Review Article

Accelerated Fibrinolysis and Its Propagation on Vascular Endothelial Cells by Secreted and Retained tPA

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Received 5 April 2012; Revised 11 July 2012; Accepted 20 July 2012

Abstract

We successfully visualized the secretory dynamics of tissue-type plasminogen activator (tPA) tagged by green fluorescent protein (tPA-GFP) from cultured vascular endothelial cells (VECs) using total internal reflection fluorescence (TIRF) microscopy and demonstrated that tPA-GFP secreted from VECs was retained on cell surfaces in a heavy-chain-dependent manner. Progressive binding of Alexa568-labeled Glu-plasminogen was also observed on the surface of active tPA-GFP expressing cells via lysine binding sites (LBS), which was not observed on inactive mutant tPA-GFP expressing cells. These results suggest that retained tPA on VECs effectively activated plasminogen to plasmin, which then facilitated the binding of additional plasminogen on the cell surface by proteolytically cleaving surface-associated proteins and exposing their C-terminal lysine residues. Thus prolonged retention of tPA appeared to play an important role in initiating and amplifying plasmin generation on VECs. LBS-dependent binding of plasminogen was also observed as a narrow band at the lytic front of the fibrin mesh formed on active tPA-GFP expressing cells, which expanded outward as the lytic area increased. This binding was not observed on inactive mutant tPA-GFP expressing cells or in the presence of aprotinin. The binding of plasminogen to partially digested fibrin appears to be indispensable for spontaneous fibrinolysis.

1. Introduction

Fibrinolysis takes place when undesirable fibrin is formed or when a hemorrhagic thrombus becomes unnecessary. Plasminogen activation by plasminogen activators (PAs) is the initial step in fibrinolysis, which is regulated by many physiological and pathological factors [1] so as to quickly dissolve unnecessary thrombi. Tissue-type PA (tPA) and urokinase-type PA (uPA) are the physiological PAs, and the former is synthesized and released from vascular endothelial cells (VECs) as an active form to initiate intravascular thrombolysis. Its characteristic of binding to fibrin is believed essential in the effective activation of plasminogen on the fibrin surface as well as in plasmin-catalyzed fibrin digestion [2]. The binding of plasminogen to fibrin through lysine-binding sites (LBS) existing in kringle domains is also believed to play an essential role in its effective activation on the fibrin surface [1]. In this paper, we introduce our recent findings on tPA secretory dynamics from VECs, and on the subsequent process of secreted tPA catalyzing plasminogen activation and fibrinolysis on vascular endothelial cells. Indispensiveness of the binding of plasminogen to partially digested fibrin in spontaneous fibrinolysis is also demonstrated.

2. Secretion of tPA from VECs and Regulation of Its Activity in Plasma

tPA is synthesized and released from VECs as an active form, either constitutively or in a regulated fashion following various kinds of stimuli from several different granules [5], and thus the amount of secreted tPA is a principal determinant of plasminogen activation potential in the vasculature. Another determinant of this potential is plasminogen activator inhibitor type 1 (PAI-1), the primary inhibitor of both tPA and uPA under physiological conditions. Since PAI-1 exists in plasma in molar excess over tPA, most of the tPA circulates in an inactive complexed form with PAI-1, and
Figure 1: Characteristics of tPA-GFP expressing EA hy926 cells. (a) Immunofluorescence images of tPA-GFP expressing EA hy926 cells were captured by either epifluorescence microscopy (epi-F) or total internal reflection fluorescence microscopy (TIRF), the latter of which more clearly detects tPA-GFP containing granules as dots. (b) Culture media of tPA-GFP expressing EA hy926 cells and control cells were analyzed by western blotting. Additional high-molecular-weight bands detected by both anti-tPA and anti-GFP were observed in tPA-GFP expressing EA hy926 cells (orange arrows), suggesting that tPA-GFP has activity to form high-molecular-weight complex with PAI-1. (c) Fibrin autography of the same culture media. The additional bands detected by western blot analysis showed a lytic band (orange arrow). This figure was originally published in [3].

only a small fraction of this total tPA circulates as an active enzyme [6].

3. tPA Secretory Dynamics

3.1. Slow Kinetics of tPA Secretion. The human vascular endothelial cell line EA.hy926 was transfected with an expression vector encoding tPA tagged by green fluorescent protein (tPA-GFP), and tPA secretory dynamics were analyzed using total internal reflection fluorescence (TIRF) microscopy (Figure 1(a)) [3]. The expressed tPA-GFP was confirmed to be active by the facts that it formed a high-molecular-weight complex with PAI-1 (Figure 1(b)) and that it developed a lytic band on a plasminogen-rich fibrin overlay (Figure 1(c)). The sizes and distribution of tPA-GFP containing granules were indistinguishable from those of intrinsic wild tPA, which was detected using an anti-tPA antibody. TIRF microscopy was used to visualize tPA-GFP-containing granules, which were detected as dots when they appeared in the so-called evanescent field, which is approximately 100 nm from the interface. The secretory dynamics of docking and opening of tPA-GFP containing granules followed by tPA-GFP secretion were appreciated by a sudden increase in fluorescent intensity followed by its gradual decrease. The most striking peculiarity in tPA secretory dynamics was that the release of tPA-GFP from the opened granule was very slow, which was shown by a slow decline in the fluorescent signal as well as by the persistence of an apparent fluorescent spot even 2-3 minutes after the opening of the granule. The mechanism for the long retention of tPA-GFP on membrane surfaces after granular opening was analyzed using several domain-deleted mutants of tPA-GFP. A mutant lacking the entire heavy chain and having only the catalytic domain disappeared quickly after granular opening, similar to the dynamics of a peptide hormone like insulin, suggesting that the prolonged retention of tPA requires the heavy chain.

4. Retention of tPA on the Cell Surface after Exocytosis and Its Release by PAI-1

Heavy-chain-dependent retention of tPA on the surface of VECs seems beneficial as it maintains an elevated plasminogen activation potential on the surface of the vasculature. The facts that in the culture medium of EA.hy926 cells tPA
exists mostly as tPA-PAI-1 complex and that free tPA rarely exists prompted us to analyze the potential effect of PAI-1 on tPA retention. The following three results suggested that PAI-1 facilitated the release of tPA from the surface of VECs. First, following suppression of PAI-1 expression in EA.hy926 cells by siRNA, the retention time of tPA-GFP was prolonged. Second, supplementation of recombinant PAI-1, in turn, shortened the retention time and increased the amount of tPA-GFP-PAI-1 complex but not of free tPA-GFP in the supernatant. Finally, a mutant tPA-GFP whose active site serine was substituted by alanine so as not to react with PAI-1 stayed on the cell surface for a longer time, and essentially no tPA-GFP was detected in the supernatant. Thus PAI-1 appeared to attenuate the fibrinolytic potential not only in plasma but also on the surface of VECs.

5. Fibrinolysis Initiated by tPA Secreted from VECs

tPA effectively activates plasminogen when it is bound to solid structures including fibrin [2], denatured proteins having a crossed beta-structure [7], and cell surfaces [2]. To achieve effective fibrinolysis, synchronized binding of plasminogen to these surfaces is necessary. When Alexa568-labeled Glu-plasminogen (568-Glu-plg) was incubated with EA.hy926 transfected with tPA-GFP, the binding of plasminogen on the cell surface was observed (orange arrow), which was suppressed when either heavy-chain-deleted tPA-GFP (tPA(CD)-GFP) or catalytically inactive tPA-GFP (tPA(S478A)-GFP) was employed. A fibrin network was formed on EA.hy926 transfected with tPA-GFP using Alexa647-labeled fibrinogen, and its spontaneous lysis was monitored by confocal microscopy. (c) Fibrinolysis initiated by tPA-GFP expressing cells, and its gradual expansion was clearly observed both focal planes of the bottom and 5 µm above the bottom. Linear binding of 568-Glu-plg was also observed at the lytic front (orange arrow), which expanded outward as the lytic area increased. A part of this figure was originally published in [4].

Figure 2: Accumulation of plasminogen and effective fibrinolysis on tPA-GFP-expressing EA hy926 cells. (a) Alexa568-labeled Glu-plasminogen (568-Glu-plg) was incubated with EA.hy926 transfected with tPA-GFP, and the binding of plasminogen on the cell surface was observed (orange arrow), which was suppressed when either heavy-chain-deleted tPA-GFP (tPA(CD)-GFP) or catalytically inactive tPA-GFP (tPA(S478A)-GFP) was employed. (b) A fibrin network was formed on EA.hy926 transfected with tPA-GFP using Alexa647-labeled fibrinogen, and its spontaneous lysis was monitored by confocal microscopy. (c) Fibrinolysis initiated by tPA-GFP expressing cells, and its gradual expansion was clearly observed both focal planes of the bottom and 5 µm above the bottom. Linear binding of 568-Glu-plg was also observed at the lytic front (orange arrow), which expanded outward as the lytic area increased. A part of this figure was originally published in [4].
front, which expanded outward as the lytic area increased. Presumably the 568-Glu-plg bound preferentially to newly exposed C-terminal lysine residues by plasmin-catalyzed partial digestion of fibrin fibers. As was shown before using many different methods [1] and was recently confirmed by analyzing the crystal structure of plasminogen [8], Glu-plasminogen seems to have undergone a conformational change from its closed conformation to its easily activatable open form, and this likely facilitates clot lysis. Thus, as has been expected for a long time, the binding of Glu-plg to fibrin that has been partially cleaved by plasmin clearly proved essential for spontaneous initiation of clot lysis.

6. Cell-Associated Fibrinolysis and Cell Function

Plasminogen activation and the associated fibrinolysis on cell surfaces are known to play a variety of functions under both physiological and pathological conditions [2, 9, 10]. Plasminogen binding-molecules play central role in these events, and several candidate molecules have been reported such as Plg-RKT, histone H2B, and annexin A2-S100A10 complex as well as alpha-enolase. The binding of plasminogen and/or its activation to plasmin on these molecules has been suggested to possibly transmit signals into the cells and to modify cellular functions [11, 12]. These specific plasminogen binding molecules, therefore, seem to play important roles in the initiation of effective plasmin generation, as well as in cell signaling, both of which are important in modifying cellular functions. Our findings, however, that the accumulation of plasminogen increased during the time course of tPA secretion from VECs suggests that other proteins also function as plasminogen binding molecules on the cell surface by providing newly exposed C-terminal lysine residues after plasmin-catalyzed cleavage (Figure 3). We believe that this augmentation of plasminogen binding by newly exposed C-terminal lysines as well as its effective activation also plays essential roles not only in the degradation of fibrin or matrix proteins, but also in protease-dependent signal transduction.

7. Summary

We have described a unique mechanism of tPA secretion and its role in effective fibrinolysis that takes place on VECs. This mechanism seems to require retention of tPA after secretion as well as the binding of plasminogen to specific molecules on the cell surface. Following the exposure of new C-terminal lysines on membrane proteins by plasmin-catalyzed cleavage, further accumulation of plasminogen on the cell surface was observed. Such initiation and augmentation of plasminogen activation seems to be important in modifying cell functions involved in a variety of physiological and pathological events. Understanding these mechanisms in greater detail will provide us with new strategies to control these events.
Acknowledgments

The research summarized here was originally published in Blood [3, 4], © the American Society of Hematology. This work was supported by a Grant-in-Aid for Scientific Research (C: 19590858) to Y. Suzuki and (C: 24590273) to T. Urano from the Japan Society for the Promotion of Science (JSPS), Japan Heart Foundation Research Grant, a grant from the Inamori Foundation to Y. Suzuki, and grants from the Smoking Research Foundation and SENSIN Medical Research Foundation to T. Urano.

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