



Effects of Post-Translational Modifications of Fibrinogen on Clot Formation, Clot Structure, and Fibrinolysis

A Systematic Review

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OBJECTIVE: Post-translational modifications of fibrinogen influence the occurrence and progression of thrombotic diseases. In this systematic review, we assessed the current literature on post-translational modifications of fibrinogen and their effects on fibrin formation and clot characteristics.

APPROACH AND RESULTS: A systematic search of Medline, Embase, Cochrane Library, and Web of Science was performed to find studies reporting post-translational modifications of fibrinogen and the effects on clot formation and structure. Both in vitro studies and ex vivo studies using patient material were included. One hundred five articles were included, describing 11 different modifications of fibrinogen. For the best known and studied modifications, conclusions could be drawn about their effect on clot formation and structure. Oxidation, high levels of nitration, and glycosylation inhibit the rate of polymerization, resulting in dense clots with thinner fibers, while low levels of nitration increase the rate of polymerization. Glycation showed different results for polymerization, but fibrinolysis was found to be decreased, as a consequence of increased density and decreased permeability of clots. Acetylation also decreases the rate of polymerization but results in increased fiber diameters and susceptibility to fibrinolysis. Other modifications were studied less or contrasting results were found. Therefore, substantial gaps in the knowledge about the effect of post-translational modifications remain.

CONCLUSIONS: Overall, post-translational modifications do affect clot formation and characteristics. More studies need to be performed to reveal the effects of all post-translational modifications and the effects on thrombotic diseases. Expanding the knowledge about modifications of fibrinogen can ultimately contribute to optimizing treatments for thrombotic diseases.

VISUAL OVERVIEW: An online [visual overview](#) is available for this article.

Key Words: fibrin ■ fibrinogen ■ fibrinolysis ■ polymerization ■ systematic review

The architecture and properties of a thrombus are an important determinant of the disease burden and mortality associated with cardiovascular diseases. One of the main determinants of thrombus characteristics is variations in the fibrinogen molecule, which affect the clotting rate, architecture of the fibrin matrix and its susceptibility to fibrinolysis. Therefore, it is essential to know the effect of variability in the fibrinogen molecule on the

coagulation cascade and the clot characteristics. Fibrinogen is a glycoprotein that is synthesized by hepatocytes and circulates in the blood of healthy individuals at concentrations between 2 and 4 g/L. Fibrinogen consists of 2 sets of 3 different polypeptide chains: 2-A α , 2-B β , and 2- γ chains, which are held together by 29 disulfide bridges.¹ The genes encoding for these chains (*FGA*, *FGB*, and *FGG*) are found in a cluster of 65 kilobases on

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Nonstandard Abbreviations and Acronyms

MI	myocardial infarction
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the human chromosome 4 (4q23-q32).² In the fibrinogen molecule, different main structural regions can be identified.³ The central E region contains the amino acid termini of the polypeptide chains and this is also the place where thrombin cleaves. The 2 distal nodules (D regions) contain the carboxyl termini of the B β and γ chains and are connected to the E region by 2 α -helical coiled-coil domains. The carboxyl termini of the A α chains form the α C-regions, which are more flexible and mobile.⁴

The final step of the coagulation cascade is the cleavage by thrombin between arginine and glycine residues in the A α and B β chain in the E region, which releases fibrinopeptides A and B.⁵ This results in the formation of fibrin monomers and initiation of polymerization. Thrombin also activates factor XIII, which cross-links γ chains or α chains from neighboring fibrin molecules, thereby enhancing the stability of the fibrin network.⁶ Fibrinolysis of the fibrin network starts when a tissue-type plasminogen activator converts plasminogen into plasmin, which cleaves fibrin after lysine residues. Susceptibility to fibrinolysis is highly influenced by the structure of the clot.⁷ Thicker fibers, reduced branching, and larger pores increase the permeability and susceptibility to fibrinolysis, which is considered anti-thrombotic. However, thinner fibers, more branching, and smaller pores make the clot less permeable and more resistant to lysis by plasmin (prothrombotic).⁸

Fibrinogen Heterogeneity

Fibrinogen heterogeneity is the result of several types of variation: genetic polymorphisms, alternative mRNA processing, proteolytic cleavage, environmental factors, and post-translational modifications of fibrinogen.^{9–11} The different combinations of these determinants lead to more than a million forms of fibrinogen within a healthy individual.^{12,13} Post-translational modifications affect the function of fibrinogen, thereby influencing clot formation, the clot structure and susceptibility to clot lysis. These effects have consequences for the occurrence and progression of thrombotic diseases. The post-translational modifications which are discussed in this systematic review will be shortly introduced below. In Figures 1 and 2, an overview of the sites of modifications in the fibrinogen chains and the biochemistry of these modifications are shown.

Post-Translational Modifications

Oxidation

One of the modifications of fibrinogen is oxidation. Oxidative stress occurs in the human body when a high

Highlights

- Variations in fibrinogen affect the occurrence and progression of thrombotic diseases.
- Post-translational modifications of fibrinogen affect clot formation, clot characteristics, and susceptibility to fibrinolysis.
- For the best known post-translational modifications of fibrinogen, conclusions regarding their effect on clot characteristics can be drawn.
- Additional research is needed to elucidate the effects of all post-translational modifications of fibrinogen on clot characteristics.

amount of reactive oxygen species are produced (eg, by immune cells, pro-oxidant enzymes, and during oxygen metabolism) which are not sufficiently detoxified by antioxidants.¹⁴ Reactive oxygen species can be produced as a consequence of exogenous insults, for example, radiation from UV-light or smoking.¹⁵ Among the reactive oxygen species are superoxide anion, hydrogen peroxide, hypochlorous acid, and hydroxyl radical.¹⁴ These different reactive oxygen species react with and cause damage to a wide range of components of the cells, for example, proteins and DNA. Consequences of oxidation of proteins are the formation of carbonyl groups and the modification of amino acids (eg, methionine is converted into methionine sulphoxide and tyrosine into dityrosine; Figure 2).^{16,17} Compared with other plasma proteins, fibrinogen is especially susceptible to oxidation. It was shown that fibrinogen is 20 times more susceptible to oxidation than albumin, the most abundant plasma protein.¹⁸ A recent article identified the amino acids in fibrinogen which are oxidized by ozone.¹⁹ Previously, other articles have identified amino acids oxidized by photo-oxidation and hypochlorous acid.²⁰ In Figure 1, we show the amino acids which were identified by at least 2 ways of oxidation or which were oxidized above 50% using 50 μ mol/L ozone.

Nitration

Oxidative stress also leads to the production of reactive nitrogen species.¹⁵ Superoxide anion reacts with nitric oxide to produce the highly reactive peroxynitrite. In addition, the reaction of oxygen with nitric oxide and peroxynitrite results in the formation of other reactive nitrogen species, for example, nitrite and nitronium ion. Nitration of fibrinogen mainly affects tyrosine residues, resulting in the formation of 3-nitrotyrosine.^{21,22} Also cysteine residues can be affected, resulting in the formation of 3-nitrocysteine (Figure 2).

Modification by Carbohydrates

Glycosylation is the enzymatic process in which sugars (glycans or polysaccharides) are attached to proteins in an ATP (adenosine triphosphate)-dependent manner.

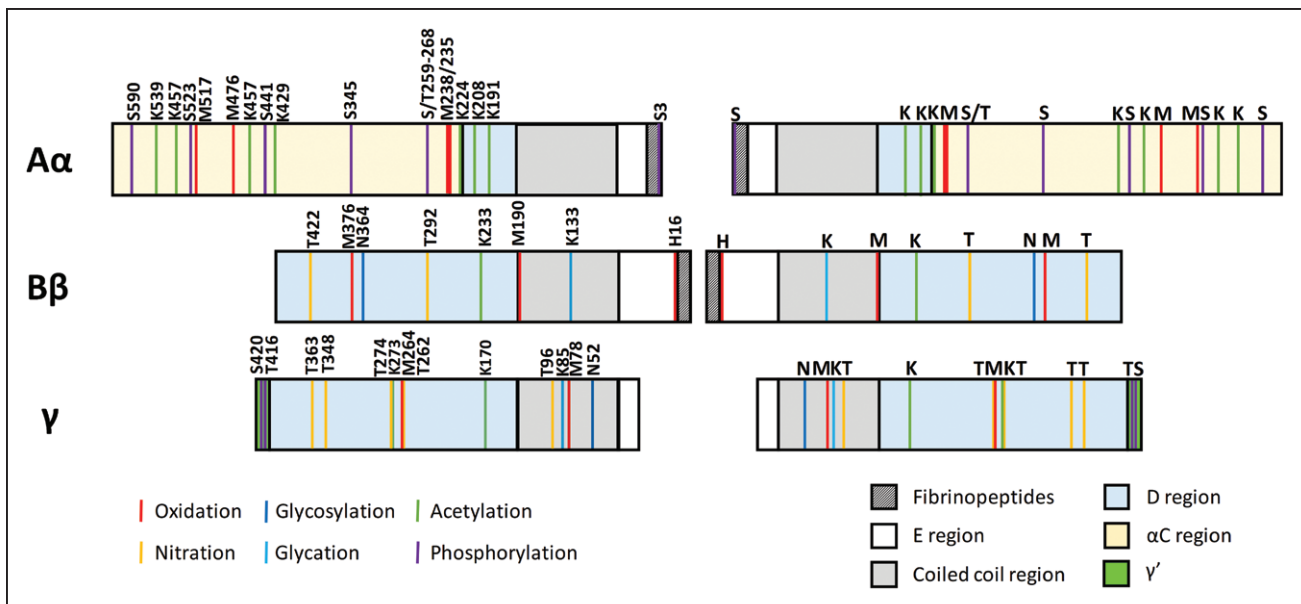


Figure 1. Known sites of post-translational modifications in the fibrinogen chains for the most studied modifications.

Schematic representation of the fibrinogen chains with the known sites of the post-translational modifications shown in lines with different colors. The letter and number indicate the type of amino acid which is modified and the position of the amino acid, respectively. The colored blocks correspond to the different regions of the fibrinogen molecule.

Glycosylation is needed at certain sites in a protein to function properly. Sialylation is a form of glycosylation in which sialic acid is bound at the end of a sugar chain of a glycoprotein. Each $B\beta$ and γ chain of fibrinogen has 1 N-glycosylation site, which can contain zero, 1, or 2 sialic acids (Figure 1).^{23,24} Normally, ≈ 6 sialic acid residues are present in each fibrinogen molecule.²⁵ Abnormal glycosylation has been related to aging and certain diseases; for example, patients with liver disease show an increased level of sialic acid content of their fibrinogen molecules.^{26,27} Glucose is known to nonenzymatically bind to proteins, often resulting in an impairment of its functions, in a process called glycation. Glycation occurs in patients with high levels of glucose in their blood, for example, in uncontrolled diabetes mellitus. Normal glucose levels are 5 mmol/L (90 mg/dL), but this can go up to 20 mmol/L (360 mg/dL) in patients with diabetes mellitus.²⁸ Also at normal glucose levels, glycation can occur. This is associated with oxidative stress, for example, observed during inflammation or in age-related diseases.²⁹ In fibrinogen, glycation is found to occur at lysine residues (Figure 1).³⁰

Acetylation

Acetylation is the attachment of an acetyl group to amino acids. In vivo, acetylation occurs at N-termini of polypeptide chains by Nt-acetyltransferases or at the ϵ -amino group of lysine residues by lysine acetyltransferases (Figure 2).³¹ Both processes have important physiological functions. Acetylation can also be caused by the intake of aspirin, which is known to exert its beneficial effects in cardiovascular disease by acetylating serine residues in the enzyme platelet cyclooxygenase. In fibrinogen and

other coagulation proteins, aspirin can acetylate lysine residues, affecting their functions (Figure 1).^{30,32}

Phosphorylation

Phosphorylation is the attachment of a phosphate group to an amino acid by protein kinases. Phosphorylation and dephosphorylation (by protein phosphatases) of proteins affect the structure and have important consequences for protein function.³³ Many enzymes are, for example, activated or deactivated by (de)phosphorylation and in many signaling pathways, phosphorylation plays a crucial role. In fibrinogen, phosphorylation occurs mainly on serine and threonine located in the $A\alpha$ chain (Figure 1).^{34,35} It is shown that fetal fibrinogen is phosphorylated to a higher degree than adult fibrinogen, suggesting a functional importance of phosphorylation.³⁴ In addition, phosphorylation of fibrinogen increases after surgery, potentially contributing to prevention of bleeding.³⁶

Other Modifications

High levels of the plasma protein homocysteine results in the nonenzymatic post-translational modification homocysteinylolation. Homocysteine has a free sulfhydryl group, with which it forms disulfide bonds with cysteine residues in proteins. Also, lysine residues can be modified, by the metabolite homocysteine thiolactone.³⁷ Homocysteine is produced in the metabolism of methionine, and it is thought that even a small increase in the plasma level is a risk factor for cardiovascular disease.³⁸ Homocysteine levels can be increased as a consequence of renal dysfunction, vitamin deficiencies, or medication affecting homocysteine metabolism.³⁸ Homocysteine levels in

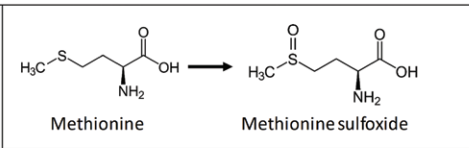
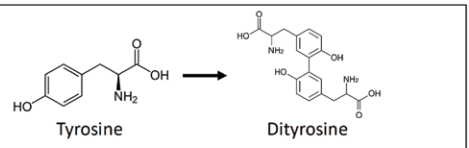
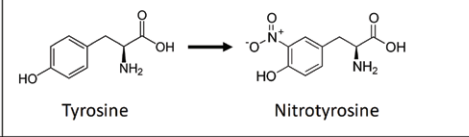
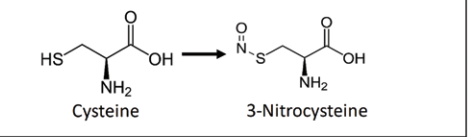
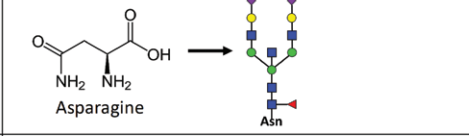
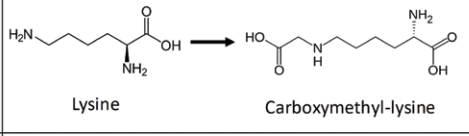
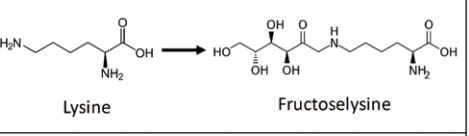
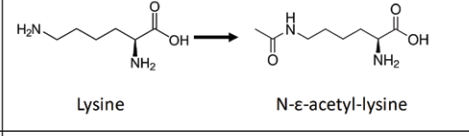
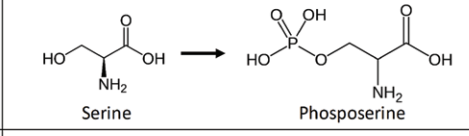
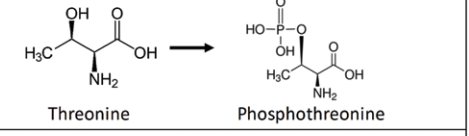
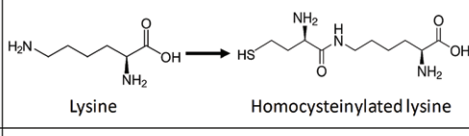
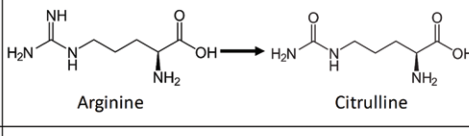
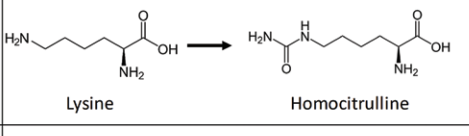
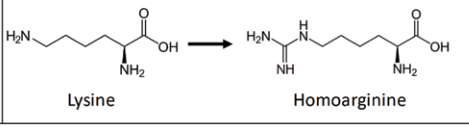
Oxidation		
Nitration		
Glycosylation		<ul style="list-style-type: none"> ◆ Sialic acid ● Galactose ■ N-Acetylglucosamine ● Mannose ▲ Fucose
Glycation		
Acetylation		
Phosphorylation		
Homocysteinylation		
Citrullination		
Carbamylation		
Guanidinylation		

Figure 2. Biochemistry of the post-translational modifications.

For each modification discussed in this review, 1 or 2 amino acids with the biochemical changes are depicted.

plasma higher than 15 $\mu\text{mol/L}$ are considered hyperhomocysteinemia and levels above 100 $\mu\text{mol/L}$ are classified as homocysteinuria.³⁹ In human fibrinogen, 3 lysine residues (Lys562 in the A α chain, Lys344 in the B β chain, and Lys385 in the γ chain) were shown to be homocysteinylation, both in vitro and in vivo.⁴⁰

Citrullination is the catalytic conversion of the amino acid arginine into citrulline by the enzyme peptidyl arginine deiminase. This post-translational modification is important in patients with rheumatoid arthritis, systemic lupus erythematosus, and other autoimmune diseases because autoantibodies in these diseases often attack

citrullinated proteins.⁴¹ Citrullination of fibrinogen potentially plays a role in the increased risk of thrombosis observed in rheumatoid arthritis.⁴² Using in vitro experiments, multiple arginine residues, mainly in the A α chain and a few in the B β chain, were identified that are citrullinated by 2 different peptidylarginine deiminases.⁴³

Carbamylation of proteins mainly occurs when amino acids interact with isocyanic acid, for example, during chronic kidney disease or atherosclerosis.⁴⁴ Isocyanic acid forms when urea decomposes into ammonium and cyanate, cyanate subsequently converts into isocyanic acid. Another source of cyanate is the thiocyanate metabolism, in which thiocyanate in the presence of hydrogen peroxide is converted into cyanate by myeloperoxidase.⁴⁴ Interaction of free amino acids or lysine residues with isocyanic acid results in the formation of α -carbamyl-amino acids or ϵ -carbamyl-lysine (homocitrulline), respectively.⁴⁴ Homocitrulline differs from citrulline by only one additional methylene group.

Finally, guanidinylation is the conversion of an amino group into a guanidine group, for example, happening in patients on hemodialysis.⁴⁵ Mostly, lysine residues are affected, which are converted into homoarginine. These last 2 post-translational modifications are studied less for fibrinogen, but because fibrinogen is a very abundant plasma protein, these modifications will most likely also affect fibrinogen.

It is important to know the effect of these post-translational modifications on clot formation, the clot structure, and susceptibility to fibrinolysis since these clot characteristics affect the development and progression of thrombosis and thromboembolic disease.^{46,47} With this knowledge, therapies to prevent or treat these diseases by normalizing clot structure or susceptibility to fibrinolysis can be optimized. During the past 70 years, multiple research groups have investigated the effects of these post-translational modifications. However, contradictory results are reported for the effects of certain modifications studied by different research groups. Therefore, the aim of this systematic review was to make an overview of the results of studies investigating the effect of post-translational modifications of fibrinogen on clot formation, clot structure, and susceptibility to fibrinolysis, to be able to draw tentative conclusions or identify the knowledge gaps.

METHODS

This systematic review was performed according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines.⁴⁸

Article Search

We conducted a systemic literature search in the Embase, Medline-Ovid, Cochrane Library, and Web of Science databases on April 18, 2019; and the search was repeated on

October 8, 2019. The search strategy included the different post-translational modifications in combination with fibrinogen and the characteristics we were interested in (eg, clotting time, clot structure, density, or permeability; see Methods in the [online-only Data Supplement](#) for the full search). No limit was set on the year of publication.

Study Selection

After deduplication, the search yielded 1278 results. Two researchers (J.J. de Vries and C.J.M. Snoek) independently screened these articles. First, relevant articles were included based on title and abstract. The included abstracts were subsequently read full text and articles that did not match the research question were excluded. In addition, conference abstracts, reviews, and articles not available in full text (in English) were excluded. Additional relevant articles identified while reading the full-text articles were also assessed for inclusion and included if there was a match with the research question. In case of disagreement, consensus about the articles was reached through discussion.

Data Extraction

To prevent bias, data were independently extracted from the articles by 2 researchers (J.J. de Vries and C.J.M. Snoek). The following data was collected: first author, publication year, method of experiments (which type of fibrinogen or plasma was used and how it was modified) and effects of the modification compared to control (clottability, cleavage by thrombin, rate of polymerization, initiation of polymerization or lag phase, maximal absorbance, diameter of fibrin fibers, stiffness, permeability, density, cross-linking, plasmin digestion of fibrinogen, and clot lysis).

RESULTS

Study Selection

A total of 1640 articles was found in the literature search. After elimination of duplication, 1278 articles were screened based on title and abstract (Figure 3). From the 183 articles read full text, we included 66 articles. In addition, while reading these articles full text, we found 39 other relevant articles in the references which were also included. The majority of these extra articles which did not show up in our literature search were old studies or did not use the terms for the modification in the title or abstract, which might explain why they did not come up in the search. In total, we identified 105 studies.

Study Characteristics

Most of the studies were performed in vitro with commercial purified fibrinogen, however, purified fibrinogen or plasma from patients was also used in a substantial number of studies. Especially studies investigating acetylation used healthy volunteers or patients taking aspirin to study the effect of acetylation on fibrinogen. Also, studies investigating the effect of other modifications

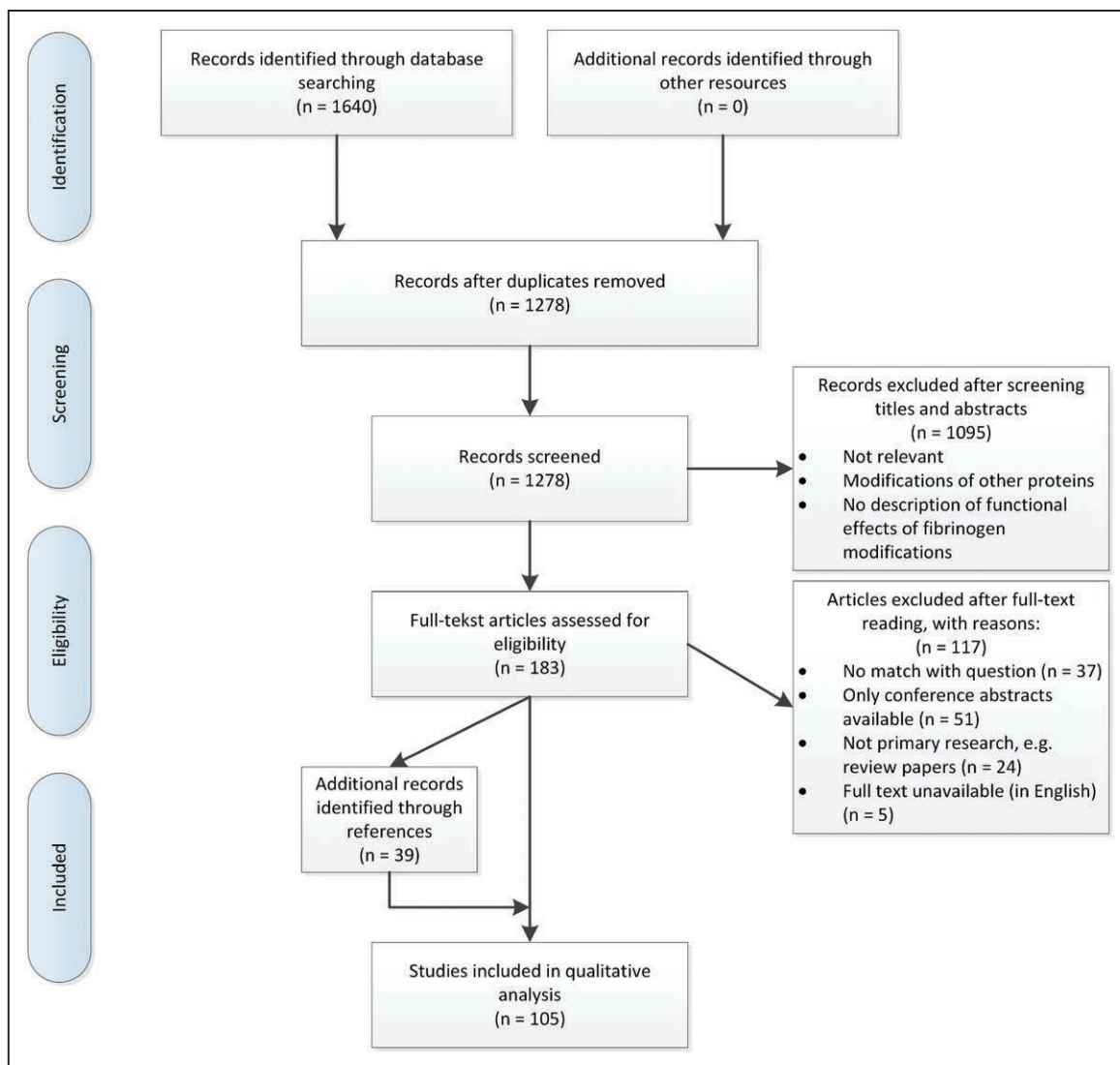


Figure 3. PRISM (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) flow diagram of study selection.

used material from patient groups in which the modification was known to be increased (eg, oxidation in myocardial infarction [MI] patients, glycation in diabetes mellitus patients, and sialylation in patients with liver disease). Twelve clot characteristics were studied in this systematic review. Clottability was used to describe the percentage of fibrinogen which is able to clot. Cleavage by thrombin describes the amount or rate of fibrinopeptide release from fibrinogen by thrombin. The rate of polymerization is reported when clotting time or the rate of aggregation was measured. Initiation of polymerization was mostly measured by the length of the lag phase, the time before clotting starts (when lag phase increases, initiation of polymerization is decreasing). Maximum turbidity means the maximum value of absorbance measured in turbidity assays. The diameter of fibrin fibers was determined using the mass-length ratio or measuring thickness of fibers on microscopy images. The stiffness of the clot was measured using rheology or thromboelastography.

Permeability describes the permeability of the clot, while density corresponds to the amount of fibers in a certain area. Cross-linking is mainly measured by measuring the amount and rate of formation of γ - or α -dimers. Plasmin digestion of fibrinogen means the breakdown of fibrinogen molecules by plasmin. Finally, clot lysis describes the degree or rate of fibrinolysis of the clot. Of these characteristics, most information was available for the rate of polymerization and maximum turbidity. The findings of all studies per modification are summarized in the Table.

Oxidation of Fibrinogen

Oxidation is the most studied post-translational modification of fibrinogen (31 articles, Table I in the [online-only Data Supplement](#)). Most studies used (human) fibrinogen and added a compound or condition that oxidizes fibrinogen (reactive oxygen species, ozone, or illumination). However, the conditions used in vitro show a lot of

Table. Overview of the Effects of the Different Modifications

Modification	Fibrin Polymerization					Clot Characteristics					Fibrinolysis		References
	Clottability	Cleavage by Thrombin	Rate of Polymerization	Initiation of Polymerization	Maximum Turbidity	Diameter of Fibers	Stiffness of Clot	Permeability	Density	Cross-Linking	Plasmin Digestion of Fibrinogen	Clot Lysis	
Oxidation	↓	=/↓	↓	↓	↓	↓	↓	↓	↑	↑		↓	16,19,49–78
Nitration		=	↑/↓	↑/↓	↑/↓						=		21,57,65,70,79–88
Glycosylation		=	↓	↓	↓	↓				=	=		25–27,87–99
Glycation	↑	=	=/↑	=	=/↑	=	=	↓	↑	=	↓	↓	28,100–114
Acetylation		=	↓		↑/↓	↑	↓	↑	↓		↑		32,71,115–125
Phosphorylation					↑/↓	↓					↓		35,36,126–132
Homocysteinylation			=/↓						↑	=		↓	39,133–137
Citrullination	↓	↓	↓		↓	↓			↓				138–140
Carbamylation			↓	↓	↓	↓			↑	↓		↓	141
Guanidinylation						↓		↓					45

variation. A few studies used fibrinogen from patients with diseases known to increase oxidative stress, and therefore the level of fibrinogen oxidation is increased^{49–53} (shown in Table I in the [online-only Data Supplement](#) below the thick line). Except for Paton et al,⁴⁹ the studies which used patient material do not show large differences compared to the in vitro studies.

The clottability of oxidized fibrinogen was decreased after oxidation in 2 studies.^{54,55} When light, radiation, or peroxyntirite was used to oxidize fibrinogen, no difference in cleavage by thrombin was found (3 studies).^{55–57} However, when fibrinogen was oxidized by reactive carbonyl compounds or ascorbate and iron, the cleavage by thrombin was decreased (2 studies).^{56,58} This suggests that the manner of oxidation influences the effects, probably by affecting different sites in fibrinogen. The rate of polymerization of fibrinogen after oxidation was mostly found to be significantly decreased compared to nonoxidized fibrinogen (in 25 of the 34 experiments). The in vitro studies reporting an increased rate of polymerization used relatively high concentrations of oxidative compounds, which might explain the discrepancy.^{59,60} In almost all experiments, the initiation of polymerization was significantly delayed (6 out of 9) and the maximum absorbance measured in turbidity assays was significantly decreased (15 out of 19). In correspondence with this, the diameter of the fibrin fibers was found to be significantly smaller in 8 of the 12 experiments. Only a few studies report contradictory results: a shorter lag phase and increased maximum absorbance or diameter.^{49,61,62} The study done by Rosenfeld et al⁶¹ found a significantly shorter lag phase and thicker fibrin fibers after oxidation of fibrinogen with ozone. The thicker fibers are also reported in⁶², where the same researchers used a similar method of oxidation. It is possible that oxidation by ozone differently affects the fibrinogen molecule than other methods of oxidation. The final study which shows contradictory results compared to the majority of the studies used purified fibrinogen from

MI patients 24 to 96 hours after the heart attack and compared clot characteristics between patients in different quartiles of plasma protein carbonyl values (representative of oxidative modifications and correlated to fibrinogen carbonyl content).⁴⁹ The authors themselves describe that the fibrinogen in other studies is probably more highly oxidized than their plasma samples, which would suggest that a low level of oxidation increases the ability of fibrinogen to clot. However, there is quite a broad range in concentrations of reactive compounds used in the other 30 studies, which gives a broad range of oxidation and most studies show a decreased ability of clotting. In addition, it can be appreciated in Table I in the [online-only Data Supplement](#) that incubating fibrinogen with relatively high concentrations of oxidative compounds shows an increased rate of polymerization, which would suggest the opposite: fibrinogen from the MI patients is oxidized to a higher degree instead of a lower degree. The other study that used fibrinogen from MI patients used blood drawn from patients 6 months after the event,⁶⁰ which is very different from 24 to 96 hours after MI. Although comparable levels of fibrinogen carbonyl content were reported in both studies, this difference in time after MI can be one of the reasons for the discrepancy between the 2 studies.

Other properties of the fibrin clot are studied in fewer articles. In general, a decreased stiffness (6 out of 7), lower permeability (4 out of 5), and higher density of fibrin clots (4 out of 8) were found after oxidation. Two studies found an increase in cross-linking after oxidation,^{63,64} while another study reported no difference.¹⁶ Digestion of fibrinogen by plasmin was found to be decreased after oxidation by concentrations of peroxyntirite higher than 100 μmol/L, whereas lower concentrations did not show significant differences.⁶⁵ Fibrinolysis was found to be significantly decreased by most studies (6 out of 9),^{16,50,51,66,67} but also no difference (one study) or an increase (2 studies) in fibrinolysis was reported.^{53,68,69}

Overall, oxidation seems to decrease the rate of clotting and results in more dense fibrin clots with thinner fibers which are less permeable. No conclusions can be drawn about the degree of cross-linking, digestion of fibrinogen or fibrinolysis, since only a few studies are done which all report different results. However, since most studies report a more dense fibrin clot with lower permeability, fibrinolysis is most likely decreased after oxidation.

Nitration of Fibrinogen

A post-translational modification that is relatively similar to oxidation and often studied in combination with oxidation is nitration of fibrinogen. We identified 12 studies on nitration of fibrinogen (Table II in the [online-only Data Supplement](#)), of which 3 are also included in the oxidation table (⁶⁵ and ⁵⁷ use peroxyxynitrite to study oxidation, which is also a nitrating agent and ⁷⁰ performed both oxidating and nitrating experiments). In Table II in the [online-only Data Supplement](#), the upper part shows the in vitro studies, arranged in order of increasing concentrations per nitrating compound used. Four studies also included experiments with fibrinogen from coronary artery disease patients,^{70,79} smokers,⁸⁰ or healthy volunteers taking lipopolysaccharides,⁸¹ all resulting in increased levels of nitration of fibrinogen (shown in the lower part of Table II in the [online-only Data Supplement](#)).

No significant effect of nitration on cleavage by thrombin was found in 2 studies.^{57,70} The rate of polymerization is generally found to be increased for nitrated fibrinogen from patients (5 out of 5 studies with patients) or fibrinogen nitrated by low levels of nitrifying agents. Except for one study, all experiments using high concentrations (>10 $\mu\text{mol/L}$ peroxyxynitrite or 100 $\mu\text{mol/L}$ nitronium fluoroborate) to nitrate fibrinogen showed a decreased rate of polymerization. Ding et al⁸² used a concentration of peroxyxynitrite slightly lower than 10 $\mu\text{mol/L}$ but also added increasing concentrations of manganese, which is known to increase fibrinogen nitration, explaining why they also find a decreased rate of polymerization. Helms et al⁸³ used 5 $\mu\text{mol/L}$ ProlinONOate to nitrate fibrinogen by the action of nitric oxide and found a decreased rate of polymerization, although not significant. There is one study that shows an increased rate of polymerization, although a high concentration of peroxyxynitrite (1 mmol/L) was used.²¹ This might be caused by a relatively low level of nitration found in this study; only 1.13 nitrotyrosine residues per fibrinogen molecule formed in contrast to levels up to 8 nitrotyrosine residues per fibrinogen molecule in other studies where high concentrations of peroxyxynitrite were used. Initiation of polymerization and maximum absorbance of turbidity measurements correspond to the results found for rate of polymerization; if the rate of polymerization was found to be increased, initiation of polymerization and maximum

turbidity were also increased. The diameter of the fibrin fibers was found to be thinner after nitration with a high concentration of peroxyxynitrite⁸⁴ or in fibrinogen from coronary artery disease patients,⁷⁰ while another study with fibrinogen from smokers reported no difference⁸⁰ and the study in which fibrinogen was nitrated by nitric oxide showed thicker fibers.⁸³ Also the stiffness of the clot showed all different results (decrease, no difference or increase in stiffness).^{70,80,83} Permeability and cross-linking were investigated by only one study, which showed both characteristics to be increased.⁷⁰ Density is shown to be higher in one study⁸⁴ and lower in another⁸³ after nitration in vivo. Digestion of fibrinogen or lysis of the fibrin clot by plasmin is shown to be similar^{65,70} or decreased after nitration of fibrinogen.^{65,80}

Overall, a low level of nitration increases the rate of fibrinogen polymerization, while increased nitration of fibrinogen decreases the rate of polymerization. The few studies studying the other clot characteristics find opposing results, probably also caused by different degrees of nitration or by the presence of other oxidative modifications that can occur upon treating fibrinogen with peroxyxynitrite.

Modification of Fibrinogen by Carbohydrates

We identified only 3 studies on the effect of glycosylation on fibrinogen (Table III in the [online-only Data Supplement](#)). One ex vivo study used fibrinogen from subjects of varying age, since aging increases the glycosylation of fibrinogen.²⁷ However, there were no differences in clot characteristics between older people (with a high level of glycosylation) and younger people.²⁷ Two other in vitro studies used another approach to study the effect of glycosylation of fibrinogen: the glycosylation of fibrinogen was inhibited during the production of fibrinogen in rabbit hepatocytes⁸⁷ or sugars were removed from fibrinogen using peptide-N-(N-acetyl- β -glucosaminy)asparagine amidase.⁸⁸ The first approach resulted in no difference in rate of polymerization, while the second study showed an increased rate of polymerization and maximum turbidity, thicker fibers and a more permeable clot, while cross-linking and fibrinogen digestion by plasmin were not affected by removal of glycosylation. Another in vitro study (not included in Table III in the [online-only Data Supplement](#)) was found in which fibrinogen was incubated with different sugars, investigating the lectin activity of fibrinogen. In addition, the effect of incubating fibrinogen with these different sugars was studied. A decreased rate of polymerization, decreased maximum turbidity and diameter of the fibers and a decreased cross-linking were found after incubation with different sugars.¹⁴²

A specific form of glycosylation, sialylation, was studied by 13 articles. Four studies with fibrinogen from patients with specific diseases which are shown

to increase the levels of sialic acid all presented a decreased rate of polymerization and initiation of polymerization.^{26,89–91} In other experiments, a different approach was used, namely removing sialic acid from fibrinogen. In addition, one article used fibrinogen from pregnant women, having a lower degree of sialylation.⁹¹ This desialylation resulted in increased rates of polymerization in 5 out of 6 studies, confirming the inhibitory role of sialic acid on polymerization.

Overall, glycosylation of fibrinogen results in decreased rates of polymerization and thinner fibers. There is not enough evidence to say something about the effect of glycosylation on the other characteristics of the fibrin clots.

Glycation of fibrinogen occurs in conditions with a high glucose concentration. We identified 16 studies investigating the effect of glucose binding on fibrinogen (Table IV in the [online-only Data Supplement](#)). Fibrinogen was incubated with glucose *in vitro*, or fibrinogen or plasma from patients with diabetes mellitus was used to assess the effects of glycation. Table IV in the [online-only Data Supplement](#) is sorted on glucose concentration used *in vitro* in the upper part of the table. Below the thick line, the studies using fibrinogen or plasma from patients with diabetes mellitus are sorted on the use of fibrinogen or plasma and type of patients with diabetes mellitus used.

After glycation, the clottability of fibrinogen was found to be similar (in one study) or increased (in 3 studies) compared to control fibrinogen. The 4 *in vitro* studies investigating the cleavage by thrombin reported no difference between glycated and control fibrinogen. However, fibrinogen from patients with type 2 diabetes mellitus showed an increased cleavage of fibrinopeptide B compared to control subjects.¹⁰⁰ The studies show contradictory results regarding the rate of polymerization, initiation of polymerization, and maximal turbidity. Different concentrations of glucose were used to glycate control fibrinogen, or fibrinogen or plasma from both patients with type 1 and type 2 diabetes mellitus were used, which might explain these differences. The diameter of the fibers did not change after glycation, but this was only investigated by 2 studies.^{100,101} The stiffness of the clot was also found to be similar for fibrinogen or plasma from patients with diabetes mellitus and controls in 2 studies.^{101,102} Three out of 4 studies measuring permeability found a decreased permeability after glycation and density is mostly shown to be increased (in 3 out of 4 studies). Most studies (7 out of 9) found no difference in cross-linking between glycated and control fibrinogen or between fibrinogen from patients with diabetes mellitus and control subjects. Finally, the majority of the results showed a decreased digestion of fibrinogen (3 out of 4) and fibrinolysis of the clot (4 out of 5) after glycation.

Overall, studies investigating glycation show different results regarding the polymerization, probably caused by the different conditions of modification used. The few

studies which are performed regarding the other characteristics of fibrin clots from glycated fibrinogen show a decreased permeability, increased density, no detectable effect on thickness of fibers, stiffness or cross-linking and decreased fibrinolysis of the clot after glycation.

Acetylation of Fibrinogen

Another modification is the acetylation of fibrinogen, which is mostly studied in the context of aspirin intake. We identified 13 studies, of which 6 did experiments with fibrinogen or plasma from patients or healthy volunteers who took aspirin (below the thick line in Table V in the [online-only Data Supplement](#)). The clottability and cleavage by thrombin were shown to be similar for control fibrinogen and fibrinogen acetylated *in vitro*.^{115–117} The rate of polymerization and maximum turbidity was mostly found to be significantly decreased after fibrinogen acetylation or ingestion of aspirin compared to nonacetylated fibrinogen (5 out of 9 studies). Only plasma from healthy volunteers taking a high dose of aspirin twice daily showed an increased polymerization.³² The other 3 studies showed no effects of acetylation on the rate of polymerization. Most experiments show a significantly increased diameter of the fibrin fibers after acetylation (7 out of 9). In correspondence with this, all studies agree that the permeability is increased (8 studies), the density is lower (3 studies) and susceptibility to clot lysis is increased (6 studies). It becomes clear that taking a low dose of aspirin is more beneficial than a higher dose since the high dose does not affect the fiber thickness and permeability of clots in 2 studies testing both doses.^{118,119} It is suggested that a high dose of aspirin can result in the formation of salicylic acid molecules, which block acetylation of fibrinogen and, therefore, inhibit the beneficial effects of aspirin on clot structure.¹²⁰ In conclusion, the acetylation of fibrinogen decreases fibrin polymerization and increases permeability and susceptibility to fibrinolysis.

Phosphorylation of Fibrinogen

The effect of phosphorylation or dephosphorylation of fibrinogen on clotting was studied by 9 research articles (Table VI in the [online-only Data Supplement](#)). These are all studies performed >20 years ago, no recent data were available. In most studies, fibrinogen was phosphorylated by incubating fibrinogen with kinases, although there is one study that uses fibrinogen from 5 patients after a hip surgery, who have elevated phosphorylation levels of fibrinogen.³⁶ It appears that phosphorylation by protein kinase A or C reduces the maximum turbidity and thickness of fibers, while phosphorylation by casein kinase II increases maximum turbidity and the fiber diameter. The patients after hip surgery also showed an increased maximum turbidity and fiber diameter, which would

suggest casein kinase II is the enzyme responsible for phosphorylation of fibrinogen *in vivo*.³⁶ However, only 5 patients were used, and more research is needed to confirm this hypothesis. The digestion of fibrinogen by plasmin was shown to be decreased in all studies, it made no difference which kinase was used. Another strategy to study the role of phosphorylation on fibrinogen was to remove the phosphate group from normal fibrinogen by alkaline phosphatase. It was found that dephosphorylation increases the maximum turbidity and diameter of fibrin fibers in 3 studies, while plasmin digestion was not affected (only one study). Finally, also the reversibility of these effects was investigated by dephosphorylating phosphorylated fibrinogen. It was shown that dephosphorylation increases maximum turbidity and diameter of fibrin fibers (and therefore normalizes back to normal after phosphorylation by protein kinase C and higher than normal in the case of casein kinase-phosphorylated fibrinogen).^{36,126,127} However, plasmin digestion of fibrinogen stayed reduced and is therefore not reversible.^{36,127,128}

In conclusion, the effect of phosphorylation on polymerization and the diameter of fibrin fibers depends on the kinase used to phosphorylate fibrinogen *in vitro*. *Ex vivo*, it was shown that increased phosphorylation of fibrinogen results in increased clot turbidity and fiber diameters, although this was only investigated by one study using a limited amount of patients.

Other Modifications of Fibrinogen

Six studies were identified in which the effect of homocysteinylation of fibrinogen was studied (Table VII in the [online-only Data Supplement](#)). Plasma from a rabbit model of homocystinuria showed an increased rate of polymerization and decreased fiber diameter,¹³³ while human fibrinogen and plasma incubated with homocysteine showed a decrease in rate of polymerization, decreased maximum turbidity, and no change in fiber diameter.³⁹ However, when plasma was incubated with higher levels of homocysteine (500 $\mu\text{mol/L}$), an increase in maximum turbidity and fiber diameter was observed.¹³⁴ These differences are probably the result of different concentrations of homocysteine, the use of fibrinogen or plasma and the difference between *in vitro* and *in vivo* situations. Clot density is observed to be increased by 3 studies and also agreement is reached over a decreased susceptibility to fibrinolysis after homocysteinylation in 5 studies. Stiffness of the clot was found to be significantly increased after incubation of plasma with 50 to 500 $\mu\text{mol/L}$ homocysteine.¹³⁵

Two studies on citrullination of fibrinogen reported an inability of fibrinogen to clot because thrombin is not able to cleave off the fibrinopeptides of citrullinated fibrinogen and, therefore, no polymerization occurs.^{138,139} (Table VII in the [online-only Data Supplement](#)). A more recent

article showed clotting after citrullination of fibrinogen and observed a decreased rate of polymerization, turbidity, fiber diameter, and density.¹⁴⁰

Carbamylation was studied in only one article in which carbamylation was found to decrease the ability of fibrinogen to clot, resulting in fibrin clots with thinner fibers, a higher density, less cross-linking, and decreased susceptibility to fibrinolysis.¹⁴¹

Finally, one article was found in which fibrinogen was modified by guanidinylation. The level of guanidinylation is increased in patients on hemodialysis.⁴⁵ Both clots formed from plasma from these patients and fibrinogen to which *o*-methylisourea-bisulfate solution was added to guanidylate fibrinogen showed clots with thinner fibers.⁴⁵ Also, the permeability of clots formed from plasma from patients on hemodialysis was decreased. The kinetics of polymerization were not investigated in this study.

DISCUSSION

In this systematic review, 105 research articles were identified that investigated the effects of post-translational modifications of fibrinogen on polymerization, clot structure, and fibrinolysis.

For oxidation, most research reported a decreased rate of polymerization, which results in more dense fibrin clots with thinner fibers and lower permeability. This results in clots that are more resistant to fibrinolysis, and therefore, prothrombotic. The mechanism by which oxidation affects clotting is thought to be oxidation of methionine residues in the αC region, which is involved in lateral aggregation. The αC region is proven to be most vulnerable to oxidation by ozone and hypochlorite.^{19,143} Weigandt et al¹⁶ identified methionine residue at position 476 in the αC region to be oxidized by hypochlorite. Conversion of Met476 into methionine sulfoxide was shown to impair dimerization of αC domains, thereby inhibiting lateral aggregation.¹⁴⁴

An interesting finding by Wang et al⁶⁴ was the observation that the stiffness of the clot measured at low frequencies is higher for oxidized fibrinogen compared to the control, while increasing the frequency at which stiffness is measured results in a lower value for oxidized fibrinogen. This difference implicates that the effect of oxidation on the stiffness can change corresponding to the conditions used in the measurement. This is an interesting finding, since blood flow and shear stress in the human body can also be variable. Therefore, it would be interesting to perform more experiments studying the effect of oxidation of clot stiffness measured during different degrees of shear stress.

It is hard to distinguish between nitration and oxidation, but in this systematic review, we separated these modifications. However, it should be noted that nitration and oxidation often occur simultaneously. In general,

nitration of fibrinogen using low concentrations of nitrating agents (low level of nitration) results in an increased rate of polymerization and higher maximum turbidity. The same results are found when fibrinogen nitrated *in vivo* is compared to control fibrinogen without nitration. However, high levels of nitration *in vitro* decrease the rate of polymerization and maximum turbidity. This shows that we need to be careful translating *in vitro* results to the *in vivo* situation. The other clot characteristics are less well studied for the effect of nitration. It is shown that nitration of fibrinogen results in the formation of nitrotyrosines in the part of the B β chain which is involved in lateral aggregation.⁸⁰ It is suggested that a low degree of nitration results in a gain of function of fibrinogen, having an increased rate of polymerization. When fibrinogen is nitrated with high concentrations of nitrating compounds, there might be too much nitrotyrosine formation which blocks lateral aggregation.

Glycosylation, and specifically sialylation, inhibits the rate of fibrin polymerization and leads to thinner fibers, which was shown both by increasing glycosylation and deglycosylation of fibrinogen. It is suggested that sialic acid inhibits polymerization by electrostatic repulsion between fibrin monomers.⁹² In normal conditions, calcium binds these sialic acid residues, thereby neutralizing this inhibitory effect. However, during liver disease, sialylation levels are increased, leading to insufficient neutralization of sialic acid by calcium and therefore inhibition of polymerization and thinner fibrin fibers.⁹² There has not been done enough research to draw conclusions about the effect of glycosylation on the other clot characteristics, such as stiffness of the clot, cross-linking, and permeability.

Glycation of fibrinogen showed contradictory effects on the polymerization, which might be caused by the different conditions in which this modification was tested (*in vitro* or fibrinogen or plasma from type 1 or type 2 diabetes mellitus patients). Therefore, more research is needed to determine the effect of glycation on fibrin polymerization and to evaluate if this effect is similar in diabetic patients. Few studies investigated other characteristics of fibrin clots and showed an increased density and decreased permeability and fibrinolysis of the clot after glycation, which corresponds to the resistance to clot lysis seen in diabetic patients.¹⁴⁵ In this systematic review, only studies are included which specifically investigated the glycation of fibrinogen (either by using purified fibrinogen or measuring fibrinogen glycation when plasma was used). However, there are multiple studies that used plasma from patients with diabetes mellitus to study clot characteristics, without quantifying fibrinogen glycation. These studies also show a decreased permeability and impaired fibrinolysis in clots from diabetic patients.^{146,147} It is known that the risk to develop major thrombotic diseases, for example, MI, stroke, or deep vein thrombosis, is increased in diabetes mellitus, due to the inflammatory and prothrombotic environment, of which

the latter can partly be ascribed to the glycation of fibrinogen.¹⁴⁸ Besides the increased density and decreased permeability, another reason for resistance to fibrinolysis might be the glycation of specific lysine residues in plasmin-sensitive coiled-coil regions of fibrinogen, thereby occupying the binding sites of plasmin.³⁰

Acetylation was shown by most studies to decrease the rate of fibrin polymerization and increase permeability and susceptibility to fibrinolysis. This is beneficial in the prevention and treatment of thrombosis and thromboembolic diseases, since a more permeable clot increases the susceptibility to fibrinolytic treatment. Acetylation as a consequence of aspirin intake is shown to target lysine residues, thereby disturbing the charge distribution of the fibrinogen molecule and affecting polymerization, resulting in clots with thicker fibers and increased permeability.¹²⁰ In addition, acetylation of lysine residues involved in cross-linking might be the cause of the decreased cross-linking seen in¹²¹, which further contributes to the increased permeability of fibrin clots.³⁰

It has been hypothesized that there is a competition between acetylation of amino acids by aspirin and glycation in diabetes mellitus, resulting in aspirin resistance in patients with diabetes mellitus.¹⁴⁹ However, identification of the lysine residues affected by both modifications showed no overlap between these sites.³⁰ Another study describes that taking aspirin can prevent the disadvantageous effects of oxidation. It was shown that acetylation of lysine residues in fibrinogen prevents the effects of oxidation of fibrinogen.⁷¹ The question whether this is due to competition remains to be answered.

The effect of *in vitro* phosphorylation of fibrinogen was different for the specific kinases used. Phosphorylation by casein kinase II showed similar results (increased maximum turbidity and thicker fibers after phosphorylation) as fibrinogen with an increased phosphorylation purified from patients, which suggests this kinase plays a role in fibrinogen phosphorylation *in vivo*. Also, the sequence specificity of type II casein kinases corresponds to the sequences around the phosphorylated amino acids.³⁴ It was reported that casein kinase II phosphorylates serine and threonine residues in the A α and γ ' chain.³⁵ Although protein kinase C also phosphorylates serine residues in the carboxyl-terminal part of the A α chain,¹⁵⁰ the effects of phosphorylation are different (decrease in maximum turbidity and fiber thickness), which suggests that amino acids at different positions are involved. To determine the effects of phosphorylation of fibrinogen, further studies are required which investigate which kinase is responsible for *in vivo* phosphorylation of fibrinogen, to use this information in *in vitro* experiments. In addition, fibrinogen phosphorylated *in vivo* can be used to investigate the effect of phosphorylation on clot characteristics. For example, no information is available on the effect of phosphorylation on clot stiffness, permeability, and susceptibility to fibrinolysis.

Studies on homocysteinylation of fibrinogen did not find consistent results in the polymerization, probably because different (mainly nonphysiological) concentrations of homocysteine were used with either fibrinogen or plasma. It would be recommended in further studies to use physiological homocysteine levels for homocysteinylation of fibrinogen to be certain about the effects on polymerization and fiber diameter. However, the studies did all show an increase in density and decrease in fibrinolysis after homocysteinylation. This can be explained by the finding that homocysteinylation of fibrinogen occurs mostly on lysines in the A α chain which are involved in plasmin binding and cleavage.^{136,137}

In 2 studies on citrullination of fibrinogen, no clots could be formed due to the citrullination of arginine residues in the N-terminus of the A α and B β chain.^{138,139} Since arginine residues are the cleavage site of thrombin, this modification inhibits the cleavage of fibrinopeptides by thrombin and formation of fibrin monomers, which explains the absence of clot formation. However, it is interesting to note that patients with rheumatoid arthritis (high levels of citrullinated fibrinogen) have an increased risk of thrombosis and do not have an increased risk of bleeding. This suggests that there is (increased) clot formation when fibrinogen is citrullinated *in vivo*. It is possible that the degree of citrullination in the performed *in vitro* studies is too high and, therefore, not relevant for the *in vivo* situation. A more recent study was indeed able to form clots with citrullinated fibrinogen and observed a decreased rate of polymerization, decreased turbidity, thinner fibers, and a decreased density when clots were formed from citrullinated fibrinogen.¹⁴⁰ Since this is only one study, these results need to be confirmed. In addition, *ex vivo* research with fibrinogen from rheumatoid arthritis patients would also be interesting.

Both for carbamylation and guanidinylation only one study was found, suggesting these modifications result in more dense clots with thinner fibers which are less susceptible to fibrinolysis. This corresponds to the *in vivo* situation seen in diseases in which these modifications occur (kidney disease or atherosclerosis).

Most studies included in this review are performed *in vitro*, sometimes with high concentrations of chemicals to induce the post-translational modifications. This makes it challenging to translate these findings to the *in vivo* situation. However, for most modifications, also *in vivo* work has been done, which mostly shows the same effects as found *in vitro* (except for nitration, as described above). Another source of variation between the studies is the variation in techniques used to study certain clot characteristics. Especially older articles used different methods to determine, for example, clotting time and clot lysis time (by visual inspection), while in more recent studies turbidity assays are used. Another example is the assessment of the diameter of the fibrin fibers, techniques used are, for example, the turbidity assay to calculate the

mass-length ratio or electron microscopy. However, we conclude that the different techniques used to measure the clot characteristics did not cause different observed effects of the post-translational modifications.

Although reviews are written about the effect of oxidative or other post-translational modifications, no recent systematic review is performed in which the effects of modifications are systematically shown as in our systematic review. A quantitative meta-analysis on the effect of post-translational modifications on clot characteristics was not possible, since there is too much variation in the way of modification and measurements of the different effects. Another limitation of this review is that studies might be missed in our literature search. However, we think that by identifying additional articles while reading our articles in full text, we covered a sufficient part of articles available on this subject.

CONCLUSIONS

In conclusion, there is knowledge about the effects of most post-translational modifications of fibrinogen on clot characteristics, especially about the best known post-translational modifications (oxidation, glycosylation, and acetylation). There are still many unanswered questions which deserve attention in upcoming research. For example, nitration shows different effects on fibrin polymerization and clot structure, potentially caused by the degree of nitration, which needs to be elucidated. Current research on glycation also shows contradictory effects on fibrin polymerization. For phosphorylation, *in vivo* studies could provide more information on which kinase is used in the body to phosphorylate fibrinogen and what the effect of this modification is on polymerization and clot structure. To investigate the effect of other (less known) modifications, for example, homocysteinylation and citrullination, only a few studies have been performed, which need confirmation. Expanding the knowledge about these modifications of fibrinogen can ultimately contribute to optimizing treatments for thrombotic diseases.

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