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PLASMA FREE HEMOGLOBIN CLEARANCE BY HEMOCOR HPH®  
HEMOCONCENTRATOR AND OTHER COMPETITIVE DEVICES

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Increased plasma hemoglobin (Hb) is often observed in bypass surgery and ECMO patients. Hb, after exceeding the capacity of its detoxifying system, triggers many pathologic reactions and poses a high risk to coronary artery endothelium. Our previous study showed ineffectiveness of cellulose-based hemodialyzers in Hb removal. The present *ex vivo* study examined free Hb clearance by clinically accepted polysulfone-based hemoconcentrators; Hemocor HPH® 1400 (Minntech Corp.), Capiox® HCO5 (Terumo Corp., Tokyo, Japan) and Senko CF11 (Senko Medical Trading Co., Tokyo, Japan). All experiments mimicked the 2-hour human bypass surgery procedure, scaled down to a 1.5 L model, using standard blood and ultrafiltrate flow rates with sampling times: 0.0, 0.25, 0.5, 1 and 2 hours. Baseline Hb level in the hemoconcentrator circuit was  $0.7 \pm 0.03$  g/dL. Results indicate that Hemocor HPH® was superior to the other tested devices. While Hemocor HPH® eliminated more than 90% of Hb after 2 hours ( $y = 1.374 \times X^{(1.022)}$ ,  $R^2 = 0.986$ ), the Capiox® HCO5 and Senko CF11 cleared less than 16% ( $p < 0.001$ ,  $y = 0.498 \times X^{(0.875)}$ ,  $R^2 = 0.982$ ) and 48% ( $p < 0.001$ ,  $y = 0.733 \times X^{(0.940)}$ ,  $R^2 = 0.993$ ), respectively. No tested polysulfone membranes affected the physico-chemical properties of Hb with respect to its tetrameric conformation, heme stability and oxidation, and ferryl-Hb species formation.