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Introduction: Autonomic dysfunction (AD) in multiple sclerosis (MS) has been recently reported, mostly affecting bladder, bowel, sexual and sweat organs. Orthostatic dizziness has been found in 50% of MS patients. Until now, studies had focused more on cardiovascular reflex testing and blood pressure responses. Autonomic manifestations may be subclinical when abnormal sympathetic skin response or decreased heart rate variation is estimated. We report 28 MS patients with discoloration in lower extremities which was not associated with vascular or other concomitant disease.

Objective: To describe a group of MS patients who developed distinctive discoloration in distal lower extremities.

Methods: Lower limb discoloration was noticed in MS patients during their follow up visit at the Maxine Mesinger Comprehensive Care Center. Epidemiologic data, treatment history, type of MS and disease duration were documented. Patients with concomitant disease (DM, vascular insufficiency) were excluded. Complete physical and neurologic exam were performed. Some patients were evaluated with arterial doppler.

<i>Demographics Table 1</i> Patients n 28		Current treatment Table 2		Graph 1 Disease duration
Women	25	Copaxone	9	
Men	3	Tysabri	5	
Caucasian	24	Avonex	3	
Hispanic	3		□ >10 years	
African American	1			■ 5-10 years ■ 3-5 years
Type of MS		Methotrexate	3	Number of patients
RRMS	21	Betaseron	2	
SPMS PPMS	6 1	Not currently treated	4	

Results: A total of 28 MS patients were enrolled, 25 women with a mean age of 52. Demographic information presented in table 1. 80% were currently on treatment with a disease modifying agent. All patients demonstrated distal discoloration of lower extremities; peripheral pulses were normal in all patients. Coloration did not change with leg raising. 11.5% had dysautonomic symptoms. 3 patients were evaluated with arterial doppler evaluation: all were normal. Current treatments are presented in table 2. Disease duration is presented in graph 1. EDSS ranged between 2 to 7 with an average of 4.5.



Conclusions: AD is seldom recognized by MS patients; it can cause paroxysmal arrhythmias, recurrent syncope, neurogenic pulmonary edema and decreased ventricular ejection fraction, leading to increased morbidity and mortality. MS can involve any part of the CNS, including critical areas subserving autonomic function causing interference of descending autonomic pathways during their course in the brainstem or spinal cord. Demyelinating lesions located among central thermoregulatory pathways may result in regional or global anhydrosis in patients with MS. Skin changes could be a manifestation of AD which may reflect an impact on the quality of life in MS patients.

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