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Near-Infrared Spectroscopy with Vascular Occlusion Test May Not Be the Adequate Tool to Explore Microcirculation in Pulmonary Arterial Hypertension

To the Editor:

Dimopoulos et al¹ likened the microcirculation of stable patients with pulmonary arterial hypertension (PAH) and congestive heart failure (CHF) and matched normal subjects. They found an effect on the microcirculation, measured by near-infrared spectroscopy with vascular occlusion test (NIRS-VOT), in patients with PAH and CHF and a deleterious action of hyperoxia (performed in the PAH group). Despite a very meticulous work and a conclusion that seems fair and pragmatic, their results should be interpreted carefully. I would like to raise several issues.

First, the authors did not comment on the role of hypoxemia in patients with PAH in their study. A partial means of moderating this would have been to assess the extraction of tissue oxygen (O_2) as a surrogate of microcirculatory function.² Elevation of the extraction before VOT would indicate microcirculatory dysfunction with decreased O_2 supply to tissue in local demand-supply dependence. This value can be obtained noninvasively and calculated as follows: microvascular oxygen extraction rate

(μOER) = $(S_{pO_2} - S_{tO_2})/0.5 \times S_{pO_2}$ (where S_{tO_2} is muscle tissue oxygenation). When fit for S_{pO_2} , μOER values were 33.6, 31, 28, and 20.3 for PAH in the basal state and hyperoxia, CHF, and normal subjects, respectively. A discussion of these results (yet more abnormally high values in PAH) and the role of chronic tissue hypoxia would have been interesting. Accordingly, a decrease in O_2 consumption is seen in tissue with chronic hypoxia, termed conformance.³ Are we confronted with a paradoxical response to hypoxia (greed for O_2 , opposite what is expected with conformance) or microvascular dysfunction? Nevertheless, the paucity of data on cardiac output and the role of chronic hypoxia make this derived value questionable. One can also imagine a different behavior of cells in chronic hypoxia, further amending the dynamic curves (ie, O_2 consumption rate, reactive hyperemia time, and time interval) compared with the princeps work during sepsis.⁴

Accordingly, elevated venous pressure has been related to impaired microvascular tissue perfusion.⁵ Thus, a possible modification of post-capillary venous vessel capacitance would obscure NIRS-VOT-derived values. In PAH patients, the role of such a right-sided outflow obstruction has been overlooked in this study. Moreover, from several intrinsic technological limiting factors, NIRS-VOT explores a derivative of the values of tissue oxygenation and may be dependent on the diffusion of O_2 in the context of hyperoxia.

I think that it may be reasonable to conclude that those patients with PAH likely had a microcirculatory dysfunction, potentially exacerbated by hyperoxia. However, a validation of NIRS-VOT to explore the microcirculation is required in this context of chronic hypoxemia and right-sided overpressure before more formal conclusions can be made. A joint (or with replacement of NIRS) assessment by sidestream dark field imaging would be interesting.

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Near-Infrared Spectroscopy With Vascular Occlusion Test May Not Be the Adequate Tool to Explore Microcirculation in Pulmonary Arterial Hypertension—Reply

In Reply:

We thank Dr Champion for his particular interest in our study and his thorough comments. However, we would like to share our concerns related to his comments.

In our point of view, we should use the microvascular oxygen extraction rate with caution, as it is not yet a valid accurate index to use. This index does not take into account all of the factors that influence oxygen delivery, such as cardiac output, serum hemoglobin levels, and hemoglobin dissociation curve, and to the best of our knowledge, it has not been investigated previously in patients with chronic hypoxemia.

Resting tissue oxygen saturation (S_{tO_2}) values depend mainly on the ratio of tissue oxygen delivery to tissue oxygen extraction. In our study,¹ pulmonary arterial hypertension (PAH) patients presented with a relatively normal cardiac index (median value of 2.5 L/min/m²) and normal serum hemoglobin levels (median value of 16 g/dL) with mild hypoxemia (S_{pO_2} median value of 94%), the latter being a possible factor that might slightly decrease tissue oxygen supply. However, tissue oxygen delivery de-