A STUDY ON LOSS FUNCTIONS AND DECISION THRESHOLDS
FOR THE SEGMENTATION OF MULTIPLE SCLEROSIS LESIONS ON SPINAL CORD MRI

${ }^{1}$ Univ Rennes, Inria, CNRS, Inserm IRISA UMR 6074, Empenn ERL U 1228, Rennes, France
${ }^{2}$ Rennes University Hospital (CHU), Department of Neurology, Rennes, France
${ }^{3}$ Rennes University Hospital (CHU), Department of Neuroradiology, Rennes, France

## Context and objectives

Multiple sclerosis patients often present hyper-intense T2-w lesions in the spinal cord. The severe imbalance between background and lesion classes poses a major challenge for Deep Learning segmentation methods. We aim at investigating the following strategies to help mitigating this issue:

- careful selection of the loss function
- adjustment of the conventional 0.5-thresholding


## Dataset

- 161 T2-w cervical and thoracic MRI were acquired in 13 different sites from 108 subjects.
- Trained neurologists and radiologists performed manual segmentations, validated by a senior expert.
- Data splits were generated by assigning each subject to a single set while balancing the lesion load (Fig.1).


Fig. 1 Boxplots for lesion number and volume in the training ( 62 MRI scans), validation (13), thr-optimization (14) and unseen set ( 26,46 with no MS lesions are not included here). Points are jittered to improve visualization.

## Methods

- Images were reoriented and resampled to 0.5 mm isotropic using linear interpolation
- The Spinal Cord Toolbox ${ }^{1}$ (SCT) was used for detecting spinal cord centerline.
- 3D patches ( $48 \times 48 \times 48$ ) were extracted along the centerline for model training using a 3D-Unet architecture ${ }^{2,3}$.
- Data augmentation techniques such as patch overlapping, random mirroring, and $90^{\circ}$ rotation were employed during the 300-epoch training.
- Overlapping patches were used for prediction with different thresholding values, and a thr-optimization subset was used to optimize binarization thresholds.
- The model was evaluated on an unseen test set and compared with the SOTA method (SCT).

Table 1. Lesion- and voxel-wise F1 scores, and patient-level sensitivity and specificity (median (mean $\pm$ std)) for models trained with the binary cross-entropy (BCE), Dice (SCT), Focal Tversky and Tversky losses. Metrics obtained on the unseen test set.

| Loss | L-F1 | V-F1 (Dice) | P-sensitivity | P-specificity |
| :--- | :---: | :---: | :---: | :---: |
| BCE | $0.27(0.28 \pm 0.3)$ | $0.33(0.29 \pm 0.24)$ | 0.58 | 0.24 |
| SCT | $0.38(0.34 \pm 0.32)$ | $0.39(0.35 \pm 0.25)$ | 0.62 | $\mathbf{0 . 3 3}$ |
| F. Tversky | $0.26(0.28 \pm 0.28)$ | $0.31(0.31 \pm 0.25)$ | 0.65 | 0.22 |
| Tversky | $\mathbf{0 . 4 ( 0 . 3 4 \pm 0 . 3 1 )}$ | $\mathbf{0 . 4 1 ( 0 . 3 3 \pm 0 . 2 4 )}$ | $\mathbf{0 . 6 5}$ | 0.26 |

## Results

- Small changes in threshold lead to a larger change in voxel-F1 score for BCE and Dice (SCT) losses (Fig.2).
- Tversky-based losses seem more stable with threshold changes.
- Highest median L-F1 and V-F1 scores (40\%) for Tversky loss compared to SOTA SCT with Dice loss (38\%, Table 1).
- Example of segmentation output between Dice and Tversky loss on unseen test data with optimized threshold (Fig.3).


Fig. 2. Boxplots of voxel-wise F1 score for BCE, SCT, Focal Tversky and Tversky losses at each decision threshold value. One point corresponds to one image.


Fig. 3. Original image (a), overlaid by a reference lesion mask (b), and segmentations with the Tversky (c) and Dice (d) losses. FN, FP and TP voxels appear in white, red and green respectively.

## Conclusion

Tversky Index-based losses and a minimal adjustment of the decision threshold can yield less dispersed and higher median scores than more standard settings (Dice loss +0.5 thr). ${ }^{1}$

[^0]
[^0]:    De Leener et al.,
    ${ }^{2}$ C. Gros et al., "Automatic segmentation of the spinal cord and intramedullary multiple sclerosis lesions with convolutional neural networks," Neurolmage, 2019
    ${ }^{3}$ O. Çiçek et al., "3d u-net: Learning dense volumetric segmentation from sparse annotation," in Medical Image Computing and Computer-Assisted Intervention - MICCAI, 2016

    This work was supported by the French National Research Agency (Agence nationale de la recherche ANR) as its 3rd PIA, integrated to France 2030 plan under reference ANR-21-RHUS-0014. Data collection was supported by a grant provided by the French State and handled by the French National Research Agency (Agence nationale de la recherche - ANR) within the framework of the France 2030 program under the reference ANR-10-COHO-002 OFSEP.

