Aspirin and clopidogrel hyporesponsiveness and nonresponsiveness in patients with coronary artery stenting

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Abstract: Patients undergoing coronary artery stenting receive an antiplatelet regimen to reduce the risk of antithrombotic complications. Current guidelines recommend the use of acetyl salicylic acid (aspirin) and clopidogrel as evidenced by large clinical trials. There has been a concern about variable responses of patients to aspirin and clopidogrel which may predispose them to subacute stent thrombosis or late stent thrombosis. Up to 25% of patients with acute myocardial infarction (AMI) undergoing percutaneous coronary intervention (PCI) were found to have hyporesponsiveness or resistance to clopidogrel which may predispose them to recurrent events. Dual antiplatelet regimen is a standard therapy in these patients and there is always a concern about variable responses to aspirin and clopidogrel predisposing them to acute coronary syndrome (ACS). Prevalence of this hyporesponsiveness or resistance may be due to noncompliance, genetic mutations, co-morbid situations and concomitant use of other drugs. This issue is of considerable importance in the era of coronary drug eluting stents when a long-term dual antiplatelet regimen is needed. This paper is a review for clinicians taking care of such patients with hyporesponsiveness or nonresponsiveness to dual antiplatelet regimen.

Keywords: aspirin, clopidogrel, acute coronary syndrome, coronary artery stenting

Background
Do we manage hypertension without monitoring blood pressure? Do we manage diabetes mellitus (DM) without monitoring glucose? Do we manage warfarin sodium (Coumadin®) without monitoring international normalized ratio (INR)? If so, why do we use an antiplatelet regimen without monitoring platelet function testing. It is not uncommon for patients with recent coronary intervention with bare metal stents (BMS) or drug eluting stents (DES) to present to the emergency department with chest pain. Just as we monitor DM with a gluco-meter or warfarin with INR, we should monitor antiplatelet regimens in patients with coronary artery stenting. It is crucial to understand the heterogeneous response of patients to aspirin and clopidogrel which can result in unstable angina, or myocardial infarction (MI).1 Trials are ongoing (Gauging Responsiveness with a VerifyNow Assay-Impact on Thrombosis And Safety [GRAVITAS] and the ARCTIC study2) to determine whether tailored antiplatelet therapy, using platelet function testing, reduces major adverse cardiovascular events after DES.

Dual antiplatelet regimen with aspirin and clopidogrel has been shown to significantly reduce the cardiovascular events.4 Although clopidogrel mostly reduces risk of cardiovascular events after coronary stenting, a significant number of events still occur in these patients.1 These events may be due to subtherapeutic responses of some patients to aspirin and clopidogrel, predisposing them to subacute stent thrombosis or
late stent thrombosis. Up to 25% of patients with acute MI (AMI) undergoing percutaneous coronary intervention (PCI) were found to have variable response to clopidogrel, predisposing them to recurrent events of acute coronary syndrome (ACS).\(^1\) Prevalence of this hyporesponsiveness or resistance may vary in certain co-morbid situations as described below. This is a critical issue in the era of DES with need of long-term dual antiplatelet regimens.

**Variable responses to antiplatelet agents and their mechanism**

The exact definition of “resistance” to antiplatelet therapy on the basis of physiology does not exist. However, there is a significant prevalence of variable response to dual antiplatelet regimens similar to different responses to anti-hypertensive therapy or warfarin therapy. Therefore, it is imperative to understand this variable response or hyporesponsiveness to aspirin and clopidogrel in these patients. A clear definition of this response should be established and, based on this, one may be able to categorize patients as responders, hyporesponders, nonresponders, or resistant and thus manage their therapeutic regimen accordingly.

The effect of aspirin is mediated by the irreversible inactivation of cyclo-oxygenase (COX-1), leading to the prevention of thromboxane \(A_2\) generation from arachidonic acid.\(^5\) Following oral administration, aspirin is effective as an antiplatelet agent within 60 minutes. COX-1 is rapidly resynthesized by nucleated cells, such as endothelial cells, and therefore the effect of aspirin on nucleated cells lasts only for a relatively short time. In contrast, the effect of aspirin on platelets (anucleate cells) lasts for the life of platelets (7–10 days). Thienopyridines inhibit platelets via a blockade of P2Y12 adenosine diphosphate (ADP) receptors.\(^6\)

As the mechanisms of aspirin and thienopyridines are complementary, the combination of both drugs has a greater degree of inhibition of platelet aggregation than either one alone. Clopidogrel is a pro-drug and is activated through sequential oxidative steps in the liver. After absorption, 85% of drug is inactivated by esterases (hydrolase enzyme) and only 15% remains for activation by the liver to its active metabolite through several cytochrome 450 (CYP-450) proteins. Therefore, genetic mutation or polymorphism of such subenzymes can affect the therapeutic response of clopidogrel.\(^7-9\)

Based on this genetic polymorphism, by means of a simple buccal swab or blood sample, 3 phenotypes can be identified as being a: poor metabolizer (PM); intermediate metabolizer (IM), or normal metabolizer (NM). Underlying genetic mutation is due to variation in alleles (nucleotide sequences). The laboratory testing for this CYP-450 2C19 DNA mutation is done by identifying 8 different kinds of alleles. The nomenclature of this CYP 2C19 mutation is reported as *1 (star 1) to *8 (star 8).\(^10\) The presence of CYP 2C19 *2 allele is associated with reduced clopidogrel responsiveness and this is related to increased risk of MI, stent thrombosis. The frequency of this mutation in Caucasians and African Americans is 30%.\(^8\) A poor metabolizer, as diagnosed by genetic testing, may exhibit different responsiveness to clopidogrel due to a failure to generate sufficient active form metabolites, which then leads to a lack of therapeutic effect. In a recent study, 162 healthy subjects were treated with clopidogrel and approximately 30% of these were found to have at least one CYP2C19 reduced-function allele that led to a relative reduction of 32.4% of the active metabolite of clopidogrel in plasma.\(^8\) Moreover, among subjects treated with clopidogrel in TRITON-TIMI 38, carriers of the *2 defective gene (CYP2C19 reducing function allele called *2 genotype) had a relative increase of 53% in the composite primary efficacy outcome of the risk of death from cardiovascular causes, MI, or stroke compared to noncarriers.\(^11\) This may lead to the need for unconventional doses of clopidogrel or an alternative oral antiplatelet drug in such patients. CYP2C19 metabolizing enzyme also catalyzes the biotransformation of many other drugs\(^12\) and the concomitant use of such drugs with clopidogrel may also change the efficacy of antiplatelet therapy. Various biological factors such as genetic polymorphism or gene mutation may account for such hyporesponsiveness or nonresponsiveness. Other causes of hyporesponsiveness of antiplatelet drugs may be as simple as noncompliance\(^13-16\) or poor absorption,\(^17\) due to abnormalities in the mechanism of action or genetic makeup.\(^18\)

Smoking has also been proposed to cause hyporesponsiveness in both aspirin and clopidogrel patients,\(^11,16,19\) whereas other researchers have reported that smokers were likely to be

<table>
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<tr>
<th>Table 1</th>
<th>Mechanisms of “resistance” to aspirin and clopidogrel</th>
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<tr>
<td>Non compliance</td>
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<tr>
<td>Poor absorption</td>
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<td>Sub optimal dose</td>
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<tr>
<td>Smoking</td>
<td>Genetic polymorphism</td>
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<td>Thrombocytosis</td>
<td>Concomitant medication</td>
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<td>Co-morbid conditions</td>
<td>Severe coronary artery disease</td>
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Platelet-platelet aggregation testing

Aggregation is the most common measure of platelet reactivity and platelet inhibition. This assay is based on platelet aggregation by stimulation with various agonists. Such aggregation testing between pretreatment and post-treatment of aspirin or clopidogrel are the most common estimates of responsiveness to aspirin or a P2Y12 receptor inhibitor like clopidogrel. 23 Historically, the “gold standard” test is LTA. 17 LTA uses an optical device that measures the rate and extent of change in light transmittance caused by platelets aggregating in a whole blood sample. This test determines the level of platelet function in response to a variety of agonists, as there are many pathways through which antiplatelet medications work. Blood samples with inhibited platelets (with antiplatelet medication) will produce low levels of light transmittance compared to normal functioning platelets which aggregate normally. This test has been widely used to measure the effect of dual antiplatelet regimens. LTA testing is tedious, time consuming and requires specialized staff, compared to rapid point-of-care (POC) assays, which are simple, less laborious and time efficient. VerifyNow® is one of the POC platelet aggregation tests that does not require sample preparation or pipetting and is also strongly correlated with LTA. 26,24

By this methodology, aspirin response is reported as aspirin reaction units (ARU). Aspirin blocks platelet activation by preventing COX-1 enzyme from converting arachidonic acid to thromboxane A2. The extent of this blockade is determined as ARUs. More than 550 ARUs is considered as diminished aspirin-induced platelet dysfunction. 27-30 Thienopyridines, like clopidogrel, block platelet activation via P2Y12 ADP receptors and the extent of this blockade is reported as P2Y12 reaction units (PRU). This test takes advantage of different receptors of platelets stimulated by different agonists. Thrombin receptors are strong platelet activators and function independently of P2Y12 ADP receptors. The base value (base PRU) is calculated by stimulating these receptors to estimate the total possible platelet aggregation irrespective of aspirin or clopidogrel responsiveness, nonresponsiveness or “resistance”

### Table 2

<table>
<thead>
<tr>
<th>Aspirin</th>
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<tr>
<td>ARU &gt; 550 (lack of ASA-induced platelet dysfunction)</td>
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<td>C/EPI-CT &lt; 193 seconds</td>
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<tr>
<td>C/ADP-CT &lt; 121 seconds</td>
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<th>Clopidogrel</th>
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<td>Responders</td>
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<td>Resistant</td>
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*Response (%) of platelet inhibition by clopidogrel.

**Abbreviations:** ARU, aspirin reaction units; C/EPI-CT, collagen/epinephrine closure time; C/ADP-CT, collagen/ADP closure time; ASA, aspirin.
of the patient taking or not taking clopidogrel. Then the extent of platelet inhibition by clopidogrel can be measured by using a selective ADP agonist to measure P2Y12 ADP receptor inhibition. The percentage inhibition of P2Y12 ADP receptors is determined by the difference between base PRU and PRU determined from ADP agonist (as illustrated in Figure 1). Less than 20% inhibition indicates low to no clopidogrel-induced inhibition of platelet function.

**VASP phosphorylation**

The surface expression of platelets can be determined by flow cytometry. Using monoclonal antibodies, platelet function can be determined by exploiting the different receptor expression of resting and activated platelets. This test (VASP) exploits the mechanism of intracellular signaling; the advantage of VASP phosphorylation testing is its specificity for the P2Y12 signaling pathway, although drawbacks include sample preparation, the need for experienced staff, and the expenses of the process.

**Tests dependent on factors released from activated platelets**

Some tests measure the factors released by activated platelets as a measure of platelet activation such as serum or thromboxane B2 and urinary 11-dehydro-thromboxane B2. These are COX-dependent and nonspecific. Moreover, urinary 11-dehydro-thromboxane depends on renal function. Another test, platelet-derived-miroparticles is expensive, and requires sample preparation and flow cytometry.

**Factors causing hyporesponsiveness or nonresponsiveness**

A major concern after interventional procedures is an ischemic event, often caused by activated platelets at the site of a coronary stent implantation, which underscores the critical role of antiplatelet agents in PCI. Most of the time, the assumption is that dual antiplatelet regimens are efficacious without any objective testing. Unfortunately, not all patients respond equally and up to one-third of patients on an antiplatelet regimen do not experience the expected results. It is very important to identify these nonresponders, especially with the increasing use of DES. Nonresponders are at 5 times greater risk of MI, stent thrombosis and death than responders. Numerous factors may contribute to this unresponsiveness which may include noncompliance, drug interactions, DM, chronic renal failure or genetic makeup.

**Factors modifying the efficacy of aspirin**

Recent observations suggest that the primary cause for aspirin resistance may be poor compliance. Hence, it is imperative to ensure compliance with an aspirin regimen prior to platelet function testing. Another important factor leading to aspirin resistance may be concomitant use of

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**Figure 1** Understanding platelet function testing.

**Abbreviations:** AA, arachidonic acid; PRU, P2Y12 reaction units; ARU, aspirin reaction units; TXA2, thromboxane A2.
nonsteroidal anti-inflammatory drugs such as ibuprofen, which interferes with COX-1 acetylation. Even though some investigators have suggested increasing the aspirin dose in such patients, in vitro studies have shown that increasing the dose of aspirin may not affect COX-1 inhibition and may work through a non-COX-1 inhibition pathway. Aspirin resistance or hyporesponsiveness may be also due to age or gender issues. Many studies have shown that women are more likely to have aspirin resistance and the majority of these subjects are postmenopausal. However, this has been disputed in other studies. Some studies have shown that advanced age may be an important factor and may be due to decreased metabolism of aspirin in old age. Therefore, it is important to test the antiplatelet efficacy of aspirin in patients with PCI.

Furthermore, this may help determine if baby aspirin is useful in such patients. Certain co-morbid conditions such as DM and chronic renal failure may cause hyporesponsiveness to aspirin. Some studies found a higher prevalence of nonresponders in smokers while other researchers have disputed this.

**Factors modifying the efficacy of clopidogrel**
The most important cause of variable platelet activity suppression may be noncompliance and inadequate dosing. Resistance to clopidogrel may co-exist with aspirin and this may be prevalent up to 50% of patients with aspirin resistance (or it may present by itself). Subacute thrombosis of coronary stents in the CREST study was found to be more common in patients with hyporesponsiveness to clopidogrel. Clopidogrel is a pro-drug and its active metabolite irreversibly inhibits the binding of P2Y12 ADP receptor on platelets. Such metabolism invokes the possibility of the concurrent use of other medications as contributing factors to hyporesponsiveness in patients with coronary artery stenting. Concomitant use of other drugs, such as calcium channel blockers (CCB), angiotensin-converting enzyme (ACE) inhibitors, or beta blockers, does not cause hyporesponsiveness to clopidogrel, although a higher number of nonresponders is seen with concomitant use of CCB and ACE inhibitors. A recent study by Siller-Matula et al showed that coadministration of CCB decreased platelet inhibition though in vitro incubation with CCB did not alter platelet aggregation in patients taking clopidogrel. This finding suggests that the negative effect in vivo may be at the level of metabolic pathways.

There has been much concern about the use of proton pump inhibitors (PPI) in conjunction with clopidogrel. A recent retrospective review of 8205 patients published in *JAMA* showed a strong association of adverse clinical outcomes when clopidogrel was used in conjunction with omeprazole, though this was recently disputed in a late breaking trial (COGENT) at the TCT Conference on September 24, 2009 in San Francisco, USA (http://www.tctmd.com/Show.aspx?id=85972). Omeprazole (Losec® or Prilosec®) is both a substrate and an inhibitor of CYP 2C19 and it may decrease the metabolism of clopidogrel to its active metabolite. Other PPIs that are CYP2c19 inhibitors are esomeprazole (Nexium®), lansoprazole (Prevacid®), and rabeprazole (Aciphex®). A recent clopidogrel medical outcome study has also suggested a similar interaction between PPI and clopidogrel.

**Management strategies**
A standard definition of hyporesponsiveness or nonresponsiveness or resistance does not exist. One may also argue that incidence of resistance or hyporesponsiveness is overestimated. But there are several studies showing that a poor response to these drugs may translate into adverse outcomes as discussed earlier. Increasing the dose of aspirin has been shown to improve the response in some patients. The impact of increased clopidogrel dosing was evaluated in patients with suboptimal response to clopidogrel in the OPTIMUS trial. This study showed that a dose of 150 mg of clopidogrel significantly decreased the platelet aggregation compared to a 75 mg dose. Likewise, Gurbel et al showed better efficacy of 600 mg compared with 300 mg loading dose. Furthermore, the combination of clopidogrel with a synergistic antiplatelet agent like dipyridamole can also improve the response. Theoretically cytochrome P 450 inducers can increase the active metabolites of clopidogrel, which could be an alternative to an increased dose, especially if there is gastrointestinal intolerance to the drug.

**New drugs on the horizon**
Other drugs on the horizon include a new thienopyridines agent (prasugrel) which has been evaluated in several large scale trials. In these studies, a loading dose of 60 mg and maintenance dose of 10 mg of prasugrel produced more consistent platelet inhibition compared to clopidogrel with a loading of 300 mg and maintenance of 75 mg. This translated into a lower rate of combined primary outcomes of death, nonfatal MI and stroke in high-risk patients with ACS. Prasugrel also demonstrated a 50% reduction in stent thrombosis in one trial. However, superior efficacy of
this agent has been somewhat offset by the increased risk of bleeding. While this drug has been approved in Europe for some time, the Federal Drug Administration (FDA) approval of this antiplatelet drug on July 10, 2009 was a major step forward in patients with ACS and PCI in the US. This may also be helpful in patients with CYP2C19 mutation, as it may not affect the pharmacokinetics and pharmacodynamics of prasugrel compared to clopidogrel.

Another highly selective, oral, nonthienopyridine drug is ticagrelor (AZD6140) which has been investigated in a large phase III trial. This also works by antagonizing ADP at the P2Y12 receptors; it does not require transformation to active metabolite and has a half-life of 7 to 8 hours. The safety and efficacy of ticagrelor were investigated in a trial named PLATO (Platelet Inhibition and Patient Outcome), the results of which were presented at European Society of Cardiology (ESC 2009) and simultaneously published in *N Engl J Med*. In this study patients were randomized for a double blind trial to a Ticagrelor group (9333 patients with loading dose of 180 mg followed by 90 mg twice daily) or a clopidogrel group (9291 patients with loading dose 300 mg followed by 75 mg daily). Death from vascular causes, MI or stroke within 12 months occurred less frequently in the ticagrelor group. The potential availability of 3 ADP receptor P2Y12 inhibitors may make it possible to individualize antiplatelet regimens rather than a “one size fits all strategy”. Ticagrelor may be preferred in acute ACS patients with unknown anatomy, in whom coronary artery bypass grafting may be anticipated, as this is a reversible ADP receptor inhibitor. There may be a potential for switching clopidogrel or prasugrel to ticagrelor in patients who need elective surgery. This may become the antiplatelet agent of choice in situations where surgical procedures cannot be deferred. Other new classes of antiplatelet agents include thrombin receptor antagonists called protease-activated receptor (PAR-1) inhibitors. PAR-1 is the main platelet receptor for thrombin, the inhibition of which may lead to the development of novel antiplatelet agents.

Conclusions

Platelets display an enormous complexity by their variety of receptors and the myriad of molecules they secrete. These receptors and molecules mediate a large number of physiologic and pathophysiologic processes and hence are a target for multiple antiplatelet agents. Variable responses to oral antiplatelet regimens are well known. Therefore, it is important to distinguish between hyporesponsiveness or nonresponsiveness or resistance (failure to inhibit platelets activity), and treatment failure (the clinical outcome of a recurrence of ischemic events). As described earlier, the prevalence of hyporesponsiveness or nonresponsiveness or resistance may be an aberration of the methodology; however, there is clearly accumulating evidence that in vivo resistance to oral antiplatelet regimens leads to a higher risk of atherothrombotic complications such as unstable angina, and MI. New developments in drugs may offer a narrow range of response variability leading to more predictive efficacy. Antiplatelet testing or genotyping may help uncover the underlying mechanisms of hyporesponsiveness or nonresponsiveness or resistance and help the development of personalized patient oral antiplatelet regimens. There is a need for large-scale studies documenting the efficacy of point of care assessment of platelet function, which will be a true departure from the “one size fits all” strategy in managing antiplatelet regimens in coronary artery stenting.

Disclosures

The authors declares no conflicts of interest.

References


