Multicenter Accuracy and Interobserver Agreement of Spot Sign Identification in Acute Intracerebral Hemorrhage

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Background and Purpose—Rapid, accurate, and reliable identification of the computed tomography angiography spot sign is required to identify patients with intracerebral hemorrhage for trials of acute hemostatic therapy. We sought to assess the accuracy and interobserver agreement for spot sign identification.

Methods—A total of 131 neurology, emergency medicine, and neuroradiology staff and fellows underwent imaging certification for spot sign identification before enrolling patients in 3 trials targeting spot-positive intracerebral hemorrhage for hemostatic intervention (STOP-IT, SPOTLIGHT, STOP-AUST). Ten intracerebral hemorrhage cases (spot-positive/negative ratio, 1:1) were presented for evaluation of spot sign presence, number, and mimics. True spot positivity was determined by consensus of 2 experienced neuroradiologists. Diagnostic performance, agreement, and differences by training level were analyzed.

Results—Mean accuracy, sensitivity, and specificity for spot sign identification were 87%, 78%, and 96%, respectively. Interobserver agreement for spot sign presence was moderate (κ=0.60). When true spots were correctly identified, 81% correctly identified the presence of single or multiple spots. Median time needed to evaluate the presence of a spot sign was 1.9 minutes (interquartile range, 1.2–3.1 minutes). Diagnostic performance, interobserver agreement, and time needed for spot sign evaluation were similar among staff physicians and fellows.

Conclusions—Accuracy for spot sign identification is high with opportunity for improvement in spot interpretation sensitivity and interobserver agreement particularly through greater reliance on computed tomography angiography source data and awareness of limitations of multiplanar images. Further prospective study is needed. (Stroke. 2014;45:107-112.)

Key Words: angiography • cerebral hemorrhage • diagnosis • multidetector computed tomography • stroke

Intrahematoma contrast density identified on first-pass computed tomography angiography (CTA), coined the spot sign, is associated with increased risk of hematoma expansion, poor functional outcome, and mortality in patients with primary intracerebral hemorrhage (ICH).1–4 The spot sign is currently being studied within 3 multicenter trials to select patients for acute hemostatic therapy. Recombinant factor VIIa is used in the Spot Sign for Predicting and Treating ICH Growth (STOP-IT)5 and Spot Sign Selection of Intracerebral Hemorrhage to Guide Hemosatic Therapy (SPOTLIGHT)6 studies and tranexamic acid in the Spot Sign and Tranexamic Acid on Preventing ICH Growth–Australasia Trial (STOP-AUST).7 Although the diagnostic performance of the spot sign is previously validated, interobserver agreement and accuracy of spot sign identification and factors associated with their improvement have not been examined in a broad cohort of physicians. To be an effective marker for guiding intervention and patient recruitment into ongoing trials, high spot sign identification accuracy and agreement among acute stroke staff is needed. We sought to evaluate the accuracy and agreement for spot sign identification among readers of various specialties and training levels through use of an online imaging certification module.

Methods

Study Design and Patient Cases

After obtaining local institutional ethics approval, an online imaging module was developed to certify clinical investigators enrolling patients based on spot sign presence in 3 clinical trials: STOP-IT (http://www.stopitstudy.com), SPOTLIGHT (http://www.spotlightstudy.org).
Statistical Analysis

Accuracy, sensitivity, specificity, positive predictive value (PPV), and negative predictive value (NPV) for true spot detection were determined for all physicians and examined by training level. The consensus agreement of 2 experienced neuroradiologists (R.I.A., S.P.S.) was used as reference standard. Differences in diagnostic performance by training level were assessed using the Wilcoxon rank-sum nonparametric test. Interobserver agreement among specialties and training level was determined using the Fleiss multirater k-statistic. Values of k of 0.21 to 0.4, 0.41 to 0.6, 0.61 to 0.8, and 0.81 to 1 were considered fair, moderate, substantial, and nearly perfect, respectively. 10 Characteristics of spots with poor interobserver accuracy (≤80%) and pair agreement (≤0.80 proportion of pairs agreeing) were compared with spots with high accuracy and agreement. Statistical significance was defined as P<0.05 for all tests. All statistical analysis was performed in SAS version 9.2 (SAS Institute, Cary, NC) and R version 3.12.3.

Results

One hundred thirty-one physicians underwent online spot sign imaging training and certification. Physicians were from 28 institutions across the United States (n=13), Canada (n=13), and Australia (n=2). One hundred eighteen (88%) physicians were neurologists (49 staff, 69 fellows), 6 (4.5%) were emergency physicians (3 staff, 3 fellows), and 10 (7.5%)were neuroradiologists (8 staff, 2 fellows). Mean time spent on the educational portion of the imaging module before certification was 17.4±14.2 minutes. Median ICH volume for all certification cases was 32 mL (range, 11–75 mL), and 70% had presence of intraventricular hemorrhage. For the 5 spot-positive cases, median (range) maximum axial spot size was 4 (3.1–10.7) mm, and spot attenuation was 203 (201–496) Hounsfield units. Three of the cases had a single spot, 1 had 3 spots, and 1 had 12 spots.

Physician Diagnostic Performance

Overall mean (95% confidence interval [CI]) for accuracy, sensitivity, and specificity for spot sign identification were 87% (85%–89%), 78% (75%–82%), and 96% (94%–98%), respectively. Overall sensitivity was lower than specificity (P<0.001), and PPV was higher than NPV (96%; [95% CI, 94%–98%] versus 84% [95% CI, 82%–86%]; P<0.001). When true spots were correctly identified, 414 of 514 (81%) responses correctly identified the presence of single or multiple spots. Diagnostic performance and agreement for spot detection are listed in Table 1. Mean (95% CI) PPV and NPV for neurologists was 96% (94%–97%) and 83% (81%–86%). PPV and NPV for limited sample of emergency physicians and neuroradiologists were 97% (92%–100%) and 85% (75%–95%) and 100% (100%–100%) and 93% (86%–99%), respectively. Accuracy, specificity, and PPV of staff compared with fellows demonstrated no significant difference (P=0.245, 0.983, and 0.916, respectively). There were weak trends toward improved sensitivity (81% [95% CI, 75%–86%] versus 77% [95% CI, 73%–81%]; P=0.123) and NPV (86% [95% CI, 82%–90%] versus 82% [95% CI, 79%–85%]; P=0.114) among staff compared with fellows.

Interobserver Agreement

Agreement for all readers and neurologists was moderate: k=0.60 (95% CI, 0.59–0.61) and k=0.59 (95% CI, 0.57–0.59), respectively. Agreement among staff was similar to fellows.

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Limited assessment of neuroradiologists and emergency physicians achieved values of $k=0.88$ and 0.68, respectively.

**Diagnostic Performance, Agreement, and Time to Spot Identification by Case**

Accuracy, pair agreement, and time to spot identification by individual examination case are listed in Table 2. Seven cases (2 spot-positive and 5 spot-negative) demonstrated high accuracy ($\geq80\%$) and pair agreement (Figures 1 and 2), whereas 3 cases (3 spot-positive) were identified as demonstrating low accuracy and agreement (Figures 3 and 4). For the 3 cases with low accuracy, incorrect responses were examined to determine whether readers may have answered the question incorrectly because of the perceived presence of a spot mimic. Of the combined 117 incorrect responses for the 3 questions with low accuracy, 111 (95%) responses also indicated the presence of a mimic. Similarly, for the 2 true spot-positive cases with high accuracy, 22 of 24 (92%) incorrect responses also indicated the presence of a mimic.

The cases demonstrating the microarteriovenous malformation and aneurysm were correctly identified as spot mimics in 69 of 131 (53%) and 48 of 131 (41%) responses, respectively.

Overall median (interquartile range) time to spot identification per case was 1.9 (1.2–3.1) minutes and was similar between staff and fellows ($P=0.673$).

**Discussion**

Previous studies of spot sign interobserver agreement have demonstrated substantial to near perfect agreement ranging from $\kappa$-statistic of 0.61 to 0.94; however, little is known about the strength of agreement among a broad range of readers, between physician specialties and training levels, or accuracy compared with a gold standard consensus. Our results demonstrate that there is an overall high accuracy (87%) for spot identification within a certification context among a large number of readers involved in acute stroke care. We have also identified high specificity (96%) and PPV (96%) for physician spot sign interpretation, and that physicians were able to perform spot interpretation rapidly $\approx$ 2 to 3 minutes. High specificity and PPV would demonstrate that physicians were highly accurate at correctly identifying the absence of a spot when not truly present and that readers had high accuracy when a spot sign was perceived. These findings are clinically important because high specificity is needed to ensure that patients without spots, and thus at lower risk of hematoma expansion, are not treated with aggressive hemostatic therapy.

<table>
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<th>Table 2. Accuracy, Agreement, and Time to Spot Identification by Case</th>
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<td><strong>Response Frequency</strong></td>
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High PPV ensures that all patients identified as having spots by physicians truly have spots and may benefit from hemo-
static therapy.

An important finding of this study, however, was the reduced sensitivity of readers compared with specificity. Reduced spot identification and, therefore, sensitivity may result in failure to treat a true spot-positive patient who potentially would benefit from therapy. Low accuracy and agreement among spot-positive cases, leading to lower sensitivity, also resulted in a lower overall \( \kappa \)-statistic \( (k=0.60) \) than previously reported. Analysis of the potential causes for reduced sensitivity demonstrated that a majority of incorrect spot identification responses in the presence of true spots were associated with the incorrect perception of spot sign mimics. In-depth examination of cases with poor sensitivity revealed that spot signs found in these cases were closely associated with adjacent lenticulostriate arteries or cortical vessels, often with only 1 to 2 mm separation between spot and vessel on axial images, as demonstrated in Figures 3 and 4, respectively. Spot-positive cases with high accuracy did not have similar adjacent vessels. A key defining feature of the spot sign definition is the absence of visible connecting vessels from outside the hematoma.\(^{1,3,9,14}\) The definition is particularly challenging for anterior basal ganglia and peripheral lobar bleeds where vessels may be in close proximity to a hematoma. Careful evaluation of the association between the focal contrast extravasation and the vessel should be made on thin-section axial CTA source images of \( \geq 0.625 \) mm to determine whether such a visible connection is present. The absence signifies a spot sign. Although review of coronal and sagittal images is helpful and often routinely provided with CTA source images, a potential pitfall of maximum intensity projection images reconstructed at 7 mm, as performed at our institution, is the inability to clearly separate spots from adjacent vessel. This emphasizes the importance of axial CTA source images for spot sign identification and mimic exclusion. The high rate of perceived mimics in the presence of true spots also demonstrates that readers were easily able to detect contrast density within hematomas but that the difficulty experienced related to deciding between a spot sign versus spot mimic.

Further improvement of spot sign interobserver agreement may be possible with multimodal CT imaging, including post-
contrast CT and CT perfusion\(^{14-17}\) visualization of extravasa-
tion improving identification. Both imaging protocols also provide a critical opportunity for improving prediction of expansion and clinical outcome and thus may represent the ideal method of spot imaging.\(^{15-18}\) Additional prospective evaluation of delayed or dynamic spot sign imaging in comparison to CTA alone is required.

Examination of accuracy and agreement between staff and fellows yielded no significant differences; however, we did observe minor trends toward improved sensitivity and NPV in staff compared with fellows. Limited data were available for neuroradiologists and emergency physicians, because the small sample size of these groups precluded meaningful comparison with the large cohort of neurologists. Online training is available and is shown to improve spot identification and diagnostic certainty and should be encouraged as a continuing quality improvement process.\(^{19,20}\)

A limitation of our study is the limited number of cases in the certification examination. Although inclusion of a greater number of cases would have been ideal, this was balanced with the time required to obtain imaging certification for a large number of investigators. Small sample size results in potentially wide variation in accuracy and agreement depending on the selected cases; however, cases were chosen by
experienced neuroradiologists and thought to be representative of acute ICH cases. The number of cases was balanced by the large number of readers of varied institutions, making our results robust for the cases included. Spot sign prevalence was set to 50%, which is at the higher range of previously reported spot prevalence in acute ICH.\textsuperscript{11,21} This was purposefully chosen to maximize reader experience with spot sign appearance. Although spot sign prevalence approaches 50% when using dynamic or delayed CT techniques,\textsuperscript{14,15} prevalence of 22% to 41% using static CTA in patients presenting <6 hours of onset in practice may alter physician diagnostic accuracy.\textsuperscript{3,21} A relatively small number of emergency medicine physicians and neuroradiologists participated in the certification process, precluding meaningful comparison with neurologists. Larger numbers of each specialty would need to be recruited to meaningfully compare performance. Because physician accuracy and agreement were assessed immediately after training, our results may represent an overestimate of true physician accuracy. Assessment with either a pretest or post-test after 3 to 6 months may have less impressive results.

To be a clinically useful predictor in the acute setting, understanding factors that affect rapid, accurate, and reliable spot sign interpretation is critical. We have demonstrated an overall high accuracy for spot identification with modest variation between specialty and training level. We have also identified opportunities for improving consensus in spot interpretation, particularly with emphasis on improving sensitivity through careful review of axial CTA source images and avoiding over-reliance on multiplanar images that may falsely suggest a spot mimic due to technical considerations. Further study in a prospective clinical cohort, ideally with inclusion of delayed or dynamic imaging, is needed. The STOP-IT study, which enrolls both spot-positive and spot-negative subjects, will provide an ideal opportunity to examine real-time accuracy and agreement of spot sign identification.

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Disclosures
Dr Flaherty serves on an advisory board and as a consultant to CSL Behring and is principal investigator of the STOP-IT study. Dr Gladstone is the principal investigator and Drs Aviv, Demchuk, and Flaherty are members of the executive committee of the SPOTLIGHT trial. Dr Davis is a principal investigator for the STOP-AUST study and Drs Meretoja and Mitchell serve in the executive steering committee. Dr Broderick is principal investigator of the SPOTRIAS Center Grant, which receives study medication for the STOP-IT study from Novo Nordisk. The other authors have no conflicts to report.
References


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