

## EXPERIMENTAL WORKS

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doi: <https://doi.org/10.15407/ubj92.03.033>THE FIBRIN B $\beta$ 125-135 SITE IS INVOLVED IN THE LATERAL ASSOCIATION OF PROTOFIBRILS

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Earlier we reported that during the human fibrinogen to fibrin transition a neoantigenic determinant was exposed in the B $\beta$ 119-133 fragment, where a hinge locus is situated. The fibrin-specific mAb FnI-3c and its Fab-fragment with epitope in this fragment inhibited the lateral association of protofibrils. We suggested that the epitope coincided with a site involved in this process. In this work we investigated the epitope location more precisely and defined a functional role for its exposure in the hinge locus of the molecule. It was found that mAb FnI-3c bound to human, horse and rabbit fibrins, all of which have Lys in the position corresponding to human B $\beta$ K130, but not to bovine and rat fibrins, which have other amino acid residues in this position, strongly suggesting that B $\beta$ K130 provides the integral part of the epitope. This fact, homology data, and structural biological analysis of the amino acid sequences around B $\beta$ K130 indicate that the site of interest is localized within B $\beta$ 125-135. The synthetic peptides B $\beta$ 121-138 and B $\beta$ 125-135, unlike their scrambled versions, bound to mAb FnI-3c in SPR analysis. Both peptides, but not their scrambled versions, inhibited the lateral association of protofibrils. The FnI-3c epitope is exposed after fibrinopeptide A cleavage and desA fibrin monomer formation. Structural biological analysis of the fibrinogen to fibrin transition showed a distinct increase of flexibility in the hinge locus. We propose that the structural transformation in the fibrin hinge regions leads to the conformation necessary for lateral association of protofibrils.

**Key words:** fibrinogen to fibrin transition, coiled-coil connector, protofibril lateral association, hinge region, neoantigenic determinant.

### Introduction

Fibrinogen is a dimer, with each subunit of the molecule being formed by three polypeptide chains: A $\alpha$ , B $\beta$  and  $\gamma$ . The molecule consists of a central E, two peripheral D, and two extended  $\alpha$ C regions [1]. The central E-region consisting of (A $\alpha$ 1-104, B $\beta$ 1-133,  $\gamma$ 1-72)<sub>2</sub> is connected to the two peripheral D-regions (A $\alpha$ 105-219, B $\beta$ 134-461,  $\gamma$ 73-411) by two long flexible coiled-coil connectors (A $\alpha$ 48-161, B $\beta$ 79-193,  $\gamma$ 23-135). The N-terminal parts of the connectors (A $\alpha$ 48-104, B $\beta$ 79-133,  $\gamma$ 23-62) belong to the E region and the C-terminal parts (A $\alpha$ 105-161,

B $\beta$ 134-193,  $\gamma$ 63-135) to the D regions. A hinge locus ( $\alpha$ 99-110,  $\beta$ 130-155,  $\gamma$ 70-100) is located in the middle part of the coiled-coil region [2, 3]. The extended  $\alpha$ C regions (A $\alpha$ 220-610) consist of unfolded flexible segments (A $\alpha$ 220-391) and more structured  $\alpha$ C-domains (A $\alpha$ 392-610).

Intermolecular binding of the fibrin polymerization sites A:a as well as putative C:c sites [4], leads to protofibril formation. The protofibrils associate laterally, initially forming fibrils and, subsequently, a three-dimensional fibrin net. At the stage of protofibrils and fibril formation, inter-protofibril binding

of the fibrin polymerization sites B:b increases the rate of protofibril lateral association. While the fibrin protofibril formation mechanism is relatively well-studied, the mechanism of their lateral association remains elusive. It has been suggested that the lateral association is realized by interactions of the DD-regions from neighboring protofibrils [5]. The likely sites of interacting regions were determined by Yang et al. [5] as  $\gamma$ 350-360 and  $\gamma$ 370-380.

Kollman et al. [6] have observed antiparallel contacts of fibrinogen coiled-coil regions in crystals of human fibrinogen. The authors proposed that fibrin molecules belonging to different protofibrils also interacted by their coiled-coil regions during the lateral association of protofibrils. It is notable that point mutations in the coiled-coil connectors of the fibrinogen molecule lead to impairment of fibrin polymerization, showing the functional significance of this region [7]. Earlier we obtained mAb FnI-3c, an antibody that reacted with human fibrin but not with fibrinogen or the D-dimer, testifying to a structural rearrangement in the epitope during the fibrinogen to fibrin transition. In a preliminary study, the epitope for this mAb in fibrinogen was localized in the coiled-coil region of the molecule B $\beta$ 119-133 because the mAb reacted with the fibrin  $\beta$ -chain, but not with the B $\beta$ 1-118 and B $\beta$ 134-461 fragments of human t-NDSK and D-dimer fragments, respectively. MAb FnI-3c and its corresponding Fab-fragment inhibited the lateral association of fibrin protofibrils. We suggested that this inhibition was the result of blocking the mAb epitope, which coincided with the site involved in the protofibril lateral association [8].

On the basis of primary structural analysis of fibrinogen chains, Doolittle [2] proposed the existence of a hinge region in the coiled-coil region of the fibrinogen molecule. Other authors considered the fragment of the coiled-coil connector consisting of A $\alpha$ 64-114, B $\beta$ 100-150, and  $\gamma$ 37-87 as a soft hinge about which the molecule adopts different conformations [9]. The crystal structure of human fibrinogen provided experimental evidence of the existence and

flexibility of hinge loci [6]. Kohler et al. [3] localized *in silico* the hinge locus in fibrinogen, although the model was lacking residues of  $\beta$ N1-57 and  $\alpha$ 201-562; they investigated the mechanism and the functional role of the hinge bending motions in fibrin polymerization, fibrinolysis and interaction with the cells. Their results define the hinge locus as including A $\alpha$ 99-110, B $\beta$ 130-155 and  $\gamma$ 70-100. Other authors assume that flexibility in the hinge allows the Y-ladder single-chain model of protofibrils to be realized, a provision not yet generally accepted.

Taking into consideration the data cited above, as well as our previous data on the inhibition of lateral association by fibrin-specific mAb FnI-3c, we investigated the localization of the mAb FnI-3c epitope at B $\beta$ 119-133, including the stage of fibrin polymerization when the epitope exposure takes place, the structure of the hinge itself, and, finally, the functional role of the structural rearrangement in the hinge locus of the coiled-coil region of the fibrin molecule.

### Materials and Methods

Tris[hydroxymethyl]aminomethan base, PMSF, aprotinin, HEPES, Tween-20, N-(dimethylamino-propyl)-N'-ethylcarbodiimide hydrochloride, 11-mercaptoundecanoic acid, 6-mercapto-1-hexanol, sodium chloride, urea, guanidine chloride, horse fibrinogen, and sheep anti-mouse IgG-HRP conjugate were obtained from Sigma-Aldrich (St. Louis, Missouri, USA). N-hydroxysuccinimide was obtained from ThermoFisher Scientific (Waltham, MA, USA).

*Preparation of fibrinogen, human fibrin, thrombin and plasmin.* Human, bovine, rat and rabbit fibrinogens and human desAB fibrin monomer were obtained as described previously [10, 11]. DesA fibrin monomer was prepared with Reptilase or thrombin [12]. Thrombin was obtained from blood plasma in accordance with published procedure [13]. Plasminogen was purified from human plasma using lysine-Sepharose [14]. Plasmin was prepared as previously described [15].

**Abbreviations:** ELISA – enzyme-linked immunosorbent assay; FpA – fibrinopeptide A; FpB – fibrinopeptide B; GPRP – Gly-Pro-Arg-Pro; HEPES – 2-[4-(2-hydroxyethyl)piperazin-1-yl]ethanesulfonic acid; HF – hydrogen fluoride; HPLC – high performance liquid chromatography; HRP – horseradish peroxidase; mAb – monoclonal antibody; MHBA – Matured Hop Bitter Acids; PAAG – Polyacrylamide Gel; PBS – 0.01 M potassium-phosphate buffer, pH 7.4 with 0.14 M NaCl; PC – principal component; PCA – principal component analysis; PMSF – phenylmethanesulphonyl fluoride; RMSF – root mean square fluctuation; SAS – solvent accessible surface; SDS – sodium dodecyl sulfate; SPPS – solid-phase peptide synthesis; SPR – surface plasmon resonance; t-NDSK – NH<sub>2</sub>-terminal disulphide knot of fibrin (A $\alpha$ 17-51, B $\beta$ 15-118,  $\gamma$ 1-78)<sub>2</sub>.

*Peptide synthesis.* Synthetic peptides B $\beta$ 121-138 and B $\beta$ 109-126 were synthesized by a solid-phase method (Fmoc chemistry) with 96% purity and identified by mass-spectrometry (MALDI) using the Bruker microflex LT (Billerica, Massachusetts, USA).

Scrambled versions of the peptides B $\beta$ 121-138 and B $\beta$ 125-135 were synthesized by a standard Boc-SPPS procedure on MBHA-modified polystyrol. The products were cleaved from the polymer by anhydrous HF, purified with HPLC and identified by mass-spectrometry (MALDI) using the Bruker microflex LT.

*Preparation and purification of mAb.* Hybridomas were obtained as described by Köhler and Milstein [16]. Human fibrin desAB in PBS with 2 M urea was used as an antigen. MAbs FnI-3c [8] was isolated from hybridoma culture medium by affinity chromatography on protein G-Sepharose 4B (Amersham, Uppsala, Sweden). The determination of immunoglobulin class and subclass was performed by ELISA using the Isotyping kit ("Clinical Credential; ICN Immunobiologicals", Lisle, IL, USA).

*Turbidity analysis of fibrin polymerization.* The effect of the mAb or its Fab-fragment on fibrin polymerization was studied spectro-photometrically at 350 nm [4, 17]. The curve of increasing turbidity during fibrin polymerization has the following parameters:  $\tau$  – the lag time representing the time of the protofibril formation;  $V_{\max}$  – the maximum rate of the fibrin polymerization defined as a tangent of the angle of the turbidity time-dependence at the point of maximum steepness; and  $h$  – the maximum turbidity of fibrin clots [4]. The polymerization of desAB fibrins was conducted at 0.1 mg/ml protein in 0.05 M ammonium acetate, pH 7.4, with 0.1 M NaCl and  $10^{-4}$  M CaCl<sub>2</sub>.

*SPR analysis.* The exposure of the neoantigenic determinant in fibrin in real time was detected by the SPR method using the Plasmon 6 device, developed in the VE Lashkarev Institute of Semiconductor Physics of the National Academy of Sciences of Ukraine (Kyiv, Ukraine). The gold surface of a chip was covered by the self-assembled layer formed of 11-mercaptoundecanoic acid and 6-mercapto-1-hexanol [18]. MAbs were immobilized covalently onto the functionalized chip using standard amino group coupling chemistry. The fibrin-specific mAb FnI-3c was put into the working cell and the fibrinogen-specific mAb FnII-4d into the control cell. To study an interaction between pep-

tides and mAb FnI-3c, each peptide was introduced into both cells at 0.5 mM concentration in 0.02 M HEPES buffer, pH 7.4, 0.15 M NaCl, and 0.005% Tween-20. The curves presenting the signals from working and control cells and their differences were recorded. To study an interaction *in situ* between fibrin and mAb FnI-3c, the latter was immobilized into working and control cells. Fibrinogen was introduced into the control cell at 1  $\mu$ g/ml concentration in 0.02 M HEPES buffer, pH 7.4, 0.3 M NaCl, and 0.005% Tween-20. In the working cell, fibrinogen was introduced in the same medium with the addition of 0.001 NIH units/ml of Reptilase. Each curve in the figures represents the typical curve for 2-3 experiments. In the SPR and ELISA experiments, we used human fibrinogen passed through a Sepacryl S-300 was obtained from Sigma-Aldrich (St.Louis, Missouri, USA) for removal of denaturated and aggregated protein molecules that appear in fibrinogen preparations after the thawing procedure.

*Electrophoresis and ELISA.* SDS electrophoresis in PAAG was performed in accordance with the Laemmli method [19]. The exposure of the neoantigenic determinant during human, horse, bovine, rabbit and rat fibrinogen transformation into fibrin was determined by ELISA. Fibrinogen was added at the concentration of 0.2 mg/ml to the medium, containing 0.02 M HEPES buffer, pH 7.4, and 0.3 M NaCl. The reaction was initiated by thrombin or Reptilase at 0.005 NIH units/ml. At a given time, the reaction was stopped by adding PMSF and aprotinin to final concentrations of 1 mM and 10  $\mu$ g/ml respectively. Aliquots were withdrawn for electrophoretic and ELISA analysis. Aliquots for ELISA were placed into microtiter plate wells. Neoantigenic determinant exposure was analyzed with mAb FnI-3c as the primary- and sheep anti-mouse IgG-HRP conjugate as the secondary antibodies.

*Lysis of fibrinogen and fibrin by plasmin.* Lysis of fibrinogen and fibrin by plasmin was performed as previously described at a protein to plasmin molar ratio of 1000:1. The plasmin activity was 15 caseinolytic units per mg of the protein [15].

*Bioinformatics analysis.* Data on fibrinogen mutations were collected with the help of the GEHT fibrinogen variants database [7] and additional literature searching. Homology models of the human fragments X of fibrinogen and fibrin were constructed using the crystallographic model, Protein Data Bank ID (PDB ID): 3GHG [6], as a template. Model 3GHG contains no coordinates for the unstructured

fragments B $\beta$ 1-57, A $\alpha$ 1-26, and A $\alpha$ 213-610 region and thus closely coincides with the structure of the fibrin(ogen) fragment X2 which lacks the residues B $\beta$ 1-53 and A $\alpha$ 221-610 [20]. The structure of the FpA (A $\alpha$ 1-16) and residues A $\alpha$ 17-26 were modeled *de novo* and optimized using Modeller9v10 software (Ben Webb, San Francisco, CA 94143, USA) [21-24]. The all-atom contacts model and the geometry evaluation were done with MolProbity (Duke, US) [25]. The dynamical properties of the fragments X of fibrinogen and fibrin were further analyzed by conformational analysis using the package CONCOORD 2.1 (Max Planck Institute for Biophysical Chemistry, Göttingen, Germany) [26]. Ensembles of 100 conformations were obtained for fibrinogen fragment X and fibrin fragment X. Principal component analysis (PCA) was performed using ProDy with NmWiz plugin [27] for VMD (University of Illinois at Urbana-Champaign, Urbana and Champaign, Illinois, USA) [28]. Solvent accessible surface (SAS) area calculations and the preparation of illustrations were carried out in PyMol (Schrodinger, New York, USA) [29]. For fast and sensitive detecting rigid blocks in large macromolecular complexes used method RigidFinder (Yale University, New Haven, Connecticut, USA) [30]. For finge prediction by network analysis of individual protein structures we used StoneHinge (Yale University, New Haven, Connecticut, USA) [31].

Statistical analysis was performed using standard Excel software (Microsoft, Redmond, USA). Mean values and their standard deviations were calculated.

## Results and Discussion

*Inhibition of protofibril lateral association by synthetic peptides.* In order to obtain direct evidence that the epitope of mAb FnI-3c is located in B $\beta$ 119-133 and coincides with the site involved in the lateral association of protofibrils, we synthesized three peptides corresponding to amino acid sequences B $\beta$ 109-126 (QTSSSSQFVMVLLKDLWQ), B $\beta$ 121-138 (LKDLWQKRQKQVKNENV) and a scrambled version of B $\beta$ 121-138 (DKWVQVELKKQKRNDLNQ).

The synthetic peptide B $\beta$ 121-138, as opposed to its scrambled version, reacted with mAb FnI-3c in SPR analyses (Fig. 1), confirming the localization of the mAb epitope in this fragment. The B $\beta$ 121-138 peptide also inhibited the lateral association of fibrin protofibrils. Based on the results of turbidity analysis

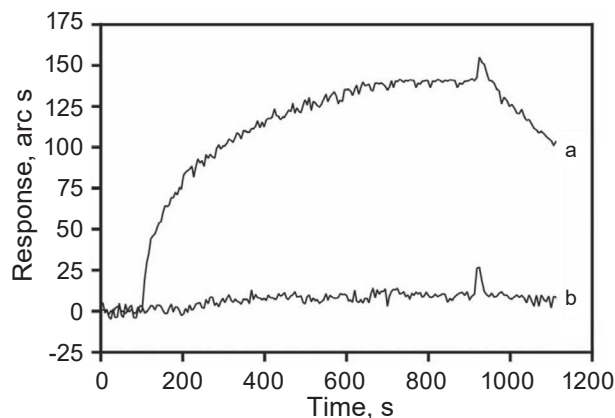


Fig. 1. Binding of synthetic peptide B $\beta$ 121-138 and its scrambled version to mAb FnI-3c as detected by the SPR method. Fibrin-specific mAb FnI-3c was immobilized into the working cell and fibrinogen-specific mAb FnII-4d into the control cell of a biosensor chamber. Each peptide was introduced into cells at 0.5 mM concentration in 0.02 M HEPES buffer, pH 7.4, 0.15 M NaCl, with 0.005% Tween-20. The curves of the difference between signals from working and control cells are presented: curve a denotes synthetic peptide B $\beta$ 121-138 and curve b its scrambled version. Each curve is an average of 2 repeats

(Fig. 2), we calculated the parameters of the polymerization process:  $V_{\max}$ ,  $\tau$  and  $\Delta h$ . The peptide at a concentration of  $2.5 \times 10^{-4}$  M caused a 6-fold decrease of the protofibril lateral association rate ( $V_{\max}$ ), increased the lag time ( $\tau$ ) by 2.5 times, and decreased the final clot turbidity ( $\Delta h$ ) by 1.4 times. In contrast, the scrambled versions of peptide B $\beta$ 121-138 (Fig. 2) and peptide B $\beta$ 109-126 (data not shown) did not inhibit fibrin polymerization, testifying to the specific inhibitory activity of peptide B $\beta$ 121-138.

The preferential inhibition of the lateral association of protofibrils by mAb FnI-3c, its Fab-fragment, and the peptide B $\beta$ 121-138 suggests this site is involved in the process of fibrin protofibril lateral association. Further localization of the FnI-3c epitope in the B $\beta$ 121-138 site was based on the observation that the peptide B $\beta$ 109-126 did not inhibit fibrin protofibril lateral association, indicating that residues B $\beta$ 127-138 must contain the epitope of mAb FnI-3c. To test this hypothesis, we studied the binding of FnI-3c mAb to fibrins of mammalian species that differ in sequence in this region: horse, rabbit, bovine, and rat. FnI-3c bound to human, horse and rabbit fibrins, but not to bovine and rat (Fig. 3, A). An alignment of the amino acid sequences shows



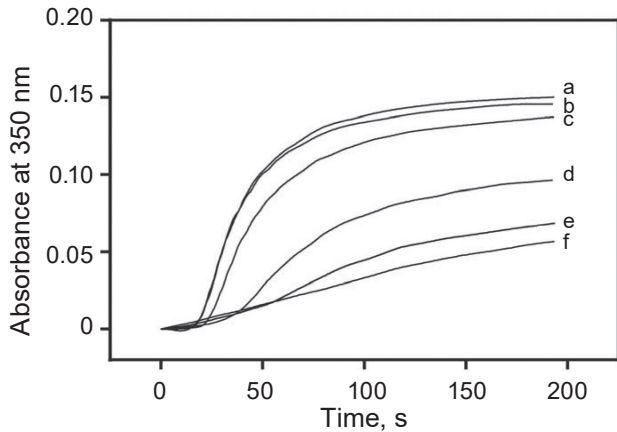


Fig. 2. Turbidity analysis of the inhibition of fibrin desAB polymerization by the synthetic peptide B $\beta$ 121-138 and its scrambled version. Curves: **a**, fibrin desAB polymerization (100  $\mu$ g/ml) without peptides, **b**, fibrin desAB polymerization with scrambled peptide (0.5 mM), fibrin desAB polymerization with **c**, 0.125 mM peptide, **d**, 0.25 mM peptide, **e**, 0.375 mM peptide, and **f**, 0.5 mM peptide. Each curve is an average of 2 repeats

that residue B $\beta$ K130 is always presents in the B $\beta$ 127-138 site in fibrins that bind to mAb FnI-3c (Fig. 3, B) but is substituted in bovine and rat fibrins by N and A, respectively, fibrins that do not bind to mAb FnI-3c (Fig. 3, A). This fact and the nature of the amino acid sequences around B $\beta$ K130 indicates that the site of interest is localized within B $\beta$ 125-138 of human fibrin.

Taking into consideration that B $\beta$ Lys130 is an integral part of the fibrin-specific mAb FnI-3c epitope, together with analysis of the crystallographic structure of this locus [6], we concluded that the epitope is formed by exposing residues B $\beta$ 125-135: B $\beta$  W125 Q126, K127, R128, K130, Q131, K133, D134, and N135 (Fig. 3, B, C) during the conversion of fibrinogen to fibrin. To verify this possibility, we synthesized an undecapeptide corresponding to fibrinogen B $\beta$ 125-135 (WQKRQKQVVDN) and analyzed its binding to mAb FnI-3c as well as its influence on fibrin polymerization. The peptide bound to mAb FnI-3c (Fig. 4), supporting the localization of the epitope in the B $\beta$ 125-135 site. The B $\beta$ 125-135 peptide at a concentration of  $2.5 \times 10^{-4}$  M caused a 2-fold decrease of the protofibril lateral association rate (Fig. 5). The scrambled version of B $\beta$ 125-135 did not bind to FnI-3c (Fig. 4) and did not inhibit fibrin polymerization (Fig. 5). The B $\beta$ 125-135 peptide did not affect the lag time ( $\tau$ ) and decreased the final

clot turbidity ( $\Delta h$ ) by 1.4 times. Probably, B $\beta$ 125-135 is the site where the structural rearrangement takes place during the transformation of fibrinogen to fibrin, which is to say the formation of the epitope of mAb FnI-3c occurs simultaneously with the lateral association of protofibrils.

*Structural rearrangements in B $\beta$ 125-135 of the monomeric fibrin desA.* The question arises, exactly which step of fibrin polymerization involves the structural rearrangement that forms the epitope in B $\beta$ 125-135? It seemed possible that the B $\beta$ 125-135 site is shielded by the  $\alpha$ C region in native fibrinogen, and that the B $\beta$ 125-135 site becomes exposed after dissociation of the  $\alpha$ C region from the bulk of the molecule. However, in ELISA experiments we did not observe FnI-3c epitope exposure during the hydrolysis of fibrinogen by plasmin, which in the initial step of digestion cleaves  $\alpha$ C regions from the molecule (Fig. 6, A, B). This indicates that the removal of the  $\alpha$ C region from the bulk of the molecule does not lead to the exposure of epitope during fibrinogen to fibrin transformation. In contrast, the results presented in Fig. 6C and 6D show that the slow cleavage of FpA from fibrinogen by small concentrations of thrombin (0.005 NIH units/ml) at the ionic strength 0.3 correlates with a gradual increase of FnI-3c binding to fibrin desA immobilized on wells of the microplate [11]. Tests with Reptilase produced the same results (data not shown). This clearly indicates that the epitope exposure correlates with FpA cleavage and the conversion of fibrinogen to fibrin desA.

Does the exposure of the epitope take place at the level of the fibrin monomer, or require fibrin oligomer formation when the interactions of the A knob and a hole are realized? To answer this question, we used a molecular system containing fibrinogen (5 mkg/ml) and Reptilase (0.001 NIH units/ml) in 0.02 M HEPES, pH 7.4, 0.3 M NaCl, 0.005% Tween-20, in the absence or presence of 1 mM GPRP peptide. The latter inhibits fibrin polymerization, keeping fibrin in solution. SPR analysis (Fig. 7) did not show any significant difference in the binding of fibrin desA formed *in situ* to mAb FnI-3c immobilized on the chip in the absence or presence of GPRP peptide. Hence, exposure of the epitope and the site involved in the lateral association of protofibrils must be in B $\beta$ 125-135 of the fibrin coiled-coil connector at the stage of the fibrin desA monomer formation.

*In silico comparison of conformational changes in the hinge loci of the fibrinogen and fibrin frag-*

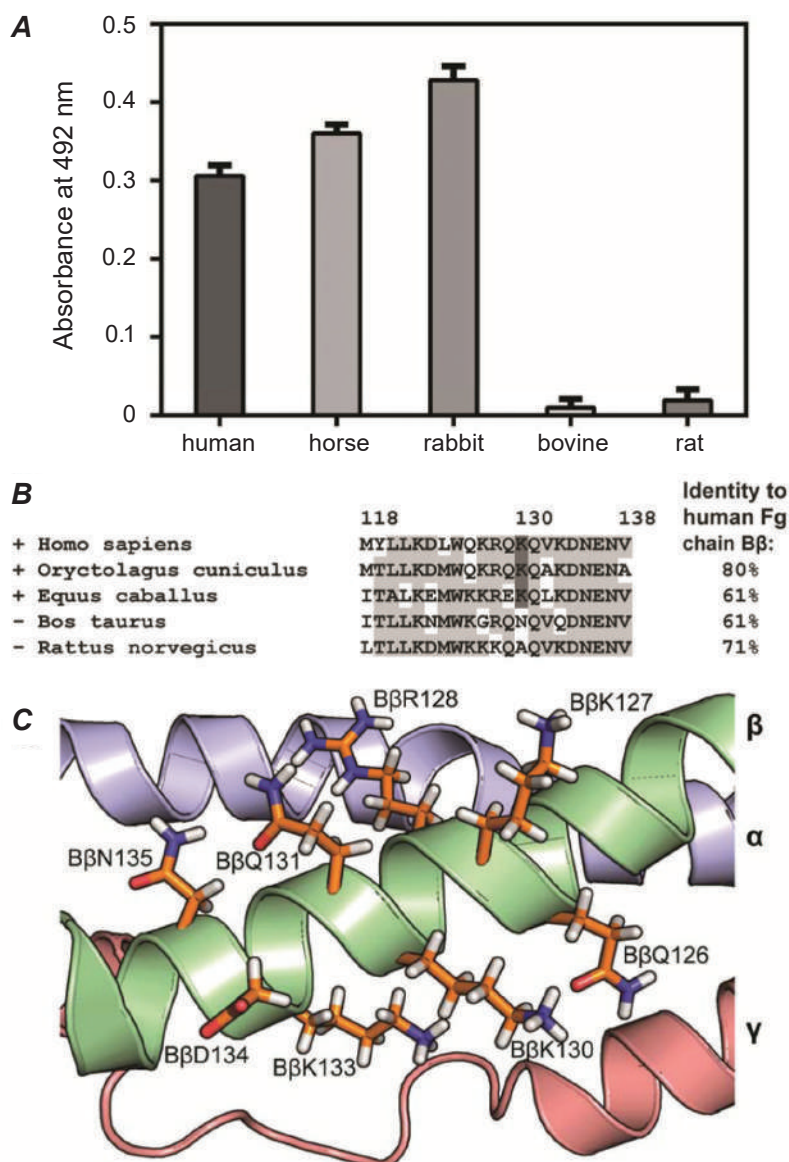


Fig. 3. Binding of mAb FnI-3c to human and animal fibrins as detected by ELISA. (A) Binding of mAb FnI-3c with human, horse, rabbit, bovine, and rat fibrins immobilized in microtiter plate wells. (B) Alignment of fragments of human, rabbit, horse, bovine, and rat fibrin(ogen) sequences homologous to the B $\beta$ 118-138 fragment of human fibrin(ogen). The percent identity with the human sequence is shown. Identical residues are shown with grey background. The sequences for fibrin(ogen)s binding and non-binding to mAb FnI-3c are marked with “+” or “-” respectively. (C) Cartoon of coiled-coil region corresponding to B $\beta$ 118-138 based on the crystallographic model of human fibrinogen (PDB ID: 3GHG). The side chains of B $\beta$  chain residues postulated to form the epitope are shown in stick form. Data represent 3 independent experiments. Error bars represent the standard deviations

ments X. An investigation of the mechanism of conformational changes in the region B $\beta$ 125-135 was performed by modeling human fibrinogen and fibrin fragments X. To this end we used the CONCOORD package to obtain ensembles of the two conformations (100 conformations each) and compared their

structural properties. We used PCA to specify the localization and the conformational properties of the hinge region, assuming that the hinge would be identified by an extremum on a plot of RMSF of C-alpha atoms. The RMSF of C-alpha atoms of the 20 slowest modes identified the principal com-

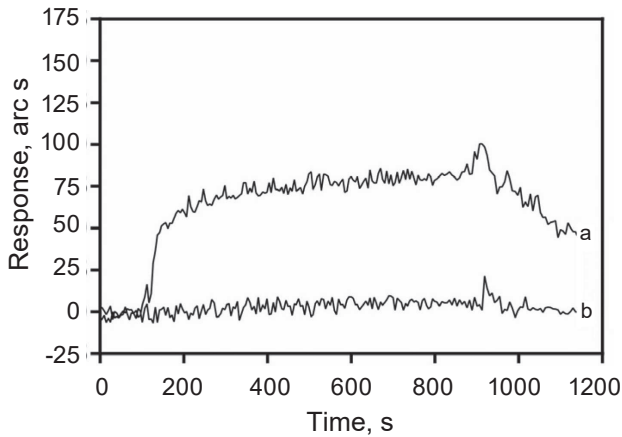


Fig. 4. Binding of synthetic peptides B $\beta$ 125-135 and its scrambled version to mAb FnI-3c as detected by the SPR method. Fibrin-specific MAb FnI-3c was immobilized into the working cell and fibrinogen-specific mAb FnII-4d into the control cell of a biosensor chamber. Each peptide was introduced into cells at 0.5 mM concentration in 0.02 M HEPES buffer, pH 7.4, 0.15 M NaCl, with 0.005% Tween-20. The curves of the difference between signals from working and control cells are presented: curve **a** denotes synthetic peptide B $\beta$ 125-135 and curve **b** its scrambled version. Each curve is typical for 3 similar experiments

ponent 7 (PC7) from the fibrinogen ensemble and PC6 from the fibrin ensemble as the slowest modes describing the hinge movement in the B $\beta$ 115-150 sector of the coiled-coil domain in one of the halves of the dimeric fibrin(ogen) molecule. Fig. 8, *A* shows that in these cases the hinge corresponds to a minimum on the RMSF of the C-alpha atoms coinciding with the residues B $\beta$ 125-135 in both fibrinogen and fibrin molecules. The correlation coefficient of the values of RMSF of the C-alpha atoms of the fragment B $\beta$ 115-150 for these two modes is 0.94. Our PCA data on the ensembles of the fibrin(ogen) conformations and the RigidFinder and StoneHinge web-server predictions allowed us to precisely identify the hinge region location in the fragment of the fibrin(ogen) coiled-coil connector comprising A $\alpha$ 91-103, B $\beta$ 125-135, and  $\gamma$ 69-77.

The RMSF data showed that the fluctuations of the C-alpha atoms on both sides of the hinge axis are higher for fibrin conformations compared to fibrinogen ones (Fig. 8, *A*), testifying to higher flexibility of the hinge region in fibrin compared to fibrinogen. This finding shows an essential increase of molecu-

lar flexibility in the coiled-coil connector in the fibrin molecule after its formation from fibrinogen.

The monoclonal antibody designated mAb FnI-3c reacts with fibrin in its monomer, oligomer and polymer forms, but not with fibrinogen [8]. This means that the transformation of fibrinogen to fibrin leads to the exposure of a stable neoantigenic determinant that retains its structure during the polymerization process. The structural rearrangement leading to neoantigenic determinant formation was found to involve residues B $\beta$ 125-135, a region thought to contain a hinge locus for the fibrin molecule.

To identify the possible mechanisms of the neoantigenic determinant exposure during the fibrinogen to fibrin transformation, we performed a comparative conformational analysis of the region B $\beta$ 125-135 in models of the fragments X of fibrinogen and fibrin. The per-residue SAS area of the fragment B $\beta$ 125-135 was analyzed and models with the lowest and highest areas were selected from ensembles of fibrinogen and fibrin fragments X, respectively (Fig. 8, *B, C, D*). This structural analysis showed (Table) that the amino acid residues B $\beta$  D134 and N135 in the  $\alpha$ -helix fragment B $\beta$ 125-135 are shielded by a nearby part of the  $\alpha$ -helix formed by B $\beta$  N137, V138, V139 in fibrinogen, and that they became

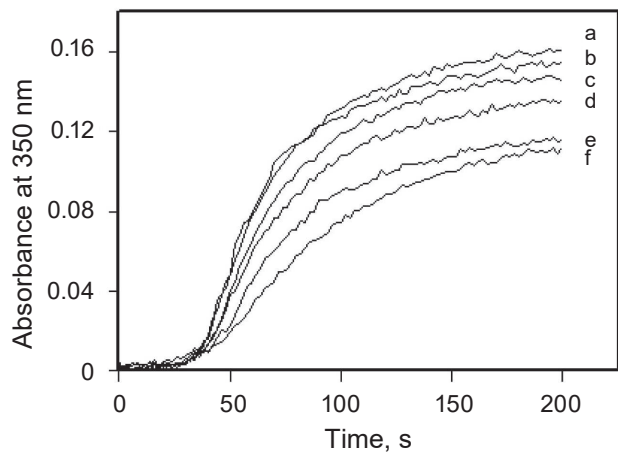


Fig. 5. Turbidity analysis of inhibition of fibrin desAB polymerization by the synthetic peptide B $\beta$ 125-135 and its scrambled version. Curves: **a**, fibrin desAB polymerization (100  $\mu$ g/ml) without peptides, **b**, fibrin desAB polymerization with scrambled peptide (0.5 mM), and fibrin desAB polymerization with **c**, 0.125 mM peptide, **d**, 0.25 mM peptide, **e**, 0.375 mM peptide, and **f**, 0.5 mM peptide. Each curve is an average of 2 repeats

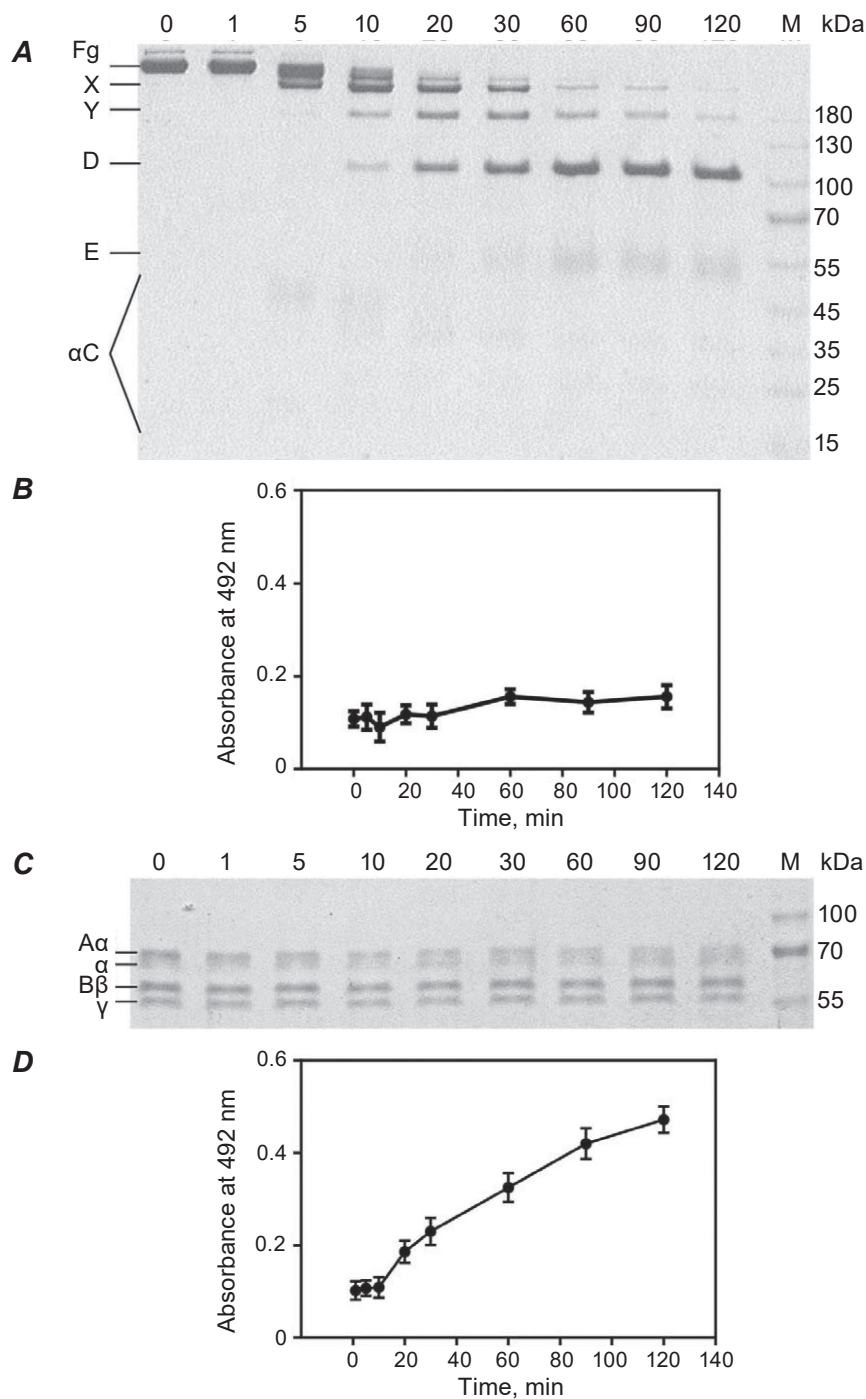


Fig. 6. Exposure of mAb FnI-3c epitope during hydrolysis of fibrinogen by plasmin and thrombin. Fibrinogen was added at the concentration of 0.2 mg/ml to the medium, containing 0.02 M HEPES buffer pH 7.4, 0.3 M NaCl, and the reaction was initiated by plasmin or thrombin/Reptilase at 0.005 NIH/ml. At a given time, the reaction was stopped by adding PMSF and aprotinin to final concentrations of 1 mM and 10 mkg/ml, respectively. Aliquots were withdrawn for electrophoretic and ELISA analysis. Aliquots for ELISA were added to microtitre plate wells. Neoantigenic determinant exposure was analyzed with mAb FnI-3c as the primary- and sheep anti-mouse IgG-HRP conjugate as the secondary antibodies. Samples of mediums of the (A, B) fibrinogen-plasmin and (C, D) fibrinogen-thrombin/Reptilase reactions were subjected to electrophoresis (A, C) and ELISA (B, D). Data represent 3 independent experiments. Error bars represent the standard deviations of the mean values



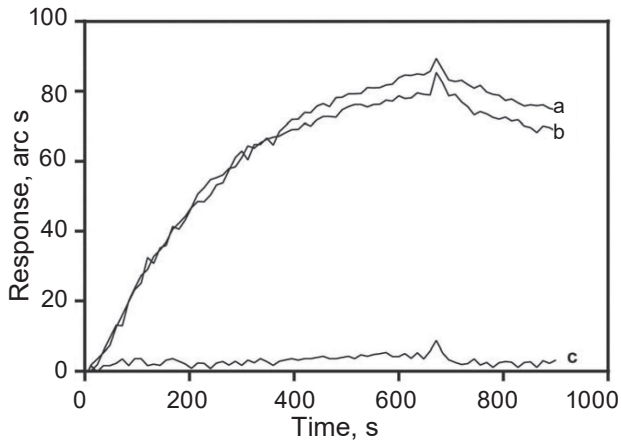


Fig. 7. Binding of fibrin desA formed *in situ* to mAb FnI-3c in the presence and absence of the GPRP peptide as detected by the SPR method. Fibrin-specific mAb FnI-3c was immobilized into working and control cells of a biosensor chamber. Fibrinogen was added to the control cell at 1  $\mu\text{g}/\text{ml}$  concentration in 0.02 M HEPES buffer, pH 7.4, 0.3 M NaCl, with 0.005% of Tween-20. In the working cell, fibrinogen was added to the same medium with the addition of 0.001 NIH units/ml of Reptilase in the presence or absence of 1 mM GPRP peptide. The interaction between fibrin desA, formed *in situ*, and the mAb immobilized on the chip was registered by the SPR method. The curves of the difference between signals from working and control cells are presented. Curves: **a**, fibrinogen + Reptilase; **b**, fibrinogen + Reptilase + GPRP; and **c**, fibrinogen as a control. Each curve represent an average of 3 repeats

more exposed in fibrin. During the fibrinogen to fibrin transformation, the surface of the B $\beta$ 125-135 site flattens due to the emergence of the  $\alpha$ -helical fragment partially buried in fibrinogen (Fig. 8, B, C, D).

The mAb FnI-3c and its Fab-fragment inhibit the lateral association of protofibrils in the process of fibrin polymerization [8]. We showed that synthetic peptides imitating the amino acid sequence of fibrin/fibrinogen fragments B $\beta$ 121-138 and B $\beta$ 125-135 also inhibit this stage of fibrin polymerization. Initially, we supposed that the fragment B $\beta$ 125-135 was the contact site of the inter-protofibril lateral binding. However, several facts now indicate that this site of the coiled-coil adopts a structure important for the subsequent association of fibrin protofibrils, and that the epitope exposure in B $\beta$ 125-135 reflects this structural rearrangement. In this regard, there are known mutations in the  $\alpha$ -,  $\beta$ -, and  $\gamma$ -constituents

Table. Per-residue surface area of the exposed residues of B $\beta$ 126-139 fragment of fibrinogen ( $S_{\text{Fg}}$ ) and fibrin ( $S_{\text{Fn}}$ ). The surface area changes ( $S_{\text{Fn}} - S_{\text{Fg}}$ ) in the models obtained by conformational analysis were calculated

Residue	$S_{\text{Fg}}$ ( $\text{\AA}^2$ )	$S_{\text{Fn}}$ ( $\text{\AA}^2$ )	$\Delta S (S_{\text{Fn}} - S_{\text{Fg}})$ ( $\text{\AA}^2$ )
B $\beta$ Q126	107.1	108.0	0.9
B $\beta$ K127	117.1	107.3	-9.7
B $\beta$ R128	141.8	144.9	3.1
B $\beta$ K130	117.7	119.1	1.4
B $\beta$ Q131	109.8	111.4	1.6
B $\beta$ K133	118.8	115.7	-3.1
B $\beta$ D134	88.4	95.0	6.6
B $\beta$ N135	87.6	98.4	10.8
B $\beta$ N137	99.8	104.2	4.4
B $\beta$ V138	89.8	95.1	5.3
B $\beta$ V139	91.9	97.0	5.1

of the coiled-coil containing B $\beta$ 111-141 that impair fibrin polymerization [31-36]. Moreover, our structural biological analysis of the B $\beta$ 125-135 site showed that conformational rearrangements take place in the hinge region during the fibrinogen-fibrin transition with the accompanying formation of the neoantigenic determinant which is preserved in the subsequent stages of fibrin polymerization.

We propose that at the start of protofibril lateral association, a specific bend and/or twist in two coiled-coil hinge regions of the dimeric fibrin molecule is required. This structural rearrangement provides a necessary mutual spatial orientation of the other regions of the molecule, which come into direct inter-protofibril contact during protofibril lateral association [6]. Our data show that the cleavage of Fp A during the fibrinogen-fibrin conversion induces the intramolecular structural rearrangement within the monomeric molecule of fibrin desA, and increases the flexibility in hinge regions. These structural changes are necessary for the subsequent protofibril association stage.

The augmentation of the flexibility in the hinge fragments of fibrin causes the increase of bending and twisting motions in the coiled-coil connector as compared to fibrinogen. Any factor affecting the flexibility of fibrin molecules in the hinge region will impair protofibril lateral association. The binding of the mAb FnI-3c or Fab-fragment to their epitope in

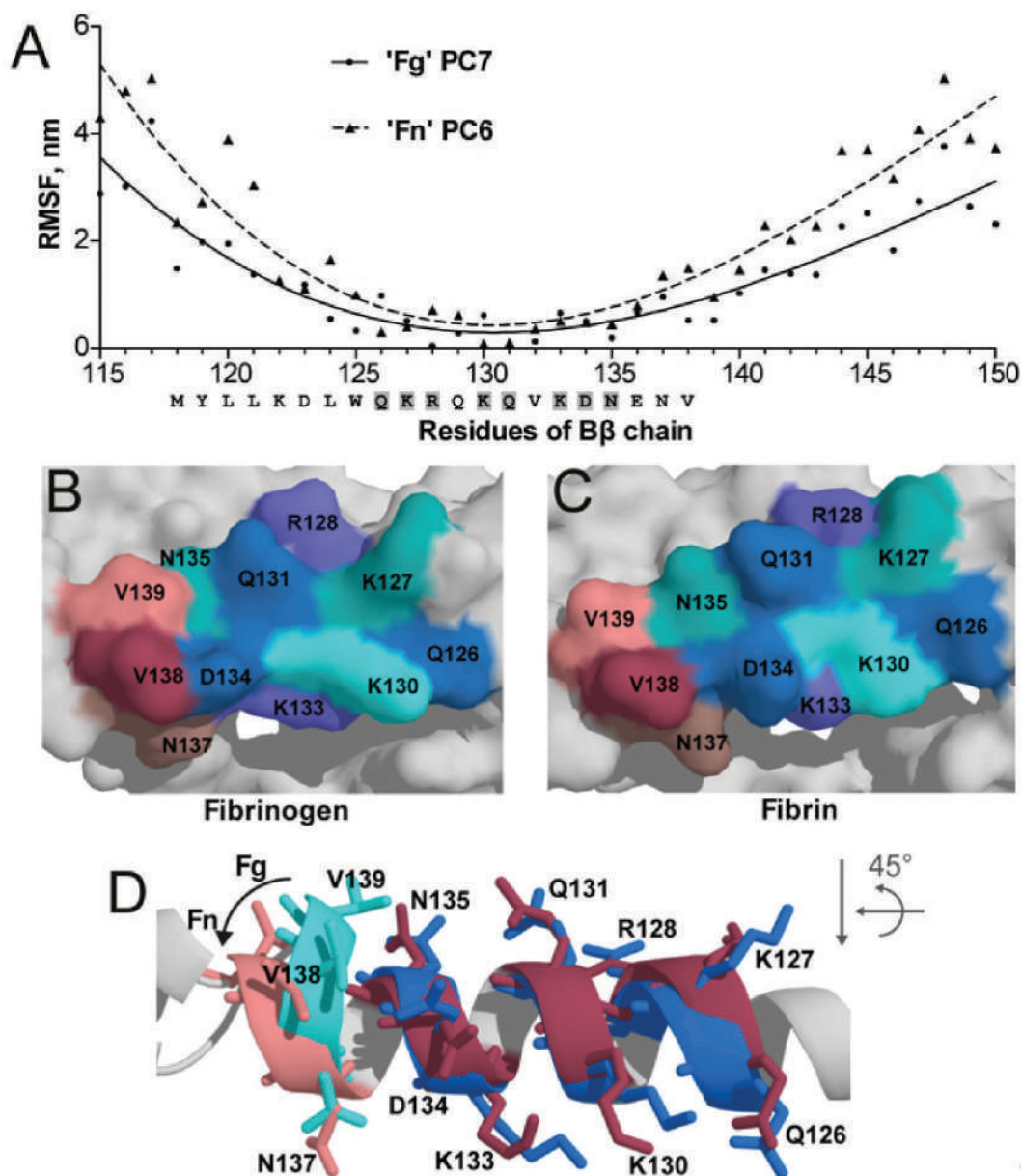


Fig. 8. Comparative RMSF analysis of conformational changes into hinge loci of fibrinogen and fibrin fragments X. (A) RMSF of the B $\beta$ 115-150 fragment C-alpha atoms for the principal component 7 for fibrinogen ('Fg' PC7, ●) and for the principal component 6 for fibrin ('Fn' PC6, ▲) ensembles of the conformations. Solid and dashed lines represent cubic polynomial approximations of the RMSF data for fibrinogen and fibrin, respectively. The coefficient of determination ( $R^2$ ) is 0.87 and 0.81 for fibrinogen and fibrin, respectively. The amino acid residues of fragment B $\beta$ 118-138 are denoted. The residues postulated to form the epitope for fibrin-specific mAb FnI-3c are marked with grey boxes. (B, C) Surface formed by the amino acid residues of fragment B $\beta$ 126-139 in models of (B) fibrinogen and (C) fibrin obtained by conformational analysis. (D) Structurally superimposed alpha-helical fragments B $\beta$ 126-139 from the selected models of fibrinogen and fibrin obtained by conformational analysis. The view in (D) is rotated by 45° around the long axis of the molecule compared to the views in (B) and (C). The residues B $\beta$  Q126, K127, R128, K130, Q131, K133, D134, and N135 of fibrinogen and fibrin are assumed to form the epitope for mAb FnI-3c. Residues B $\beta$  N137, V138, and V139 shield the residues B $\beta$  D134 and N135 in fibrinogen. The latter become more exposed in fibrin

the hinge region inhibited the polymerization process. The impairment of fibrin polymerization by the synthetic peptides B $\beta$ 121-138 and B $\beta$ 125-135 may be caused by decreased flexibility at the hinge loci of separate fibrin molecules in forming protofibrils. These peptides may bind with the corresponding  $\alpha$ - or  $\gamma$ -remnants of the hinge region in the fibrin coiled-coil and block the bending motions. This mechanism may be similar to the self-assembly of the coiled-coil from peptides described by Jing et al. for the synthetic peptides imitating the human fibrinogen site  $\gamma$ 23-57 and its modified form [37].

### Conclusions

Cleavage of fibrinopeptide A exposes a stable neoantigenic determinant in the fibrin desA B $\beta$ 125-135 site. B $\beta$ 125-135 is a constituent of the fibrin hinge consisting of A $\alpha$ 91-103, B $\beta$ 125-135, and  $\gamma$ 69-77. The flexibility of the hinge is increased during the fibrinogen to fibrin transformation. We assume the increase in flexibility in the fibrin molecule around the hinges allows the structural reorganization of the fibrin desA molecule during the self-assembly of protofibrils and their lateral association into fibrils.

*Conflict of interest.* Authors have completed the Unified Conflicts of Interest form at [http://ukrbiochemjournal.org/wp-content/uploads/2018/12/coi\\_disclosure.pdf](http://ukrbiochemjournal.org/wp-content/uploads/2018/12/coi_disclosure.pdf) and declare no conflict of interest.

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### В $\beta$ 125-135 ДІЛЯНКА МОЛЕКУЛИ ФІБРИНУ БЕРЕ УЧАСТЬ У ЛАТЕРАЛЬНІЙ АСОЦІАЦІЇ ПРОТОФІБРИЛ

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Раніше ми показали, що в процесі перетворення фібриногену людини у фібрин на фрагменті В $\beta$ 119-133, який розташований у шарнірній ділянці молекули, експонується неантигенна детермінанта. Фібринспецифічне mAb FnI-3c та його Fab-фрагмент, епітоп для яких знаходиться в цьому фрагменті, гальмують латеральну асоціацію протофібрил. Ми припустили, що епітоп співпадає із сайтом, який бере участь у цьому процесі. У цій роботі ми більш точно визначили розташування епітопу для mAb FnI-3c у молекулі фібрину та встановили його роль у функціонуванні шарнірного локусу молекули. Було виявлено, що mAb FnI-3c зв’язується з фібрином людини, коня та кроля, які мають лізин у положенні, що відповідає такому В $\beta$ K130 фібрину людини, але відсутній у фібрині бика та щура, у яких в цьому положенні є інші амінокислотні залишки, що вказує на те, що залишок В $\beta$ K130 є невід’ємною складовою епітопу. Цей факт, дані гомології та структурно-біологічний аналіз амінокислотних послідовностей навколо В $\beta$ K130 свідчать про те, що досліджуваний сайт локалізований у межах В $\beta$ 125-135 молекули фібрину. Синтетичні пептиди В $\beta$ 125-135 та В $\beta$ 121-138, на відміну від їхніх неупорядкованих аналогів, зв’язуються з FnI-3c в ППР аналізі. Обидва пептиди на відміну від їхніх неупорядкованих версій, інгібували латеральну асоціацію протофібрил. Епітоп



mAb FnI-3c експонується після відщеплення фібринопептиду А та утворення мономеру desA фібрину. Структурний біологічний аналіз переходу фібриногену у фібрин показав чітке підвищення рухливості в шарнірному локусі. Ми вважаємо, що така структурна перебудова в шарнірних областях фібрину спричинює формування конформації молекули, яка необхідна для латеральної асоціації протофібрил фібрину.

**Ключові слова:** трансформація фібриногену у фібрин, суперспіралізований конектор, латеральна асоціація протофібрил, шарнірна ділянка, неоантигенна детермінанта.

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