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Primary and secondary Raynaud's phenomenon: does small-neuropathy remain the same?

Dear Sir,

We took great interest in reading Manek and Coll paper's [1], who hypothesize the presence of a small-fiber neuropathy in primary Raynaud's phenomenon (RP). Indeed, we reached similar results testing 12 patients with secondary RP in systemic sclerosis (SSc) when compared to a group of healthy controls [2]. In these patients, cardiovascular reflexes tested by tilt-test, Valsalva manoeuvre, deep breathing and sustained handgrip were normal but the cold face test was abnormal in almost all patients. By this test we explored the function of afferent cutaneous C and A δ fibres instead of using the cold pressor test, which is painful and is difficult to perform in SSc patients with severe sclerodactyly. In healthy people a cold stimulus applied to the face during spontaneous breathing evokes bradycardia and peripheral vasoconstriction as measured by increased systolic and diastolic blood pressure [3]. Our SSc patients with secondary RP showed blunted bradycardic and hypertensive response to cold-face test, suggesting that a dysfunction of epidermal small fibers (C-fibers) may be involved in SSc.

Manek observations as well as ours are very preliminary and the number of patients is too low to definitely prove a unique and characteristic small-fiber neuropathy in the RP.

However, assuming there is a small-fiber neuropathy in both primary and secondary RP, the question arises as to whether the same fibers are involved or to the same extent in both conditions. There are several observations supporting the idea that this may not be the case.

Autonomic c-fibers around sweat glands are deep, mostly located between dermis and subcutaneous tissue. Skin biopsy in SSc with secondary RP showed loss of sensory

neuropeptidic fibers sprouting from dermal papillary layer into the epidermis, whereas a similar picture but with less severe loss of fibers was described in primary RP [4,5].

Peripheral neuropathy is intriguingly common evidence of very different diseases such erythromelalgia [6], connective tissue diseases [2,5,7,8], and RP [1,2,4,5,9]. It seems unlikely that the same postganglionic neuropathy is involved in all these conditions, unless the small fiber neuropathy is considered as a sign of neuropeptidic dysfunction that may manifest in different pathophysiologic ways thereby leading to a different clinical presentation.

Further studies are needed to clarify this issue although the number of converging results not due to chance is increasing, possibly opening up new avenues of therapies for these diseases.

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