Reversible exposure of hydrophobic residues on albumin as a novel strategy for formulation of nanodelivery vehicles for taxanes

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Background: We report herein a novel strategy for the preparation of protein-based nanodelivery vehicles for hydrophobic active pharmaceutical ingredients.

Methods: The procedure consisted of three steps, ie, exposure of hydrophobic residues of a protein to a pH-induced partial unfolding: interaction between hydrophobic residues on the protein and the hydrophobic active pharmaceutical ingredient, and a final step where the structure of the protein was reversed to a native-like state by returning to neutral pH. As proof of concept, the interaction of paclitaxel with partially unfolded states of human serum albumin was evaluated as a potential method for the preparation of water-soluble complexes of the taxane with albumin.

Results: We found that paclitaxel readily binds to pH-induced partially unfolded albumin, leading to the formation of optically clear water-soluble complexes. The complexes thus formed were more stable in solution when the albumin native state was at least partially restored by neutralization of the solution to a pH around 7. It was also observed that the hydrodynamic radius of human serum albumin was only slightly increased after the cycle of pH changes, remaining in a monomeric state with a size according to paclitaxel binding. Furthermore, paclitaxel binding did not affect the overall exposure of charged groups of human serum albumin, as evaluated by its interaction with an ionic exchange resin.

Conclusion: The in vitro biological activity of the complexes formed was qualitatively equivalent to that of a Cremophor®-based formulation.

Keywords: human serum albumin, paclitaxel, unfolded states, solubility

Introduction
Paclitaxel (Taxol®), Bristol-Myers Squibb, New York) is a very potent anticancer agent currently approved as first-line treatment for advanced carcinoma of the ovary, adjuvant treatment of breast carcinoma and nonsmall cell lung cancer, and as a second-line treatment for Kaposi’s sarcoma related to acquired immunodeficiency syndrome.1,2 Because of its extremely poor aqueous solubility, paclitaxel has been formulated in ethanol and polyethoxylated castor oil (Cremophor EL®), a vehicle that has been associated with bronchospasm, hypotension, and various hypersensitivity reactions, in particular after rapid administration or 10 minutes after initiating drug infusion.3,4 Premedication with corticosteroids and antihistamines is mandatory to reduce the incidence of serious hypersensitivity reactions. However, milder reactions still occur in 5%–30% of treated patients.5 To avoid the toxicities associated with the cosolvents required for taxane administration, and also to improve the solubility of paclitaxel, a number of alternative formulation strategies have been investigated.6–8 In this context, polyethylene glycol...
was evaluated as a biocompatible polymer which improves water solubility of paclitaxel. However, the polyethylene glycol–paclitaxel complex thus formed showed a decrease in antitumor activity. Other polymers, such as hyaluronic acid, polyglutamic acid, and nucleic acids, have been evaluated as carriers to improve paclitaxel solubility. Although these complexes had a highly stable composition, the biological activity of the pharmaceutical principal was slow.7,9,10

Proteins have been used as carriers for hydrophobic drugs, especially human albumin. In this context, Desai et al described a composition for in vivo delivery of water-insoluble drugs, such as paclitaxel.11,15 In their report, paclitaxel was incorporated in a polymeric shell of albumin with a size no bigger than 10 µm, which was substantially crosslinked through disulfide bonds induced in the protein by sonication. Desai et al then described a new formulation to deliver paclitaxel. In this case, the agent was delivered in the form of suspended particles coated with a stabilizing protein. The complex of protein with drug is subjected to high shear, in the absence of conventional surfactants or any polymeric core, a procedure that yielded amorphous nanoparticles with a diameter of less than 200 nm. These results were the basis for the development of Abraxane®, a formulation approved for the development of Abraxane®, a formulation approved for metastatic breast cancer. However, the formulation developed requires the use of a percentage of organic solvent in order to deal with the very low solubility of paclitaxel in aqueous media.

**Materials and methods**

Paclitaxel was obtained from Yunnan Smandbet Co Ltd (Kunming, China). Stock solutions of paclitaxel were prepared by dissolving the drug in either ethanol or dimethylsulfoxide in a final concentration of 11.1 mg/mL. Further dilutions in organic solvent were prepared as necessary when assaying for different ratios and concentrations.

HSA 20% (w/v) with sodium caprylate (0.04 M) and N-acetyl tryptophan (0.04 M) as stabilizers was obtained from the Laboratorio de Hemoderivados, National University of Córdoba, Argentina. To remove the excipients, albumin was dialyzed against distilled water or saline solution, and diluted with distilled nonpyrogenic water to the final concentration used in each assay. Defatted albumin was prepared by adsorption of fatty acids onto charcoal as previously described. Sodium dodecyl sulfate and urea were from Sigma Chemical Co (St Louis, MO). All other chemicals used were of analytical grade.

**Turbidimetry**

Optical density at a wavelength of 600 nm was used as a measure of the optical clarity of the samples.

**Determination of HSA concentration**

Protein concentration was determined using a Coomassie Brilliant blue assay or by direct absorbance at 280 nm, as previously described.23

**Determination of paclitaxel concentration**

Paclitaxel concentration was measured on a Curosil B C18 column (250 × 3.20 mm ID, particle size 5 µm) and a Curosil B C18 guard column (30 × 4.60 mm ID, particle size 5 µm) supplied by Phenomenex (Torrance, CA). The mobile phase was 60% (v/v) acetonitrile and 40% (v/v) biodistilled water. Flow rate was 0.7 mL/min and the eluent was monitored at 227 nm. Chromatography was performed at ambient temperature (20 ± 2°C).

**Protein chromatography**

Reaction mixtures and controls were chromatographed on an Äkta Explorer 100 system (GE Healthcare, Barrington, IL) fitted with a Superdex 200 or Mono Q column previously equilibrated with 50 mM phosphate buffer (pH 7.0) and 100 mM NaCl (phosphate-buffered saline). The elution
profile was followed using an ultraviolet detector at 280 nm, and the total protein levels were quantified using a Coomassie brilliant blue assay. Molecular weight markers in concentrations of 3 mg/mL were run similarly, and fractions analyzed for protein content using the Coomassie brilliant blue assay. HSA was eluted from the anion exchange resin with 50 mM sodium phosphate pH 7.0, 0.5 M NaCl.

**Water-soluble paclitaxel noncovalently bound to albumin**

Paclitaxel was solubilized in an organic solvent, such as ethanol or dimethylsulfoxide, and the solutions were then centrifuged to remove any potential particles. The solutions were slowly added under gentle agitation to the solution of HSA previously adjusted to the different conditions to be tested. After 1 hour of stabilization at the selected experimental conditions (temperature, pH, and ionic strength), the pH was slowly adjusted to 7.0 ± 0.2 with 1 M Tris-HCl pH 7.0, if necessary, and the paclitaxel precipitated was removed by centrifugation (14,000 × g, 10 minutes) at a series of time intervals. Finally, an aliquot of supernatant (300 µL) was filtered through 0.2 µm and the concentration of paclitaxel in solution (bound and unbound to HSA) was determined by reverse-phase high-pressure liquid chromatography. The amount of insoluble paclitaxel was determined from the precipitates by reverse-phase high-pressure liquid chromatography after its solubilization in ethanol.

**Human larynx epithelioma cell culture conditions**

Human larynx epithelioma cells (ATCC CCL-23) were propagated in minimum essential medium supplemented with 10% irradiated fetal bovine serum (Natocor, Córdoba, Argentina), 100,000 IU/L penicillin (Life Technologies, Rockville, MD), and 2 mM L-glutamine (Sigma). Cell monolayers were incubated in the above medium in the presence or absence of different concentrations of paclitaxel: HSA or paclitaxel in Cremophor (control) over 24 hours. Cell viability was assayed by trypan blue exclusion. Cell monolayers were photographed without any staining in an Axiovert 135 M Carl-Zeiss microscope at 50 × (Oberkochen, Germany).

**Results and discussion**

**Solubilization of paclitaxel by interaction with reversibly denatured HSA**

It is known that HSA undergoes reversible conformational isomerization with changes in pH of the solution containing the protein. At neutral pH, HSA assumes the normal form, which abruptly changes to a highly charged fast migrating form at pH values lower than 4.3 because this form moves “fast” on gel electrophoresis. In this condition, albumin is in a compact partially denatured state, with a significant amount of secondary structure but a largely disordered tertiary structure. Upon further reduction in pH to lower than 2.7, the protein structure changes to the fully extended form. The α-helix content decreases to a minimal value of 25%, and HSA is in an expanded form and increases the exposure of its hydrophobic surface. On the other hand, an increase in pH to 10 also induces a reversible denaturation of HSA that exposes part of its hydrophobic residues. We hypothesized that these reversible partially denatured structures could contribute to the binding of a highly hydrophobic molecule such as paclitaxel, leading to the formation of soluble complexes. As shown in Table 1, not only does paclitaxel form clearer solutions when incubated with denatured acid than with native HSA, but also a proportion of the complexes thus formed remains soluble even after pH is reversed to 7. The differences between the preparations are also reflected in the dramatic differences in their turbidity which, when measured before clarification by centrifugation, turned out to be a good predictor of the outcome in terms of paclitaxel solubilization.

<table>
<thead>
<tr>
<th>Ptx: HSA Molar ratio</th>
<th>Sample incubated at pH 2.7 (1 h) adjusted pH to 7.0</th>
<th>Sample incubated at pH 7.0 (1 h)</th>
<th>Sample incubated at pH 10.0 (1 h) adjusted pH to 7.0</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Pre-centrif. OD (600 nm)</td>
<td>Soluble Ptx (µg · mL⁻¹)</td>
<td>Pre-centrif. OD (600 nm)</td>
</tr>
<tr>
<td>9:1</td>
<td>0.543</td>
<td>355</td>
<td>1.824</td>
</tr>
<tr>
<td>4:1</td>
<td>0.596</td>
<td>446</td>
<td>1.483</td>
</tr>
<tr>
<td>2:1</td>
<td>0.664</td>
<td>455</td>
<td>1.386</td>
</tr>
</tbody>
</table>

**Note:** Samples of HSA were adjusted to pH 2.7, 7.0 or 10.0 and kept at 25°C during 1 hour, afterwards, 0.1 volume of Ptx in ethanol was added to reach a final Ptx concentration of 1 mg · mL⁻¹. The samples were incubated at 25°C during 1 hour and then the pH was adjusted to 7.0 with 0.1 N NaOH or HCl. Finally all samples were clarified by centrifugation at 14,000 × g during 10 minutes and the concentration of soluble Ptx was determined by RP-HPLC as described under materials and methods. In all cases OD at 600 nm after centrifugation was below 0.1. Results are the means of three different experiments performed in duplicates. RSD were all <10%.

**Abbreviations:** BDL, below detection limit; Ptx, Paclitaxel; OD, optical density; RSD, relative standard deviation.
It was also observed that the mixtures prepared at a low incubation temperature remained clearer and more soluble for at least a week at 4°C, indicating that the complexes formed had greater stability (data not shown). Upon increasing the temperature from 4°C to 37°C, unacceptable formulation mixtures were obtained due to increased precipitation of paclitaxel and HSA. Similar results were obtained at alkaline pH, except that at a pH of 10, the ester bond in paclitaxel was rapidly hydrolyzed at high temperatures (data not shown).

Effect of ionic strength on formation of water-soluble paclitaxel–HSA complexes

Given the zwitterionic character of HSA and the hydrophobicity of paclitaxel, the ionic strength of the aqueous media is almost certainly a variable that will affect the formation of water-soluble complexes between paclitaxel and HSA. First, an increase in ionic strength could reinforce and stabilize the mainly hydrophobic interaction of paclitaxel with HSA. On the other hand, using a high protein concentration, increasing ionic strength could promote interaction among albumin molecules, leading to a decrease in stability with a concomitant increase in turbidity of the solution. Although not statistically significant, a slight improvement in paclitaxel solubility was apparent with the addition of NaCl up to 0.4 M, suggesting that the addition of salt reinforces and stabilizes the interaction of paclitaxel with HSA (Table 3).

Table 2 Effect of incubation temperature on the turbidity and solubility of Ptx–HSA complexes

<table>
<thead>
<tr>
<th>Ptx:HSA Molar ratio</th>
<th>Incubation temperature</th>
<th>4°C</th>
<th>25°C</th>
<th>37°C</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Pre-centrif. OD (600 nm)</td>
<td>Soluble Ptx (μg mL⁻¹)</td>
<td>Soluble Ptx (μg mL⁻¹)</td>
<td>Soluble Ptx (μg mL⁻¹)</td>
</tr>
<tr>
<td>9:1</td>
<td>0.123</td>
<td>328</td>
<td>0.352</td>
<td>240</td>
</tr>
<tr>
<td>4:1</td>
<td>0.154</td>
<td>257</td>
<td>0.386</td>
<td>213</td>
</tr>
<tr>
<td>2:1</td>
<td>0.203</td>
<td>252</td>
<td>0.686</td>
<td>179</td>
</tr>
</tbody>
</table>

Note: Samples of HSA were adjusted to pH 2.7 and kept at 4, 25 or 37°C during 1 hour, afterwards, 0.1 volume of Ptx in ethanol was added to reach a final Ptx concentration of 0.5 mg mL⁻¹. The samples were incubated at 4, 25 or 37°C during 1 hour and then the pH was adjusted to 7.0 with 0.1 N NaOH. Finally, all samples were clarified by centrifugation at 14,000 × g during 10 minutes and the concentration of soluble Ptx was determined by RP-HPLC as described under materials and methods. Results are the means of three different experiments performed in duplicates. RSD were all <10%.

Abbreviations: Ptx, Paclitaxel; OD, optical density; RSD, relative standard deviation.

Table 3 Effect of ionic strength on the optical clarity and solubility of Ptx–HSA complexes

<table>
<thead>
<tr>
<th>NaCl (M)</th>
<th>Sample incubated at pH2.7*</th>
<th>Sample incubated at pH 2.7 (1 h) adjusted pH to 7.0**</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Pre-centrif. OD (600 nm)</td>
<td>Soluble Ptx (μg mL⁻¹)</td>
</tr>
<tr>
<td>0</td>
<td>0.266</td>
<td>BDL</td>
</tr>
<tr>
<td>0.2</td>
<td>0.196</td>
<td>869</td>
</tr>
<tr>
<td>0.4</td>
<td>0.196</td>
<td>890</td>
</tr>
<tr>
<td>0.6</td>
<td>0.223</td>
<td>891</td>
</tr>
<tr>
<td>0.8</td>
<td>0.425</td>
<td>465</td>
</tr>
<tr>
<td>1.0</td>
<td>&gt;2</td>
<td>BDL</td>
</tr>
</tbody>
</table>

(Precipitated) (Precipitated)

Note: *Samples of HSA (20 mg mL⁻¹) were adjusted to pH 2.7 and kept at 4°C during 1 hour, afterwards, 0.1 volume of Ptx in ethanol was added to reach a final Ptx concentration of 1 mg mL⁻¹. The samples were incubated at 4°C during 1 hour and then samples labeled * were adjusted to pH 7.0 with 0.1 N NaOH. All samples were incubated 1 h at 4°C, clarified by centrifugation at 14,000 × g during 10 minutes and the concentration of soluble Ptx was determined by RP-HPLC as described under materials and methods. Results are the means of three different experiments performed in duplicates. RSD were all <10%.

Abbreviations: BDL, below detection limit; RSD, relative standard deviation; Ptx, Paclitaxel; OD, optical density.
As expected, we found that a gel was formed at a high ionic strength and an acidic pH, and this led to marked precipitation of protein and turbid mixtures (Table 3). This effect was even more sensitive to salt concentration when the HSA concentration was 40 mg/mL, at which a gel was formed with 0.6 M NaCl, whereas at 60 mg/mL a gel was formed with 0.4 M NaCl.

**Chromatography of paclitaxel–HSA complexes**

To explore the structural properties of the complexes formed between paclitaxel and acid-denatured HSA, samples of the soluble complexes formed were run on size exclusion and ionic exchange chromatography columns. As determined by reverse-phase high-pressure liquid chromatography analysis of the fractions, paclitaxel coeluted with HSA from a size exclusion resin indicating that there is a physical association between them (data not shown). It was also observed that the elution volume of HSA changed slightly upon binding of paclitaxel, suggesting that it remains monomeric and

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**Table 4 Binding of Ptx–HSA to anion exchange resin**

<table>
<thead>
<tr>
<th>Paclitaxel: HSA Molar ratio</th>
<th>Nonbound to Mono Q HSA (%)</th>
<th>Paclitaxel (%)</th>
<th>Eluted from Mono Q HSA (%)</th>
<th>Paclitaxel (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>9:1</td>
<td>5.0</td>
<td>22</td>
<td>95</td>
<td>80</td>
</tr>
<tr>
<td>4:1</td>
<td>8.0</td>
<td>14</td>
<td>92</td>
<td>82</td>
</tr>
<tr>
<td>2:1</td>
<td>7.0</td>
<td>16</td>
<td>93</td>
<td>85</td>
</tr>
</tbody>
</table>

**Note:** Results are the means of three different experiments performed in duplicate. RSD were all <10%.

**Abbreviations:** RSD, relative standard deviation; Ptx, Paclitaxel.
globular, with a hydrodynamic radius in agreement with that expected for HSA complexed with 4–6 molecules of paclitaxel (Figure 1). In agreement with this finding, more than 90% of the albumin and paclitaxel were bound and coeluted from an ionic exchange resin, with a profile similar to that of pure HSA (Table 4 and Figure 2). Altogether, these results confirm that paclitaxel is solubilized in water through a physical interaction with HSA, and also provide evidence suggesting that the complexes formed do not substantially modify the hydrodynamic radius or the net charge of HSA.

Evaluation of biological activity of water-soluble paclitaxel–HSA complexes

The test used to evaluate the biological activity of the paclitaxel–albumin complex was the inhibition of division of a tumoral cell line. As seen in Figure 3, control samples show a clear monolayer of human larynx epithelioma cells covering the surface of the Petri dishes. On the other hand, cell samples treated with paclitaxel either in Cremophor or complexed with HSA show few cells, indicating that albumin does not impair the biological effect of paclitaxel on cell division. Furthermore, at least at the dose tested, there was
no statistical difference in the number of viable cells between paclitaxel in Cremophor and those complexed with HSA.

**Conclusion**

Standard formulations of taxanes require use of solvents, such as Cremophor or Tween, which contribute to some of the toxicities associated with paclitaxel-based therapy. To overcome these drawbacks, nanoparticle HSA-bound paclitaxel has been recently approved and is available commercially as Abraxane®. However, this formulation is prepared by high-pressure homogenization of albumin and paclitaxel, leading to the formation of a colloidal suspension with very limited stability. We hypothesized that the exposure of hydrophobic domains of albumin via a partial opening of its structure could improve its interaction with a hydrophobic molecule like paclitaxel. The results obtained confirm that, as previously known, the binding affinity of paclitaxel with native HSA is very low (Table 1). In agreement with our hypothesis, we found that paclitaxel readily binds to pH-induced partially unfolded HSA, leading to the formation of optically clear water-soluble complexes. The complexes thus formed appear to be more stable in solution when the native state of HSA is at least partially restored by neutralization of the solution to a pH around 7. In agreement with this, we observed that the hydrodynamic radius of HSA is only slightly affected after the cycle of pH changes, and is consistent with paclitaxel binding. Furthermore, paclitaxel binding does not affect the overall exposure of charged groups of HSA, as evaluated by its interaction with an ionic exchange resin. The in vitro biological activity of the complexes formed was qualitatively equivalent to that of a Cremophor-based formulation.

In conclusion, our results not only show that HSA-paclitaxel complexes can be prepared by a simple method that does not require use of high-pressure homogenization, but also open up the possibility to extend this strategy for the design of novel protein-based nanodelivery vehicles for other hydrophobic active pharmaceutical ingredients. Further work is in progress in order to characterize the structure, as well as energetic and dynamic parameters of the complexes formed by the binding of paclitaxel to partially unfolded HSA and to evaluate their activity.

**Acknowledgments**

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**Disclosure**

The authors report no conflicts of interest in this work.

**References**
