Hemostasis in the Lab, Approaching to the Correct Diagnosis in the Coagulopathies

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To cite this article:

Received: October 2, 2020; Accepted: February 24, 2020; Published: March 6, 2020

Abstract: Coagulation is the result of a coordinated interaction of blood proteins, circulating cells, endothelium cells and extracellular matrix proteins in the vessel wall. This process works in conjunction with other mechanisms that have to keep the blood in an effective flow, without hemorrity when damage occurs or without a thrombus when clotting begins. Most of the time, there is not so much information that a clotting test gives us, however, in the face of a real clinical scenario that concurrent with a personal haemorrhage, this is essential to take action. In addition, there is a thrombophilic scenario that has to be investigated. In all of them, it is important to note that doctors have to take into account that the pre-analytical phase plays a fundamental role, so it is very important to look around any situation that may impact the patient's results. There is a wide range of laboratory tests to evaluate different mechanisms of hemostasis, according to the history and characteristic of the clinical patient. With this in mind, there is a right time to ask for the right test, to get a better result for diagnosis and treatment. Analytical quality is the set of actions that are carried out in the laboratory with the aim of obtaining an accurate result, reflecting the real situation of the patient. Grossly, platelets, clotting factors and the fibrinolytic system could be studied, not all patients need all tests, they could be solved with a simple test, but may require a deep focus; that is why the patient with a history of mucocutaneous bleeding has to be studied for thrombocytopenies or VWD; a patient with muscle or joint bleeding has to be a factor deficiency test. We propose an initial approach to detecting the pathological situations essential to make decisions. Each person has different symptoms and different approaches, which is why identification and the right tests give us the greatest opportunity to get the right diagnosis.

Keywords: Blood Coagulation, Hemostasis, Blood Test, Clinical Laboratory, Coagulopathy

1. Introduction

1.1. Initial Approach

Coagulation [1] is the result of a coordinated interaction of blood proteins, circulating cells, cells of the vasculature and proteins of the extracellular matrix in the vessel wall. This complex mechanism makes it difficult to evaluate it in the laboratory, which only restricts itself to measuring the coagulation proteins and circulating cells, while the vascular elements are not measurable.

When is necessary study a patient with abnormal coagulation test, the preanalytical phase plays a fundamental role, if this takes wrong way, the results will not be assessable, affecting the efforts from hemostasis laboratory team that are aimed providing excellence in care, which means reporting an accurate results, in the shortest time and the lowest possible cost according with the International standard care.

1.2. Preanalytical Phase

The preanalytical phase involves all those factors that affect the quality of the sample (identification, patient condition, adequate sampling collection, correct conservation and transport of the sample until it will be used).
The analytical quality is the set of measurements that are carried out in the laboratory with the aim of obtaining an accurate and precise result. [2]

Currently, laboratory tests that are available to evaluate different phases, depends on the sintomatology, is the test that should be ordered.

Figure 1. Shows the coagulation pathways.

aPTT: activated partial thromboplastin time, PT: prothrombin time, TT: Thrombin time.

2. The Correct Approach

2.1. The Platelet Phase; The Beginning

The platelet disorders symptoms are characterized by mucocutaneous bleeding; The principles causes of thrombocytopenia is show in Figure 2.

Figure 2. Study a patient with mucocutaneous hemorrhage.

For the platelet study used in the past, the bleeding time, by the Ivy's or Mielke’s technique. Bleeding time (BT) was a global test of primary hemostasis, its greatest utility was evaluate the platelets function; it can be altered in certain circumstances such as uremia, the use of some drugs such as antiplatelets, non-steroidal anti-inflammatory drugs (NSAIDs), ticlopidine among others, the same happens with low platelet counts, likewise in the alteration of platelet glycoproteins as in Glanzmann thrombasthenia, or von Willebrand's disease, but this test has not enough sensibility/specificity, so its use is not recommended actually. [3] Have fallen into disuse, due to the difficulty in its standardization.

2.2. Platelet Evaluation & Thrombocytopathies

To evaluate the platelet function, exist different tests but it is essential first of all, obtained the platelet count and the platelet size by CBC and the volume platelet mean (VPM), normally goes from 5-12 femtoliters (fL), and a normal count goes between 150 - 450,000 cells / µL.

Thrombocytopenia may be secondary to an increase in peripheral blood destruction, which is often associated with increased VPM, while lack of production in bone marrow (BM) is associated with decreased VPM. The decrease in platelets number can be the consequence of an antibody sensitive to EDTA that promotes the agglutination of platelets, known as pseudothrombocytopenia.

The most frequent cause of elevation in the number of circulating platelets is iron deficiency, followed by some other reactive causes such as infection, however, if thrombocytosis persists for more than 6 months, the possibility of a myeloproliferative disorder should be considered.

In the past, the clot retraction test was widely used [4], the test has the basis that the retraction of the clot is produced through the interaction of glycoprotein αIIbβ3 (GIIb-IIIa) with the actin of the cytoskeleton, a process that takes the rearrangement of the platelet membrane. Retraction requires ATP as an energy source.

It depends on several factors:
- a) Quantity and quality of the platelet,
- b) Fibrinogen concentration,
- c) Hematocrite.

This test has low sensitive and its influenced by the cleaning of the material [1] is not recommended.

The test that offers a good sensitivity and specificity to assess platelet defects is platelet aggregometry, described by Born in 1962, which has been revised and modified at the present [5]. It consists of measuring in real time the aggregation of platelets in a sample of platelet-rich plasma (PRP) by optical clearance. It requires a spectrophotometer in which the PRP sample is deposited in an incubation cuvette at 37°C, which is between a light source and a photocell that calculates the optical density or turbidity of the PRP. Several substances known for their agonist effect on platelets are added, in order to induce aggregation of platelets and simulate in vitro what happens in vivo. However, it must be remembered that during the neonatal period it can be a confuse test since it has been shown that neonatal platelets normally respond less to agonists, by the other side we have to consider is necessary a great blood volume. [6] So the flow cytometry would be much more helpful during this period. [7]

Something that we must have to considerer in the first instance for the aggregometry, is the platelet count, since in patients with thrombocytopenia the reported values will not be accurate, so international recommendation is a platelet count at least over 100×10⁹ cells/L.

Like all the rest of coagulation test, the preanalytical phase it’s crucial, some food and drugs can affect platelet function;
as garlic, chocolate, different kind of oils, alcohol, onions, peppers, ginger, NSAIDs, antibiotics, including antacids, so we must be cautious, for an accurate result.

The most commonly used agonist agents are ADP, epinephrine, collagen, ristocetin and arachidonic acid. The aggregation cinetica interpretation should always be done by an hematologist trained.

In Figure 3 the normal aggregation pattern is shown.

If a pattern of abnormal aggregation is observed in a patient, it is advisable to repeat the evaluation at least once, to verify that the abnormality persists, since the collection time, the centrifugation time, the platelet count, the pH, mixing speed, hematocrite, temperature, lipemia, air bubbles among others factors can interfere with the results.

The test helps us to diagnoses some thrombocitopaties like: Bernard Soulier disease, Glanzmann thromboastenia, some types of von Willebrand Disease (vWD), as well as cyclooxygenase alterations, and in patients with antiplatelet drugs it helps us to monitor their use and stabish resistance.

Flow cytometry [8] is the most accurate, quick, objective and quantitative method to study platelet membrane glycoproteins, the expression of some glycoproteins, like Ib-IX and V is diminished, it consists in the evaluation of the constitutive glycoproteins of the platelet membrane using markers such as CD41, CD42, CD61, CD62; as we describes:

CD41: recognises the platelet membrane glycoprotein GPIIb (the integrin alpha IIb chain) which is non-covalently associated with GpIIIa (the integrin beta 3 chain) to form the GpIIb/IIIa complex.

CD61: recognises the platelet membrane glycoprotein GpIIia (the integrin beta 3 chain).

CD42b: reacts with GPIb on megakaryocytes and platelets. CD42b also inhibits ristocetin-dependent binding of Von Willebrand Factor to platelets and ristocetin-induced platelet agglutination [8].

CD62p: P-selectin is found in the α-granules of platelets. Presence of CD42 on the platelet surface indicates platelet activation, while their absence could be implicated a granules disease. [9]

2.3. Point of Care Test [10]

Point of Care tests, such as the platelet function test (PFA-100) is not considered as a screening test for platelet dysfunction, it was originally thought for vWD, however emerge as an alternative for the monitoring of the various antiplatelet agents, other tests as point of care can be mention are: [11].

2.3.1. The Impact Cone and Plate (let) Analyzer

Is shear-induced platelet adhesion/aggregation onto surface in whole blood. Is use for screening of congenital primary hemostasis abnormalities with low sensitivity ans specificity; as well for evaluation of platelet response to aspirin and clopidogrel.

2.3.2. The VerifyNow System

Is due Fibrinogen-platelet agglutination in response to agonist in whole blood, to monitoring of the platelet response to antiplatelet agents.

2.3.3. The Plateletworks

Is due platelet counting pre- and post-activation in whole blood, to monitoring of the platelet response to antiplatelet agents.

However the results are not encouraging and these test are not recommended as a routine for the diagnosis of congenital thrombocitopaties, very useful for monitoring antiplatelet agents. [12]
3. Coagulation Factors, the Second Phase

On the other hand coagulation tests such as: prothrombin time (PT) and activated partial thromboplastin time (aPTT) are used as screening tests to evaluate the coagulation pathways, and hemorrhage risk, traditionally describes, intrinsic and extrinsic pathway of coagulation in a didactic way, however this is explained better in the coagulation cellular theory. [13]

The factors involved in the intrinsic pathway of coagulation are evaluated by the aPTT, while the PT evaluates the extrinsic pathway, both coincide in the common pathway (Figure 1).

Characteristically, patients with factor deficiency present joint or muscle hemorrhages. Figure 4, shows a quick review but no more than 110%, it is important that if the amount of anticoagulant. However, an error that is frequently not respected, so that will have to adjust the amount of citrate anticoagulant relation), the tubes must be filled up to 90% siliconized sodium citrated tubes in a 9:1 (total blood: citrate relationship) is important to remember the reference values of each laboratory and the age of the patients.

The result of the prothrombin time can be expressed in seconds, percentages (%) or in ratios.

The report is internationally recommended in seconds and the reason depend strongly on the reagent and the instrument used; to report in (%) a calibration curve should be made with a pool of normal plasmas or a commercial calibrator [14]. Current recommendations express their concern to continue using the percentage of activity given that it does not have a linear relationship and can fall into interpretation errors.

Depending on the type of thromboplastin (human placenta, rabbit brain, mouse or monkey and currently the recombinant) the result is added can vary widely, so a standardized method has been developed to express these variations: International Normalized Ratio (INR), the importance of this parameter lies in its usefulness to evaluate the effectiveness of anticoagulation with vitamin K antagonists, but it has little use in other conditions of coagulopathy such as liver failure.

\[
\text{INR} = \left( \frac{\text{PT}_{\text{pat}}}{\text{TP}_{\text{NGM}}} \right)^{1\text{ISI}}
\]

Therefore, it is suggested internationally that INR should only be reported for patients using vitamin K antagonists (VKA), currently there is a capillary TP which facilitates the monitoring of patients and their use with VKA.

Some causes of PT prolonged are hepatopathy, vitamin K deficiency, factor deficiency VII.

3.2.2. Activated Partial Thromboplastin Time (aPTT)

The aPTT evaluates the intrinsic pathway of coagulation. For this reaction to the citrated plasma, phospholipids, and a contact factor initiator such as kaolin, silica, celite or ellagic acid are added, then the incubation and finally Ca\(^{2+}\) is added. Thus, the sensitivity of the reagent depends on the origin, concentration of phospholipids, type of activator, ionic strength and stabilizers.

The normal result varies from 25-45 seconds; However, it is important to remember the reference values of each laboratory and the age of the patients.

The most frequent cause of the alteration of aPTT is the deficiency of one of the factors of the intrinsic pathway (II, V, VIII, IX, XI or XII), although this must be a place with an activity <50% to modify it, the most frequent cause is the use of unfractionated heparin (UFH) when the blood sample is taken through the catheters that are heparinized even with many people and after "washing" the catheter, we may have falsely prolonged aPTT; however, it is recommended internationally for the monitoring of UFH [15], as well as the
presence of acquired inhibitors such as aPTT, and the most frequent of these is the lupus anticoagulant, one way to differentiate them is by mixing with normal plasma, this is done with a volume of the patient's plasma plus a normal plasma volume, the correction should be observed in the first dilution since we remember that we only need 50% of the concentration of factors to present a normal coagulation and this directs the clinician to look for a deficiency of the factors of this way, the correction must be valued by the Rosner index, which is calculated in the following way:

\[
\frac{(\text{aPTT mix} + \text{aPTT pool})}{\text{aPTT patient}} \times 100
\]

Where each laboratory should establish the correction values, and if don’t get the correction index, considered an inhibitor.

Is a different history for the patient with thrombosis, the Figure 5 shows the approach of prolonged aPTT & thrombosis, where a inhibitor could be present, or a contact factor low level as Factor XII [16] has to be test.

Figure 5. aPTT & Thrombosis.

3.2.3. Thrombin Time (TT)

The thrombin time allows to evaluate the fibrin formation stage, by measuring the time it takes to coagulate the citrated plasma in the presence of thrombin, its reference value is between 15 to 20 sec. The reference values vary with the concentration of thrombin used. Thrombin time is prolonged with low levels of fibrinogen or dysfibrinogenemia, treatment with UFH (however, it is not used for monitoring), high levels of fibrin digestion products (FDP), the presence of paraproteins, as well as inhibitors of thrombin. [17]

Another screening test is reptilase time (RT). It is a coagulation time like the thrombin time, but where the clot is produced by the action of an enzyme of viper venom called reptilase. Reptilase transforms fibrinogen. It differs from the thrombin action on fibrinogen in that it releases only fibrinopeptide A. Reptilase time is not inhibited by heparin and can be used in place of the thrombin time in the evaluation of fibrinogen in heparinized patients. When evaluating hypercoagulability states, a normal result of the reptilase time excludes the presence of abnormal fibrinogen. If the effect of heparin is the only cause of prolonged prothrombin time, the RT will be normal.

The simultaneous alteration of PT and aPTT frequently indicates an alteration of both the intrinsic and extrinsic pathways, such as the case of liver disease, coagulopathy due to consumption, in less frequent cases. (Figure 1). Under these conditions, a utility will be obtained to measure FDP, D Dimer (DD) and fibrinogen levels that allows to orient the origin of a simultaneous alteration, of course in the correlation with the clinic manifestations.

4. Fibrinogen, the Sticky Phase

The fibrinogen [18], the last protein of the coagulation cascade, can be measured by chemical or immunological methods, and is found in concentrations ranging from 200-400mg / dL.

In fibrinogen deficiency o dysfibrinogenemia, patients traditionally show muco-cutaneous hemorrhages.

The fibrinogen level can increase up to 800 mg/dL in acute inflammatory process.

4.1. Measurement

Currently, its measurement is recommended by the Clauss method, which is functional and is based on the clottable fibrinogen in the presence of thrombin excess. The fibrinogen concentration in g/L is obtained by comparing the clotting time of the plasma sample with serial dilutions of a reference plasma with known fibrinogen concentration. [19]

As it is a semiautomatic coagulometric method, half of the indicated volumes can be used, this decreases the amount of sample needed, the presence of heparin or high levels of FDP give falsely diminished results.

4.1. Breaking Bad

D-dimer is a soluble fibrin digestion product from the plasmin action on fibrin formed, thus is a coagulation and fibrinolytic marker.

The measurement of DD [20] that is carried out by means of a specific monoclonal antibody against the D regions of fragmented fibrin, is a biomarker of fibrin formation, which not only rises in thrombosis, it is normally elevated in pregnancy, inflammatory processes, neoplastic, with age and others. It is more specific and sensitive test in patients with DIC, which values arise over than 500ng / mL.

Remember only values below 50mg of fibrinogen alter PT/aPTT.

5. Viscoelastic Test, the Vintage Era

There is an assay modality in the patient with coagulopathy, the so-called viscoelastic tests such as thromboelastography (TEG) and thromboelastometry (ROTEM) which describe in a graph, the global interaction of different components of the hemostatic system, such as coagulation factors, fibrinogen, platelets and fibrinolytic system, since whole blood is used and the kinetic and viscoelastic characteristics of the clot are evaluated in real time. [21]

Heparinase to deactivate heparin is now employed in the TEG (r) technology to prevent interference from heparin in samples.
Which allow us to evaluate hemostasis in a global way and allows us to make decisions immediately in the critical patient, just what the patient need, low risk better results.

Nomenclature Differences for ROTEM. [22]

Although TEG and ROTEM measure the same coagulation functions there is a different naming convention for each device.

5.1. Features of a Teg Curve

5.1.1. Initiation (R)
Represents period of time of latency from start to initial fibrin formation due to effects of Factor VIIa and Tissue Factor.

5.1.2. Amplification (K)
Represents time taken to achieve a certain level of clot strength due to thrombin and activation of platelets (where r-time = time zero)

5.1.3. Propagation (α-Angle)
Measures the speed at which fibrin build-up and cross-linking takes place (clot strengthening), and hence assesses the rate of clot formation.

5.1.4. Maximum Amplitude (MA)
A function of the maximum dynamic properties of fibrin and platelet bonding via GPIIb/IIIa and represents the strongest point of fibrin clot and correlates to platelet function: 80% platelets; 20% fibrinogen.

5.1.5. Clot Stability (LY 30%)
This is percentage decrease in amplitude 30 minutes post-MA and gives measure of degree of fibrinolysis.
In Figure 6, shows an example of the curve that is generated in a patient without pathology.

5.2. Rotem
ROTEM offers some advantages as portable device, and dinamyc test, for the platelet, factors and fibrinogen; more accurate viscoelastic test.
But both are incredible useful in the critical bleeding patient.
A normal TEG/ROTEM shows in Figure 6.

6. Conclusions
Once an adequate clinical evaluation of the patient with hemorrhage has been carried out, we will be able to guide the type of studies to be requested, which will also finally allow us to define the treatment to be followed according to the proposed approaches in this review, that is a quick form to evaluate a patient, depending of the symptoms and the correct test in the correct time to get a better diagnosis and treatment for our patients.

References


