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Impact of training dataset size on technical performance of a deep learning model for detection and quantification of lymphomatous disease on ¹⁸F-FDG PET/CT

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Objective

FDG PET/CT is widely used for staging high-grade lymphoma. Artificial intelligence has the potential to improve efficiency and enable use of advanced quantification methods in a clinical setting. Here we investigate the impact of the amount of data used to train a deep learning (DL) model on detection and segmentation performance.

Materials & Methods



Pre-treatment FDG PET/CT scans of 420 patients with a total of 6150 lymphoma lesions segmented as ground truth by experienced PET- reporters were randomly split into training (300) and test sets (120).

A DL model, consisting of an ensemble of patch-based 3D DenseNet, was trained using various dataset sizes: N = 50, 100, 150, 200, 250 and 300, randomly sampled from a total of 300 cases.



Lesion detection performance was assessed using sensitivity and false positives (FPs) per patient, and true positives to false positives ratio (TPs/FPs) across the test set.

Segmentation and quantification performance were evaluated using sensitivity, positive predictive value (PPV), Dice score and non-parametric Bland Altman analysis for SUV_{max} and SUV_{mean} per lesion, and total metabolic volume (TMV) and total lesion glycolysis (TLG) per patient.

Results

Lesion detection sensitivity varied between 82% to 88%, whilst FPs per patient decreased with more training data (see **Table 1**). TPs/FPs improved as the training dataset size increased.

 Table 2 shows the segmentation performance for the six models: voxel-wise sensitivity, PPV and Dice score.

Bland Altman analysis showed improvement in Limits of Agreement (LoA) for lesion volume, TMV and TLG with more training data (See **Figure 2**).

Dataset size	50	100	150	200	250	300
Sensitivity (%)	82	83	88	83	83	86
FPs	9	4	4	4	3	3
TPs/FPs	0.73	1.42	1.40	1.43	1.67	1.69

Table 1: Lesion detection performance obtained for different dataset sizes. Median sensitivity was similar across models. FPs decreased with increasing dataset size. TPs/FPs ratio improved with more training data.

Dataset size	50	100	150	200	250	300
Sensitivity (%)	91	93	92	91	89	93
PPV (%)	75	82	83	86	88	88
Dice (%)	78	83	84	85	85	86

Table 2: Segmentation performance. Median sensitivity, PPV and Dice obtained for different dataset sizes.



Figure 1: Examples of predictions for N=50 and N=300. Maximum intensity projections (MIP) of PET images with overlaying contours, showing true positives (green), false positives (orange) and false negatives (red).



Figure 2: Statistics from Bland Altman analysis for SUVmean (A), Volume (B), TMV (C) and TLG (D). Median difference, lower and upper limits of agreement (LoA) calculated as 5^{th} and 95^{th} percentile are reported for the six models trained with different dataset sizes.

Conclusion

A deep learning model was relatively unaffected by the size of the training dataset in its ability to detect lymphoma lesions on PET/CT scans. However, more training data reduced FP rate, and improved agreement between prediction and ground truth segmentations for lesion volume $\rm SUV_{mean}, TMV$ and TLG.

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