Nebuilsed exogenous natural surfactant after cardiac surgery

SIR—Pulmonary changes after cardiac surgery (CS) with cardiopulmonary bypass (CPBP) are similar to those in adult respiratory distress syndrome (ARDS). Exogenous natural surfactant (ENS) seems a promising approach for the treatment of ARDS.¹ We investigated nebulised ENS in 6 patients (mean age 66 [SD 5]; 2 valve replacement, 4 coronary revasularisation) after C9 with CPBP. All patients were mechanically ventilated and under continuous nebulisation for almost 6 hours with ENS (30 mg/kg bodyweight) via a jet nebuliser. An arterial catheter monitored mean blood pressure and the arterial oxygen pressure. The arterial/alveolar oxygen ratio (a/A PO₂) was calculated. Intravenous volume of colloidal fluid transfused was estimated in mL/m². The prenebulisation (PreNeb) and 2 hours post-nebulisation (PostNeb) values were compared:

Patients	a/A PO ₂ *		Blood preassure (mm Hg)†		Colloidals
	PreNeb	PostNeb	PreNeb	PostNeb	(mL/m²)
1	0.45	0.72	80	58	1818
2	0.60	0.83	80	71	700
3	0.26	0.55	71	61	633
4	0.39	0.61	61	79	523-25
5	0.67	0.58	78	94	1111 11
6	0.72	0.76	115	73	1052

Paired t test: *p = 0.02, †p = not significant.

Because of a slight hypotensive effect of ENS in this group of unstable postoperative patients, colloidal fluids were perfused. The fluid requirements of these 6 patients during the ENS nebulisation was 806 (581) mL/m²—therefore higher than the mean 595 (225) mL/m² used in postoperative patients under CS with CPBP not receiving ENS.

Walmrath et al² described a redistribution of blood flow from pulmonary shunt areas to regions with normal ventilation with epoprostenol (prostacyclin, PGI₂) nebulisation. The effect of ENS is to open unventilated areas and augment the functional residual capacity with a better static compliance and arterial oxygenation.³ This effect could be compared with the benefit of positive end-expiratory pressure application on mechanically ventilated patients.⁴ Newly ventilated areas generated by ENS prevent pulmonary hypoxic vasoconstriction and facilitate the recruitment of pulmonary vessels. The consequence of pulmonary blood redistribution is hypotension, also described by Hallman et al in an animal experiment.⁵

We consider both ENS and epoprostenol as useful tools for ARDS treatment. Nebulising the two agents together will be the subject of future research.

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