



N5
School Year
2008/2009
Training Period

**Institut Suprieur de l'Electronique et
du Numrique**

Tel. : +33 (0)2.98.03.84.00

Fax : +33 (0)2.98.03.84.10

CS 42807 - 29228 BREST Cedex 2 -
FRANCE

Parameter space exploration and tools for fast
visualization in EM segmentation and MRI bias
field correction in Slicer 3

From April 6th to Septemner 11th
At
Surgical Planning Laboratory (SPL)



Brigham & Women's Hospital
Harvard Medical School
75 Francis St.
Boston, MA 02115

Supervisors:

- Ron KIKINIS, Surgical Planning Laboratory, Harvard Medical School,
Boston, MA - USA
- Sylvain JAUME, CSAIL, Massachusset's Insitut of Technologies, Boston,
MA - USA

Referring teachers:

- Dominique MARATRAY, Institut Supérieur de l'Electronique et du Numerique, Brest - FRANCE
- Christine CAVARO-MENARD, Universite de Angers, Angers - FRANCE

Student:

- Nicolas RANNOU, Institut Supérieur de l'Electronique et du Numerique, Brest

Abstract

A couple of sentences on three or four lines to summarise your work.

This is a L^AT_EX template for undergraduate project reports.

Its detailed contents evolve to reflect FAQs.

Expectation-maximization is very popular for segmentation but it can be tricky to understand and to use. A full description of the EMS algorithm is done in this report. Different methods for fast parameters exploration are described. As part of the research, preprocessing methods like MRI bias field correction will be explained. The results obtained will be presented. Following the new workflow should allow the user to segment more datasets, more accurately.

Keywords: segmentation, expectation, maximization, correction, bias.

Resumé

Quelques phrases pour resumer mon travail.

C'est un template L^AT_EX pour les rapport.

Le contenu peut evoluer.

Mots clés :segmentation, expectation, maximisation, correction, biais.

Contents

1	Introduction	4
1.1	Context and motivation	4
1.2	Contents	5
2	Expectation-maximization applied to brain segmentation	6
2.1	Presentation of the EM segmentation	6
2.2	Fundamentals	7
2.2.1	Statistical model used for the brain	7
2.2.2	Gaussian mixture model	8
2.2.3	Maximum likelihood	8
2.3	Expectation maximization algorithm	11
2.4	Expectation maximization algorithm used in Slicer 3	15
2.4.1	Probabilistic atlas	15
2.4.2	Multichannel segmentation	15
2.4.3	Bias field correction	16
2.4.4	Hierarchical information	18
2.5	Workflow in Slicer 3	19
2.5.1	User interface	19
2.5.2	Algorithm	20
2.5.3	Summary	20
2.6	Limitations	20
3	The contributions	21
3.1	MRI Bias Field correction	21
3.1.1	Interest	21
3.1.2	Our approach	21
3.1.3	Results	21
3.2	Intensity Normalization	21
3.2.1	Interest	21
3.2.2	Our approach	21
3.2.3	Our implementation	22
3.3	Class Distribution selection	22
3.3.1	Interest	22

3.3.2	Method used	22
3.4	Class Distribution visualization	22
3.4.1	Interest	22
3.4.2	Method used	24
3.4.3	Results	25
3.5	Global Prior Estimation	25
3.5.1	Presentation of the problem	25
3.5.2	Our approach	25
3.5.3	Results and discussion	26
4	Results and discussion	28
4.1	Results	28
4.1.1	Bias correction	28
4.1.2	Global Prior estimation	30
4.1.3	Class Selection	30
4.2	Limitations	30
4.3	Future work	30
	Acknowledgements	31
	Bibliography	32

List of Figures

2.1	A simple tree structure of the brain	19
3.1	Axial view of a T1 volume without the label map.	23
3.2	Axial view of a T1 volume with the label map.	23
3.3	Axial view of a T2 volume without the label map.	23
3.4	Axial view of a T2 volume with the label map.	23
4.1	Module created for the bias correction.	29
4.2	Sagittal view of a biased T1 volume.	29
4.3	Sagittal view of the T1 volume after bias correction.	29

Chapter 1

Introduction

1.1 Context and motivation

Nowadays, medical image processing is becoming a major field of research in most of the laboratories. Indeed, because of the increasing complexity of the data they have to deal with, physicists need something to help. Help must be provided in many different ways. Before the surgery, to establish a fast and accurate diagnosis. During the surgery to prevent the physicist from errors and to help him to proceed to more precise moves. After the surgery, to see if the surgery succeed, or to follow the pathology of a patient. Nevertheless, the informations bring to the physicist by the tool must be accurate, robust and provide a fast feedback.

In this context of pre and post operation, plenty of work as already been done. Thus, there is still a lot of work to achieve. Regarding data storage and exchange, the increasing among of informations leads us to find other and more appropriate methods. Another interesting contribution is images segmentation. New methods have to be developped for a better diagnosis, or to detect new pathologies. A lot of methods appears like level-set segmentation, region growing or texture based segmentation. Each one is adapted for a specific problem like vessels segmentation, tumors detection or SDFS lungs detection. Another remarquable contribution is the segmentation based on expectation maximization (EM), which is very well suited for brain segmentation.

For the MR images segmentation purpose, the Surgical Planning Laboratory (SPL), Harvard Medical School, has developped an EM algorithm to segment brain's MR images. The results obtained are very good until we segment small structures and we select the optimum parameters. The approach used for the intensity inhomogeneities estimation appears less efficient for the particular purpose of large structures segmentation WHY. Moreover, the implementation is not widely used so far, regarding the complexity of the segmentation process. In this report we will present an approach to

enhance the segmentation for large structures, correcting the intensity inhomogeneities in large structures and providing the end-user tools for an easier segmentation process.

1.2 Contents

The main body of this report is divided as follows.

Chap. 2 deals with the EM segmentation. Fundamentals will be reminded and the algorithm used will be described. We will also present in this chapter the limitations of the current implementation. Chap. 3 describes our contribution. It explains the solution we choose to enhance the segmentation and to improve the usability of the current framework. Chap. 4 shows the results achieved. It also discusses about what has been done, the limitations of the current module and the next work which has to be done.

Chapter 2

Expectation-maximization applied to brain segmentation

Here we get going with theory of the expectation-maximization (EM) applied to brain segmentation. We show firstly a simple approach of the problem, in the particular case of Gaussian Mixture Models (GMM) followed by a more generalistic approach. The simple approach will give you an intuitive understanding of the problem then the general approach will formalize it in order to adapt it to most of the segmentation problems. Finally, there will be a presentation of the algorithm used in Slicer 3¹.

2.1 Presentation of the EM segmentation

The EM algorithm was originally described in 1977 by Arthur Dempster, Nan Laird, and Donald Rubin[1]. They generalized and developed a method used in several times by authors, for particular applications. It is widely used to solve problems where data are "missing". The EM algorithm is an iterative algorithm which works in two steps: Expectation and Maximization. It can be used to solve a lot of image processing's problems like classification, restoration[3], motion estimation[2], etc.. Since the generalization of the algorithm, a lot of related papers were proposed. Most of them bring algorithms derived from the original one to adapt it to particular problems using additional informations like spatial or structural information.

Nowadays, EM algorithms have become a popular tool for classification problems. It is particularly well suited for brain MR images segmentation. A lot of algorithms already exist. They present complex frameworks using spatial information, neighborhood or intensity inhomogeneities to enhance

¹open source software developed in the SPL for biomedical engineering purpose

the classification.

In the SPL, the algorithm developed uses spatial, structural and intensity inhomogeneities informations to segment the brain.

2.2 Fundamentals

Here we get going with a presentation of all the fundamentals you need to have a good understanding of EM segmentation. We begin with a description of the statistical model used for the brain. Then we present briefly the widely used GMM. Finally we will introduce you to what the maximum likelihood function. This part is mainly inspired from [4], [5] and [6]

2.2.1 Statistical model used for the brain

We define the voxel intensities of a MR image as $Y = \{y_1, \dots, y_n\}$ when the image consisted in n voxels. Each y intensity is called *observed data* because this is the data we see when we observe the image. Each y is a realization of the random variable Y . The real labelling of the image is Z . Z is called *hidden data* because we don't know the value of each label. This is precisely the purpose of the segmentation: estimating the *hidden data* from the *observed data*. We assume that the *observed data* is generated from the *hidden data* and a parameter Φ . The parameter Φ can either be a probability density function, noise, bias field, etc., depending on the model.

Y and Z can be viewed as n -dimensional random variables $Y = \{Y_1, \dots, Y_n\}$ and $Z = \{Z_1, \dots, Z_n\}$ then each y_i is a realisation of Y_i and each z_i is a realization of Z_i . The conditional probability function describing Y_i is $p(Y_i|Z_i, \Phi)$.

The easiest model assumes that each intensity in one class is the same, but this intensity is corrupted by factors like noise, with a Gaussian Distribution. We can describe the relationship as below:

$$y_i = \mu_k + n_i$$

where μ_k is the mean intensity of the k^{th} tissue and n_i a random sample generated by the corrupting factor(s). Let's say that n_i is generated by a Gaussian probability distribution function $G(., 0, \sigma)$, with 0 mean and σ variance. That means that y_i is a random sample generated by Gaussian probability density function $G(., \mu_k, \sigma)$. Let's assume that each class has a different variance, $G(., \mu_k, \sigma)$ becomes $G(., \mu_k, \sigma_k)$ and it leads to:

$$p(Y_i = y_i|Z_i = k, \Phi) = G(y_i, \mu_k, \sigma_k) \quad (2.1)$$

As the labelling is not known, it is useful to express the probability density function (PDF) of Y_i only depending on parameter Φ with the total probability theorem:

$$p(Y_i|\Phi) = \sum_{k=1}^K p(Y_i = y_i|Z_i = k, \Phi)p(Z_i = k|\Phi) \quad (2.2)$$

$p(Z_i = k|\Phi)$ is the *prior probability*. It expresses the probability that a voxel i belongs to a class k . $p(Y_i = y_i|Z_i = k, \Phi)$ is the *likelihood*. In our case, we will assume that the *prior probability* is constant. The new model we obtain for the labelling, via a gaussian distribution, is a widely used one: the *Gaussian mixture model*.

2.2.2 Gaussian mixture model

Let's remind the first hypothesis: the conditional probability function for each tissue to segment is Gaussian equation (2.1). Moreover, we will assume that *prior probability* is a constant c_k for each class k . c_k is the *weight* of the class k .

$$p(Z_i = k|\Phi) = c_k \quad (2.3)$$

The last assumption will be that Φ contains unknown means, variances and weights for each tissue. Then we can express Φ as $\Phi = (\mu_1, \sigma_1, c_1, \dots, \mu_K, \sigma_K, c_K)$.

Using equations (2.1) and (2.3), equation (2.2) becomes:

$$p(Y_i = y|\Phi) = \sum_{k=1}^K G(y, \mu_k, \sigma_k)c_k \quad (2.4)$$

In the case of *Gaussian mixture model*, each voxel is considered to be independent. That means that each voxel will have his own probability density function. Consequently, the normalized histogram of the whole volume can be interpreted as an approximation of the sum of all the probability density functions.

2.2.3 Maximum likelihood

In our case, we know the intensity of each observed pixel y_i . Φ has to be found. The best estimation of Φ will be obtain using the maximum likelihood principle. $p(Y = y_i|\Phi)$ is called likelihood function and for it returns the value of the likelihood for y_i , given Φ . We can generalize it to the whole image with $p(Y|\Phi)$.

As we said in SDFSDF, the voxels are considered to be independent through the whole volume. It leads us to:

$$p(Y|\Phi) = \prod_{i=1}^n p(Y_i|\Phi) \quad (2.5)$$

We must keep in mind that the objective is to find the parameter Φ which will maximize the likelihood of the observed volume. We can note this parameter:

$$\hat{\Phi} = \arg \max_{\Phi} p(Y|\Phi) \quad (2.6)$$

Therefore, it is more convenient to work with logarithm because the product from equation (2.5) will be converted into a sum. Equation (2.6) becomes:

$$\hat{\Phi} = \arg \max_{\Phi} \log p(Y|\Phi). \quad (2.7)$$

Let us denote:

$$\begin{aligned} L(\Phi) &\triangleq \log p(Y|\Phi) \\ &= \sum_{i=1}^n \log \sum_{k=1}^K p(Y_i = y|Z_i = k, \Phi) p(Z_i = k|\Phi) \end{aligned}$$

Finally, in case of GMM, with eq. (2.4), $L(\Phi)$ becomes:

$$L(\Phi) = \sum_{i=1}^n \log \sum_{k=1}^K G(y, \mu_k, \sigma_k) c_k \quad (2.8)$$

The maximized *log* likelihood can then be computed using partial derivatives for each parameter of Φ . When the partial derivative of $L(\Phi)$ is 0 for a parameter, we found the maximum likelihood for the parameter.

For example, to find the maximum likelihood for μ_k , we have to find when:

$$\frac{\partial}{\partial \mu_k} (L(\Phi)) = 0 \quad (2.9)$$

Then we compute the partial derivative of $L(\Phi)$ over μ_k :

$$\begin{aligned}
\frac{\partial}{\partial \mu_k}(L(\Phi)) &= \frac{\partial}{\partial \mu_k} \left(\sum_{i=1}^n \log \sum_{k=1}^K G(y, \mu_k, \sigma_k) c_k \right) \\
&= \sum_{i=1}^n \frac{G(y, \mu_k, \sigma_k) c_k}{\sum_{j=1}^K G(y, \mu_j, \sigma_j) c_j} \frac{\partial}{\partial \mu_k} \left(-\frac{(y - \mu_k)^2}{2\sigma_k^2} \right) \\
&= \sum_{i=1}^n \frac{G(y, \mu_k, \sigma_k) c_k}{\sum_{j=1}^K G(y, \mu_j, \sigma_j) c_j} \left(-\frac{(y - \mu_k)}{\sigma_k^2} \right) \\
&= \sum_{i=1}^n \frac{p(Y_i = y | Z_i = k, \Phi) p(Z_i = k | \Phi)}{\sum_{j=1}^K p(Y_i = y | Z_i = j, \Phi) p(Z_i = j | \Phi)} \left(-\frac{(y - \mu_k)}{\sigma_k^2} \right) \quad (2.10)
\end{aligned}$$

Using Bayes formula, we notice that:

$$p(Z_i = k | Y_i = y_i, \Phi) = \frac{p(Y_i = y_i | Z_i = k, \Phi) p(Z_i = k | \Phi)}{\sum_{j=1}^K p(Y_i = y_i | Z_i = j, \Phi) p(Z_i = j | \Phi)} \quad (2.11)$$

Thus, setting the denominator to 0 in equation (2.10) and using equation (2.11) yields:

$$\sum_{i=1}^n p(Z_i = k | Y_i = y_i, \Phi) (y - \mu_k) = 0 \quad (2.12)$$

Let us denote

$$p_{ik} = p(Z_i = k | Y_i = y_i, \Phi) \quad (2.13)$$

Equation (2.12) leads us to:

$$\mu_k = \frac{\sum_{i=1}^n y_i p_{ik}}{\sum_{i=1}^n p_{ik}} \quad (2.14)$$

Proceeding the same way as we did for equation (2.12), we can get similar equations for variance σ_j and weight c_j .

We find that:

$$\sigma_k^2 = \frac{\sum_{i=1}^n (y_i - \mu_k)^2 p_{ik}}{\sum_{i=1}^n p_{ik}} \quad (2.15)$$

$$c_k = \frac{1}{n} \sum_{i=1}^n p_{ik} \quad (2.16)$$

Equations (2.14), (2.15), and (2.16) provides us an equation for soft segmentation. This equation expresses that a voxel i belongs to the class k .

$$p_{ik} = \frac{G(y_i, \mu_k, \sigma_k) c_k}{\sum_{j=1}^K G(y_i, \mu_j, \sigma_j) c_j} \quad (2.17)$$

In the case of GMM, the segmentation can now be done following an iterative algorithm called *expectation maximization algorithm*.

2.3 Expectation maximization algorithm

The EM algorithm is a method to find the maximum likelihood for a given set of parameter (Φ in our case). Here we first get going with an intuitive description of the algorithm in the particular case of GMM then we will present a more general definition.

Algorithm in case of Gaussian mixture data model

Let's assume that we can find the maximum likelihood of the hidden data by a direct differentiation (because of the GMM). The EM algorithm is an iterative process of two steps: the expectation step (E-Step) and the maximization step (M-Step). At each iteration, the maximum likelihood will be increased until convergence is reached.

E-step

In this step, we calculate an estimation of soft segmentation $p^{(m+1)}$ with equation (2.17) as below. We know all the variables needed for the calculation from the observed data and the current parameter estimate $\Phi^{(m)}$. Note that an initialization is necessary for the first iteration.

$$p_{ik}^{(m+1)} = \frac{G(y_i, \mu_k^{(m)}, \sigma_k^{(m)}) c_k^{(m)}}{\sum_{j=1}^K G(y_i, \mu_j^{(m)}, \sigma_j^{(m)}) c_j^{(m)}}$$

M-step

In this step, we estimate the maximum likelihood for parameter $\Phi^{(m+1)}$. We do it with equations (2.14), (2.15), and (2.16) as below. We know all the variables needed for the calculation from the observed data and the current estimate $p^{(m+1)}$ of hidden data.

$$\begin{aligned} \mu_k^{(m+1)} &= \frac{\sum_{i=1}^n y_i p_{ik}^{(m+1)}}{\sum_{i=1}^n p_{ik}^{(m+1)}} \\ (\sigma_k^{(m+1)})^2 &= \frac{\sum_{i=1}^n (y_i - \mu_k^{(m+1)})^2 p_{ik}^{(m+1)}}{\sum_{i=1}^n p_{ik}^{(m+1)}} \end{aligned}$$

$$c_k^{(m+1)} = \frac{1}{n} \sum_{i=1}^n p_{ik}^{(m+1)}$$

The problem is simple as long as we are working with GMM. In the other case, the log-likelihood can not be maximized by direct differentiation and a more generalized approach must be used.

Generalized algorithm

Now we assume that we are no longer working with GMM. Thus, we must use a more general algorithm. To explain the general algorithm, we will start from the log-likelihood $L(\Phi)$. As presented in the previous subsection:

$$L(\Phi) = \log p(Y|\Phi)$$

Since \log is a strictly increasing function, the value of Φ which will maximizes $p(Y|\Phi)$ will also maximizes $L(\Phi)$. We want to maximize $L(\Phi)$. Thus, after the m^{th} iteration, we want an estimated Φ_n such as:

$$L(\Phi) > L(\Phi^{(m)})$$

In other words, we want to maximize the difference $L(\Phi) - L(\Phi_n)$. For convenience, we introduce a new variable z_{ik} which means that $Z_i = k$. Using the new notation, we can transform this difference as below:

$$\begin{aligned} L(\Phi) - L(\Phi_n) &= \log p(Y|\Phi) - \log p(Y|\Phi_n) \\ &= \sum_{i=1}^n \left\{ \log \sum_{k=1}^K p(Y_i|z_{ik}, \Phi) p(z_{ik}|\Phi) - \log p(Y_i|\Phi_n) \right\} \\ &= \sum_{i=1}^n \left\{ \log \sum_{k=1}^K p(Y_i|z_{ik}, \Phi) p(z_{ik}|\Phi) \cdot \frac{p(z_{ik}|Y_i, \Phi_n)}{p(z_{ik}|Y_i, \Phi_n)} - \log p(Y_i|\Phi_n) \right\} \\ &= \sum_{i=1}^n \left\{ \log \sum_{k=1}^K p(z_{ik}|Y_i, \Phi_n) \cdot \frac{p(Y_i|z_{ik}, \Phi) p(z_{ik}|\Phi)}{p(z_{ik}|Y_i, \Phi_n)} - \log p(Y_i|\Phi_n) \right\} \\ &\geq \sum_{i=1}^n \left\{ \sum_{k=1}^K p(z_{ik}|Y_i, \Phi_n) \log \frac{p(Y_i|z_{ik}, \Phi) p(z_{ik}|\Phi)}{p(z_{ik}|Y_i, \Phi_n)} - \log p(Y_i|\Phi_n) \right\} \end{aligned}$$

We can deduce this inequality from Jensen's inequality since $p(z_{ik}|Y_i, \Phi^{(m)})$ is a probability measure:

$$p(z_{ik}|Y_i, \Phi_n) > 0$$

$$\sum_k p(z_{ik}|Y_i, \Phi_n) = 1$$

Finally, we use the fact that $\sum_k p(z_{ik}|Y_i, \Phi^{(m)}) = 1$. In this case, it leads to $\log p(Y|\Phi^{(m)}) = \sum_k p(z_{ik}|Y_i, \Phi^{(m)}) \log p(Y|\Phi^{(m)})$. This allows us to bring $\log p(Y|\Phi^{(m)})$ into the summation. We will also use a new variable e_k to express the difference through the whole volume without a summation. e_k is defined as $e_k = \{z_{1k}, \dots, z_{nk}\}$ when the image consisted in n voxels. For example, $z_{ik} = e_k = \{0, \dots, 0, 1, 0, \dots, 0\}$ means that voxel i belongs to class k .

$$\begin{aligned} L(\Phi) - L(\Phi_n) &\geq \sum_{i=1}^n \left\{ \sum_{k=1}^K p(z_{ik}|Y_i, \Phi^{(m)}) \log \frac{p(Y_i|z_{ik}, \Phi)p(z_{ik}|\Phi)}{p(z_{ik}|Y_i, \Phi^{(m)})} - \log p(Y|\Phi^{(m)}) \right\} \\ &= \sum_{i=1}^n \left\{ \sum_{k=1}^K p(z_{ik}|Y_i, \Phi^{(m)}) \log \frac{p(Y_i|z_{ik}, \Phi)p(z_{ik}|\Phi)}{p(z_{ik}|Y_i, \Phi^{(m)})p(Y_i|\Phi^{(m)})} \right\} \\ &= \sum_{k=1}^K p(e_k|Y, \Phi^{(m)}) \log \frac{p(Y|e_k, \Phi)p(e_k|\Phi)}{p(e_k|Y, \Phi^{(m)})p(Y|\Phi^{(m)})} \\ &\triangleq \Delta(\Phi|\Phi_n) \end{aligned}$$

We can then conclude that:

$$\begin{aligned} L(\Phi) &\geq L(\Phi^{(m)}) + \Delta(\Phi|\Phi^{(m)}) \\ &\geq l(\Phi|\Phi^{(m)}) \end{aligned}$$

where $l(\Phi|\Phi^{(m)}) \triangleq L(\Phi^{(m)}) + \Delta(\Phi|\Phi^{(m)})$.

We have now a function $l(\Phi|\Phi^{(m)})$ which is bounded above by $L(\Phi)$. The only time when the two functions are equals is when $\Phi = \Phi^{(m)}$. Our objective is to find the value of Φ which will maximize $L(\Phi)$.

We can formalize this research. $\Phi^{(m+1)}$ is the updated value which is get after maximization of Φ using $\Phi^{(m)}$. Once $\Phi_n = \Phi^{(m+1)}$, the algorithm has converged.

$$\begin{aligned}
\Phi^{(m+1)} &= \arg \max_{\Phi} \{l(\Phi|\Phi^{(m)})\} \\
&= \arg \max_{\Phi} \{L(\Phi^{(m)}) + \sum_{k=1}^K p(e_k|Y, \Phi^{(m)}) \log \frac{p(Y|e_k, \Phi)p(e_k|\Phi)}{p(e_k|Y, \Phi^{(m)})p(Y|\Phi^{(m)})}\} \\
&= \arg \max_{\Phi} \{\sum_{k=1}^K p(e_k|Y_i, \Phi^{(m)}) \log p(Y|e_k, \Phi)p(e_k|\Phi)p(e_k|Y, \Phi^{(m)})p(Y|\Phi^{(m)})\} \\
&\text{using probaSDSFSFSDFSDF} \\
&= \arg \max_{\Phi} \{\sum_{k=1}^K p(e_k|Y, \Phi^{(m)}) \log \frac{p(Y, e_k, \Phi)p(e_k, \Phi)}{p(e_k, \Phi)p(\Phi)}\} \\
&= \arg \max_{\Phi} \{\sum_{k=1}^K p(e_k|Y, \Phi^{(m)}) \log p(Y, e_k|\Phi)\} \tag{2.18} \\
&= \arg \max_{\Phi} \{E_{Z|Y, \Phi^{(m)}} \{\log p(Y, Z|\Phi)\}\}
\end{aligned}$$

We notice from equation (2.18), that , for a given voxel i :

$$\sum_{k=1}^K p(e_k|Y_i, \Phi^{(m)}) \log p(Y_i, e_k|\Phi) = \sum_{k=1}^K p_{ik}^{(m+1)} \log p(Y_i, e_k|\Phi) \tag{2.19}$$

Now, both expectation and maximization steps are apparents.

E-step

This this the expectation step. During this step, we estimate the probability that the pixel i belongs to class k regarding $\Phi_{(m)}$. This equation is obtained from equation (2.19) with Bayes formula.

$$p_{ik}^{(m+1)} = \frac{p(Y_i|e_k, \Phi^{(m)})p(e_k|\Phi^{(m)})}{\sum_{j=1}^K p(Y_i|e_j, \Phi^{(m)})p(e_j|\Phi^{(m)})} \tag{2.20}$$

Using this probability, $E_{Z|Y, \Phi^{(m)}}$ returns the expected value of the parameter Φ regarding $\Phi_{(m)}$.

M-step

This this the maximization step. During this step $\arg \max_{\Phi}$ maximizes $E_{Z|Y, \Phi^{(m)}}$ for the parameter Φ . It returns $\Phi_{(m+1)}$

$$\arg \max_{\Phi} \{E_{Z|Y, \Phi^{(m)}} \{\log p(Y, Z|\Phi)\}\} \tag{2.21}$$

We are no familiar with the theory behind the EM algorithm. The logic appears clearly and we can summarize the algorithm will the simple schema below: SDFSDFGSDFGSDF

2.4 Expectation maximization algorithm used in Slicer 3

In this part, we present the EM algorithm which has been integrated in Slicer 3 and the workflow in which it has been integrated as well. Finally, we briefly discuss the limitations of this one.

The EM algorithm which is used in the SPL is a derived one, from the original one. It enhances the original algorithm adding informations like a probabilistic atlas, a multichannel segmentation, a bias correction and a structure information. In Slicer 3, the GMM is used to describe the tissues to segment. Thus it will simplify the problem and the notations.

2.4.1 Probabilistic atlas

The EM algorithm is very sensitive to initialization since it only finds local extremums during the maximization step. A solution to enhance the initialization is to use atlases. There is an atlas for each class you want to segment. For each voxel i of the volume, it returns the probability that this voxel belongs to class k . This probability can be used as initialization.

$$p_{ik}^{(0)} = p_{ik}^{atlas} \quad (2.22)$$

From this value, it estimates $\Phi^{(1)}$ and the algorithm iterates until convergence is reached. The probabilistic atlases are not only used to initialize the process. It is also used to get a more robust algorithm. Indeed, we can use the spatial information given by the atlases. Voxels will be classified not only based on intensity but regarding spatial position as well. Van Leemput *et al.* ([8] and [9]) used the spatial prior at each iteration. It is constant and we then have a spatial information. The probability that a pixel i belongs to class k , in the E-Step changes. From equation (2.20), it becomes:

$$p_{ik}^{(m+1)} = \frac{p(Y_i|e_k, \Phi^{(m)})p_{ik}^{atlas}}{\sum_{j=1}^K p(Y_i|e_j, \Phi^{(m)})p_{ik}^{atlas}} \quad (2.23)$$

2.4.2 Multichannel segmentation

Most of the time, several modalities are used to process brain segmentation. Indeed, the best suited modality to use depends on the tissue you want to segment. For example, T1 MR images are well suited to segment white matter (WM) but is really bad for cerebrospinal fluid (CSF). On the contrary, T2 MR images are well suited for CSF and not for WM. To formalize it, we will change the definition of y_i and of μ_k we did at the beginning. Let $y_i = \{y_{i1}, y_{i2}, \dots, y_{iR}\}$ and $\mu_i = \{\mu_{i1}, \mu_{i2}, \dots, \mu_{iR}\}$ when we use R images, from different modalities to do the segmentation. The equations will remain the same.

2.4.3 Bias field correction

A major issue in MR modality is that the images can be corrupted by a low field bias field. It is due to equipment limitations or/and to patient induced electrodynamic interactions.

Principle

Let $I = (I_1, \dots, I_n)$ the observed intensities in an image, $I^* = (I_1^*, \dots, I_n^*)$ the ideal intensities and $F = (F_1, \dots, F_n)$ the bias field. Then, the degradation at each voxel can be expressed as:

$$I_i = I_i^* F_i$$

Let $Y = (Y_1, \dots, Y_n)$ and $Y^* = (Y_1^*, \dots, Y_n^*)$ be the log-transformed observed and ideal intensities. $B = (B_1, \dots, B_n)$ the log-bias field. This transform makes the bias field becomes additive instead of multiplicative without the log-approach.

$$Y_i = Y_i^* + B_i$$

We can model the PDF of the voxel intensity with a gaussian distribution

$$p(y_i | e_k, \Phi, B) = G(y_i - b_i, \mu_k, \sigma_k)$$

The low frequency characteristic of the bias field B can be modeled by a linear combination of smooth basics functions (ANNEXES) $\Psi_l(x)$. Let b_i be the realisation of the random variable B_i

$$b_i = \sum_{l=1}^L a_l \Psi_l(pos(i))$$

$pos(i)$ returns the 3D position (x, y, z) of the voxel i . a_i is the i^{th} value of the vector $A = (a_1, \dots, a_L)$. A represents the bias field parameters.

In the GM model, bias field can then be estimated using EM framework. The bias field parameter A will be used during the E-Step to estimate the probability that a voxel i belongs to class j . It will be computed during the M-Step, in parallel with the maximization of the parameters (mean and variance). Van Leemput formalised the two steps as below ([8] and [9]):

- **E-step**

$$p_{ik}^{(m+1)} = \frac{G(y_i - b_i, \mu_k^{(m)}, \sigma_k^{(m)}) p_{ik}^{atlas}}{\sum_{j=1}^K G(y_i - b_i, \mu_j^{(m)}, \sigma_j^{(m)}) p_{ij}^{atlas}}$$

- **M-step**

- Gaussian distribution parameters estimation

$$\mu_k^{(m+1)} = \frac{\sum_{i=1}^n y_i p_{ik}^{(m+1)} - b_i}{\sum_{i=1}^n p_{ik}^{(m+1)}}$$

$$(\sigma_k^{(m+1)})^2 = \frac{\sum_{i=1}^n (y_i - \mu_k^{(m+1)} - b_i)^2 p_{ik}^{(m+1)}}{\sum_{i=1}^n p_{ik}^{(m+1)}}$$

- Bias field correction

$$(A^{(m+1)})^T = (F^T W^{(m+1)} F)^{-1} F^T W^{(m+1)} R^{(m+1)}$$

with:

$$\mathbf{F} = \begin{pmatrix} \Psi_1(pos(1)) & \Psi_2(pos(1)) & \dots & \Psi_L(pos(1)) \\ \Psi_1(pos(2)) & \Psi_2(pos(2)) & \dots & \Psi_L(pos(2)) \\ \vdots & \vdots & \ddots & \vdots \\ \Psi_1(pos(N)) & \Psi_2(pos(N)) & \dots & \Psi_L(pos(N)) \end{pmatrix}$$

$$\mathbf{W}^{(m+1)} = \begin{pmatrix} \sum_{k=1}^K w_{1k}^{(m+1)} & 0 & \dots & 0 \\ 0 & \sum_{k=1}^K w_{2k}^{(m+1)} & \dots & 0 \\ \vdots & \vdots & \ddots & \vdots \\ 0 & \dots & 0 & \sum_{k=1}^K w_{Nk}^{(m+1)} \end{pmatrix}$$

$$w_{ik}^{(m+1)} = \frac{p_{ik}^{(m+1)}}{(\sigma_k^{(m+1)})^2}$$

$$\mathbf{R} = \begin{pmatrix} y_1 - \tilde{y}_1^{(m+1)} \\ \vdots \\ y_N - \tilde{y}_N^{(m+1)} \end{pmatrix}$$

$$\tilde{y}_i^{(m+1)} = \frac{\sum_{k=1}^K w_{ik}^{(m+1)} \mu_{ik}^{(m+1)}}{\sum_{k=1}^K w_{ik}^{(m+1)}}$$

The bias field correction step can be interpreted as follows: the estimated soft segmentation and gaussian distribution parameters are used to reconstruct the image $\tilde{Y} = (\tilde{y}_1, \dots, \tilde{y}_n)$. The new image is supposed not to be corrupted by the bias field. We then subtract the reconstructed image

\tilde{Y} from the observed image Y . We obtain the residual image R . From R , we estimate the bias field. F represents the discretized geometry of the bias field. W is an inverse covariance matrix.

The approach used in Slicer 3 is based on the same principle but differs regarding the parameters which are known and estimated, in the EM algorithm.

Variation used in Slicer 3

In this method, we are working with a GMM. Moreover the parameters of this gaussian distribution are assumed to be known. The idea of estimating the field in EM framework was originally proposed by Wells *et al.* ([10]) The EM is used to estimate the bias field using the *maximum a posteriori principle* (MAP) instead of the maximum likelihood principle:

$$\hat{\Phi} = \arg \max_{\Phi} p(Y|\Phi) = \arg \max_{\Phi} \frac{p(\Phi|Y)p(\Phi)}{p(Y)} = \arg \max_{\Phi} p(Y|\Phi)p(\Phi)$$

The estimation of Φ , during the E-Step, can no longer be done with E We must include the fact $\log(\Phi)$. In Wells' method, the only parameter to be estimate is the bias field:

$$p(\Phi) = p(B) = G(B, 0, \Sigma_B)$$

The equation for the bias field will change. We add the smoothness constraint in the gaussian distribution: Σ_B^{-1} . We also set F to unit matrix as no parametric model for the bias field is assumed. We also define the mean residual image $\bar{M}^{(m+1)}$

$$\bar{M}^{(m+1)} = W^{(m+1)} R^{(m+1)}$$

The equation for the bias field, in the maximization step $(A^{(m+1)})^T$ will then be replaced by $(B^{(m+1)})^T$:

$$(B^{(m+1)})^T = (W^{(m+1)} + \Sigma_B^{-1})^{-1} \bar{M}^{(m+1)}$$

2.4.4 Hierarchical information

The last modification of the original EM algorithm is the addition of hierarchical information in the iterative process. The algorithm was described by Pohl *et al.* ([?]) The idea was to describe the structures we want to segment as a tree. At the first iteration, the highest leaves of the tree, the parents leaves, will be segmented. Then, following an iterative process, the child of each parent will be segmented, then the child of the child, etc..

At the first iteration, the MR image will be segmented into the background (BG) and the intracranial cavity (ICC). At the second iteration, the

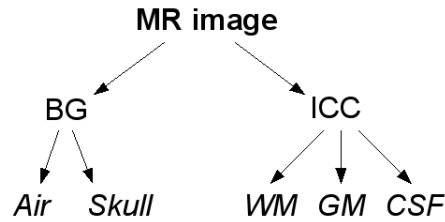


Figure 2.1: A simple tree structure of the brain

BG will be segmented into the air and the skull. Finally, at the last iteration, the ICC will be segmented into white matter (WM), grey matter (GM) and cerebrospinal fluid (CSF).

We incorporate H , a set of structure-specific information in SDFSDF. Add H , (re) define clearly W . EM segmentation done for each set of childs ALGO..?

2.5 Workflow in Slicer 3

We will now present the whole workflow used in Slicer 3, for the EM segmentation module. This will describe all the initialisation steps done by the user, via the graphi user interface. It will also present how the whole algorithm works. We will remind why is each initialisation step important and where is this information used in the pre-processing treatments or in the EM algorithm.

2.5.1 User interface

Step 1: Tree structure creation

We first create the tree structure we want to use for the segmentation. It will be used to define H .

Step 2: Atlas assignement

We assign for each node of the tree, the related atlas. It will be used to define H .

Step 3: Multimodal segmentation

We choose the images we want to use for the segmentation. As discussed in DSFSD, it is usefull because some tissues are not clearly visible in some modalities. It helps us to define Y , the observed images.

Step 4: Intensity normalization

We normalize the intensity of the images to be segmented, regarding the atlas. It is usefull if we want to use the command-line module. Doing an intensity normalisation, the initialisation value will be the same for each class, using the same atlas. It is a convenient way to run a lot of segmentation. Indeed, the initialisation values are supposed to remain the same.

Step 5: Class definition

We define mean value, variance and covariance for each class and modality. Moreover, it is a more precise way to initialise the algorithm. It is usefull because the EM algorithm used only estimates the bias field but still requires informations about the tissues to be segmented.

Step 6: Prior weights

We define the importance of each image and of the atlas for each node. It is will be usefull to define H . For each tissue to segment, H knows usefull informations as which modality is the more relevant and the size that the tissue is supposed to be.

Step 7: Registration selection

We choose the type of registration we want. Different kind of registrations are available.

2.5.2 Algorithm

Step 1: Intensity normalisation

Step 2: Images registration

Step 3: EM Algorithm in the tree structure

2.5.3 Summary

In Slicer 3, the whole segmentation workflow

2.6 Limitations

Sensitive to initialization.(vizu) Not reproducible (labelmap) problem of registration (bias correction) at things in the workflow

We are now aware of the whole

After the presentation of the main issues, we will now show you some solution we brought to try to solve the problems.

Chapter 3

The contributions

3.1 MRI Bias Field correction

Bias correction is not efficient so we propose an other method, as part of the segmentation process.

Rescaling

3.1.1 Interest

algorithm is only accurate for small region bias correction, because au the gaussian model he uses. Correction the bias in the whole volume first could enhance results.

3.1.2 Our approach

Gaussian distrib.... ITK filter Automatic

3.1.3 Results

corrected not corrected parameters explanation

3.2 Intensity Normalization

3.2.1 Interest

usefull if you need to do a lot of segmentations enhance the rescaling..?

3.2.2 Our approach

background/brain

3.2.3 Our implementation

show results... corrected image...

3.3 Class Distribution selection

During parameters initialization, the user has to define each class distribution. The current methods presents some troubles so we present here a new approach to estimate each class distribution.

3.3.1 Interest

So far, the user haf two choices to define each class distribution. He could enter manually the intensities mean value and variance for each class, for each volume to be processed. This way, the user can be very precise and accurate when he defines each class. But it is very hard for the user to found the good mean value and variance for each class for each volume. Moreover, each time he wants to process a new volume, he will have to redefine mean values and variances. This is not convenient and it can takes a ot of time. The next approach consisted in defining a class model by manual sampling. For each class, the user clics in the related part of the volume. With this methos, the problem is that you compute your mean value and variance using a few samples. Then, your mean values and variances are not accurate. That's why we proposed a new approach using a label map, to estimate each class model.

3.3.2 Method used

Create a label map. colors we choose a the same as the one we defined for each class. then run algorithm which will estimate the mean value and covariance values.

Algorithm..?

Explain each colors

3.4 Class Distribution visualization

An important contribution is a tool which allows to visualize the classes to be segmented distribution, through 2 volumes.

3.4.1 Interest

As we discussed in the EMS algorithm requires variance and mean values for each class to be segmented. So far, the user defines these parameters by selecting samples on the dataset. exampleSDFSF A main issue is that user

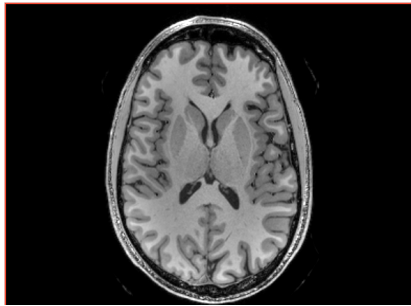


Figure 3.1: Axial view of a T1 volume without the label map.

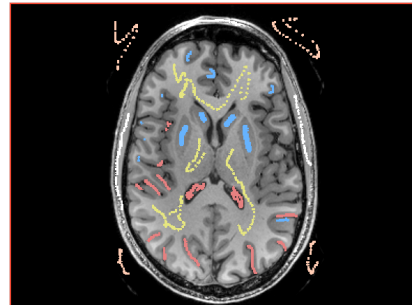


Figure 3.2: Axial view of a T1 volume with the label map.

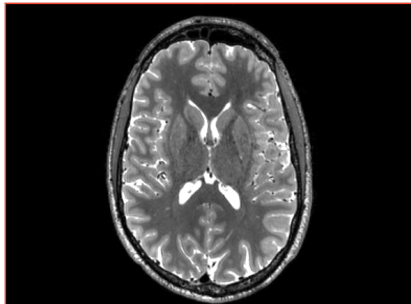


Figure 3.3: Axial view of a T2 volume without the label map.

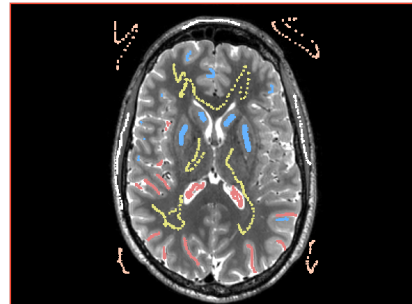


Figure 3.4: Axial view of a T2 volume with the label map.

user can know if the pixels he selected are really representative or not of the class he wants to select. Thus, our objective was to find a way to give the user if his samples are representative or not of a tissue.

3.4.2 Method used

Problem's description

Here is a brief description of the situation to let you understanding the problem. As input we have N images, K classes to segment, and for each class a mean value μ , a variance σ and covariance ζ . The covariance is estimated for each couple of images. We will come back on it later. where Using these informations, we had to find a way to give the user usefull informations about the class distribution.

Choices

Our first idea was to display gaussian curves in 3D, for each class. The X and Y axes would be the intensity ranges for the two images we want to use. The problem with this representation is that it does not contain any information about the number of pixels. 3d too sophisticated...? To solve this problem, we decided to divide the problem in two parts. The first part will consist in creating a "temperature map" as background. The second part will consists in displaying ellipses over this map. This will give the user a good information about how good his selection was.

Background creation

First of all, the background has to be created. This background is a "temperature map". We can present it as a two dimensions array. The height and the length of the array depend of the intensity range of the 2 volumes we want to process. Then, we fill the array in an easy but useful way. Each case of the array contains the number of occurrence of an intensity pair at the same position, through the 2 volumes.

Algorithm 1: BACKGROUND_CREATION($V1, V2$)

```

define RangeI1  $\leftarrow$  intensity range in  $V1$ 
define RangeI2  $\leftarrow$  intensity range in  $V2$ 
define  $A(\text{RangeI1}, \text{RangeI2}) \leftarrow$  background
for each pixel  $i$  in  $V1$ 
for each pixel  $j$  in  $V2$ 
   $\triangleright A(i, j) = A(i, j) + 1$ 
return  $A$ 

```

Ellipse representation

We will first start with a brief rappel.. The covariance matrix is
Representing ellipse associated to this covariance matrix.
Maximum variance axe.Eigen values
Algorithm used. Big and small axes of the ellipse

3.4.3 Results

show result

3.5 Global Prior Estimation

The last contribution to the EMS is a tool which provides the user an easy and fast way to estimate the global prior weights (GPW).(ref ch2 ...)

3.5.1 Presentation of the problem

This contribution is usefull in many different ways. When you run the segmentation process,at the 6th step of the process, you have to provide to the algorithm an estimation of the GPW for each node in the tree. First of all if there are a lot of structures to segment, they user can spend a lot of time during this step. Indeed, for each part of this tree strcuture, they have to define the GPW. Moreover, the user may not know at all which weights to choose. This new approach will provide the users a good estimations of the weights to use. We must also keep in mind that the end users are physicists. They might don't understand what the parameters meanings and providing them a visual feedback could help them a lot.

3.5.2 Our approach

We divided the problem in two parts. The first part will be about providing the user a real-time feedback regarding the GPW estimation. The second part will consist in developping an algorithm which fills automatically the tree.

Fast user feedback

We can divide the feedback part in 3 steps: the histogram computation and utilisation, the multicolumn list and the labelmap generated. The histogram allows the user to manual segment classes based on intensity. The multicolumn list allows the user to change the order of the classes in the histogram. The labelmap provides to the user a visual feedback, base on the segmentation realized in the histogram. Using these three complementary tools, the

user, even if he is not initiated can estimate easily, accurately and rapidly the GWP.
 schema

Global priors evaluation

The algorithm used to estimate the weight of each node is iterative. It starts from the root and goes to the leaves. It evaluates the weight of the childs of the active node at each iteration. Here is a description of the algorithms used to compute the GPW of each node.

Description algo 1

Algorithm 1: TREEWEIGHTESTIMATION(R, W)

```

define C = CHILD(R)  $\leftarrow$  set of childrens of root R
define LEAF(C)  $\leftarrow$  set of leaves of tree with roots C
define H  $\leftarrow$  set of structure-specific information defined by LEAF(C) for
each leaf
update W in childrens of root R with the results of WEIGHTESTIMA-
TION(C,LEAF(C),H)
for each node R' in CHILD(R) that is not a leaf
 $\triangleright$  TREEWEIGHTESTIMATION(R', W)

```

Description algo2

The algorithm used estimates the global prior of the leaves of the current node, based on the number of pixel which belong to the child classes. This number of pixels is calculated from the segmentation computed in the histogram.(CF ..)

Algorithm 2: WEIGHTESTIMATION(C,LEAF(C), W,H)

```

define T  $\leftarrow$  set of total weight of leaves in LEAF(C). Leaves weights are
contained in H
define E  $\leftarrow$  set of weight for each node of C
for each node of C
 $\triangleright$  E=E+H : Get the total weight of each node
W=E/T
return (W)

```

3.5.3 Results and discussion

As we can see, the results obtained are good and accurate. The time of processing is fast and most of the people are now able to understand these parameters, even they are not familiar with EMS. This is an intuitive way

to get an estimation of the GPW parameters.

Of course some sfsdfs must be done. If there is a strong bias in the images we process, the pixels of a same class will have different values. Then, a segmentation based on intensity will provide bad results. Moreover, if the user wants to segment 2 classes which have the same color, just using the spatial information for example, they won't be able to use it properly. Thus, the user must always keep in mind that it is just an estimation and that he has to check if the values are accurate.

Chapter 4

Results and discussion

This chapter will review what has been done and mentions the main open questions.

Results using different tools
limitations
improvement (in relation to limitations)

4.1 Results

Tests has been performed to show the utility of the work done. The results are presented below. The testings are comparaisons between the results obtained with the previous workflow and the new one. The results obtained will then be reviewed by a specialist to evaluate the previous and the new segmented datasets.

4.1.1 Bias correction

Here we get going with the intensities inhomogeneities correction.

Testing process

we create a mask.

Results

Here are the results

more comments tttttttttttttttttt ttttttttttt tttttttttt tttttttttt
 tttttttttt tttttttttt tttttttttttttttttt ttttttttttt tttttttttttttttttt ttttttttttt ttttttttttt

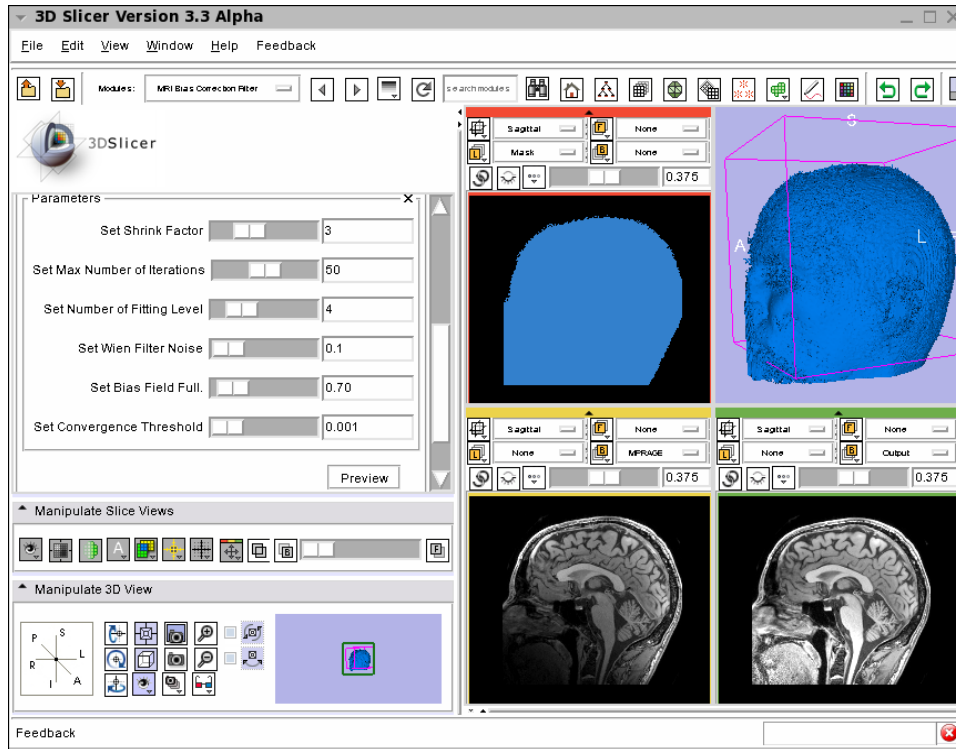


Figure 4.1: Module created for the bias correction.

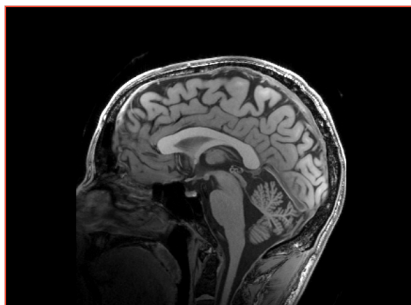


Figure 4.2: Sagittal view of a biased T1 volume.



Figure 4.3: Sagittal view of the T1 volume after bias correction.

Specialist's point of view

4.1.2 Global Prior estimation

Testing process

Results

Specialist's point of view

4.1.3 Class Selection

Testing process

Results

Specialist's point of view

4.2 Limitations

4.3 Future work

Acknowledgements

Ron Kikinis who gave me the opportunity to carry out my intersnship in the SPL. Sylvain Jaume who supervises me during all my work. Andryi, Daniel, Steve?

Bibliography

- [1] A.P. Dempster, N.M. Laird, and D.B. Rubin, "Maximum likelihood from incomplete data via the em algorithm", *Journal of the Royal Statistical Society: Series B*, vol. 39, pp. 1-38, November 1977.
- [2] Y. Weiss, "Bayesian motion estimation and segmentation", *PhD thesis*, Massachusetts Institute of Technology, May 1998.
- [3] R.C. Hardie, K.J. Barnard, and E.E. Armstrong, "Joint MAP registration and high-resolution image estimation using a sequence undersampled images", *IEEE Transaction on Image Processing*, vol. 6, pp. 1621-1633, December 1997.
- [4] M. Murgasova, "Tutorial on Expectation-Maximization: Application to Segmentation of Brain MRI", May 2007.
- [5] S. Borman, "The Expectation Maximization Algorithm", January 2009.
- [6] Wikipedia, "Expectation-Maximization algorithm", http://en.wiki.org/wiki/Expectation-maximization_algorithm, June 2009.
- [7] G. McLachlan, and T. Krishnan, "The EM Algorithm and Extensions", *John Wiley & Sons*, New York, 1996.
- [8] K. V. Leemput, F. Maes, D. Vandermeulen et al., "Automated model-based tissue classification of MR images of the brain", *IEEE Transaction on Medical Imaging*, 18(10), pp. 857-908, 1999.
- [9] K. V. Leemput, F. Maes, D. Vandermeulen et al., "Automated model-based bias field correction of MR images of the brain", *IEEE Transaction on Medical Imaging* 18(10), pp. 885-896, 1999.
- [10] W. M. Wells III, W.E.L. Grimson, R. Kikinis et al., "Adaptive segmentation of MRI data", *IEEE Transaction on Medical Imaging* 15(4), pp. 429-442, 1996.