

Commentary

Colloid-induced kidney injury: experimental evidence may help to understand mechanisms

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See related research by Hüter *et al.*, <http://ccforum.com/content/13/1/R23>

Abstract

Fluid resuscitation is widely used, and many patients are therefore exposed to plasma volume expanders. Among these, colloids, particularly hydroxyethyl starches, have been shown in recent experiments and clinical studies to induce acute kidney injury. The mechanisms of colloid-induced acute kidney injury remain incompletely elucidated. The risks associated with colloid osmotic pressure elevation *in vivo* and the high incidence of osmotic nephrosis lesions in experimental models and clinical studies indicate that hydroxyethyl starches can no longer be considered safe.

Plasma volume expansion is often required in the operating room, emergency department, or intensive care unit. The safety of plasma volume expanders therefore deserves careful consideration. Low renal perfusion is a major risk factor for acute kidney injury (AKI), and plasma volume expansion is therefore crucial for its prevention. On the contrary, among plasma volume expanders, colloids can induce kidney injury, as shown many years ago [1-5]. Recent experiments and clinical studies have supplied further information on the renal toxicity of some colloids, particularly hydroxyethyl starches (HESs) [6-9].

Colloid-induced AKI with morphological abnormalities of the proximal tubular cells, or osmotic nephrosis, has been reported after the infusion of low-molecular-weight dextran or, more recently, HES. The tubular lesions reflect the accumulation of proximal tubular lysosomes due to pinocytosis of exogenous osmotic solutes (for example, mannitol, sucrose, iodinated contrast media, or colloids) [10]. The tubular cells swell because they contain numerous lysosomes and endocytotic vacuoles. Furthermore, the oncotic force of colloids may induce further renal function impairment by decreasing the renal filtration pressure [3]. The exact mechanisms of

colloid-induced AKI remain incompletely elucidated, and controversy exists regarding the relative roles for morphological and functional changes [4].

Because HESs are widely used and AKI is strongly associated with decreased survival, the risk of AKI associated with various HESs needs to be determined. Three generations of HES have been developed over time. Newer HESs have lower molecular weight and lower degree of substitution, two changes that should decrease accumulation and toxicity [11]. Third-generation HESs have molecular weights lower than 200 kDa and degrees of hydroxyl substitution lower than 0.5, the most common combination being 130/0.4. First-generation and second-generation HESs have been found to induce AKI in heart surgery patients, in brain-dead organ donors, and in patients with sepsis [6,7,9,12]. The most recent randomized controlled trial, which included a large number of patients, showed a higher incidence of AKI with 10% HES 200/0.5 than with Ringer lactate solution in intensive care unit patients with sepsis [6]. Both the administration of large volumes and high *in vitro* colloid osmotic pressure (COP) may contribute to renal toxicity. No large randomized controlled trial establishing the safety of third-generation HESs is available.

In the previous issue of *Critical Care*, Hüter and coworkers report an interesting experiment aimed at improving our understanding of HES-induced AKI [1]. Using hemodilution in a model of isolated kidney perfusion, they assessed the renal effects of one second-generation HES solution and one third-generation HES solution comparative to a crystalloid. Although their model was very different from the clinical situation, and the number of studied animals was limited, morphological studies of the kidneys yielded useful infor-

AKI = acute kidney injury; COP = colloid osmotic pressure; HES = hydroxyethyl starch.

mation. After *in vivo* hemodilution, creatinine clearance was higher with the crystalloid than with either HES. As expected, the glomerular filtration pressure was much higher with the crystalloid. For a similar mean arterial pressure, the COP was considerably lower after crystalloid infusion and was similarly increased with both HESs. Although the two HESs had different *in vitro* COP values, their *in vivo* COP effect was similar. The rapid *in vivo* degradation of HES 130/0.4 usually results in a high plasma COP, as illustrated here. After isolated kidney perfusion, the plasma COP remained similar in the two HES groups, but creatinine clearance was lower with the second-generation 10% HES 200/0.5. This result suggests an additional role for a delayed decrease in glomerular filtration, independent of filtration pressure. Morphological examination showed that the lesions of osmotic nephrosis were more severe in the two HES groups, despite the limited volumes infused. Tubular lesions appeared as early as 6 hours after exposure and were associated with greater severity of the interstitial inflammation and tubular dysfunction in the 10% HES 200/0.5 group.

The results of this experiment have implications for clinical practice. First, the *in vivo* COP of the fluid used may have an early effect on the glomerular filtration pressure, as recently suggested in patients resuscitated for shock [8]. With polydispersed macromolecules such as HES, the *in vivo* COP differs from the *in vitro* COP. Also, third-generation HESs can induce osmotic nephrosis similar to that seen with older compounds, within a few hours of exposure. Recent systematic reviews have alerted clinicians to the renal toxicity of HES [13,14]. In the absence of large randomized controlled trials, doubts about the safety of third-generation HES persist, and the results reported by Hüter and colleagues leave room for concern about the safety of widespread use of third-generation HES. Furthermore, HESs have been reported to induce not only AKI, but also irreversible chronic renal failure [10,15,16].

The question therefore is should HESs or other colloids ever be used for fluid resuscitation? There is no evidence from randomized controlled trials that colloids improve patient outcomes [17]. Thousands of patients included in randomized controlled trials have been safely resuscitated using only crystalloids [17]. Furthermore, the study by Hüter and coworkers shows that colloid toxicity and the risk of colloid-induced AKI can be assessed experimentally before colloids are considered for use in humans.

Competing interests

The authors declare that they have no competing interests.

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