

# An automatic method for change detection in serial DTI-derived scalar images

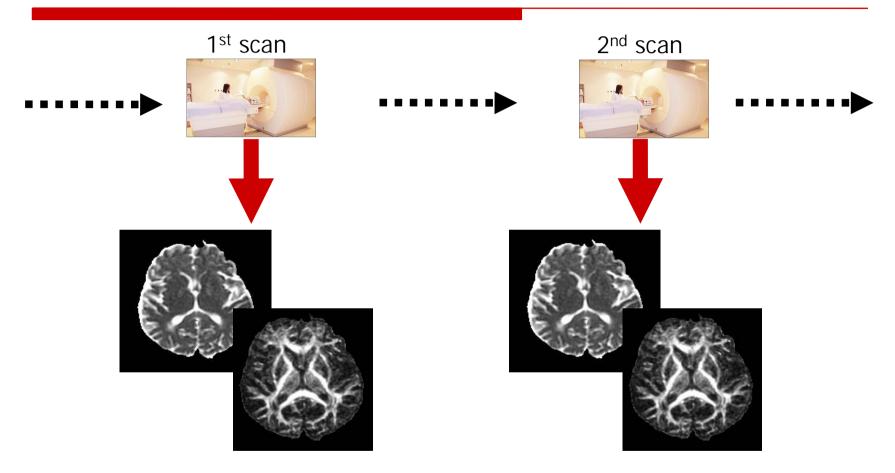
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Work supported by



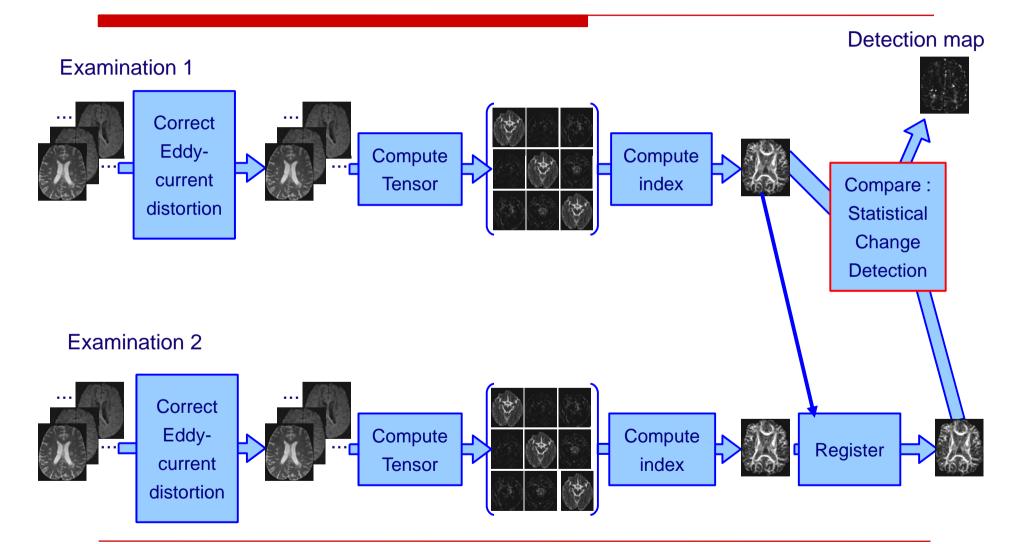
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# Objective



=> Detect significant changes between diffusion-weighted scalar images

## **General scheme**



## Statistical change detection

We consider the Generalized Likelihood Ratio Test (GLRT): Likelihood ratio under 2 hypothesis:

- $H_0$ : there is no significant change between images  $I_1$  and  $I_2$ .
- $H_1$ : there is a significant change between images  $I_1$  and  $I_2$ .

$$GLRT = \frac{p(I_1|\hat{\theta}_1) \ p(I_2|\hat{\theta}_2)}{p(I_1|\hat{\theta}_0) \ p(I_2|\hat{\theta}_0)}$$

 $I_1$  and  $I_2$  are considered as realizations of random variables drawn according to parametric probability density functions:

- with the same parameters  $\theta_0$  under assumption H<sub>0</sub>
- with two sets of different parameters  $\theta_1 \neq \theta_2$  under assumption H<sub>1</sub>

## Bosc's method

Bosc *et al* (Neuroimage, vol 20 (2), pp 643-656, 2003) use a similar framework for detecting MS lesion evolution in conventional multimodal MRI sequences

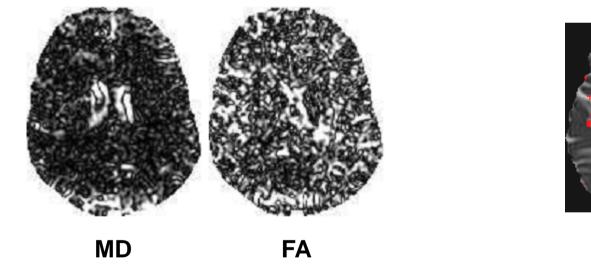
Hypothesis:

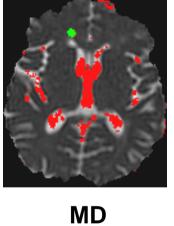
- Intensities are modeled as a constant value  $\mu$  in a window W<sub>s</sub> of 3x3x3 voxels
- additive stationary Gaussian noise

$$\log GLRT(s) \propto rac{\left(\mu_2^{W_s}-\mu_1^{W_s}
ight)^2}{\sigma^2}$$

### Bosc's method

However, Bosc's approach fails when coping with DTI-derived scalar images because of the non-stationarity of noise.





## **Proposed approach**

We propose to account for the non stationarity of the noise by estimating a variance at each voxel:

$$GLRT(s) = \frac{\sigma_0^2(s)}{\sqrt{\sigma_1^2(s) \ \sigma_2^2(s)}} \exp \frac{(I_1(s) - \mu_0(s))^2 + (I_2(s) - \mu_0(s))^2}{2 \ \sigma_0^2(s)}$$

Estimation of the variance is done using the method described in Chang *et al*, Magnetic Resonance in Medicine, vol 57(1), pp 141 - 149, 2007

## Variance estimation (1/2)

Variance on the tensor elements is obtained as a by-product of least squares estimation:

$$Y = AX \qquad X = [D_{xx}, D_{yy}, D_{zz}, D_{xy}, D_{xz}, D_{yz}]^{t}$$
$$Y = AX \qquad Y = [1/b \ \log(S_{0}/S_{1}), \dots, 1/b \ \log(S_{0}/S_{N})]^{t}$$
$$A(i, :) = \left[g_{i;x}^{2}, g_{i;y}^{2}, g_{i;z}^{2}, 2g_{i;x}g_{i;y}, 2g_{i;x}g_{i;z}, 2g_{i;y}g_{i;z}\right]$$

.

$$\widehat{X} = (A^{t}A)^{-1}A^{t}B$$

$$\Sigma_{X} = \sigma^{2}(A^{t}A)^{-1} \text{ with } \sigma^{2} = \frac{1}{N}(Y - A\widehat{X})^{t}(Y - A\widehat{X})$$

## Variance estimation (2/2)

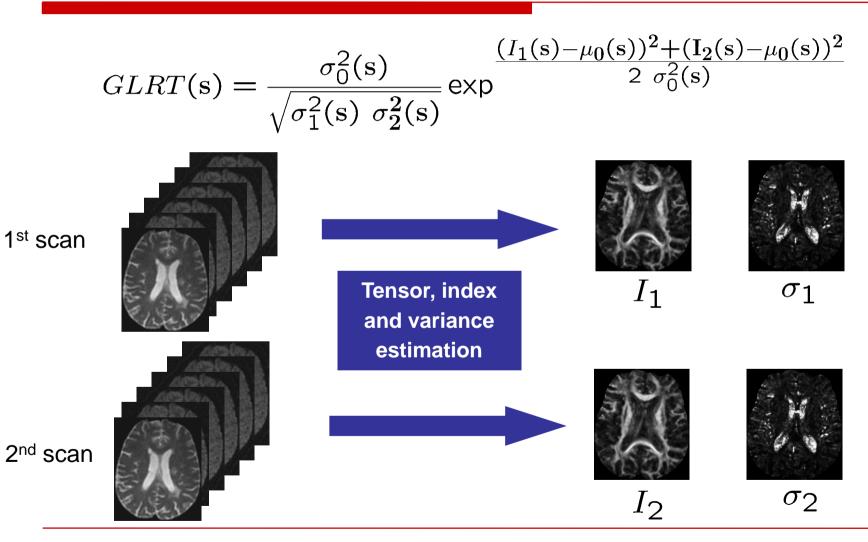
The covariance matrix is propagated to the eigenvalues of X

$$\Sigma_{\lambda} = R \ \Sigma_X \ R^t$$

and then to FA or MD

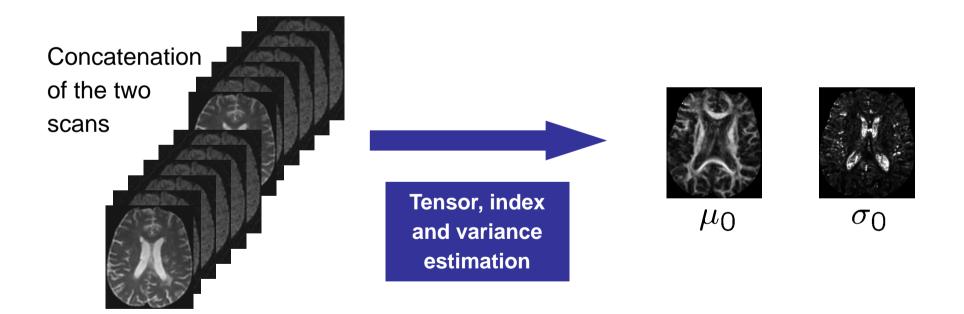
$$\begin{split} \sigma_{MD}^{2} &= \sigma_{\lambda_{1}}^{2} + \sigma_{\lambda_{2}}^{2} + \sigma_{\lambda_{3}}^{2} + 2 \sigma_{\lambda_{12}}^{2} + 2 \sigma_{\lambda_{13}}^{2} + 2 \sigma_{\lambda_{23}}^{2} \\ \sigma_{FA}^{2} &= \sigma_{\lambda_{1}}^{2} \left(\frac{\partial FA}{\partial \lambda_{1}}\right)^{2} + \sigma_{\lambda_{2}}^{2} \left(\frac{\partial FA}{\partial \lambda_{2}}\right)^{2} + \sigma_{\lambda_{3}}^{2} \left(\frac{\partial FA}{\partial \lambda_{3}}\right)^{2} \\ &+ 2 \sigma_{\lambda_{12}}^{2} \frac{\partial FA}{\partial \lambda_{1}} \frac{\partial FA}{\partial \lambda_{2}} + 2 \sigma_{\lambda_{13}}^{2} \frac{\partial FA}{\partial \lambda_{1}} \frac{\partial FA}{\partial \lambda_{3}} + 2 \sigma_{\lambda_{23}}^{2} \frac{\partial FA}{\partial \lambda_{2}} \frac{\partial FA}{\partial \lambda_{3}} \end{split}$$

#### GLRT: H<sub>1</sub> hypothesis

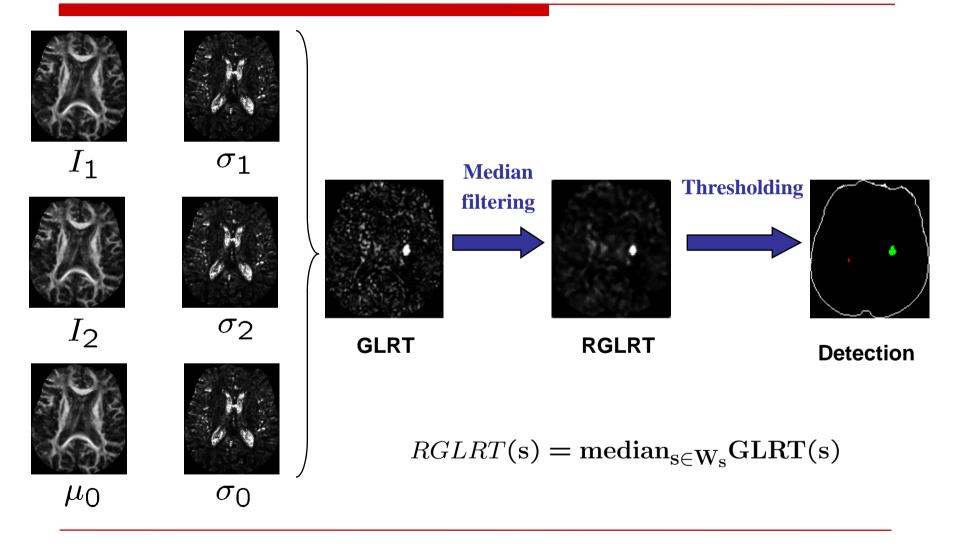


#### GLRT: H<sub>0</sub> hypothesis

$$GLRT(s) = \frac{\sigma_0^2(s)}{\sqrt{\sigma_1^2(s) \ \sigma_2^2(s)}} \exp \frac{(I_1(s) - \mu_0(s))^2 + (I_2(s) - \mu_0(s))^2}{2 \ \sigma_0^2(s)}$$



# **Spatial information**



# Validation

- Validation of such method is difficult due to the lack of ground truth
- Even for experts, it is difficult to detect changes in DT-images
- □ We resort to simulation of MS lesion appearance

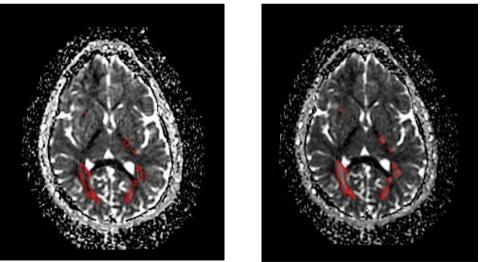
# **MS** lesion simulation

Experimental studies have shown that demyelination leads to an increase of the radial diffusivity  $(D_{\perp})$  in DW-images.

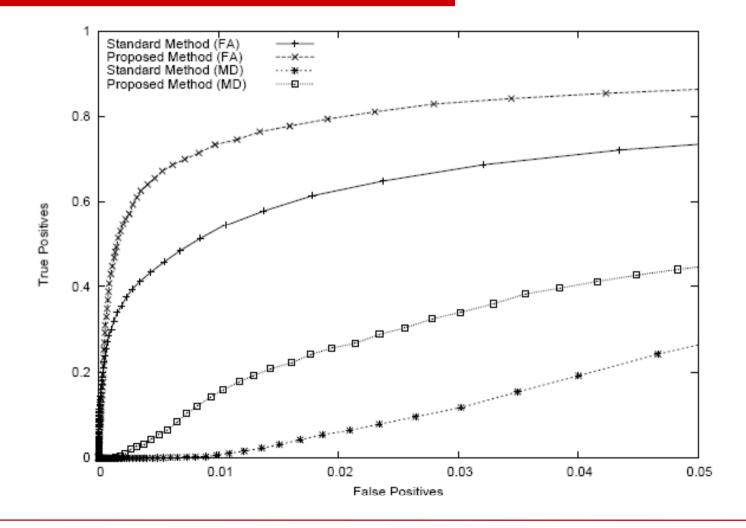
[Harsan et al, Journal of neuroscience research, vol 83(3), pp 392-402, 2006]

• Two scans of a healthy subject are considered

• The two last eigenvalues (corresponding to the radial diffusion) of the second examination are increased in several regions of Interest to simulate lesion apparitions

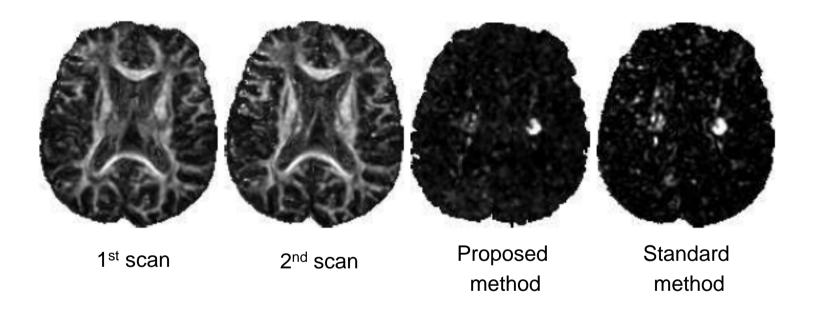


## **MS** lesion simulation



MIAMS'08: MICCAI workshop on "Medical Image Analysis on Multiple Sclerosis"

# Follow-up of MS patients



# **Conclusion and perspectives**

- **Further validation should be done on real cases**
- Automatic threshold selection: p-value estimation, multiple comparison problem
- Extension for multimodal detection
- Extension to tensor images and to other model of diffusion