Solvent/Detergent-Treated Plasma: A Virus-Inactivated Substitute for Fresh Frozen Plasma

By Bernard Horowitz, Richard Bonomo, Alfred M. Prince, Sing N. Chin, Betsy Brotman, and Richard W. Shulman

Fresh frozen plasma (FFP) is prepared in blood banks worldwide as a by-product of red blood cell concentrate preparation. Appropriate clinical use is for coagulation factor disorders where appropriate concentrates are unavailable and when multiple coagulation factor deficits occur such as in surgery. Viral safety depends on donor selection and screening; thus, there continues to be a small but defined risk of viral transmission comparable with that exhibited by whole blood. We have prepared a virus sterilized FFP (S/D-FFP) by treatment of FFP with 1% tri(n-butyl) phosphate (TNBP) and 1% Triton X-100 at 30°C for 4 hours. Added reagents are removed by extraction with soybean oil and chromatography on insolubilized C18 resin. Treatment results in the rapid and complete inactivation of \(10^8\) infectious doses (ID\(_{50}\)) of vesicular stomatitis virus (VSV) and \(10^6\) ID\(_{50}\) of sindbis virus (used as marker viruses), \(10^6\) ID\(_{50}\) of human immunodeficiency virus (HIV), \(10^6\) chimp infectious doses (CID\(_{50}\)) of hepatitis B virus (HBV), and \(10^6\) CID\(_{50}\) of hepatitis C virus (HCV). Immunization of rabbits with S/D-FFP and subsequent adsorption of elicited antibodies with untreated FFP confirmed the absence of neoimmunogens formation. Coagulation factor content was comparable with that found in FFP. Based on these laboratory and animal studies, together with the extensive history of the successful use of S/D-treated coagulation factor concentrates, we conclude that replacement of FFP with S/D-FFP, prepared in a manufacturing facility, will result in improved virus safety and product uniformity with no loss of efficacy.

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MATERIALS AND METHODS

Reagents. Chemicals were reagent grade unless otherwise stated. TNBP and Tween 80 were obtained from Fisher Scientific (Springfield, NJ); Triton X-100 was obtained from Serva Feinbiochemica (Heidelberg, Germany); and Prep C18 resin was obtained from Millipore Corp (Milford, MA).

Virus-inactivation procedure and virus assays. Studies of the inactivation of vesicular stomatitis virus (VSV) and Sindbis virus were performed as described previously. Both viruses are lipid enveloped. Previous studies have shown that solvent/detergent mixtures do not inactivate protein-enveloped virus. The reactions were stopped by 100-fold dilutions of virus into medium (Minimal Essential Medium Eagle; GIBCO Laboratories, Grand Island, NY) containing 5% fetal calf serum. After sterile filtration and storage at −70°C, virus titer was determined by endpoint dilution and cellular cytopathology, using human A549 cells with VSV and primary chicken embryo cells with Sindbis virus. HIV-III strain was used to assess the inactivation of HIV added to plasma, as described previously. The reaction was stopped by adding 0.12 g of Waters Prep C18 resin to 1 mL of the treated plasma, mixing for 3 minutes, removing the resin by centrifugation, and repeating the process once. HIV infectivity was assayed by endpoint dilution, measuring reverse transcriptase after cultivation of virus exposed CEM cells for 14, 21, and 28 days.

The inactivation of HBV (strain NYBC 75-564) and NANBH (strain 59, a passage from the Hutchinson inoculum) on treatment of plasma was studied as described. Plasma from one carefully selected donor who was free of hepatitis B surface antigen (HBsAg), hepatitis B surface antibody (HBsAb), and hepatitis B core antibody (HBcAb), who had an alanine aminotransferase (ALT) level \(\leq 25\) IU/L, and who came from a group with a low risk of exposure to hepatitis viruses (ie, middle class, white, and...
heterosexual) was contaminated with either HBV to an end concentration of $10^5$ chimpanzee-infectious doses (CID$_{50}$) per milliliter or NANBH to an end concentration of $10^5$ CID$_{50}$ per milliliter. The virus-spiked plasma solutions were incubated in a shaking water bath at 30°C for 4 hours. After 1 hour of TNBP treatment, the plasma-virus mixture was transferred to a second tube to ensure that every droplet of plasma was contacted by the TNBP. The chimpanzees were immediately inoculated without further processing of the samples.

Four chimpanzees were injected with treated plasma: two received plasma spiked with $10^5$ CID$_{50}$ of HBV and two received $10^6$ CID$_{50}$ of NANBH. Chimpanzees used for the HBV inactivation study had no previous exposure to blood products and were free of hepatitis B markers. Chimpanzees for the NANBH inactivation study had been previously used to assess the safety of HBV vaccine.

After 9 months, the chimpanzees who received treated HBV-containing plasma were injected with 1 mL of a dilution of the untreated inoculum containing $10^5$ CID$_{50}$ of HBV. After 6 months, the chimpanzees injected with treated NANBH-containing plasma received 1 mL of the untreated inoculum containing $10^5$ CID$_{50}$ of NANBH. All chimpanzees were observed for an additional 6 months. Weekly sera were assayed for ALT, aspartate aminotransferase, HBsAg, anti-HBs, and anti-HBc. Bimonthly Menghini needle liver biopsies were examined by both light and electron microscopy.

Preparation of S/D-plasma. FFP was thawed rapidly and treated with stirring for 4 hours with 1% (vol/vol) TNBP and 1% (vol/vol) Triton X-100 at 30°C. After treatment, soybean oil (5% vol/vol) was added, mixed gently for 30 minutes, and then removed by centrifugation at 10,000g for 20 minutes. The clarified plasma was then applied to a column of Waters Prep C18 resin such that the ratio of plasma to column volume was 6 and the contact time was 3 minutes. The column eluate was filtered on a 0.2-µm filter.

Neoinmunogenicity. New Zealand white rabbits were immunized with S/D-plasma. Each animal was injected subcutaneously on three occasions at 2-week intervals; the first inoculum was emulsified with Freund’s complete adjuvant and the subsequent ones with Freund’s incomplete adjuvant. To determine whether S/D-plasma elicited antibody nonreactive with untreated plasma, antisera from each animal were collected immediately preceding each injection (bleeds 1 to 3) and for additional times over the subsequent 3 months. Antisera were analyzed by the Ouchterlony technique and by neutralizing crossed immunoelectrophoresis. In the latter case, anti-S/D-plasma was incubated with plasma in a test tube. After removal of the precipitate, remaining soluble antibody was incorporated into a gel to be used as the second dimension of a crossed immunoelectrophoresis. As a control, chicken albumin was added to the S/D-plasma used as the sample for electrophoresis and antichicken albumin was added at the outset of the experiment to the anti-S/D-plasma samples.

Assays. Coagulation factor activity for FVIII, FIX, and FXI was determined by one-stage activated partial thromboplastin (APTT) time clotting assay, which measures the degree of correction of the clotting time of factor-deficient plasma (George King Biomedical, Overland Park, KS) in the presence of APTT reagent (Organon-Teknika, Durham, NC). FV activity was assessed similarly, except thromboplastin with calcium (Sigma Diagnostics, St Louis, MO) replaced the APTT reagent.

TNBP was quantitated after hexane extraction by gas chromatography using a 0.25-in by 2 mm ID by 4-ft glass column packed with 10% SP-1000 on an 80/100 mesh Supelcoport (Supelco, Bellefonte, PA). Triton X-100 was assayed by first extracting it from plasma using a Prep C18 column, eluting with 75% isopropanol, concentrating the eluate with a Speed Vac concentrator (Savant, Farmingdale, NY), diluting the concentrate fivefold with water, and subjecting this sample to high performance liquid chromatography (HPLC) on a 3.9 x 300 mm Bondapak C18 column (Waters) with a Hewlett Packard HP1090 HPLC apparatus (Palo Alto, CA) and the UV detector set at 230 nm. The column was eluted with a water/isopropanol linear gradient.

RESULTS

Marker virus inactivation. The inactivation of all detectable VS and Sindbis virus added to plasma on treatment with 1% TNBP and 1% Triton X-100 at 30°C was very rapid, occurring within 15 minutes (Fig 1). Total demonstrated inactivation was $10^3$ tissue culture infectious doses (TCID$_{50}$) for VSV and $10^6$ TCID$_{50}$ for Sindbis virus. Use of 0.3% TNBP or a reaction temperature of 24°C resulted in a substantially slower rate of inactivation of

![Fig 1. Rate of inactivation of marker viruses added to plasma on treatment with TNBP and Triton X-100. VSV (upper panel) and Sindbis virus (lower panel) were added to human blood plasma (solid circles) or to an AHF concentrate prepared at the New York Blood Center and sampled at the stage in manufacture at which TNBP and detergent are normally added (open circles and triangles). Plasma was treated with 1% TNBP and 1% Triton X-100 at 30°C (solid circles), and plasma was treated with 0.2% TNBP together with either 0.2% sodium cholate at 30°C (open circles) or 1% Tween 80 at 24°C (open triangles). The titer of infectious virus was determined at the indicated times after a 100-fold dilution, used to stop the reaction.](image-url)
VSV (data not shown). Inactivation of each of these marker viruses on treatment of plasma under our selected conditions was faster than that observed on treatment of an antihemophilic factor (AHF) concentrate with 0.3% TNBP and 0.2% sodium cholate at 30°C or 0.3% TNBP and 1% Tween 80 and 24°C, the conditions in common use today for antihemophilic factor (AHF) concentrate with 0.3% TNBP and 0.2% sodium cholate at 30°C or 0.3% TNBP and 1% TNBP and 1% Triton X-100 at 30°C for 4 hours, not after 30 minutes of treatment (data not shown).

HIV. HIV, >10^6 TCID50, was inactivated on treatment of plasma with 1% TNBP and 1% Triton X-100 at 30°C for 4 hours (Table 1). A separate experiment showed that HIV inactivation was complete to the limit of detection on exposure to the S/D reagents at the earliest timepoint taken, 1 hour. Additionally, HIV (10^5 TCID50) was removed from plasma on exposure to Waters C18 Prep gel, the resin used in the removal of both TNBP and Triton X-100 during routine processing of S/D-plasma. Thus, the combined efficacy was ≥10^3 TCID50.

HBV. HBV (10^8 CID50/mL) added to plasma was treated with 1% TNBP and 1% Triton X-100 at 30°C for 4 hours, after which 10 mL was injected immediately into each of two animals. Neither animal developed signs of hepatitis through the initial period of follow-up: 28 weeks for chimpanzee 326, which died of an anesthesia-related incident, and 36 weeks for chimpanzee 327 (Fig 2). The low level of anti-HBs reactivity observed is not considered significant. Following challenge of chimpanzee 327 with the untreated inoculum, 10^3* CID50 of HBV, a sharp increase in serum ALT level and the appearance of anti-HBs were observed.

HCV. HCV (10^8 CID50/mL) added to plasma was treated in the manner described for HBV, after which 10 mL was immediately injected into two chimpanzees each. The two animals receiving HCV spiked plasma were monitored for indications of hepatitis transmission for 32 weeks. Throughout this period, serum ALT levels remained below 2 times the upper limit of normal, antibody to the C-100 protein of HCV was undetectable, and liver histology (light and electron microscopy) was normal (Fig 3). To confirm the susceptibility of the animals to HCV, an untreated sample was injected, using an HCV dose of 10^9 CID50/animal. Both animals developed signs of HCV infection, indicating susceptibility. Chimpanzee 298 had evidence of a transient, though sharp increase in serum ALT level, and Pfeiffer structures types I to IV were observed on examination of liver biopsies by electron microscopy. Anti-HCV (C100) was not observed. Chimpanzee 299 exhibited Pfeiffer structures 4 to 14 weeks after challenge and subsequently developed detectable anti-HCV (C100).

Coagulation factor activity. All coagulation factors measured remained at or near the level observed in the control, start sample when FFP plasma was thawed and processed at the 15- to 20-L scale (Table 2). In processing four separate plasma pools, comparison of the start and end concentrations of coagulation factors V, VIII, IX, and XI.
Absence of neoimmunogenicity. S/D-plasma injected into each of three rabbits did not elicit an antibody reactive with S/D-plasma that did not react with untreated plasma, as judged by the Ouchterlony technique (Fig 4) or by neutralizing crossed immunoelectrophoresis (Fig 5).

Removal of TNBP and Triton X-100 by C18 chromatography. S/D-treated plasma was extracted with 5% soybean oil, subjected to centrifugation at 10,000g for 10 minutes to remove lipid, and then applied to a column (14 x 14 cm) of Waters C18 Prep resin. With a contact time of 3 minutes, <2 µg/mL of either TNBP or Triton X-100 was present in the eluate when as much as 6 column volumes of plasma showed a recovery of 87%, 88%, 99%, and 108%, respectively. Prothrombin times and APTT values were normal, varying in four consecutive lots from 10.5 to 12.1 seconds and 32.0 to 33.8 seconds, respectively. These results suggest that contact factor activation is not encountered during processing. Moreover, factor XII levels were normal (0.78 to 0.88 U/mL; n = 3) in S/D-plasma prepared by Octa-

### Table 2. Recovery of Selected Coagulation Factors

<table>
<thead>
<tr>
<th>Factor</th>
<th>Start (U/mL)</th>
<th>End (U/mL)</th>
<th>Recovery (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Factor V</td>
<td>0.83</td>
<td>0.07</td>
<td>0.72</td>
</tr>
<tr>
<td>Factor VIII</td>
<td>0.93</td>
<td>0.06</td>
<td>0.82</td>
</tr>
<tr>
<td>Factor IX</td>
<td>0.98</td>
<td>0.07</td>
<td>0.97</td>
</tr>
<tr>
<td>Factor XI</td>
<td>0.86</td>
<td>0.05</td>
<td>0.93</td>
</tr>
</tbody>
</table>

FFP (15 to 20 L) as thawed, treated with 1% TNBP and 1% Triton X-100 at 30°C for at least 4 hours, and passed through a column of Waters C18 Prep Gel as described in Materials and Methods. Coagulation factor activities of the freshly thawed plasma and the C18 eluate were measured and compared (n = 4; SD = standard deviation).
DISCUSSION

Approximately 2 million units of FFP are used in the United States each year. While it is generally conceded that much of this usage is inappropriate, a Consensus Conference sponsored by the National Heart Lung Blood Institute concluded that FFP is useful in the treatment of coagulation factor disorders where appropriate concentrates are unavailable and when multiple coagulation factor deficits occur, such as in surgery. Since that meeting, FFP usage in the United States has remained essentially constant. If FFP is to be transfused, it should be made as safe as possible even while efforts continue to eliminate inappropriate usage.

We have now shown that treatment of pooled plasma with 1% TNBP and 1% Triton X-100 for 4 hours at 30°C inactivates $10^6$ CID$_{50}$ of HBV, $10^5$ CID$_{50}$ of HCV, and $10^6$ TCID$_{50}$ of HIV. Subsequent exposure to the C18 column used to remove TNBP and Triton X-100 was shown to remove an additional $10^3$ TCID$_{50}$ of HIV. Furthermore, because the rate of inactivation of VSV is faster in this system than observed on treatment of an AHF concentrate with either 0.3% TNBP and 0.2% sodium cholate at 30°C or 0.3% TNBP and 1% Tween 80 at 24°C (conditions in use worldwide), it seems likely that S/D-plasma will enjoy the same record of virus safety.

While considerable data support the effectiveness of the S/D approach against lipid-enveloped viruses, nonenveloped viruses, should they be present, would not be inactivated. The example most frequently raised in this context is parvovirus B19. Fortunately, B19 infections are self-limiting and largely asymptomatic. While not considered to be a serious threat in most transfusion settings, antiparvovirus antibody expected to be present in more than half of the units might effectively inactivate the rare unit containing parvovirus, depending on the respective titers of antibody and antigen. This conclusion seems plausible because routinely prepared IVIG preparations have been shown to neutralize parvovirus in vitro and in vivo.

The protein content of S/D-plasma prepared from large pools of FFP appears to be normal in all respects. Initial preparation at the 20-L scale resulted in excellent maintenance of the levels of coagulation factors V, VIII, IX, and XI, and of PT and APTT values. Neoimmunogens were absent when analyzed in a rabbit model. Additionally, the considerable worldwide experience with S/D-treated products indicates that the proteins present in S/D-plasma will circulate and function normally in vivo.

Finally, it should be noted that TNBP and Triton X-100 were efficiently removed by the described process. Residual values were $\leq 2 \mu g/mL$ of each when processing at the 20-L scale, and always $\leq 10 \mu g/mL$, when processing was performed at larger scale. If present at $10 \mu g/mL$, a 70-kg man receiving 2 L of S/D-plasma could potentially be exposed to 0.29 mg/kg body weight. This is more than 2 orders of magnitude lower than the reported lowest effect dose on intraperitoneal injection of TNBP into mice and intravenous injection of Triton X-100 into mice. Results on the acute toxicity of TNBP and Triton X-100 in mice and rats from studies jointly sponsored by the New York Blood Center and Octapharma (Dusseldorf, Germany) conducted by the Laboratory of Pharmacology and Toxicology (Hamburg, Germany) are in accord with these earlier, published studies (unpublished results, October 1991). Soybean oil is...
used as a principal ingredient in lipid emulsions designed for total parenteral nutrition. It consists almost entirely of triglycerides naturally occurring in humans, largely those of linoleic, oleic, and linolenic acid, and daily intravenous administration of more than 2 g/kg body weight of lipid emulsion has proven to be safe.26,27 By contrast, we have found no more triglyceride in the treated solution than that normally found in plasma. Toxicity should not be associated with the use of the Prep C18 resin. It is prepared by reaction between octadecyldimethyl silanol and the silica backbone. Information from the manufacturer (Millipore Corp, Waters Chromatography Division) indicates that leakage of this functional group should be very low, and none was normally found in plasma. Toxicity should not be associated with this preparation (M.S. Horowitz, personal communication, October 1991).

REFERENCES

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