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**Clopidogrel Hyper-Response and Bleeding Risk in Neurointerventional Procedures**

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**ABSTRACT**

**BACKGROUND AND PURPOSE:** Antiplatelet therapy is associated with decreased ischemic events after neurointerventional procedures. Antiplatelet resistance negates the protective effects of antiplatelet medication, leading to a higher incidence of ischemic events. A possible link between antiplatelet hyper-response and increased hemorrhagic complications has been inadequately investigated. We aimed to examine the correlation between antiplatelet hyper-response and the risk of hemorrhagic complications.

**MATERIALS AND METHODS:** Patients who were treated with antiplatelet medications and underwent neurointerventional procedures were prospectively recruited. We collected the following data: demographics, vascular risk factors, antiplatelet and anticoagulation treatment, antiplatelet responsiveness, coagulation profile, and hemorrhagic complications. P2Y12 receptor–mediated platelet inhibition was tested by using the VerifyNow assay device. The primary end points were postprocedural major and minor hemorrhagic complications. Receiver operator characteristic analysis was used to evaluate the percentage of platelet inhibition as a diagnostic tool for bleeding events. The association between hemorrhage and percentage of platelet inhibition was investigated by using logistic regression modeling.

**RESULTS:** Forty-seven patients were enrolled. The mean age was 56 ± 12 years, and 28% were men. Ten patients (21.3%) developed hemorrhagic complications. Clopidogrel response was higher in patients with a major bleeding complication compared with those with minor or no bleeding (median, 94% versus 24% platelet inhibition; \( P = .0084 \)). Of the 7 patients (14.9%) defined as hyper-responders with ≥72% platelet inhibition, 42.8% had a major bleeding complication.

**CONCLUSIONS:** Hyper-response to clopidogrel is associated with increased risk of hemorrhagic complications. Larger studies are urgently needed to validate a clinically useful threshold to define clopidogrel hyper-response and to examine the clinical effects of antiplatelet dosage adjustment.

**ABBREVIATIONS:** ACT = activated clotting time; aPTT = activated partial thromboplastin time; CI = confidence interval; IQR = interquartile range; ROC = receiver operating characteristic
The study protocol received ethics committee approval.

Patients were excluded if they refused consent, were younger than 18 years of age, or had a history of an acute event (transient ischemic attack, ischemic infarct, or acute intracranial hemorrhage) within the 24 hours before the procedure. Demographic and clinical data collected included age, sex, and cardiovascular risk factors (hypertension, dyslipidemia, diabetes mellitus, smoking history, and history of previous cerebrovascular ischemic events).

Forty-seven patients were enrolled in the study, 34 women and 13 men, with a mean age of 56 ± 12 years (range, 30–78 years). There was a history of hypertension in 46.8%, dyslipidemia in 25.5%, and diabetes mellitus in 4.3% of patients, and 29.8% were current smokers. Previous TIA or ischemic stroke was documented in 12% of patients, with a history of hypertension in 46.8%, dyslipidemia in 25.5%, and diabetes mellitus in 4.3% of patients, and 29.8% were current smokers.

The VerifyNow assay is a point-of-care device using a 2-channel disposable cartridge, with platelet aggregation measured by increasing light absorbance through the sample. The first chamber gives a measure expressed as P2Y12 reaction units (PRU), representing platelet activation by the adenosine diphosphate-P2Y12 pathway. Percentage of inhibition is calculated from the formula \( \frac{1 - (\text{PRU/BASE})}{100} \). The VerifyNow assay has been shown to correlate well with the current criterion standard light transmittance aggregometry and with other point-of-care assay devices, and it has been validated in a head-to-head comparison of assay methods based on outcome measures.

Outcomes were measured by clinical assessment during the hospital admission and at the 6-week review appointment. The primary end point was Thrombolysis in Myocardial Infarction major or minor periprocedural bleeding events. The need for investigation with Doppler sonography or CT or MR imaging of the brain was guided by clinical assessment.

Statistical analysis was performed by using the commercial statistical software STATA, Version 11IC (StataCorp, College Station, Texas). Platelet inhibition is reported as median IQR due to non-normal distribution as assessed by the Shapiro-Wilk test. The Wilcoxon-Mann-Whitney test was used to compare the percentage of platelet inhibition by clopidogrel between bleeding and nonbleeding groups with the corresponding effect size estimated by using median regression. The receiver operator characteristic analysis curve was used to evaluate the percentage of platelet inhibition measured by the VerifyNow assay device as a diagnostic tool for bleeding events. Association between the presence of periprocedural bleeding and the percentage of platelet inhibition was investigated by using logistic regression modeling (both unadjusted and adjusted for mean aPTT and intra-procedural ACT). The statistical significance threshold for all the tests performed was set as \( P = .05 \).

### RESULTS

#### Study Population

Forty-seven patients were enrolled in the study, 34 women and 13 men, with a mean age of 56 ± 12 years (range, 30–78 years). There was a history of hypertension in 46.8%, dyslipidemia in 25.5%, and diabetes mellitus in 4.3% of patients, and 29.8% were current smokers. Previous TIA or ischemic stroke was documented in only 0.9%.

#### Procedure

All endovascular procedures were performed by 1 of 3 experienced neurointerventionalists. Arterial access was via a 6F femoral sheath, with hemostasis obtained postprocedure by use of an Anglo-Seal device (St. Jude Medical, Minnetonka, Minnesota). Systemic anticoagulation with intravenous heparin during the procedure was titrated to an ACT twice normal. The postprocedural unfractionated heparin infusion rate was adjusted on the basis of aPTT.

Whole blood was obtained for testing at the time of the initial femoral puncture; following discard of the first 5 mL, citrated Vacette tubes (VWR, Arlington Heights, Illinois) were filled and inverted 5 times with blood analyzed by using the VerifyNow assay device (Accumetrics, San Diego, California).

The VerifyNow assay is a point-of-care device using a 2-channel disposable cartridge, with platelet aggregation measured by increasing light absorbance through the sample. The first chamber contains a strong agonist for platelet activation independent of aspirin or clopidogrel therapy, providing a baseline platelet function measurement (BASE). The second chamber gives a measure expressed as P2Y12 reaction units (PRU), representing platelet activation by the adenosine diphosphate-P2Y12 pathway. Percentage of inhibition is calculated from the formula \( \frac{1 - (\text{PRU/BASE})}{100} \). The VerifyNow assay has been shown to correlate well with the current criterion standard light transmittance aggregometry and with other point-of-care assay devices, and it has been validated in a head-to-head comparison of assay methods based on outcome measures.

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intravenous heparin following the procedure: 21 patients for up to 24 hours and 17 patients for 48 hours. One patient who had a Factor V Leiden deficiency received 10 hours of intravenous unfractionated heparin postprocedure and then low-molecular-weight heparin until therapeutic anticoagulation with warfarin had been achieved.

Treatment of 3 patients deviated from the standard antiplatelet regimen. Two received a single loading dose of 450 mg of clopidogrel at 3 hours or at 5 hours before the procedure. The other had a history of Factor V Leiden deficiency and had been receiving subcutaneous low-molecular-weight heparin due to cessation of his usual long-term anticoagulation with warfarin. The decision was made to give him 3 days of clopidogrel loading at 75 mg daily, rather than dual antiplatelet therapy.

**Clopidogrel Response and Bleeding Events**

Response to the antiplatelet effects of clopidogrel varied widely, with the percentage of platelet inhibition ranging between 0% and 99% (median, 30%; IQR, 12%–61%). Figures 2 and 3 illustrate the distribution of platelet inhibition and bleeding events among the study population.

Eleven patients had a bleeding event during the procedure or in the 24 hours postprocedure. Thrombolysis in Myocardial Infarction major bleeding events occurred in 3 patients (6.4%). Two of these had documented periprocedural intracranial hemorrhages. A 5-mm left temporal intracerebral bleed occurred in a patient with 99% platelet inhibition and PRU 5, whose right posterior communicating artery aneurysm was treated by placement of a flow-diverting stent. MR imaging of the brain showed no evidence of an underlying cavernoma, mass, or infarct. A tiny subarachnoid bleed within a sulcus at the right vertex occurred in a patient with 94% platelet inhibition and PRU 21, who had a right supracerebral carotid aneurysm treated by placement of a flow-diverting stent. The other patient had a large retroperitoneal hemorrhage in 1 patient was related to on-table aneurysm perforation, so it was not included in the bleeding group for statistical analysis. Minor bleeding events in 7 (15%) patients were small groin hematomas of little clinical significance.

Clopidogrel response was greater in the 10 patients with a bleed of any size (median platelet inhibition, 63.5%; IQR, 36%–75%) compared with those with no bleeding complication (median platelet inhibition, 19%; IQR, 11%–47%), resulting in a median difference between the degree of platelet inhibition in bleeding and nonbleeding groups of 38% (95% CI, 10.6%–65.4%; P = .015).

Comparison between clopidogrel response in the 3 patients with a major bleed and those with minor or no bleeding demonstrated a more pronounced difference, with median platelet inhibition of 94% (IQR, 72%–99%) versus 24% (IQR, 11.5%–55.5%), resulting in a median difference between percentage platelet inhibition in these 2 groups of 69% (95% CI, 23%–100%; P = .008).

ROC analysis was undertaken to identify an optimal threshold to define hyper-response to clopidogrel. Using 72% or greater platelet inhibition as the criterion, 7 patients (14.9%) were defined as clopidogrel hyper-responders; 42.8% of these patients had a major bleeding complication, while all non-hyper-responders were major bleeding complication-free. If we used a threshold of ≥72% inhibition to define hyper-response, the percentage of platelet inhibition by clopidogrel as a diagnostic tool for clinically significant bleeding events demonstrated high diagnostic capacity (ROC area = 0.96; 95% CI, 0.89–1), resulting in sensitivity of 100% and specificity of 90.9%. Due to the small sample size and low number of events, caution is required in the extrapolation of these results into clinical practice; larger studies would be required to define and validate a robust threshold for clinical use.

Including minor with major bleeding in the ROC analysis results in a lower diagnostic capacity (ROC area = 0.75; 95% CI, 0.58–0.93), with a threshold of ≥53% inhibition corresponding to a sensitivity of 70% and a specificity of 75.7%, and is of limited clinical utility. The incidents of minor bleeding in our study were all small groin hematomas, a complication that may be influenced by puncture technique and by achieving inadequate hemostasis at the end of the procedure. Also, because the cutoff below which patients are deemed to be low responders has variably been nominated as anywhere between 23% and 40% platelet inhibition, use of this lower threshold would necessarily result in a narrow therapeutic window.

**Systemic Anticoagulation with Intravenous Heparin and Bleeding Events**

When divided by bleeding outcome, mean ACT was similar among all groups with 392.2 (95% CI, 354.32–430.15) in those with no bleeding complication, 373.3 (95% CI, 315.67–431) in those with any bleed, and 375 (95% CI, 207.9–542.1) in those with clinically significant bleed.
In the patients whose intravenous heparin was continued in the ward, the mean aPTT was also similar between each of the bleeding outcomes: the postprocedure aPTT was 63.3 (95% CI, 51.09–75.57) in those with no bleeding complications, 60.4 (95% CI, 49.38–71.38) in those with any bleed, and 60.6 (95% CI, 54.66–66.47) in those with a clinically significant bleed.

FIG 3. Percentage of platelet inhibition for patients with major or minor hemorrhagic complications, compared with those with no hemorrhagic complications.

While the relatively high intraoperative ACT and the use of heparin anticoagulation in the postprocedural period may well potentiate the effects of antiplatelet medication on bleeding risk, they do not account for the preponderance of bleeding events among the patients with a high level of response to the antiplatelet effects of clopidogrel.

The possibility that the type of intervention performed may impact bleeding risk is another consideration. The 2 small intracranial hemorrhages in this study group occurred in patients who had undergone placement of a flow-diverting stent. However, it is difficult to implicate traction on arteries or altered flow dynamics related to stent placement in the etiology of these events, given that the bleeding site was distant from the treated vessel in both cases. The early experience of this method of aneurysm treatment emphasizes the potential thrombotic complications of parent vessel occlusion, small artery occlusion, and in-stent thrombosis. A multicenter study of 70 patients treated with the Silk flow-diverting stent (Balt Extrusion, Montmorency, France) reported 3 extracranial hemorrhages but no intracranial hemorrhage.22 A multicenter study of 31 patients treated with the Pipeline flow-diverting device (Covidien, Irvine, California) reported no hemorrhagic complications.23

A more likely mechanism by which flow-diverting stents could affect perioperative bleeding risk is in their requirement for postprocedural anticoagulation with intravenous heparin, which accounts for the high number of such patients in our study. In addition, these patients continue to receive at least 6 months of antiplatelet medication, usually with clopidogrel. While the administration of heparin did not significantly affect the results of this study, the need for ongoing clopidogrel administration highlights the potential for clopidogrel hyper-response to increase the cumulative risk of adverse bleeding events.

Differences between patients undergoing neurointerventional procedures and the cardiac patients who have been studied in greater numbers may influence the clinical impact of interindividu-
vidual variability in the clopidogrel response. In concentrating on elective patients and excluding those with a recent ischemic or hemorrhagic event, our study population is strongly weighted toward patients undergoing treatment for a cerebral aneurysm. These patients tend to be younger with fewer cardiovascular risk factors than those undergoing cerebrovascular or coronary revascularization, so the background risk of thromboembolic events may be reduced. Whether this is offset, or perhaps overshadowed, in the periprocedural period by the need for catheterization of intracranial arteries is uncertain, but regardless, it may have implications for clopidogrel hyper-responders during the 6–12 months postprocedure when patients with intracranial stents will receive maintenance dual antiplatelet therapy. It is possible that a given antiplatelet dose in a neurointervention patient will be associated with an increased bleeding risk compared with patients undergoing cardiac percutaneous coronary intervention or cerebrovascular stent placement.

Authors of future research into the identification and management of clopidogrel hyper-response should bear in mind the lessons from previous research into hypo-response. Despite the large body of evidence available linking low clopidogrel response to an increased risk of adverse cardiovascular outcomes, differences in assay devices and the choice of statistical methods to define “hyporesponse” mean that researchers have struggled to achieve consensus on the optimal cutoff defining low response.

A different approach to understanding the impact of variability in the clopidogrel response relies on identifying genetic polymorphisms in the CYP2C19 allele that are associated with high or low response. The CYP2C19*2 variant is the most common of several loss-of-function alleles. Conversely, the CYP2C19*17 variant allele causes increased metabolism and function of clopidogrel and has been shown to reduce the incidence of major adverse cardiovascular events in patients with acute coronary syndrome.

A target of future research would be a prospective study comparing genetic testing with measures of platelet aggregation to determine which is best able to predict the risk of an adverse bleeding outcome. The high specificity of genetic testing may prove to be a drawback, given the multifactorial etiology of clopidogrel response variability, and the time required to perform genetic testing may reduce its clinical utility in comparison with point-of-care assay devices. In addition, genetic testing will need to include the full panel of CYP2C19 alleles because there is evidence that the combination of high- and low-function variants will result in a low-to-normal function phenotype.

CONCLUSIONS

Hyper-response to the antiplatelet effects of clopidogrel is associated with a significant increase in bleeding risk in the setting of endovascular neurointervention. Any proposed threshold value for a clinically significant increased risk of major bleeding will require validation with a larger study population. Testing for genetic polymorphisms associated with increased platelet inhibition by clopidogrel is a promising avenue of further research. However, point-of-care assay devices retain the advantage of speed and simplicity of use and will include the effects of nongenetic determinants of platelet function. Standardization of study methodology in future research protocols is of critical importance to provide statistically robust and clinically useful thresholds defining high and low response to clopidogrel.

REFERENCES

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