Timing and modality of the sclerosing agents binding to the human proteins: laboratory analysis and clinical evidences

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Abstract

Sclerosing agents (SA) are blood inactivat-ed. Nevertheless, investigations concerning the interaction among SA and blood components have never been deeply investigated. Aim of the study is to precisely identify SA blood ligands, to determine their binding time and to highlight the clinical consequences. Thirty-one blood samples were collected from chronic venous disease patients and tested by capillary and agarose gel (AGE) electrophoresis before and after adding polidocanol (POL) and sodium tetradecyl sulphate (STS). The two different types of electrophoresis allowed an evaluation of the blood proteins binding with the sclerosing agents, with a reaction time lower than 8 seconds for the AGE. Subsequently six patients underwent foam sclerotherapy and then were subdivided in group A (4 patients) and B (2 patients). In group A blood sample was obtained from the ipsilateral brachial vein immediately before (T0) and repeated 1, 3, 5, and 10 minutes after injection of STS 3% injection into the GSV. In group B, the same procedure was performed subsequently compared after addition of sclerosing agents. The study population mean age was 54±8 years, with a female/male ratio of 5/1. No significant co-morbidities were present.

In the first one, 31 blood samples were collected following 12 h or fasting in CVD patients (C1-6 Ep As Pr) and tested both by capillary and agarose gel (AGE) electrophoresis, in order to obtain a normal control curve, to be subsequently compared after addition of sclerosing agents. The assessment was performed at 5, 8, 12, 14 and 16 min by CE evaluation and at 8 h by AGE. Subsequently, the same 31 patients blood samples were evaluated in new CE and AGE runs adding both 3% STS and 3% POL (200 µL of SA+400 µL of blood). In CE the proteins fractions separations occurred in a fused silica capillary, which was previously filled with a buffer solution. The main components of the system are a vial with the sample, a source and a destination recipient, electrod, a high voltage generator, a shallow capillary (in silica), a recorder, a data collector and the support. The migration of the particle is created by the high voltage that was applied to the same solution, thus reducing the migration time to 5 min. The proteins were then assessed and quantified by direct measurement of their 214 nm wave length absorbance.[10]

In AGE, proteins fractions separation is based on the competition between he applied electric field and the specific protein affinity to the media. In AGE, an agarose gel matrix is used to deposit the particles. The band quantification, together with the standard diagram production, occurs by swartz starch or other substances coloration. Subsequently, a cleaning and media diaphanization are performed. At the end of the processing, the colored bands film is created. A densitometer is used for the assessment. The particles separation is obtained following their different electrophoretic mobility.[10]

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STS 3% injection into the GSV (T0). The sampling was repeated 1, 3, 5, 10 min later.

In group B (2 patients), the same procedure was performed with the same timing on the ipsilateral femoral vein.

Free STS (fSTS) and total protein-bound STS (bSTS) were measured in both groups.

The foam was produced according to the Tessari technique, thus mixing 1 cc of 3% STS with 4 cc of gas.21

The blood samples were immediately spin-dried and the plasma was stored in another vial. Everything was frozen at □26° and subsequently sent to the laboratory.

fSTS and total protein-bSTS were measured.

In group B (2 patients, 1 male and 1 female), the same procedure was performed with the same timing on the ipsilateral femoral vein.

An informed consent was obtained from all the patients, both for the clinical and the laboratory procedures.

**Results**

All the set-up tests were correctly performed and reproduced reliable outcomes. All the proteins fractions absolute values were physiological (Figure 1).

In the first set of investigations, the human plasma proteins displayed a weak bond with POL, mainly in the β1 an β2 range (only about 11% of the plasma proteins pool) (Figure 2A). Conversely, the STS-human plasma protein interaction was especially represented in the albumin (62.6%), α1 (10.7%) and α2 (20.0%) fractions (Figure 2B).

In the binding time analysis, the CE evaluation at 5, 8, 12, 14 and 16 min highlighted a total overlap of STS 3%-plasma proteins (1:2 ratio respectively) with the basal plasma proteins electrophoretic run (Figure 3).

The AGE evaluation demonstrated an 8-s drug-protein binding peak, that was equally maintained at the 12 and 60 s evaluation (Figure 4).

The second investigation, concerning fSTS amount, demonstrated that in the brachial vein, the average bSTS concentration in μg/mL was 0, 0.568, 5.98, 6.91 and 7.2, respectively at T0, 1, 3, 5, 10 min.

In the femoral vein, bSTS average concentration in μg/mL was 0, 1.62, 13, 24.6 and 8.67, respectively at T0, 1, 3, 5, 10 min.
Both in the brachial and in the femoral vein ISTS was always 0 (Figure 5).

Discussion

Nowadays, FS is one of the most popular CVD therapeutic tools worldwide. Some articles have reported possible threatening side-effects. Nevertheless, recent reviews have demonstrated that FS complications and side effects remain uncommon, thus confirming FS efficacy and safety.

On the other hand, the pathogenesis of FS-induced side effects still needs deeper investigation.

According to recent papers, the gas used to produce foam can be important in modification of the incidence of side effects, with a better safety profile for carbon dioxide-oxygen foam. But the side effects incidence isn’t related only to the chosen gas.

A deeper analysis of the biochemical interaction between the sclerosing agents and the blood components is mandatory to find out the possible new pathogenetic mechanisms of complications during sclerotherapy.

The present paper evaluates the sclerosing drug interaction with the human blood proteins, suggesting clinical consequences that improve the FS safety profile.

The first issue in the drug-protein binding assessment is identifying the exact timing in which it occurs.

A bias in the drug-protein CE assessed binding is the possible interaction between the two substances inside the test tube, during the same CE required processing time.

Therefore two different kind of electrophoresis have been used: CE and AGE.

CE is a more sophisticated analysis, designed to separate species and, based on their size, to charge ratio in the interior of a small capillary that is filled with an electrolyte. This characteristic brings a higher sensitivity, but requires some minutes to have the automated process started.

To the contrary, AGE is a totally manual procedure, bringing less precise measurement, but providing the test result in a few seconds.

The overlapping drug-protein peaks of CE and AGE demonstrated how the bond, and thus the inactivation, occurs after just a few seconds by the SA contact with the human blood.

Consequently, all the sclerosing drug circulation within the lungs is bounded to the proteins, and thus inactive.

Moreover, the FS safety profile is confirmed by the lack of detectible free sclerosing drug not only in the upper limb venous circulation, but also in the ipsilateral femoral vein.

Further considerations regarding coagulation modification were reported by Parsi last year, demonstrating how low concentration sclerosants inhibit platelet aggregation because of GPIIb/IIIa activation. The present study suggests that proteins like antiplasmin could be also involved. This is a c2 globulin which when bound to STS the lock on the plasmin is removed, thus enhancing fibrinolysis. This may represent a possible rationale for fewer post-sclerotherapeutic deep venous thromboses with the STS use.

It has recently been suggested that FS complications and side effects may be related to new pathogenetic mechanisms, based on
endothelial release of vasoactive molecules, such as endothelin-1, histamine or serotonin.22-28

Further analysis should be aimed to elucidate both the post-sclerotherapic endothelial catabolite identification and deeper insight into the drug-protein kinetic.

Conclusions

Foam sclerotherapy safety remains a hot topic in the phlebology community, despite everyday clinical use which demonstrates an extremely low complication and side effect rate. No data are available concerning the in vivo binding and neutralization of the SA whenever inside the blood stream.

The present paper offers two investigations demonstrating the SA binding features when in contact with human blood and the consequent kinetics within the venous network.

The results demonstrate in vivo the rapid inactivation of the SA in circulating blood, thus their high safety profile.

References

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