Platelet dysfunction in the perioperative and critical care setting

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\textbf{Abstract:} Achieving haemostasis and, simultaneously, preventing venous thromboembolism (VTE) in the perioperative and critical care setting is challenging. Although the traditional coagulation parameters, including international normalised ratio (INR), activated partial thromboplastin time (aPTT) and platelet count, are important, evidence suggests that far more subtle aspects of platelet dysfunction may occur without antiplatelet therapy, and play a pivotal role in critical bleeding and the pathogenesis of VTE. With advances in our ability to measure platelet activity and function beyond platelet count alone, we are making substantial progress in our understanding of how and when platelets interact and cross-talk with coagulation factors to achieve haemostasis and, conversely, contribute to the development of VTE after major surgery or trauma. With the ability to quantify the ability of platelets to aggregate, it is clear that platelet function tests have the potential to assist prediction of bleeding after cardiac surgery, especially for those who are treated with P\textsubscript{2}Y\textsubscript{12}-blockers. In addition, unregulated platelet activation or, conversely, platelet dysfunction in the absence of antiplatelet therapy, is increasingly being recognised as one of the key missing elements in the pathogenesis of VTE or critical bleeding, respectively. Our current understanding of the coagulation system—in particular the significance of platelet dysfunction and activation—may not be adaptive to benefit survival in all patients, especially after traumatic brain injury (TBI). Indeed, cardiolipin release from damaged neuronal mitochondria into the systemic circulation may explain why coagulopathy after TBI is associated with increased VTE instead of bleeding. Similarly, in patients with acquired coagulopathy due to liver disease, sepsis or trauma, excessive platelet activation has been observed explaining why these patients are not immune to developing VTE and, hence, VTE prophylaxis is still needed. While many drugs can inhibit platelet function through one of the three major platelet activation pathways—cyclooxygenase/thromboxane A\textsubscript{2}, P\textsubscript{2}Y\textsubscript{12}-ADP and thrombin—currently we do not have many interventions that can improve platelet function. Recent evidence suggested that an antioxidant polyphenol, resveratrol, may improve platelet function by reducing apoptosis and improving platelet mitochondrial function. We are now entering the exciting stage of precision-medicine in the area of coagulation in perioperative and critical care. Ultimately, we would be able use information beyond abnormal platelet count, INR and aPTT to guide platelet transfusion including withholding unnecessary platelet transfusion, and also avoid omission of VTE prophylaxis in the critically ill.

\textbf{Keywords:} Bleeding; coagulation; platelet function; transfusion; thrombocytopenia; thrombosis

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Background

Approximately $1\times10^{11}$ platelets are produced by the cytoplasmic fragmentation of megakaryocytes each day (1). It has been argued that platelets—in many ways—behave more like cells than fragments. Studies suggest that thrombopoiesis can occur in the circulation, and platelets are capable of turning buds and extrusions into pro- and then pre-platelets, a process not normally expected for a simple cell fragment (2). Platelets can produce phospholipid microvesicles (100–1,000 nm), also called platelet microparticles (PMPs) or extracellular vesicles, with the stimulation of thrombin, collagen, or high shear stress with turbulent blood flow and during the apoptosis process. Unlike microparticles secreted by other cells, PMPs are not released through exocytosis, and have the distinct property of expressing the same antigens as the (parent) platelets (2). In addition, platelets can sense injured vascular subendothelium structures or infectious pathogens and respond to soluble molecules—including thrombin or thrombin-derived peptides—or bind to pathogens, either directly or indirectly, as immune complexes with antibodies or complement factors (2,3). Platelets are considered as one of the most short-lived ‘cells’ in human by a process called senescent platelet death (or apoptosis by others) that is highly regulated by intra-platelet signalling mechanisms (2,4) through thrombin activation (5), resulting in release of PMPs (6). Functionally, platelets play a pivotal role to control bleeding by adhering to an injured vessel wall and activating the coagulation factors.

Achieving haemostasis and, simultaneously, preventing arterial and venous thromboembolism (VTE) in the perioperative and critical care setting is challenging. With advances in our ability to measure platelet activity and function beyond platelet count alone, we are making substantial progress in our understanding of when and how platelets interact and cross-talk with the coagulation factors to achieve haemostasis and, conversely, contribute in the development of thrombosis after major surgery and trauma. In addition, platelets are increasingly recognized to play a role, as well as being affected by a variety of pathophysiological processes including inflammation and host defence (7). In this narrative review, we aimed to summarize the significance of platelet dysfunction and the possible roles of measuring platelet function in the perioperative and critical care setting.

Current state of managing platelet transfusion & function

To a large extent, bleeding and thrombosis are the different sides of the same coin. Current transfusion and thromboprophylaxis guidelines, including the National Blood Authority critical bleeding management tool, recommended platelet transfusion and withholding anticoagulants using a certain platelet count threshold in an attempt to reduce bleeding and unnecessary platelet transfusion (8-10). These guidelines stated that these recommendations were based on weak evidence with an emphasis that more research is needed. The obvious evidence gap in these guidelines is whether the platelet function—beyond the count alone—has any important clinical relevance and, therefore, should be considered in the medical decision-making in perioperative medicine.

Our understanding of platelet physiology has dramatically improved in the last few decades, primarily driven by the need to support the advances in percutaneous cardiac interventions. It is clear now that platelet activity or function cannot be considered as an “all or none” phenomenon. Three distinct major platelet activation pathways are well established (7,11), and each of these can be specifically inhibited by an antiplatelet agent, so much so that it is not rare to have patients treated with dual antiplatelet therapy (DAPT) and, in some acute cardiac care situations, even triple antiplatelet therapy. There is strong evidence to support such approach in reducing in-stent thrombosis and subsequent acute coronary events, especially when reperfusion is not satisfactory (12). More interestingly, large randomised controlled trials (RCTs) showed that aggressive DAPT or triple therapy can also have an ‘antiplatelet intensity-related’ protective effect on VTE (12), confirming the pivotal role of platelets in the pathogenesis of both venous and arterial thrombosis. Nevertheless, there is also a problem of variability in the pharmacokinetics and pharmacodynamics of anti-platelet therapy between individuals (e.g., aspirin resistance and clopidogrel CYP-2C19 enzyme pharmacogenomic poor-metaboliser in 30% and 40% of the population, respectively) (13). This has led to widespread use of platelet function testing in an attempt to individualise antiplatelet therapy, by adjusting the dose or type of antiplatelet drug between different patients as well as within the same individuals over the time, in cardiology practice.
Counterintuitive results of the antiplatelet therapy and platelet transfusion trials in the perioperative and critical care setting

Traditionally, most clinicians would rely on how long a patient has stopped taking antiplatelet therapy as a way to determine the degree of residual antiplatelet effect that may still exist, and whether the patient would be safe to undergo invasive procedures. Two recent landmark studies have challenged the validity of this simple intuition. The ATACAS RCT involving 2,100 patients tested the hypothesis that a single-dose of preoperative aspirin before coronary artery surgery would reduce a composite endpoint of death and thrombotic complications (non-fatal myocardial infarction, stroke, pulmonary embolism, renal failure, or bowel infarction) within 30 days after surgery. This trial did not find any significant differences in all clinical outcomes between the placebo and aspirin groups (14). The second study assessed the hypothesis that platelet transfusion for patients who were taking aspirin before haemorrhagic stroke could limit the extension of cerebral haemorrhage with better neurological outcomes. Surprisingly, the odds of death or dependence at 3 months were actually higher in the platelet transfusion group than in the standard care group (adjusted common odds ratio 2.1, 95% CI: 1.2–3.6; \( P=0.01 \)) (15). In addition, 40 (42%) participants who received platelet transfusion had a serious adverse event during their hospital stay, compared to only 28 (29%) who received standard care. The ‘empirical’ nature of initiating an antiplatelet agent or platelet transfusion—without considering whether significant antiplatelet therapeutic effect already existed—could at least in part explain the rather counterintuitive results of these two landmark trials. Theoretically, we would like to know that both groups were well-balanced in their degree of platelet inhibition (by a platelet function analyser) at baseline; and in retrospect, platelet transfusion should only be reserved for those with a certain degree of antiplatelet activity on board at the time of stroke, instead of assuming that any patients who had been taking aspirin within 7 days of stroke would benefit from platelet transfusion without harms.

Viscoelastic blood tests in perioperative and critical care setting

Whole-blood viscoelastic blood tests are increasingly used to assess bleeding risk and guide transfusion in the perioperative and critical care setting (16-19). The initial clotting time (CT) on ROTEM® (or r-time on the thromboelastography®) is useful to detect the effect of unfractionated heparin (UFH) or low-molecular-weight-heparin (LMWH) or coagulation factor deficiency, while the maximum clot firmness (MCF) or maximum amplitude (MA) is more useful to delineate platelet activity or fibrinogen effect on the ROTEM® or thromboelastography®, respectively. The evidence to support their role, in particular the MCF or MA (17), to guide transfusion in cardiac surgery as well as to assess thrombotic risk (20) is gathering strength, and has now been incorporated in some official perioperative and critical care transfusion guidelines (18,21).

It is, however, important to understand the limitations of viscoelastic tests. These may include: (I) it does not detect the effects of hypothermia on coagulation factors activities and platelet dysfunction (as the sample is usually measured at 37 °C); (II) it does not measure the effects of hypocalcaemia on clot strength and platelet function when citrated blood samples are used for analysis; (III) it does not measure platelet adhesion (e.g., von Willebrand disease) or platelet aggregation problems (including the thrombotic risk of microangiopathic thrombotic (22), and—most important of all—it does not detect the effects of aspirin and clopidogrel in cardiac surgical patients (unless a TEG® Platelet Mapping module is used), because the activator used in viscoelastic tests would generate enough thrombin to activate phospholipase C (the dominant platelet activation pathway) overriding the inhibition of minor platelet activation pathways through the P\(_i\),Y\(_2\) receptors and cyclooxygenase-1 (23). As such, whole-blood viscoelastic blood tests cannot be considered as the ‘panacea’ in the management of haemostasis and thrombosis and should also be used in conjunction with full blood count, coagulation blood tests and possibly also platelet function tests (23).

Platelet function tests in perioperative and critical care setting

Excessive platelet activation (which may result in increased thromboembolism) can be implied by a number of platelet bioassays. These platelet-related assays and biomarkers are primarily used for research purposes currently and hence, we will not discuss their utility further in this review. Interested readers can refer to some of other reviews on this topic (24-26). Conversely, platelet function analysers that can detect suboptimal platelet function or effect of
antiplatelet agents are much more widely used in a clinical environment. These platelet function analysers can be broadly divided into measuring the overall platelet function or the integrity of a specific platelet activation pathway. The most important limitation for almost all of these analysers is that the findings can be affected by thrombocytopenia and hence the results must be interpreted with the platelet count. The principles behind these analysers, their possible applications and limitations are summarised in Table 1 (23,27-29).

Platelet function tests in cardiac and non-cardiac surgery

Cardiac surgical patients are often treated with one or more antiplatelet agents before surgery, and bleeding requiring blood product transfusion is common. The use of platelet function test, in both preoperative and postoperative periods, to predict blood loss and transfusion requirements in cardiac surgery has been reported by a number of studies. While heterogeneity between multiple small studies (n<200 in most studies) exists, there is a reasonable signal to suggest that quantitative assessment of preoperative platelet inhibition (or dysfunction) has an association with perioperative blood loss and risk of requiring allogeneic blood transfusion [platelet function analyzer (PFA): area under the receiver-operating-characteristic curve (AUROC) 0.66; multiple electrode aggregation (MEA): AUROC 0.64–0.77] (30).

In addition, there is also a signal to suggest that there is a quantitative relationship between the risk of severe bleeding after cardiac surgery and the degree of platelet inhibition by ADP-blockers (31) (Figure 1). Although using platelet function assessment to assess risk of bleeding appears promising, there is substantial uncertainty about its utility, in particular whether only one, two or all three platelet activation pathways are equally important in determining perioperative blood loss and transfusion. Currently, platelet function test has been recommended (class II-b recommendation, grade-b evidence derived from a single randomized clinical trial or large non-randomized studies) by the European Association for Cardio-Thoracic Surgery (EACTS) and the European Association of Cardiothoracic Anaesthesiology (EACTA) to guide timing of cardiac surgery if patients are treated with DAPT or P(Y12) blockers (32). Because most existing studies are relatively small, whether platelet function test can be used to predict blood loss and transfusion accurately, and hence can be used to guide platelet transfusion when bleeding occurs remains scientifically unproven. Although it makes clinical sense, the validity of using platelet function tests to guide timing of other invasive procedures, including epidural catheter insertion or neurosurgical procedures (33,34), in anaesthesia and perioperative medicine has also yet to be firmly supported by high-grade evidence.

Despite the promising hypothetical reasons to measure and optimize platelet function in both cardiology and cardiac surgical studies, the evidence to demonstrate the benefits of platelet function tests remains elusive (30,35). There are a few reasons why it is premature to abandon platelet function test in cardiac surgery completely. In the first instance, many studies have only assessed one specific type of platelet function analyser; it is possible that the other platelet function analysers may be better under different circumstances. Second, although still not completely proven, preliminary data suggests that platelet function test (supplemented by a viscoelastic test) may play a more important role to rationalize the choice of blood products during the period of active bleeding (36), when international normalised ratio (INR), activated partial thromboplastin time (aPTT), fibrinogen concentration and platelet count all relatively normal and the cause of ‘medical bleeding’ is uncertain. Third, platelet function test may also be considered in some special situations when any avoidable causes of bleeding or transfusion should be minimised, including those with difficulties in cross-match due to antibodies, and people who would not accept any form of allogeneic transfusion for a variety of reasons. Finally, it is most likely that platelet function test may be more accurate in predicting blood loss and transfusion in emergency cardiac surgical care or for those who are only treated with an antiplatelet agent stronger than aspirin alone (e.g., on clopidogrel or DAPT) and thus more indicated in emergency cardiac surgery when DAPT cannot be ceased prior to surgery (32). Indeed, the most promising studies on use of platelet function test to predict cardiac surgical bleeding appeared to involve primarily patients who were treated with ADP blockers or thienopyridines (37,38).

Improving platelet function during the perioperative period

While many drugs can inhibit platelet function through one of the three major platelet activation pathways (cyclooxygenase/thromboxane A2, P(Y12)-ADP and thrombin), currently we do not have many interventions that can improve platelet function. Resveratrol, a
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<tr>
<td>VerifyNow®</td>
<td>Point-of-care closed system coated with fibrinogen measuring platelet reactivity by the rate and extent of light transmission as aggregates form. Measures platelet αIIb/β3 aggregation with fibrinogen (αIIb/β3 blockers); ADP (P2Y12 receptor inhibitors) and thromboxane-A2 activation (aspirin detection)</td>
<td>Can measure response to antiplatelet agents including aspirin, P2Y12 inhibitors and αIIb/β3 inhibitors</td>
<td>Monitor antiplatelet therapy: anti-αIIb/β3 (abciximab or eptifibatide) in cardiac surgery. Quick, results provided in &lt;10 minutes. Requires very little sample processing and hands on time. Can detect platelet reactivity. Most accurate measure of response to clopidogrel</td>
<td>Has no proven efficacy in large randomised trials in guiding antiplatelet therapy based on platelet reactivity in patients with acute coronary syndromes (ACS) undergoing percutaneous coronary intervention (PCI)</td>
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<tr>
<td>Multiplate®</td>
<td>Measures platelet activity in response to different platelet agonists: arachidonic acid (APTimet), thrombin (TRAPTest), collagen (COLtest), ADP (ADPtest and ADPtest HS—for high sensitivity testing)</td>
<td>Can detect various antiplatelet activity: aspirin, P2Y12 receptor inhibitors (ADP blockers) and αIIb/β3 inhibitors. Uses whole blood and allows for platelet surface adherence (to sensors), mimicking in vivo platelet activation, adhesion and aggregation mechanisms. User friendly—machine has in-built pipette and instructional prompts</td>
<td>Effective in detecting platelet reactivity with a number of antiplatelet agents. Potentially useful for de-escalation of antiplatelet therapy in patients with ACS undergoing PCI</td>
<td>Conflicting results from numerous small studies in its ability to guide clinical practice or predict clinical outcomes. Not effective in predicting platelet transfusion requirements in perioperative setting for patients undergoing coronary artery bypass graft (CABG) surgery</td>
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<td>Vasodilator-stimulated phosphoprotein (VASP) phosphorylation test</td>
<td>A flow cytometry-based method that measures the magnitude of platelet activation. ADP and prostaglandin reagents are added as activators and the fluorescence intensity is calculated as an index</td>
<td>Sensitive analysis for detection of P2Y12 receptor inhibitors</td>
<td>Most accurate measure of the peak response to clopidogrel</td>
<td>Time consuming. Requires specialised equipment, staff training and processing procedures. Most laboratories do not typically perform these tests routinely or “after-hours”</td>
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<tr>
<td>Platelet Function Analyser (PFA)-100®/Innovance PFA-200®</td>
<td>Measures coagulation in the form of closure-time of an apparatus within a collagen coated cartridge under high-shear flow. Two cartridges are run simultaneously, one with an ADP activator and the other with epinephrine (adrenaline). Results represented in time (seconds). Normal ranges are advised to be established within house, theoretically however, closure times should be &lt;110 s for ADP and &lt;150 s for epinephrine</td>
<td>Detect platelet dysfunction caused by von Willebrand factor (VWF) deficiency on board antiplatelet activity. Ease of use: citrated whole blood, minimal set up</td>
<td>Excellent for detecting VWF deficiency (prolonged closure of both cartridges in absence of an antiplatelet agent) and low dose aspirin activity (prolonged epinephrine cartridge, but normal ADP cartridge)</td>
<td>Falsely prolonged closure times with low haemoglobin—low viscosity increases vacuum suction through capillary not allowing adequate flow of platelets through aperture. Times &gt;300 in both cartridges may be due to thrombocytopenia or high dose aspirin, but cannot differentiate between different mechanisms. Limited application for intraoperative use</td>
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<td>Light Transmission Platelet Aggregometry (LTA)</td>
<td>Platelet rich plasma is continuously stirred and measured as an increase in light transmission as platelets form aggregates with the addition of an exogenous platelet agonist (ADP, epinephrine, collagen, ristocetin or arachidonic acid)</td>
<td>Can measure a number of platelet functional disorders and detect the effects of many antiplatelet agents</td>
<td>Gold standard for detecting disorders of platelet dysfunction</td>
<td>Time consuming. Specialised processes to prepare and test sample are required and specialised training of staff to perform and interpret results. Affected by pre-analytical conditions (lipaemic samples, concentration of agonists). Heavily dependent on platelet count and must be adjusted in vitro, thereby potentially negating in vivo results</td>
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<td>TEG® platelet mapping &amp; ROTEM® Platelets™</td>
<td>TEG6S: provides measure of platelet function in four different tests (maximum response to thrombin, thrombin blocker, ADP and TXA2). Results are calculated in reference to the maximum potential to thrombin as a degree of inhibition</td>
<td>User friendly and quick to perform</td>
<td>TEG6S® lw platelet function can predict platelet transfusion requirements during perioperative for patients undergoing CABG. Can detect clopidogrel activity and measure overall hyper/hypocoagulability</td>
<td>ROTEM® Platelets™ not superior to Multiplate® in detecting platelet reactivity after discontinuation of P2Y12 inhibitors</td>
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Polyphenol with strong antioxidant properties, is commonly found in plants and plant products such as berries, grapes and red wine (~14 mg/L). Its actions on sirtuins (silent information regulator—SIRT1, SIRT3, and SIRT4) gene expression is similar to caloric restriction, resulting in activation of telomerase to maintain telomere length (39). It is increasingly consumed as a nutritional supplement for cardiovascular, metabolic, and possibly anti-ageing benefits (39,40). In a large number of small-animal studies, intraperitoneal resveratrol administration has been shown to restore SIRT1 activity, attenuate hepatocyte injury, improve cardiac contractility, and ameliorate hypoxia-induced liver and kidney mitochondrial dysfunction following haemorrhagic injuries (41). A recent human in-vitro study showed that resveratrol can preserve function of stored platelets by improving mitochondrial function and reducing apoptosis (42).

Using a double-blinded RCT design in a large animal (greyhound) model, our recent work showed that use of oral micronized resveratrol (10 mg/kg/d) for 1 week was associated with a substantial improvement in the animals’ tolerance to development of shock, requiring a much larger volume of haemorrhage to induce mean arterial blood pressure <40 mmHg compared to controls (67 vs. 51 mL/kg, Δ =16 mL/kg, 95% CI: 10–21) and there were significant improvement in platelet activity [measured by the maximal clot firmness (MCF) by the rotational thromboelastometry—ROTEM®] and thrombin generation (43) (Figure 2).

In addition, small phase-I human trials involved patients with colorectal cancer with hepatic metastases scheduled to undergo hepatectomy and colonic cancers preparing for surgery, oral resveratrol (at 5 and 1 g/d for 14 days before surgery, respectively) was well tolerated and associated with an increase in tumour apoptosis and a reduction in tumour proliferation (44,45). Therefore, there is a clinical need to confirm the potential benefits of resveratrol on coagulation by a phase II clinical trial before it can be tested as a perioperative therapy to prevent cancer recurrence in subsequent phase III cancer surgery trials.

**Platelet dysfunction and thromboembolism in the critically ill**

**Cardiolipin release, platelet dysfunction and coagulopathy after traumatic brain injury (TBI)**

TBI is a leading cause of death and disability in young people and accounts for approximately one-third of all trauma-related deaths. Coagulopathy is common in TBI and associated with poor clinical outcomes but the underlying pathogenesis remains poorly understood (46). Recent studies clearly showed that inhibition of platelet, in one or more of the three activation pathways, is common after severe TBI (47,48); and the degree of platelet dysfunction bears a quantitative relation with the severity of TBI in the absence of haemorrhagic shock or multisystem injury and antiplatelet therapy.

Cardiolipin is an anionic phospholipid primarily located in the leaflet of inner mitochondrial membrane of all cells and plays an important role in maintaining mitochondrial

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**Figure 1** Relationship between risk of severe bleeding and degree of platelet P<sub>2</sub>Y<sub>12</sub> inhibition.

**Figure 2** Changes in thrombin generation with and without resveratrol pre-treatment before development of haemorrhagic shock in dogs.
membrane fluidity. It regulates various mitochondrial processes including electron transport chain and programmed cell death (49). Cardiolipin maintains brain homeostasis, hence any alteration of its chemical structure or decrease in overall levels are implicated in the pathogenesis of various neurodegenerative diseases like Alzheimer’s and Parkinson’s disease. Recent experimental study using a murine TBI model showed that cardiolipin release from the brain could cause disruption of blood-brain-barrier and systemic and consumptive coagulopathy through activation of platelets (50-52). However, there are hardly any human clinical studies available that systematically investigated the effects of cardiolipin on coagulation, including its relationship to platelet dysfunction, after TBI. In addition, whether cardiolipin release leading to excessive platelet and thrombin activation is related to a reduction in plasma anti-thrombin levels (53,54)—which determines effectiveness of both UFH and LMWH—is also unknown. Understanding the mechanisms behind the development of TBI-induced coagulopathy is paramount to test and develop appropriate strategies to manage thrombocytopenia, in terms of platelet transfusion and thromboprophylaxis, after severe TBI. Further studies are therefore needed to determine whether the thrombocytopenic state after TBI is indeed prothrombotic, pro-bleeding, or both—changing with the time course of the disease.

Relationship between VTE and activation of platelets in patients with acquired coagulopathy

Acquired coagulopathy with thrombocytopenia, prolonged INR or aPTT is common in the perioperative setting. Yet, despite their underlying apparent bleeding propensity, the risk of thrombosis is high (55,56). In patients with chronic liver diseases, platelets appear to play a pivotal role in inducing both bleeding and thrombosis (57). In patients with coagulopathy in the perioperative setting (e.g., surgical emergencies and trauma) without chronic liver diseases, the contrition of excessive activation of platelets to development of bleeding requiring transfusion or thromboembolism is largely unknown. As such, the optimal haemostatic management including whether platelet transfusion is contraindicated for such patients remains highly controversial, and is one of the most important unanswered questions in VTE prophylaxis. Our recent work suggests that an increased clot strength on whole-blood viscoelastic test is associated with an increased risk of thromboembolic diseases in a variety of clinical situations (20), including patients with a prolonged INR or aPTT. We also found that excessive platelet activation in patients with acquired coagulopathy is associated with an increased risk of subsequent risk of clinical thromboembolic events (manuscript in preparation), consistent with the other evidence supporting the pivotal role of platelet in both arterial and venous thrombosis (12). In contrast to whole-blood viscoelastic tests, the currently available platelet function analysers are not useful to identify which patients at increased risk of thromboembolic events—with the exception in patients with heparin-induced thrombocytopenia syndrome (HITS) (58).

Conclusions and future research direction

Platelets are now known to process a variety of different physiological roles other than haemostasis alone, even though this latter function will remain as the main focus for most critical care physicians (2). Coagulation is an important area of perioperative and critical care medicine; and it is clear that the current practice of using platelet count and standard coagulation parameters alone to determine bleeding and thrombotic risk is inadequate, resulting in both false positives and negatives with important patient outcome and financial implications compared to in combination with viscoelastic and platelet function tests (15,16,59-61). Advances in quantitative assessment of platelet activity and/or function have allowed us to assess not only the effects of antiplatelet therapy, but also demonstrate the existence of platelet dysfunction without antiplatelet therapy in patients with severe TBI which has prognostic significance (46,52,62). Cardiolipin release from the brain appears to be responsible for the circulating microparticles that can explain why coagulopathy and thrombosis may coexist in patients with severe TBI; and excessive platelet activation in response to trauma and sepsis contribute to a high risk of thrombosis despite thromboprophylaxis in these patients (6).

Current evidence suggests that platelet function tests may assist clinicians in predicting their risk of bleeding especially in cardiac surgical patients treated preoperatively with thienopyridines, and platelet function may be improved using antioxidant polyphenol. Many of these hypotheses and emerging evidence do need further confirmation by adequately-powered phases II and III clinical trials. We are now entering the exciting stage of precision-medicine in the area of coagulation in perioperative and critical care (28). Ultimately, we would be able to use individualised-patient information—
beyond abnormal platelet count, INR or aPTT alone—to avoid unnecessary platelet transfusion and omission of anticoagulant VTE prophylaxis in the critically ill.

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Footnote

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References

2. Garraud O, Cognasse F. Are Platelets Cells? And if Yes, are They Immune Cells? Front Immunol 2015;6:70.


