. Cherry, and J. C. Mazziotta. Rapid automated algorithm for aligning and reslicing PET images. J. Computer Assisted T omgraphy, $16(4):620-633, 1992.$