

Improved detection of clinically relevant MRI findings in Multiple Sclerosis radiology reports using FDA-approved quantitative software

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DISCLOSURES



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Contribution to this study was as a paid consultant, and was not part of his Stanford University duties or responsibilities

BACKGROUND

Lesion metrics relate to disease progression in multiple sclerosis (MS)

• Quantitative metrics (e.g. lesion count and brain volume) provide objective data of disease progression.



- Semi-automated software can improve lesion detection, which can impact treatment-related decision-making.¹
- Implementation of FDA-cleared quantitative software in clinical practice requires significant effort and investment.
- Not yet established **if quantitative software can consistently improve detection of clinically relevant MRI findings** in MS-specific radiology reports.

¹ Van Heerden J et al. *AJNR*, 2015

OBJECTIVE

Can using FDA-cleared quantitative software improve lesion detection?

• To characterize clinically relevant findings in MS-specific radiology reports generated by a neuroradiologist after visual interpretation of images alone and with the use of FDA-cleared software.

Clinically relevant findings



Generate quantified MRI metrics and images

- Identify patients from anonymized retrospective MRI dataset between 2013-2017
- 2 MRIs/patient: 3D T1, 3D T2 FLAIR, T1-post
- 3D T1 and 3D T2 FLAIR images processed using FDA-cleared software (NeuroQuant & LesionQuant 3.0.1) to generate:
 - \circ lesion count and volume
 - brain volume
 - color-coded map that highlighted lesion changes between MRI timepoints
 - "Hot (red)": new or enlarging lesion
 - "Cold (blue)": shrinking lesion



METHOD

Compare qualitative vs quantitative radiology reports

- Board-certified neuroradiologist visually compared MRIs and reported the number of new, enlarging, and shrinking T2 lesions, gad-enhancing lesions, and brain atrophy assessment in a **qualitative report (qual)**.
- To avoid recall bias, studies were randomized and re-anonymized, and one month later, the neuroradiologist used the post-processed data and images to generate a second **quantitative report (quant)**.
- Recorded time of interpretation and report generation during both sessions.
- Two neuroimaging experts coded report differences in the following categories:
 - Presence or absence of MS finding
 - Change in count or location
 - Change in qualitative descriptor

Improved lesion detection on quantitative report

- 26 MS patients who had <u>></u>2 brain MRIs (mean MRI time interval: 1 year)
- In 13 patients we detected a difference between qual vs quant reports, where quant reports had on average 1 more new, enlarging and/or shrinking lesion (SD= 2.4).
- Brain atrophy descriptions changed with the addition of quantitative metrics in 7 cases, where 5 cases were upgraded and 2 cases were downgraded in severity.
- No significant difference (p=.16, paired t-test) was found in interpretation time (qual 12.3 minutes; quant 11.4 minutes).

Comparison of Lesion Counts

	Qual	Quant
New	44	50
Enlarging	10	25
Shrinking	5	8
Gad+ Enhancing	3	3

RESULTS

Examples: T2 Lesion Lesions missed on qualitative assessment



New

Baseline MRI Follow-up MRI Software Detection

Shrinking T2 Lesion

DISCUSSION

More clinical relevance in quantitative reports

- In addition to the metrics reported, all quant reports provided information not available on qual reports, such as T1 white matter hypointensity volume, and regional brain volumes over time
- Use of FDA-approved quantitative software improved detection of clinically relevant neuroimaging findings in MS patients, without adding time to image interpretation, suggesting its potential value in clinical practice

NEXT STEPS

Evaluate impact on clinical management

- Compare MS-specific radiology reports and lesion detection rates between general radiologists, neuroradiologists using visual assessment only, and neuroradiologists using combined visual assessment and FDA-cleared software
- Assess MS Neurologists' satisfaction of reports and evaluate potential changes to treatment plan and disease management

Questions?

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