Iron-induced parafibrin formation in tumors fosters immune evasion

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Although efficacious in vitro, it is well known that adoptive immunotherapeutic modalities lose their potency when applied in vivo. Furthermore, malignant cell exposure to blood platelets attenuates the anticancer activity of natural killer (NK) cells. We argue that upon contact with redox iron, fibrinogen is converted to a hydrophobic fibrin-like polymer that coats tumor cells and provides protection from immune-mediated destruction.

Introduction

Placke et al. have recently presented a view that blood platelets impair natural killer (NK) cells ability to recognize cancer cells by forming a physical barrier around the tumor cells. In connection with this it should be remembered that platelet require for their proper functions the presence of fibrinogen, an essential component of blood coagulation cascade. Fibrinogen (FBG) is a large molecular weight (340 kDa) multi-chain plasma protein that is converted with thrombin to fibrin monomer(s) (FM), an essential component of the clotting assembly that spontaneously polymerizes to form fibrin clots. However, prior to the formation of solid clots FMs remain soluble by forming complexes with the intact fibrinogen molecules. Thus, in addition to inducing platelet aggregation, FM complexes have been shown to cause red blood cell (RBC) agglutination and increased sedimentation, as well as cell aggregation of the bacterium Staphylococci. Under normal hemostatic conditions fibrin clots are susceptible to degradation by the blood fibrinolytic system that secures proper wound healing and growth of connective tissue at the site of vessel wall injury. This system is activated by the release of tissue plasminogen activator (tPA) resulting in the generation of active plasmin that under normal conditions effectively degrades fibrin into soluble fragments. However, it is not understood why degradation of fibrin in certain types of cancer is incomplete, despite increased tPA production as in the case of prostate cancer.

Mechanism of Parafibrin Formation

We have previously reported that the exposure of plasma proteins to disulfide-reducing agents resulted in the formation of huge insoluble aggregates, which when adhered to tumor cell membranes have been proposed to act as a barrier to tumor recognition by the innate immune system. Subsequently, it was further documented that similar aggregates could be induced with another non-enzymatic agent, free iron ions. In contrast to thrombin-generated fibers, those formed in the presence of iron ions exhibit dramatically different physicochemical properties. The first step in this conversion involves the interaction of trivalent iron ions (Fe³⁺) with the hydroxyl groups of water to form highly reactive hydroxyl radicals (HO⁻) according to the following reaction: HO⁻ + Fe³⁺ → HO⁻ + Fe²⁺.

Of note, our first observance of this reaction took place without the involvement of hydrogen peroxide required for the classic Fenton reaction. The hydroxyl...
radicals generated by the interaction of hydroxyl and iron cleaves intra-molecular disulfide bonds in the fibrinogen molecules causing their unfolding with the exposure of hydrophobic groups normally buried inside the tridimensional structure of the polypeptide chains. In the absence of specific chaperons the exposed hydrophobic epitopes form scrambled intermolecular linkages resulting in the formation of fibrin-like fibrils (parafibrin). The most important feature of parafibrin is its hydrophobicity and total resistance to proteolytic degradation. This unusual phenomenon is due to the fact that the hydrophobic forces holding together the polypeptide chains are purely physical and do not involve peptide bonds. Consequently, once attached to the surfaces of various cells, parafibrin induces a permanent state of inflammation by eliciting the release of cytokines and proteases from macrophages that impair their normal functions.

The damaging effect of parafibrin has recently been extended to the pathogenesis of cardiovascular and Alzheimer disease. These pathological conditions share a common mechanism by which native FBG molecules become hydrophobic by a single event of the conversion from the folded to the unfolded state. This phenomenon is very similar to that occurring when native prion protein PrP becomes converted to its toxic form PrPSc simply by unfolding, without any alteration in its primary structure. Interestingly enough, this concept has been met for many years with great skepticism from mainstream protein scientists, until its author was awarded with a Nobel Prize.

Iron and Cancer

Accumulating evidence suggests that a correlation exists between increased blood concentration of unbound iron and the incidence of cancer in humans. Furthermore, iron level reduction may prevent cancer morbidity and mortality. It should be noted that it is only trivalent iron (Fe³⁺) and not divalent (Fe²⁺) which participate in the generation of hydroxyl radicals and subsequent formation of insoluble parafibrin from soluble plasma fibrinogen. However, when hemoglobin is released from the hemolyzed erythrocytes, the divalent ferrous ions are enzymatically converted to ferric ions. Thus, any pathologic condition in which erythrocyte membranes are damaged, e.g., in response to infections and/or exposure to environmental toxins, may contribute to the excessive body storage of trivalent iron. It should be noted that this form of iron accumulates with age due to the fact that there is no mechanism for its physiologic elimination, and may therefore underlie prior observations of associations between cancer incidence and aging.
Current Issues of Cancer Immunotherapies

It is of extreme importance to remember that the presence of fibrin-like deposits, identified as thrombosis, has been reported for almost a century.20-22 However, no explanations have been offered why such deposits are not being removed by the fibrinolytic enzymes frequently found to be dramatically activated in tumors, particularly in prostate cancer. Apparently, there must be some post-translational modifications of fibrinogen/fibrin structure that render the tumor thrombi resistant to the proteolytic degradation. The same phenomenon may also be responsible for the disappointing results of chemotherapy for the most types of solid tumors. In the past decade, attempts have been made to develop alternative methods using adaptive and/or adoptive immunotherapies.1 It is of interest to note that while some the adoptive modalities work perfectly well in artificially contrived systems in vitro they lose their efficacies in the physiologically relevant environment in vivo. Obviously, the difference is in the exposure of tumor and immune cells to blood and its components as emphasized by Placke et al.1 As mentioned before another important hemostatic factor is plasma fibrinogen that, in the presence of free iron, is converted to parafibrin. We propose that this insoluble polymer forms a protective barrier around tumor cells by means of the hydrophobic interaction, similar to those of a detergent react with the non-polar regions of the fat with the exposure of hydrophobic groups and subsequently inhibit protein unfolding and scrambled refolding.20 Finally, it should be emphasized that the present concept of parafibrin formation and its role in tumor evasion would not be possible without the pioneering work on the protein structure-function relationships elucidated by the great American protein chemist, Chris Anfinsen.30

The Role of Polyphenols and Other Dietary Agents

In view of the known cancer protective effect of Mediterranean diet23,24 it is quite possible that certain polyphenolic substances present in this diet, such as EGCG,25 ferulic acid,26 and curcumin27 may exert their preventive and/or therapeutic effects by helping remove the protective barrier from the tumor membranes. The proposed mechanism of the action of polyphenols is based on their amphiphilic character that allows parafibrin to be displaced in a zipper-like manner (Fig. 2). This mechanism is similar to that operating in the removal of fat by detergents, in which the hydrophobic groups of a detergent react with the non-polar regions of the fat with the exposure of hydrophilic sites to the aqueous milieu. In addition to these organic substances suggested here to displace parafibrin from the tumor cell surfaces, there are two minerals that may also play an important role in this phenomenon. Thus, magnesium ions contained in green leaves and vegetables may exert their anticancer properties28 by inhibiting the intrinsic blood coagulation and in this way limiting the formation of parafibrin as indicated in the pathogenesis of Alzheimer disease29 The other essential, albeit not generally recognized mineral, is selenium that in a specific chemical form as sodium selenite blocks the protein thiol groups and subsequently inhibit protein unfolding and scrambling refolding. Finally, it should be emphasized that the present concept of parafibrin formation and its role in tumor evasion would not be possible without the pioneering work on the protein structure-function relationships elucidated by the great American protein chemist, Chris Anfinsen.30

Conclusions

Our perspective is that a singular electron transfer event from the redox iron ion into the hydroxyl group of water can initiate a chain of events that can explain the enigmatic phenomenon of immune evasion. This novel concept offers an additional scenario accounting for improper tumor immunoediting, as well as fostering a defective cellular immune system. However, a philosophical question arises whether such a simple idea can find its place in this day-and-age of the enormous complexity and technical sophistication of the current cancer research.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

References