



Department of Neuroradiology  
Klinikum rechts der Isar  
Technical University of Munich



# Comparing myelin-sensitive markers MWF, ihMTR, and MTsat in healthy and normal-appearing brain tissue and multiple sclerosis lesions

Ronja C. Berg<sup>1</sup>, Viola Pongratz<sup>2</sup>, Markus Lauerer<sup>2</sup>, Thomas Amthor<sup>3</sup>, Guillaume Gilbert<sup>4</sup>, Aurore Menegaux<sup>1</sup>, Claus Zimmer<sup>1</sup>, Christian Sorg<sup>1</sup>, Mariya Doneva<sup>3</sup>, Irene Vavasour<sup>5</sup>, Mark Mühlau<sup>2</sup>, Christine Preibisch<sup>1</sup>

Poster Session **Multiple Sclerosis II**

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Computer # **10**

Program # **0890**

<sup>1</sup> Technical University of Munich, School of Medicine, Department of Neuroradiology, Munich, Germany

<sup>2</sup> Technical University of Munich, School of Medicine, Department of Neurology, Munich, Germany

<sup>3</sup> Philips Research Europe, Hamburg, Germany

<sup>4</sup> MR Clinical Science, Philips Healthcare, Mississauga, ON, Canada

<sup>5</sup> University of British Columbia, Department of Radiology, Vancouver, BC, Canada





JOINT ANNUAL MEETING ISMRM-ESMRMB

ISMRT 31<sup>ST</sup> ANNUAL MEETING

07-12 MAY 2022 | LONDON, ENGLAND, UK

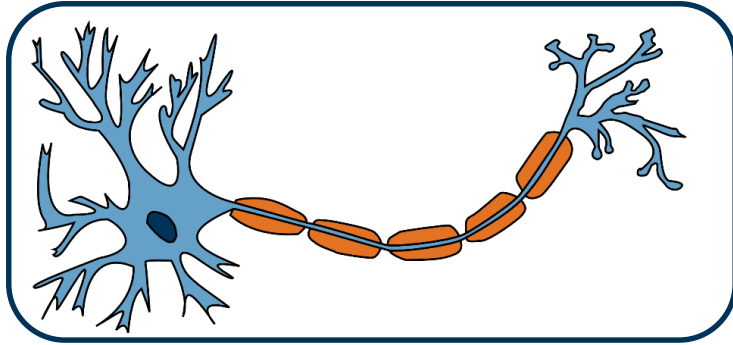
A HYBRID EXPERIENCE



# Declaration of Financial Interests or Relationships

Speaker Name: **RONJA BERG**

I have no financial interests or relationships to disclose with regard to the subject matter of this presentation.



MS

Demyelination



Aim

Monitoring demyelinating diseases

Imaging methods

Myelin water imaging



Myelin-sensitive markers

MWF

$$= \frac{\text{Myelin water}}{\text{Total water}}$$

Magnetization transfer



ihMTR

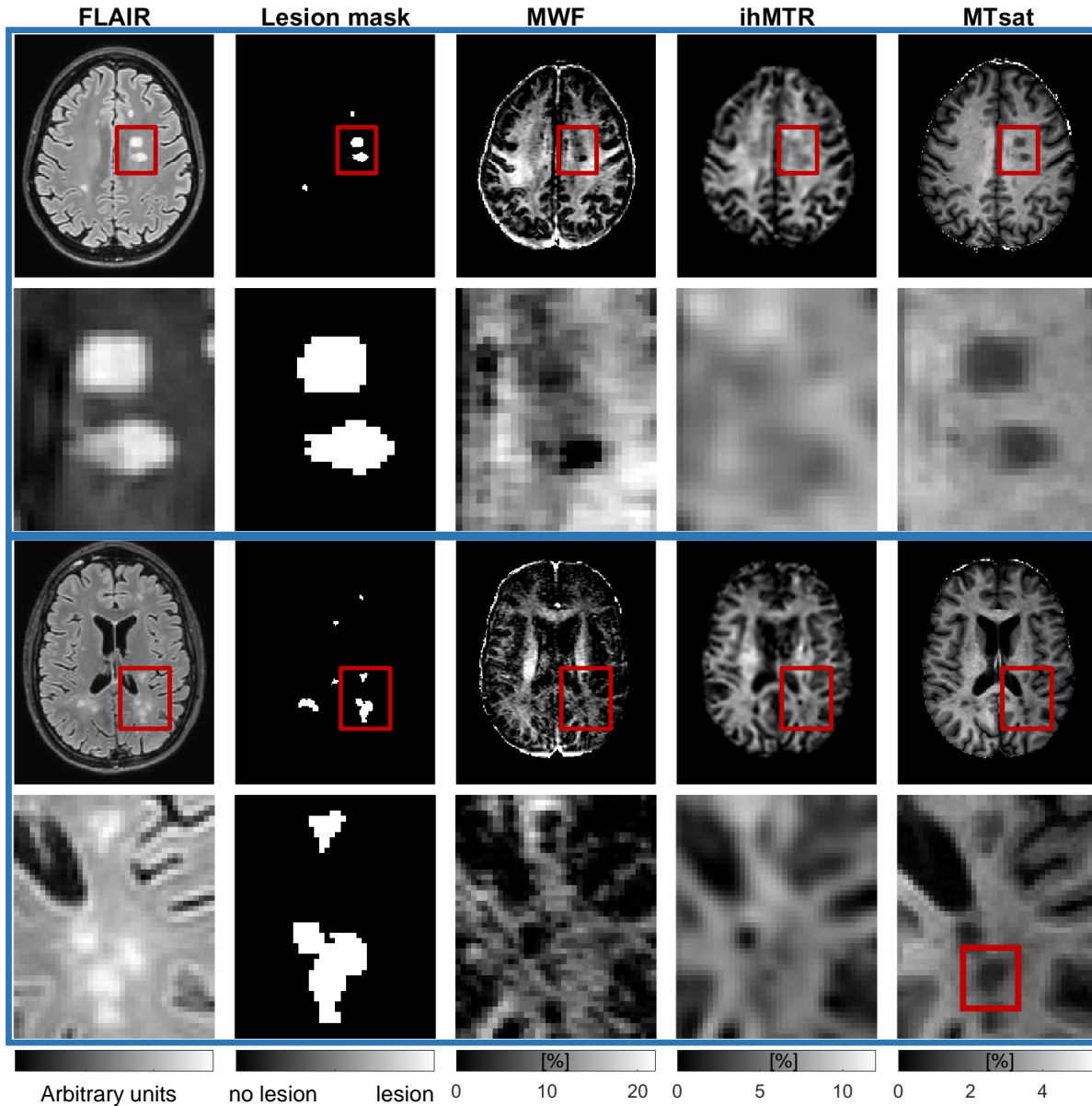
$$= \frac{1}{S_0} (MT^+ + MT^- - MT^{+-} - MT^{-+})$$



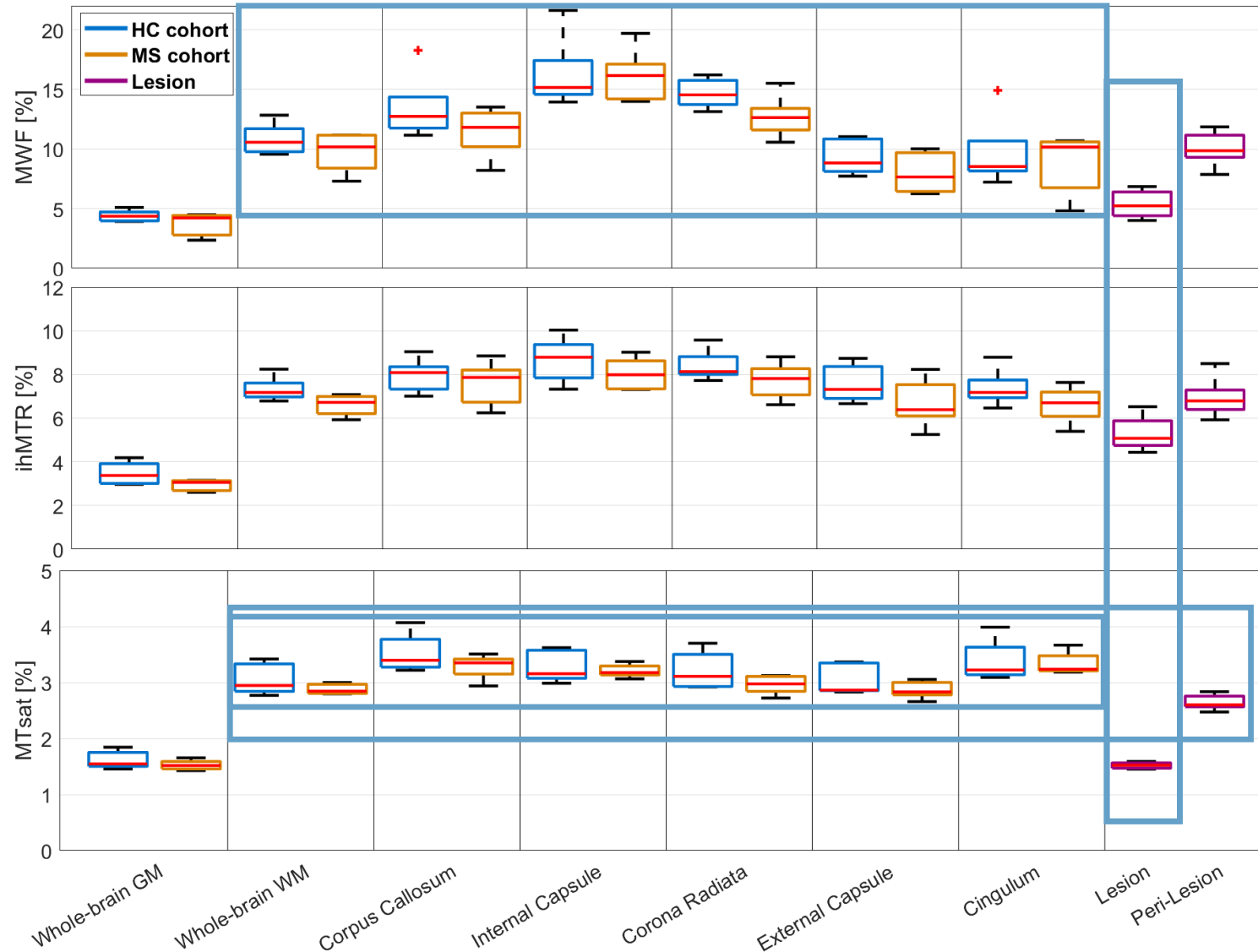
MTsat

$$= \left( \frac{A_{app} * \alpha}{S_{MT}} - 1 \right) * R_{1app} * TR - \frac{\alpha^2}{2}$$

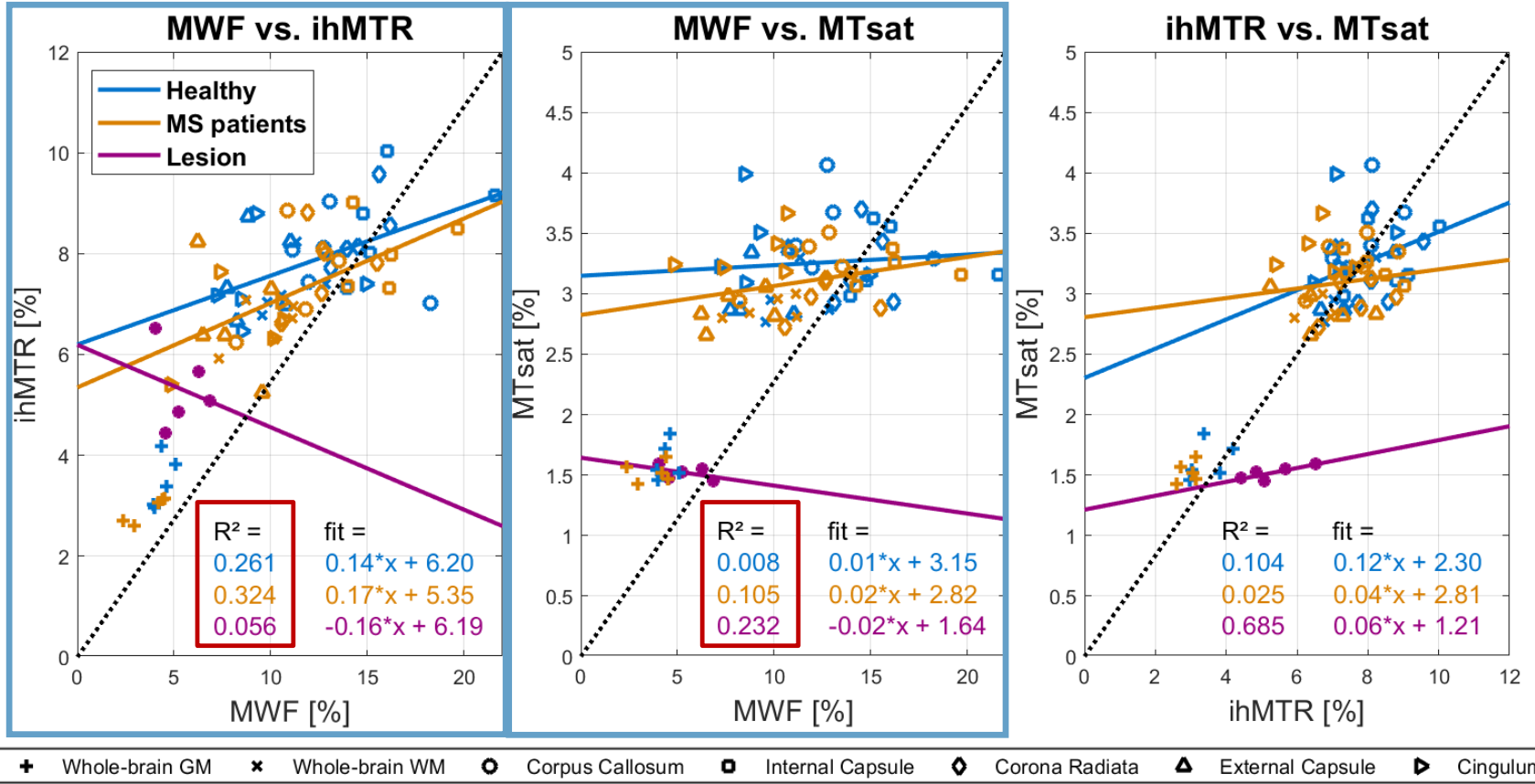
<b>Participants</b>	<b>3 T, Philips</b>	<b>MWF</b>	<b>Lesions</b>	<b>Brain regions</b>
<p><u>Healthy:</u></p> <ul style="list-style-type: none"> <li><math>n = 5</math>, 3f / 2m</li> <li><math>32 \pm 3</math> years</li> </ul> <p><u>MS patients:</u></p> <ul style="list-style-type: none"> <li><math>n = 5</math>, 2f / 3m</li> <li><math>33 \pm 6</math> years</li> <li>4 RRMS, 1 CIS</li> <li>Disease duration: 3-15 y, (avg.: 9.4 y)</li> <li>EDSS: 0-1.5, (avg.: 1.1)</li> </ul>	<p><u>MWF:</u></p> <ul style="list-style-type: none"> <li>3D gradient- and spin-echo (GRASE) sequence</li> <li><math>TE1/\Delta TE = 8/8</math>ms,</li> <li>48 echoes, 20 slices</li> <li>res: <math>1 \times 2 \times 5</math> mm<sup>3</sup></li> </ul> <p><u>ihMTR:</u></p> <ul style="list-style-type: none"> <li>3D gradient-echo (GE)</li> <li><math>TE1/\Delta TE = 3.5/5.7</math>ms</li> <li>res: <math>2.2 \times 2.2 \times 2.</math> mm<sup>3</sup></li> <li>10 MT pulses, <math>\alpha_{MT} = 90^\circ</math>, <math>t_{MT} = 0.9</math> ms</li> </ul> <p><u>MTsat:</u></p> <ul style="list-style-type: none"> <li>3 x 3D GE: 1) <math>\alpha = 4^\circ</math>, 2) <math>\alpha = 25^\circ</math>, both TR = 18ms; 3) MT-w: <math>\alpha = 6^\circ</math>, TR = 35ms</li> <li>All: <math>1 \times 1 \times 1</math> mm<sup>3</sup>, 6 echoes</li> <li><math>TE1/\Delta TE = 2.4/2.4</math> ms</li> <li>MT pulse: <math>\alpha_{MT} = 540^\circ</math>, <math>t_{MT} = 12.8</math> ms, <math>f_{MT} = 2200</math> Hz</li> <li>4) B1-map for bias field correction</li> </ul>	<p><u>MWF:</u></p> <ul style="list-style-type: none"> <li>Using the Sparsity Promoting Iterative Joint Non-negative least squares (SPIJN) algorithm</li> </ul> <p><u>ihMTR:</u></p> <ul style="list-style-type: none"> <li>Combination of single and dual frequency offset saturation MT images</li> </ul> <p><u>MTsat:</u></p> <ul style="list-style-type: none"> <li>Parameter map calculation via the hMRI toolbox</li> </ul>	<p><u>Segmentation:</u></p> <ul style="list-style-type: none"> <li>Lesion growth algorithm, lesion segmentation tool for SPM12</li> <li>Based on FLAIR &amp; MPRAGE</li> </ul> <p><u>Lesion VOI:</u></p> <ul style="list-style-type: none"> <li>Lesion probability &gt; 0.5</li> </ul> <p><u>Peri-lesion:</u></p> <ul style="list-style-type: none"> <li>3-voxel wide shell surrounding lesions within NAWM</li> </ul>	<p><u>Whole-brain:</u></p> <ul style="list-style-type: none"> <li>GM and WM segmentation SPM12's segment module (tissue prob. &gt; 0.5)</li> </ul> <p><u>Anatomical:</u></p> <ul style="list-style-type: none"> <li>Several tracts from the ICBM-DTI-81 WM labels atlas</li> </ul> <p><u>Evaluation:</u></p> <ul style="list-style-type: none"> <li>In subjects' native spaces (MPRAGE data space)</li> </ul>



- Visual similarity
- Appearance of lesions
  - Lower values
  - Some differences visible
- White matter
  - Stronger variation in MWF
  - MTsat most homogeneous

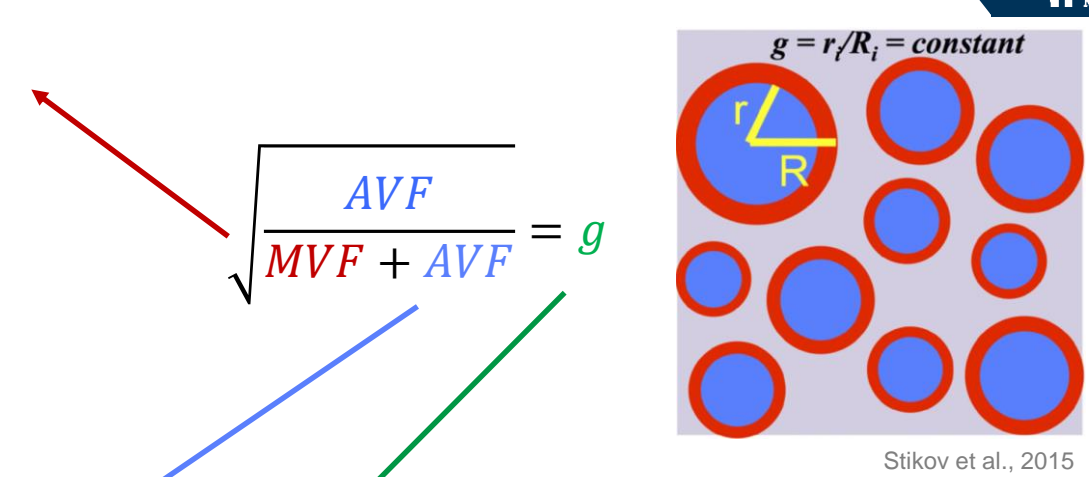
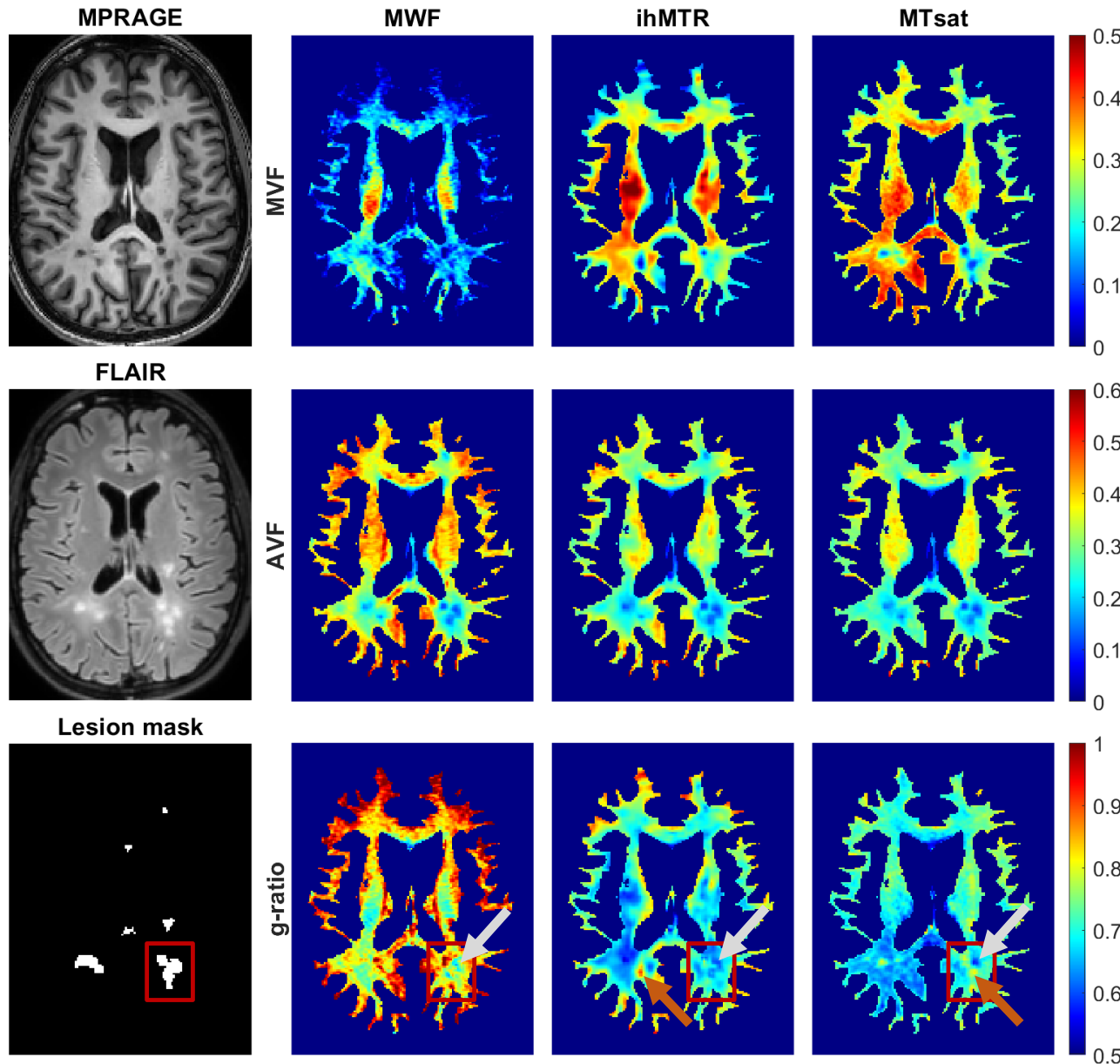


- MWF varies most strongly across WM VOIs
- MTsat most homogeneous
- Within MS lesions:
  - Clearly reduced values
  - MTsat values comparable to GM
  - For MWF and ihMTR, differences to MW less prominent
- Peri-lesion:
  - Largest difference to NAWM in MTsat



## Correlation between VOI-average myelin marker values in WM

- Highest between MWF and ihMTR
- Lowest between MWF and MTsat
  - → Rely on different contrast mechanisms



- MVF estimation from myelin-sensitive markers
- AVF calculation based on diffusion data
- g-ratio evaluation
- g-ratio values within WM strongly depend on the myelin-sensitive marker
- g-ratio values within lesion quite diverse
  - sometimes > WM, sometimes < WM

Stikov, Nikola, et al. "In vivo histology of the myelin g-ratio with magnetic resonance imaging." Neuroimage 118 (2015): 397-405. [www.doi.org/10.1016/j.neuroimage.2015.05.023](http://www.doi.org/10.1016/j.neuroimage.2015.05.023)



MWF

- Largest differences between various WM structures

ihMTR

- Good correlation with MWF

MTsat

- Largest difference between peri-lesion and NAWM

g-ratio

- g-ratio values quite diverse within lesions

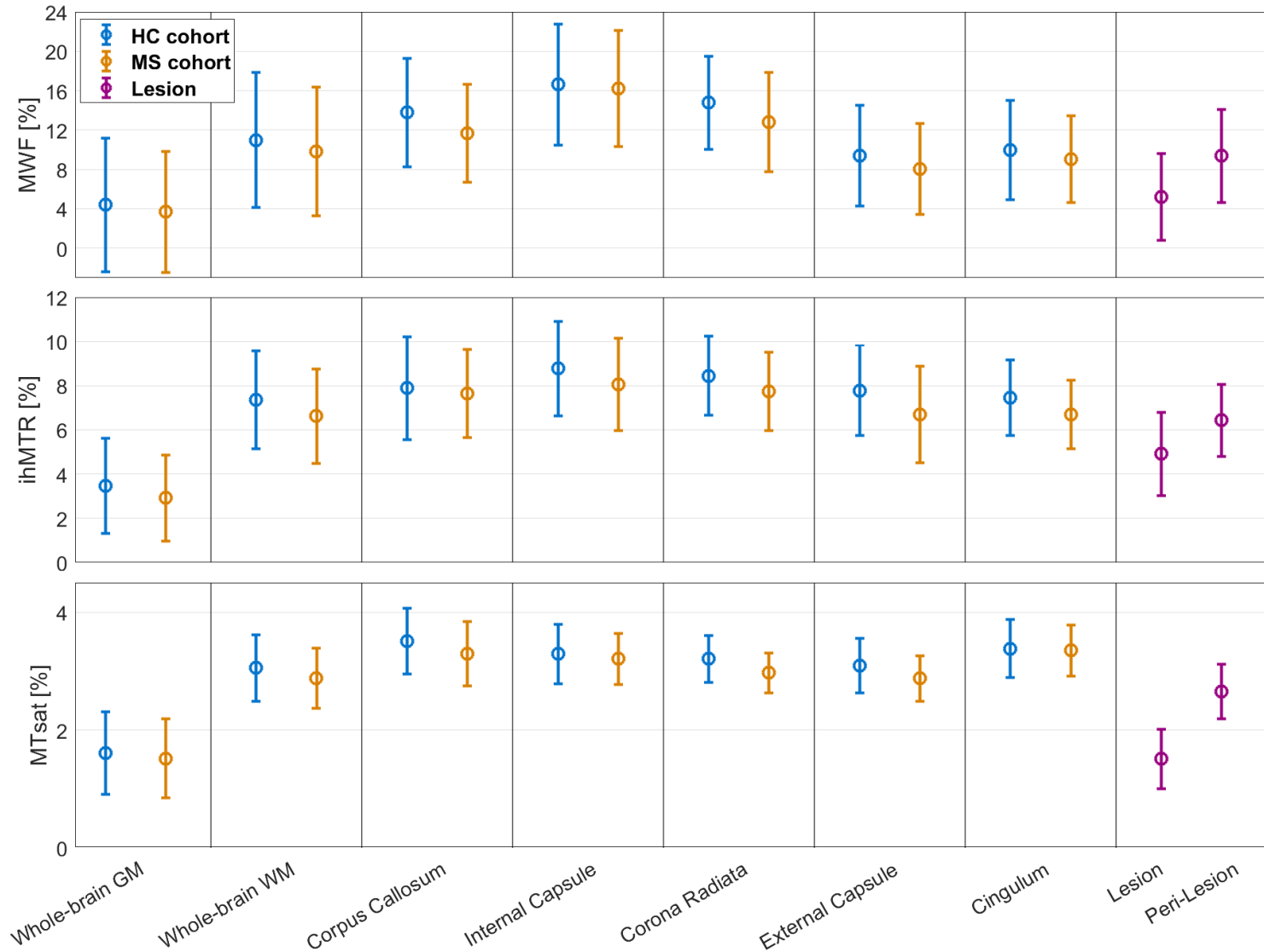
→

- Combined use of several myelin-sensitive markers
  - Disentangling microstructural effects
- Further studies needed



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Thank you for your attention!



Pooled standard deviation, averaged across participants:

- Highest for MWF
- Lowest for MTsat
- Often slightly lower for normal-appearing than for healthy tissue