Letter

Removal of Triton X-100 and SDS from Protein Solutions with Zeolite Y

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Detergents are commonly used in biochemistry. They are amphiphilic molecules containing a hydrophobic and a hydrophilic part. The hydrophilic part can be charged or uncharged and detergents often are characterized as being an-, cat- or non-ionic. In biochemistry detergents are mainly used for solubilizing cell membranes and protein aggregates.

Purification of membrane components and intracellular molecules is often carried out in presence of detergents. However, detergents may interfere with the purification method used. Thus ion-exchangers may bind anionic or cationic detergents and non-ionic detergents, that form large micelles, may cause an increased apparent molecular weight upon gel filtration.

The development of cDNA technology and the increased production of recombinant proteins, which often form intracellular protein aggregates, ‘inclusion bodies,’ has increased the demand for fast and large-scale purification methods of intracellularly formed proteins. So far, the most commonly used methods for the removal of detergents from protein solutions are dialysis, ion exchange chromatography and adsorption onto hydrophobic matrices. In this letter we describe the use of a hydrophobic zeolite as an adsorbing matrix for detergents.

Zeolites, framework aluminium silicates or tectosilicates, are characterized by their chemical composition and pore size.1 The chemical composition is conveniently expressed in terms of the Si:Al ratio. High-silica zeolites carry less framework charge and are commonly referred to as hydrophobic; the opposite holds for high-alumina zeolites which are labelled hydrophilic. Pore sizes vary typically in the range 3–7.5 Å and the accessibility of the porous system is dimensionality dependent. Of the suitable zeolites, zeolite Y, mordenite, and silicalite can be obtained essentially free from aluminium either through direct synthesis (silicalite) or by means of post-synthetic manipulations (mordenite, zeolite Y). Zeolite Y and mordenite have pore sizes in the upper range, 7.5 and 7.0 Å respectively, while silicalite is a middle-range zeolite with 5.5 Å pore size. The porous systems in zeolite Y and silicalite are readily available owing to the three-dimensional arrangement of the channels; mordenite is somewhat less efficient since the porous system is one-dimensional.

Experimental

Zeolites were added to phosphate-buffered saline (PBS, 10 mM phosphate, 150 mM NaCl, pH 7.4) containing 10 mg ml⁻¹ bovine serum albumin (BSA, Cohn fraction V, Sigma Chemicals Co., USA) and 10 mg ml⁻¹ Triton X-100 (Merck, Darmstadt, FRG) or 10 mg ml⁻¹ sodium dodecyl sulfate, SDS (specially pure, BDH Chemicals Ltd., Poole, England). The suspensions were vortex mixed for 60 s and centrifuged for 10 min at 12,000 × g before assaying of the detergent remaining in the supernatant. Zeolites were obtained from Tosoh Co., Japan and silica (SiO₂·xH₂O) used in the control experiments was obtained from Merck, Darmstadt, FRG.

1H-Triton X-100 (NET-556 lot number 2481-229) used as a tracer was purchased from NEN Research Products, Boston, USA. Before the detergent was dissolved in PBS, 1H-labelled Triton X-100 was mixed with unlabelled Triton X-100, dissolved in methanol and evaporated to dryness. This procedure reduced the amount of radioactivity by 30%.

Table 1. Adsorption of detergents onto zeolites.*

<table>
<thead>
<tr>
<th>Zeolite</th>
<th>Pore size /Å</th>
<th>Si:Al</th>
<th>Triton X-100 remaining in the supernatant (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Zeolite Y</td>
<td>7.5</td>
<td>&gt;1000</td>
<td>0.7</td>
</tr>
<tr>
<td>Silicalite</td>
<td>5.5</td>
<td>&gt;1000</td>
<td>87</td>
</tr>
<tr>
<td>Mordenite</td>
<td>7.0</td>
<td>46</td>
<td>86</td>
</tr>
</tbody>
</table>

*Zeolites (100 mg ml⁻¹) were added to a solution of 10 mg ml⁻¹ Triton X-100 in PBS. After vortex mixing for 60 s, the zeolites were pelleted by centrifugation and the detergent remaining in the supernatant was determined by adsorption at 276 nm.

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**Adsorption of Triton X-100 onto zeolite Y and onto silica particles.** Triton X-100 (10 mg ml⁻¹), and ³H-labelled Triton X-100 (1.4×10⁶ dpm ml⁻¹) were dissolved in PBS (10 mg ml⁻¹ BSA) as described above. Various amounts of zeolite Y or silica particles were added to the detergent solution and this was vortex mixed for 60 s. The adsorbents were removed by centrifugation and 100 µl of the supernatants were counted in a β-counter.

**Result and discussion**

Zeolites are commonly used as adsorbers of low molecular weight substances. A recent proposition as to the mechanism of adsorption in zeolites in general, and especially in hydrophobic zeolites, is based on the geometry of the framework structure.² In a hydrophobic zeolite carrying little or no framework charge, the adsorptive properties are described as a result of the focusing of the total van der Waals forces.³ The focusing efficiency is a consequence of the particular geometry involved and hence is different for different zeolites. In an attempt to find a zeolite that adsorbs detergents, three zeolites of known hydrophobic properties were examined (Table 1). The initial experiments were performed with an anionic detergent, sodium dodecyl sulfate (SDS), and a non-ionic detergent, Triton X-100. Significant adsorption of both detergents was only obtained with zeolite Y.

Binding of ³H-labelled Triton X-100 to zeolite Y and to silica particles is shown in Fig. 1. After the addition of 50 mg ml⁻¹ zeolite Y, 1.7% of the radioactivity remained in the supernatant. This gives a concentration of 0.27 mM Triton X-100 which is close to its critical micelle concentration (cmc), 0.24 mM. Furthermore, no lysis of red blood cells could be obtained with Triton X-100 solutions treated with 50 mg ml⁻¹ zeolite Y, which is in agreement with a detergent concentration below its cmc. A concentration below cmc was also obtained when BSA was omitted from the detergent solution and the amount of detergent remaining after treatment with zeolite Y was assayed by absorption at 276 nm. However, if the mixture of ³H-labelled and non-labelled Triton X-100 was not dissolved in methanol and evaporated to dryness before dissolution in PBS, 30% of the radioactivity remained in the supernatant after treatment with 50 mg ml⁻¹ zeolite Y. The manufacturer states their ³H-labelled Triton X-100 to be 99% pure. We agree with the manufacturer after removal of the volatile radioactivity from the ³H-Triton X-100 preparation.

Adsorption of Triton X-100 to zeolite Y is achieved after vortex mixing for less than 60 s. Compared with other hydrophobic matrices, such as BioBeads SM-2,⁴ the adsorption onto zeolite Y is much faster and indicates an adsorption mechanism based on direct binding of the detergent micelles.

A solution of 10 mg ml⁻¹ SDS treated with 100 mg ml⁻¹ zeolite Y showed no lysis of red blood cells. SDS is an anionic detergent and hence the adsorption of detergent to zeolite Y is independent of charge, instead the adsorption is dependent on the size of the hydrophobic part of the detergent molecules. Removal of detergents from solutions by the use of zeolite Y⁵ is fast and efficient and should have numerous applications in biochemistry and biotechnology.

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**References**


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