Role of Ticagrelor in Clopidogrel Nonresponders: Resistance Is Futile?

Luke Tapp, MRCP; Eduard Shantsila, MD; Gregory YH Lip, MD

Dual antiplatelet therapy with aspirin and a thienopyridine drug that blocks the platelet adenosine diphosphate receptor P2Y12 is the cornerstone of management in patients presenting with acute coronary syndrome (ACS). Clopidogrel is the most widely used thienopyridine, with evidence of benefit in non–ST-elevation myocardial infarction (MI),1 percutaneous coronary intervention (PCI),2 and ST-elevation MI treated with thrombolysis or primary PCI.3,4 Improvement in Therapeutic Outcomes by Optimizing Platelet Inhibition With Prasugrel-Thrombolysis In Myocardial Infarction 38 (TRITON-TIMI 38) patients allocated to prasugrel. CYP polymorphisms did not attenuate the pharmacological response to prasugrel and were not associated with adverse cardiovascular outcomes. These important observations may contribute to the differences in clinical response to these drugs.

The TRITON-TIMI 38 trial6 demonstrated a relative risk reduction of 19% in the primary efficacy endpoint (death from cardiovascular causes, nonfatal MI, and nonfatal stroke) with prasugrel compared with clopidogrel in ACS patients scheduled for PCI. This benefit was amplified in high-risk groups, such as patients with diabetes mellitus or ST-elevation MI, and it was apparent as early as day 3. Additionally, the frequency of both early and late stent thrombosis was reduced by approximately 50%.8 These important findings validate the hypothesis that a more potent antiplatelet regimen can reduce ischemic events. However, the sacrifice is an increased risk of bleeding. Prasugrel was associated with a statistically significant increase in non-coronary artery bypass grafting-related TIMI major bleeding (the key safety endpoint), coronary artery bypass grafting-related TIMI major bleeding, and a small increase in fatal bleeding. More important, a prespecified analysis of net clinical benefit showed statistical significance in favor of prasugrel.8 Avoiding prasugrel (or reducing its dose) in groups with a higher risk of bleeding and less evidence of clinical efficacy (age ≥75 years, weight <60 kg, and previous stroke or transient ischemic attack) may attenuate the increased bleeding risk.

What are the alternatives? Ticagrelor is an oral, direct, reversible, nonthienopyridine P2Y12 receptor antagonist that has a rapid, powerful, and consistent antiplatelet effect. Because ticagrelor is not a prodrug, a loading dose of 180 mg achieves effective inhibition of platelet aggregation within 30 minutes. Platelet function returns to normal 2 to 3 days after discontinuation.8 The Study of Platelet Inhibition and Patient Outcomes (PLATO) trial compared ticagrelor to clopidogrel in ACS patients managed invasively or noninvasively.10 Ticagrelor was associated with a 16% relative risk reduction in the primary endpoint (composite of death from vascular causes, MI, and stroke) and a 22% relative risk reduction in all-cause mortality. There was no significant difference in the primary safety endpoint (overall rate of major bleeding) between the ticagrelor and clopidogrel groups (11.6% and 11.2%, respectively; P = 0.43), but ticagrelor was associated with a higher rate of major bleeding not related to coronary artery bypass grafting (4.5% versus 3.8%; P = 0.03), including more instances of fatal intracranial bleeding and less fatal bleeding of other types.10 Additionally, ticagrelor reduced...
Resistance to antiplatelet drugs

The definition of aspirin resistance is controversial. Lab response to antiplatelet agents appears to be normally distributed, and it therefore depends on where the cutoff of sufficient platelet inhibition lies. Moreover, there may not be a linear relationship between enhanced antiplatelet effect and a relevant reduction in ischemic events. Clinically, so-called resistance broadly encompasses individuals with recurrent atherothrombotic events despite appropriate medication and lab-assessed failure to sufficiently inhibit platelet aggregation. This incorporates both intrinsic (eg, P2Y12 receptor density and affinity, internal cell signaling variability) and extrinsic factors (eg, absorption, biotransformation, drug interactions, underdosing, increased platelet turnover, and, ultimately, noncompliance). However, atherosclerosis is a multifactorial disease, and therefore some patients will inevitably experience recurrent atherothrombotic events despite optimal platelet inhibition. Accordingly, a clinical definition of platelet resistance is also problematic.

Noncompliance with antiplatelet therapy may be as high as 22%. Thus, resistance may be overestimated if noncompliance is not eliminated. Clopidogrel resistance may become increasingly relevant as it moves to the generic market given that it may become increasingly widely prescribed.

Identification of clinically meaningful in vivo pharmacological resistance in a reproducible, cost-effective, and timely manner, using an in vitro test, is challenging. Several tests of antiplatelet resistance lack acceptable sensitivity, specificity, and reproducibility. Platelet reactivity is only measured via a single pathway, whereas a synergistic blockade of multiple pathways is likely to be important in vivo. A more complex approach to the assessment of platelet function may improve the accuracy of predicting thrombotic events.

Quo vadis?

Rapid decision making about the optimal antiplatelet regimen is essential in ACS management, especially in patients requiring primary PCI. Identification and management of clinically relevant clopidogrel nonresponsiveness is therefore a priority. Ideally, antiplatelet therapy could be individualized, incorporating a validated assessment of ischemic and bleeding risks and perhaps CYP2C19 genotyping. A point-of-care test determining responsiveness could facilitate appropriate drug selection. Subsequently, the dose may be adjusted to optimize therapeutic effect while minimizing bleeding risk. In the RESPOND study, nonresponders had consistently lower clopidogrel-induced platelet inhibition with all assays studied. This result suggests that screening with a single 300 mg dose can perhaps identify clopidogrel nonresponders. An alternative agent may be an effective strategy in this scenario. Evaluation of whether this strategy translates into long-term benefit in a prospective clinical trial will be necessary to move beyond mere speculation.

However, if ticagrelor is to become a mainstream antiplatelet drug, certain caveats should be heeded. Ticagrelor requires twice-daily dosing because of its rapid reversibility. This requirement necessitates scrupulous compliance to minimize the risk of recurrent ischemic events and stent thrombosis. Additionally, ticagrelor is associated with dyspnea and bradycardia, which may lead to discontinuation. For example, dyspnea was more common with ticagrelor in the PLATO study (13.8% versus 7.8% with clopidogrel). Few
patients discontinued ticagrelor because of dyspnea (0.9% versus 0.1% with clopidogrel). Discontinuation of the study drug because of adverse events occurred slightly more frequently with ticagrelor (7.4% versus 6.0% with clopidogrel; \(P<0.001\)) in the PLATO study.\(^{10}\) The dyspnea and bradycardia-related could be related to the inhibition of adenosine reuptake by erythrocytes, but paradoxically this has been hypothesized to contribute to the therapeutic effect of ticagrelor by reducing blood pressure, improving microcirculatory function, and protecting against reperfusion injury.

### Conclusions

Strong evidence supports the hypothesis that drugs which enhance platelet inhibition reduce the risk of recurrent ischemic events.\(^8,10\) If this is combined with an absence of overall increased bleeding\(^8\) or is directed toward subgroups likely to benefit most,\(^9\) then the benefits may be even greater. Moreover, strategies that directly reduce the incidence of bleeding complications, such as radial artery access, are intrinsically linked with enhanced outcomes of more potent antiplatelet drugs. The antplatelet armamentarium continues to evolve rapidly, affording clinicians an increasing choice in drug therapy in patients with ACS. Long-term results and comparisons between prasugrel and ticagrelor in prospective randomized clinical trials are awaited with great interest.\(^19\) With the availability of novel antiplatelet agents with more potent platelet inhibition, aspirin and clopidogrel resistance may soon be irrelevant.

### Disclosures

Dr Lip was an investigator in the PLATO study and has received funding for research, educational symposia, consultancy, and lecturing from different manufacturers of drugs used to treat thrombosis.

### References


Key Words: Editorials, antiplatelet therapy resistance, aspirin, clopidogrel, ticagrelor
Role of Ticagrelor in Clopidogrel Nonresponders: Resistance Is Futile?
Luke Tapp, Eduard Shantsila and Gregory YH Lip

Circulation. 2010;121:1169-1171; originally published online March 1, 2010;
doi: 10.1161/CIR.0b013e3181d8d929
Circulation is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2010 American Heart Association, Inc. All rights reserved.
Print ISSN: 0009-7322. Online ISSN: 1524-4539

The online version of this article, along with updated information and services, is located on the
World Wide Web at:
http://circ.ahajournals.org/content/121/10/1169

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published
in Circulation can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial
Office. Once the online version of the published article for which permission is being requested is located,
click Request Permissions in the middle column of the Web page under Services. Further information about
this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to Circulation is online at:
http://circ.ahajournals.org/subscriptions/