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Microbial invasion or severe inflammatory or nutritional conditions may cause the activation of various serine proteases such as neutrophil proteases, trypsin, elastase, and plasmin, which may lead to the augmentation of fibrinolytic systems and subsequent inhibition of the fibrinolytic system in patients with sepsis [10]. Thus, the activation of coagulation and fibrinolytic systems is thought to be associated with the pathogenesis of sepsis. The role of fibrinolytic systems is not limited to the control of fibrin deposition, and they can also modulate the activation of inflammatory mediators and various components of the immune system. Recent data suggest that there is a tight connection between coagulation and fibrinolytic systems in sepsis [11]. There is a complex molecular-pathophysiological relationship between coagulation and fibrinolysis in sepsis. It is possible that these two physiological systems are cross-linked to achieve favorable clinical outcomes. There are 4 things that distinguish the disease-specific treatment from treatments that are used to treat any type of infection. First, of the disease-specific treatments are directed at the specific pathogen (that is, directed at a drug that targets a particular drug-resistant microorganism). Second, most disease-specific treatments are given by IV infusion instead of oral administration. Third, they are usually short courses of treatment-9 hours or less. Finally, and most importantly, disease-specific treatments are beneficial and do not adversely affect treatment of any other condition or disease. [click here to download my Cheat Sheet: What is a disease-specific treatment](#)

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