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Pentasaccharide Enhances the Inactivation of Factor Xa by Antithrombin through Promoting the Assembly of a Michaelis-type Intermediate Complex. Demonstration by Rapid Kinetic, Surface Plasmon Resonance and Competitive Binding Studies

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Abstract

It has been demonstrated that a unique pentasaccharide fragment of heparin (H5) activates AT by exposing an exosite on the serpin that is a recognition site for interaction with the basic autolysis loop (residues 143–154) of fXa. In support of this, the substitution of Arg-150 of fXa with Ala (R150A) impaired the reactivity of the mutant with AT by an order of magnitude specifically in the presence H5. To understand the mechanism by which heparin activation of AT improves the reactivity of the serpin with fXa, the H5-catalyzed reaction of AT with fXa, fXa R150A and fXa S195A was studied using rapid kinetic, surface plasmon resonance and competitive binding methods. The pseudo-first-order rate constants for the H5-catalyzed AT inhibition of both fXa and fXa R150A exhibited a linear dependence on the serpin concentration, thereby yielding second-order rate constants of $1.0 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$ and $1.5 \times 10^5 \text{ M}^{-1} \text{ s}^{-1}$, respectively. On the other hand, an ~70-saccharides long, high affinity heparin-catalyzed AT inhibition of both fXa derivatives showed a saturable dependence on the inhibitor concentration, yielding an identical rate constant of $\sim 20 \text{ s}^{-1}$, but different ternary fXa-heparin-AT dissociation constants ($K_{E,ATH}$) of ~130 nM and ~1780 nM for wild-type and R150A fXa, respectively. Competitive kinetic and surface plasmon resonance binding studies using the catalytically inactive S195A mutant of fXa yielded dissociation constants of 255 nM and 610 nM, respectively, for the mutant protease interaction with the AT-H5 complex. These results suggest that H5 enhances the reactivity of AT with fXa primarily through lowering the $K_{E,ATH}$ for the formation of a Michaelis-type serpin-protease encounter complex.

Antithrombin (AT) is the primary serpin inhibitor in plasma that regulates the activities of the trypsin-like serine proteases of both the intrinsic and the extrinsic blood coagulation cascade (1–5). Similar to other inhibitory serpins, AT inactivates its target proteases by binding to their active-sites through an exposed reactive center loop, and undergoing a conformational change which traps the enzymes in inactive, stable complexes (5,6). However, AT is a slow inhibitor of coagulation proteases unless it binds to heparin-like glycosaminoglycans, similar to those found on the endothelium surface (1,7). A full-length heparin accelerates the inhibition of coagulation proteases by AT by 3 to 4 orders of magnitude, thus the rate of protease inhibition by the serpin is only limited by diffusion (3,8). This is the basis for the extensive use of heparin for prophylaxis and treatment of venous thrombosis (9). In the case of fXa, the dramatic increase in the rate of protease inhibition by AT in the presence of heparin has been demonstrated to arise from 1) the ability of a unique pentasaccharide fragment of heparin to

change the conformation of the serpin to facilitate its optimal recognition by the protease (activation mechanism) (10,11), and 2) the ability of a full-length heparin to bridge the serpin and enzyme in one complex in the presence of physiological concentrations of Ca^{2+} , thereby promoting the initial interaction between the two proteins (template mechanism) (12,13). It has been demonstrated that the cofactor effect of a full-length heparin (~70 saccharides) accelerates the AT inhibition of fXa ~300-fold through the activation of serpin (10) and ~200–300-fold through a template mechanism in the presence of Ca^{2+} (13).

Previous rapid kinetic studies have indicated that AT inactivates fXa and other target coagulation proteases by a two step reaction mechanism in which an initial Michaelis-type enzyme-inhibitor complex, formed in the first reaction step, is converted to an irreversible covalent complex in the second step of the reaction (14,15). The dissociation constant ($K_{E,AT}$) for formation of the initial non-covalent serpin-protease complex is very high for the reaction of AT with fXa (and other coagulation proteases) in the absence of heparin (14,15). A full-length heparin consisting of ~70 saccharides (H70) has been demonstrated to lower the dissociation constant for formation of the noncovalent serpin-protease complex by ~4–5 orders of magnitude with both fXa and thrombin at physiologic Ca^{2+} , ionic strength and pH (13). In the case of thrombin, it is well established that the heparin bridging of both the serpin and the protease in one complex is solely responsible for rate enhancement of the polysaccharide, which lowers the dissociation constant for formation of a non-covalent AT-thrombin encounter complex (15). Although the 200–300-fold rate accelerating effect of H70 in catalyzing the AT inhibition of fXa is also mediated through the polysaccharide lowering the dissociation constant for the non-covalent serpin-protease encounter complex formation (13), the mechanism of the H5-mediated interaction of the activated AT with fXa is not known.

It has recently been demonstrated that Arg-150 of the autolysis loop of fXa (chymotrypsin numbering (16)) is a recognition site for specific interaction with the activated conformation of AT (17). The substitution of this residue with Ala has decreased the reactivity of the mutant protease with AT by an order of magnitude in the presence of H5 (17). The decrease in the reactivity of the fXa mutant with AT has been shown to be specific for the activated conformation of the serpin since the mutant protease reacted normally with the native conformation of the serpin (17). The important role of the autolysis loop of fXa in reaction with the activated conformation of AT has been confirmed by the observation that activated protein C and thrombin fusion mutants containing the autolysis loop of fXa both exhibited high reactivity with AT specifically in the presence of pentasaccharide (18). In the current study, rapid kinetic measurements of the H5 and H70-catalyzed reactions of AT with both wild-type and R150A fXa together with surface plasmon resonance and competitive binding methods have been employed to determine the mechanism by which activated AT effectively inhibits fXa.

Materials and Methods

Preparation of recombinant proteins and other reagents

Construction and expression of the Arg-150 to Ala (R150A) substitution mutant of fX was previously described (17). Gla-domainless fX (GD-fX) and a mutant in which the catalytic residue Ser-195 has been replaced with Ala (S195A) were constructed by PCR mutagenesis methods and expressed in human embryonic kidney cells 293 using the RSV-PL4 expression/purification vector system as described previously (19). The fX derivatives were activated by the factor X-activating enzyme from Russell's viper venom (Haematologic Technologies Inc., Essex Junction, VT) and purified on a SBTI-agarose column as described previously (20). Human plasma fXa and AT were purchased from Enzyme Research Laboratories (South Bend, IN). Active-site concentrations of fXa derivatives were determined by titrations with AT assuming a 1:1 stoichiometry (13). The therapeutic pentasaccharide fragment of heparins,

fondaparinux sodium (H5, MW=1.728 kD) was from Organon Sanofi-Synthelabo (France). The full-length high-affinity heparin containing ~70-saccharides (H70) isolated from commercial heparin by size and AT affinity fractionation, was a generous gift from Dr. Steven Olson from the University of Illinois-Chicago. Concentrations of heparins were based on AT binding sites and were determined by stoichiometric titrations of AT with the polysaccharides, with monitoring of the interaction by changes in protein fluorescence as described (21). The chromogenic substrate Spectrozyme FXa (SpFXa, MeO-CO-D-Chg-Gly-Arg-pNA) was purchased from American Diagnostica (Greenwich, CT). Polybrene and the fluorogenic substrate N-t-Boc-Ile-Glu-Gly-Arg-7-Amido-4-Methylcoumarin (Boc-IEGR-amc) were purchased from Sigma (St. Louis, MO).

Kinetic Methods

The rate of inactivation of fXa derivatives by AT in both the absence and presence of H5 and H70 was measured under pseudo-first-order conditions by a discontinuous assay method as described previously (13). In the absence of heparin, 1 nM fXa was incubated with 250–1000 nM AT in 0.1 M NaCl, 0.02 M Tris-HCl, pH=7.5, (TBS buffer, ionic strength = 0.12) containing 2.5 mM CaCl₂ (TBS/Ca²⁺), 0.1 mg/mL bovine serum albumin (BSA) and 0.1% polyethylene glycol (PEG) 8000. In the presence of H5, 0.5 nM of either wild-type or R150A fXa was incubated with 25–200 nM AT in the presence of excess H5 (1 μM) in the same buffer system. In the presence of H70, the experimental conditions were the same except that each protease was incubated with 200 nM AT and catalytic levels of heparin (1–10 nM). All reactions were carried out in 50 μL volumes in 96-well vinyl plates at room temperature. After a period of time (10 sec to 10 min depending on the rate of the reactions), 50 μL of the chromogenic substrate, (200 μM SpFXa), in TBS containing 0.1% PEG 8000 and 1 mg/mL Polybrene was added to each well and the remaining enzyme activity was measured with a V_{max} Kinetics Microplate Reader (Molecular Devices, Menlo Park, CA). The observed pseudo-first-order rate constants (*k*_{obs}) and the second-order association rate constants for uncatalyzed and catalyzed reactions were obtained as described previously (13). All values are the average of at least 3 independent measurements ±S.D.

Rapid kinetic analysis was carried out to resolve the two-step reaction of the AT-heparin complex with wild-type and R150A fXa in TBS/Ca²⁺ containing 0.1% PEG 8000 at 25 °C. In this case, the inactivation reaction was followed by a continuous assay in an Applied Photophysics SX.18MV-R stopped-flow instrument using the fluorogenic substrate, Boc-IEGR-amc as described previously (13). The excitation wavelength was set to 380 nm and fluorescence detected with a 420 nm emission cutoff filter. For the AT reactions in the presence of H70, AT concentrations ranged from 100 to 5,000 nM, H70 concentrations from 5 to 4,000 nM, fXa derivatives from 2 to 300 nM and Boc-IEGR-amc was present at 50 or 100 μM. For the reactions in the presence of H5, AT concentrations ranged from 200 to 12,000 nM, H5 from 10 to 10,000 nM, fXa derivatives from 2 to 300 nM, and substrate was present at 50 or 100 μM. Pseudo-first-order conditions were obtained by employing a greater than 10-fold molar excess of AT-heparin complex over the enzymes. To maintain relatively constant fluorescence amplitude in these reactions, the ratio of heparin to enzyme was kept constant. The *k*_{obs} values were determined by computer fitting of fluorescence progress curves to a single exponential with linear endpoint equation. The dependence of *k*_{obs} (the average of 5–10 progress curves) on the AT-H70 complex concentration was computer fit by the hyperbolic equation 1 (13),

$$k_{\text{obs}} = k_{\text{lim}} [\text{AT} - \text{heparin}] / K_{\text{E,ATH}} (1 + [S_0] / K_M) + [\text{AT} - \text{heparin}] \quad (1)$$

where *k*_{lim} represents the limiting rate constant for conversion of the intermediate AT-H70-fXa ternary encounter complex to a stable AT-fXa complex with release of heparin, [AT-heparin] is the concentration of the AT-heparin complex which was calculated based on the

dissociation constant for the AT-heparin interaction using the quadratic equation (fixed to the values of 5 and 23 nM for H70 and H5, respectively (13)), $K_{E,ATH}$ is the dissociation constant for the Michaelis-type fXa-AT-heparin ternary complex, $[S]_0$ is the concentration of the fluorogenic substrate, and K_M is the Michaelis constant for substrate hydrolysis by the enzymes. In both cases, K_M was fixed at measured value for fXa hydrolysis of Boc-IEGR-amc of 300 μ M in TBS buffer containing 2.5 mM Ca^{2+} as described (13). In the case of H5 where the dependence of k_{obs} on [AT-H5] was linear, the linear regression slope according to equation 2 provided the overall second-order rate constant ($k_{lim}/K_{E,ATH}$) after multiplying by the factor, $1 + [S]_0/K_M$, in order to correct for the competitive effect of the fluorogenic substrate.

$$k_{obs} = k_{lim} [AT - heparin] / K_{E,ATH} (1 + [S]_0 / K_M) \quad (2)$$

Competitive binding

The competitive effect of the catalytically inactive S195A mutant of GD-fXa on the wild-type GD-fXa inhibition by AT both in the absence and presence of H5 was studied using the same discontinuous assay described above. In this case, the inhibition of GD-fXa (1 nM) by AT (50 nM) was monitored in the presence of increasing concentrations of GD-fXa S195A (0–10 μ M) in the absence and presence of a fixed concentration of H5 (25 nM) in TBS/ Ca^{2+} containing 0.1 mg/mL BSA and 0.1% PEG 8000. Following 2–3 min incubation at room temperature, 50 μ L SpFXa was added to each well and the remaining activity of GD-fXa was measured as described above. Under these experimental conditions more than 90–95% of the enzyme activity in the absence of the competitor was inhibited. K_D for the non-covalent interaction of the activated conformation of AT with GD-fXa was determined from the nonlinear regression analysis of the saturable S195A concentration dependence for recovery of the enzyme activity monitored by the hydrolysis of SpFXa at 405 nm as described above.

Surface plasmon resonance (SPR)

The interaction of AT with GD-fXa S195A immobilized on a dextran-coated CM5 sensorchip was analyzed by SPR using the BIAcore X biosensor instrument (Uppsala, Sweden). The inactive fXa mutant was coupled to the CM5 sensorchip to obtain a resonance unit (RU) of ~500 using an amine coupling kit according to the manufacturer's instruction. The HBS buffer containing 10 mM HEPES (pH 7.4), 150 mM NaCl, 3.4 mM EDTA and 0.005% surfactant P20 was used as the running buffer. For analysis of the binding interactions, human AT was diluted to different concentrations of 2.5–1280 nM in the same buffer containing 4 μ M H5 and was injected into the flow cell 2, housing the immobilized S195A GD-fXa sensorchip. The binding sensogram was generated with a flow rate of 40 μ L/min with an injection time of 2 min. Under these conditions an RU value of ~150 was obtained at the highest concentration of the serpin. Control experiments included injection of 1280 nM AT alone or 4 μ M H5 alone in the same buffer for 2 min with the same flow rate. No regeneration step was necessary since the running buffer lacking H5 completely dissociated AT from the sensorchip as evidenced by RU returning to a zero baseline following the completion of sample injection. The flow cell 1 reference deducted sensogram data output obtained with different concentrations of the AT-H5 complex were analyzed using BIAevaluation software provided by the manufacturer. The global fitting of the binding data to a 1:1 Langmuir binding model yielded rate constants for association (k_a) and dissociation (k_d) and the equilibrium dissociation constant (K_D) for the non-covalent interaction of AT with the inactive fXa mutant in the presence of pentasaccharide.

Results and Discussion

AT inhibits fXa by a two-step reaction mechanism in which a non-covalent enzyme-inhibitor encounter complex, formed in the initial step, is converted to a stable covalent complex in the second step of the reaction (14). Rapid kinetic analysis was carried out to determine the step

which was affected in the reaction of R150A fXa with AT in the presence of pentasaccharide. Under the experimental conditions described under the legend of Fig. 1, the pseudo-first-order rate constant (k_{obs}) for inhibition of both wild-type and mutant fXa by the AT-H5 complex increased linearly, suggesting that the ternary dissociation constant ($K_{\text{E,ATH}}$) for forming the intermediate non-covalent complex is higher than 10 μM . In this case, only second-order rate constants could be calculated from the slope of the linear dependence of k_{obs} values on the AT-H5 complex concentration according to equation 2 (Table 1). Since a full-length ~70-saccharides heparin (H70) has been demonstrated to decrease $K_{\text{E,ATH}}$ of the fXa-AT interaction to ~100 nM in the presence of a physiological concentration of Ca^{2+} (13), rapid kinetic analysis was also carried using increasing concentrations of the AT-H70 complex in TBS containing 2.5 mM Ca^{2+} . In this case, a saturable dependence of k_{obs} on the AT-H70 complex concentration was observed for both wild-type and mutant fXa (Fig. 2), indicating the saturation of an intermediate heparin-AT-fXa encounter complex prior to formation of the stable serpin-protease complex. Non-linear regression analysis of data by the hyperbolic equation 1 yielded values of 130 ± 20 nM for $K_{\text{E,ATH}}$, and 19 ± 1 s^{-1} for the limiting rate constant (k_{lim}) for the stable complex formation for wild-type fXa and 1780 ± 260 nM for $K_{\text{E,ATH}}$, and 21 ± 2 s^{-1} for k_{lim} for the mutant protease (Table 1). These results indicate that the impairment in the reactivity of the fXa mutant with the H5-activated conformation of AT is localized to formation of the Michaelis-type non-covalent complex in the first reaction step. The calculated ratio of k_{lim} to $K_{\text{E,ATH}}$, corresponding to the overall second-order rate constant, agreed reasonably well with the directly determined value by a discontinuous assay, and was an order of magnitude lower for the mutant than for wild-type fXa (Table 1). Noting that the lower reactivity of the fXa mutant with AT is specific for the activated conformation of AT (k_2 values of 3×10^3 $\text{M}^{-1} \text{s}^{-1}$ and 2×10^3 $\text{M}^{-1} \text{s}^{-1}$ were obtained for wild-type and mutant fXa reactions with AT in the absence of heparin, respectively), these results indicate that the stimulating effect of H5 on AT inhibition of fXa is primarily due to the pentasaccharide lowering the dissociation constant for formation of the Michaelis-type non-covalent encounter complex.

Similar to its reaction with true substrates, the fXa catalytic residue, Ser-195, is required to form a covalent protease-serpin complex in the second step of the reaction. Thus, the inactive S195A mutant of fXa is expected to undergo only the first reaction step to form a reversible Michaelis-type complex with AT. To provide further support for the proposal that the primary role of pentasaccharide is to lower the dissociation constant for the formation of a non-covalent serpin-protease complex, the ability of the S195A mutant of GD-fXa to compete with wild-type GD-fXa for interaction with AT was evaluated in both the absence and presence of a fixed concentration of H5. As shown in Fig. 3, the inactive fXa mutant effectively competed with its wild-type counterpart for interaction with AT in the presence of H5. Under these experimental conditions, S195A enzyme did not influence the reactivity of wild-type enzyme in the absence of H5 (data not shown). Non-linear regression analysis of data yielded K_{D} of 255 ± 22 nM ($n = 6 \pm \text{S.D.}$) for the mutant fXa interaction with the activated conformation of the serpin. These results clearly indicate that pentasaccharide improves the dissociation constant for formation of the non-covalent complex of fXa with AT. The ability to measure K_{D} for interaction of the mutant fXa with the AT-H5 complex by this approach was in apparent contradiction to that by rapid kinetic approach in which no dissociation constant for the serpin-protease Michaelis-type complex could be measured for up to 10 μM of the AT-H5 complex (Fig. 1). To investigate this question further, we evaluated the direct binding of S195A GD-fXa to AT by SPR analysis in both the absence and presence of H5. In agreement with the competition kinetic data, H5 lowered the K_{D} for the non-covalent interaction of AT with the mutant protease. As shown in Fig. 4, AT interacted with the mutant protease with a K_{D} of 610 ± 70 nM in the presence of H5, whereas no interaction of AT with the protease was detected in the absence of pentasaccharide at the highest concentration of the serpin (Fig. 4, baseline). SPR analysis yielded association (k_{a}) and dissociation (k_{d}) rate constants of $(8.2 \pm 0.3) \times 10^4$ M^{-1}

s^{-1} and $0.05 \pm 0.004 s^{-1}$, respectively for the mutant fXa interaction with the AT-H5 complex. Similar results were obtained in both TBS and HBS buffers. The reason for an ~2-fold difference between K_D values as determined by either competition kinetic or SPR approach is not known. Nevertheless, the results clearly indicate that the activation mechanism of AT involves H5 lowering K_D for formation of a non-covalent serpin-protease encounter complex in the first step of the reaction.

The observation that the dissociation constant for interaction of the AT-H5 complex with the inactive S195A fXa could be determined by both competition kinetic and SPR approaches, but not for interaction with wild-type fXa using the rapid kinetic method may be explained by the Michaelis-type fXa-AT-H5 complex not establishing equilibrium at the initial binding step of the reaction. This kinetic behavior is characteristic for the reaction of enzymes with “sticky substrates” and occurs when the dissociation rate constant for the formation of a reversible Michaelis complex is slower than the rate constant for the subsequent formation of an acylated intermediate complex (22,23). Under these conditions, association of the enzyme with the substrate is diffusion-controlled and acylation is the rate-determining step in the reaction. This also appears to be the case for the reaction of fXa with AT-H5 since as discussed below the rate of covalent complex formation is much faster than the rate constant of the non-covalent complex dissociation (k_d). The resolvable rate constants for the reaction of fXa with the AT-heparin complex either by rapid kinetic or SPR method as determined in this study are presented in Scheme 1.

Although it was not possible to resolve the two-step reaction of the AT-H5 complex with fXa, it is expected that the rate constant for the stable covalent complex formation in this reaction to be higher than $19 s^{-1}$, the rate constant for fXa reaction with the AT-H70 complex. This contention is derived from the previous observation that in the absence of a template effect of heparin, fXa exhibits both high $K_{E,ATH}$ and k_{lim} values in reaction with the AT-heparin complex (14). Thus, previous rapid kinetic analysis has estimated a $K_{E,ATH}$ of ~200 μM for the non-covalent reaction step and a limiting rate constant of $140 s^{-1}$ for fXa reaction with AT in complex with a 26-saccharides high affinity heparin in the absence of Ca^{2+} (14). Calcium affects the reactivity of fXa with AT in complex with longer chain heparin by allowing the polysaccharide to bind to a basic exosite on fXa, thereby enhancing the reactivity of the protease with the serpin by a template mechanism (12,24). With the exception of its role in binding to the Gla-domain and allowing the longer chain heparins to interact with a basic exosite of fXa (24), Ca^{2+} is not known to influence any other step of the AT-fXa reaction. In support of this proposal, fXa exhibits similar reactivity with the AT-H5 complex in both Ca^{2+} and EDTA (13), and under both experimental conditions, the Gla-domainless fXa reacts with the AT-H70 complex with $K_{E,ATH}$ and k_{lim} values similar to those obtained for wild-type fXa in the presence of Ca^{2+} (13). Thus, the rate constant for the covalent reaction step of fXa with the AT-H5 complex should approximate to the same value ($\sim 140 s^{-1}$) reported for the reaction of fXa with AT in complex with 26-saccharides high affinity heparin in EDTA (14). In this case, the rate constant of covalent complex formation is 2800-fold higher than the dissociation rate constant of non-covalent complex formation ($k_d=0.05 s^{-1}$), as determined by SPR analysis. Because of such a large difference between reverse and forward rate constants, upon association of AT with the protease, the non-covalent complex is rapidly converted to a covalent complex without establishing equilibrium with free proteins. Under these conditions, the Michaelis-type complex [$K_{E,ATH} = (k_{lim} + k_d)/k_a = k_{lim}/k_a$] would not represent K_D and the association rate of fXa and AT-H5 would be diffusion-limited, with acylation being the rate-determining step in the reaction (25). This kinetic behavior can explain the inability of the rapid kinetic method to resolve the Michaelis constant for the non-covalent interaction of fXa with AT-H5 in the concentration ranges studied. On the other hand, in the case of fXa reaction with AT in complex with the longer chain heparin, the ability of heparin to simultaneously bind to both the serpin and the protease in the presence of Ca^{2+} dramatically lowers the dissociation constant for the

non-covalent step at the expense of limiting the rate constant for the covalent step of the reaction. It is not known how the bridging function of heparin, mediated by Ca^{2+} , limits the rate of covalent protease-serpin complex formation. Nevertheless, it has been speculated that the bridging function of heparin may impede the conformational freedom of the serpin's trapping mechanism at the covalent step of the reaction (25,26). This reaction mechanism is analogous to that observed for the reaction of tissue plasminogen activator with plasminogen activator inhibitor-1. In this case, the interaction of a basic insertion loop of the protease with two negatively charged residues on the primed side of the reactive site loop of the serpin limits the rate constant for the acylation step of the reaction (25,27).

In summary, the results presented above, for the first time, demonstrate that pentasaccharide activates AT by decreasing the dissociation constant for the formation of a non-covalent encounter complex between AT and fXa in the first-step of a two-step resolvable reaction. This was evidenced by the observation that 1) that the equilibrium dissociation constant for the ternary AT-heparin-fXa Michaelis-type complex formation was impaired by ~an order of magnitude for the R150A mutant of fXa, 2) the catalytically inactive S195A mutant of fXa effectively competed with wild-type fXa for binding to AT in the presence, but not in the absence of H5, and 3) the SPR analysis suggested that AT can bind to S195A fXa only in the presence of H5.

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Abbreviations

AT	antithrombin
fXa	active factor Xa
Gla	γ -carboxyglutamic acid
GD-fXa	Gla-domainless factor Xa, factor Xa from which the amino-terminal residues 1–45 have been removed
GD-fXa S1950A	Gla-domainless factor Xa derivative in which the catalytic residue Ser-195 (in the chymotrypsin numbering system of Bode et al. (16)) has been substituted with an Ala
fXa R150A	factor Xa mutant in which Arg-150 has been replaced with an Ala
BSA	bovine serum albumin
PEG	polyethylene glycol

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Figure 1.

Dependence of k_{obs} for wild-type and R150A fXa inhibition on the AT-H5 complex concentration. The k_{obs} values were determined from the exponential decrease in the rate of hydrolysis of the fluorogenic substrate, Boc-IEGR-amc, due to inhibition of the wild-type fXa ($\circ$) or R150A fXa ($\bullet$) by the increasing concentrations of the AT-H5 complex (x-axis) in TBS/ Ca^{2+} containing 0.1% PEG 8000. All reactions used a molar excess of AT over H5 to saturate the pentasaccharide, with exact concentrations of the AT-H5 complex calculated based on the known dissociation constant for the interaction. The solid lines are least squares computer fits of data by the linear equation 2 as described under "Materials and Methods".

Figure 2.

Dependence of k_{obs} for wild-type and R150A fXa inhibition on the AT-H70 complex concentration. The k_{obs} values for wild-type (○) and R150A fXa (●) inhibition by AT were determined for ~70-saccharide heparin-catalyzed reactions in TBS/ Ca^{2+} containing 0.1% PEG 8000 as in Fig. 1. The solid lines are least squares computer fits of data to the hyperbolic equation 1 as described under “Materials and Methods”.

Figure 3.

Competitive effect of inactive S195A GD-fXa on inhibition of wild-type GD-fXa by the AT-H5 complex. One nM GD-fXa was incubated with 50 nM AT in complex with 25 nM H5 in TBS/Ca²⁺ containing 0.1 mg/mL BSA and 0.1% PEG 8000 at room temperature for 2.5 min followed by monitoring the amidolytic activity of GD-fXa using Spectrozyme FXa as described under “Materials and Methods”.

Figure 4.

Biacore sensogram for binding of the AT-H5 complex to S195A mutant of GD-fXa. The fXa mutant was immobilized on CM5 sensorchip followed by the injection of different concentrations of AT (2.5–1280 nM) diluted in the running buffer (HBS) containing 4 μ M pentasaccharide as described under “Materials and Methods”. The rate constants for association (k_a) and dissociation (k_d) of the AT-H5 interaction with fXa was calculated from global fitting of the binding data (solid straight lines) to a 1:1 Langmuir binding model and presented in Scheme 1.

Scheme 1.

Table 1
Kinetic constants for the heparin-catalyzed reactions of AT with wild-type and R150A fXa as measured by continuous (stopped-flow) and discontinuous (end-point) assay methods.

	Heparin	$K_{E,ATH}$ (10^{-6} M)	k_{lim} (s^{-1})	k_2 ($10^6 M^{-1} s^{-1}$)
<i>^aStopped-flow</i>				
fXa	H70	130 ± 20	19 ± 1	146 ± 30
fXa R150A	H70	1780 ± 260	21 ± 2	12 ± 3
fXa	H5	ND	ND	0.7 ± 0.01
fXa R150A	H5	ND	ND	0.1 ± 0.001
<i>^bEnd-point</i>				
fXa	H70	ND	ND	110 ± 10
fXa R150A	H70	ND	ND	15 ± 1
fXa	H5	ND	ND	1.0 ± 0.1
fXa R150A	H5	ND	ND	0.15 ± 0.02

^a The kinetic constants were determined by rapid kinetic analyses of the observed pseudo-first-order rate constant (k_{obs}) for wild-type and R150A fXa inactivation by the AT-heparin complexes as a function of the AT-heparin complex concentrations in TBS buffer containing 0.1% PEG 8000 and 2.5 mM Ca^{2+} . AT-heparin complexes were formed by mixing full-length (H70) or pentasaccharide heparin (H5) with a molar excess of AT sufficient to saturate the polysaccharides. $K_{E,ATH}$ and k_{lim} values were obtained from computer fits of the saturable dependence of k_{obs} on the AT-heparin complex concentrations according to equation 1, after correction for the competitive effect of the reporter substrate as described under "Materials and Methods". Overall second order rate constants (k_2) were calculated from the ratio of $k_{lim}/K_{E,ATH}$ or were determined from the slope of the linear regression line fitted to plots of k_{obs} vs. AT-heparin complex concentration according to equation 2 after correction for substrate competition. ND, not determinable.

^b The heparin-catalyzed second-order association rate constants were determined from the residual chromogenic activities of fXa derivatives (0.5 nM) after incubation at room temperature with AT (25–200 nM) in complex with 1 μ M H5 for 30 sec in TBS buffer containing 0.1 mg/mL BSA, 0.1% PEG 8000 and 2.5 mM Ca^{2+} . The same methods were used to measure the k_2 values in the presence of H70 except that 0.5 nM enzyme was incubated with 1–10 nM heparin and 200 nM AT for 10–20 sec. The second-order inactivation rate constants were calculated as described under "Materials and Methods".