Cognitive impairment screening in a cohort of multiple sclerosis outpatients: demographic characteristics, psychological features and relation to hospital care.

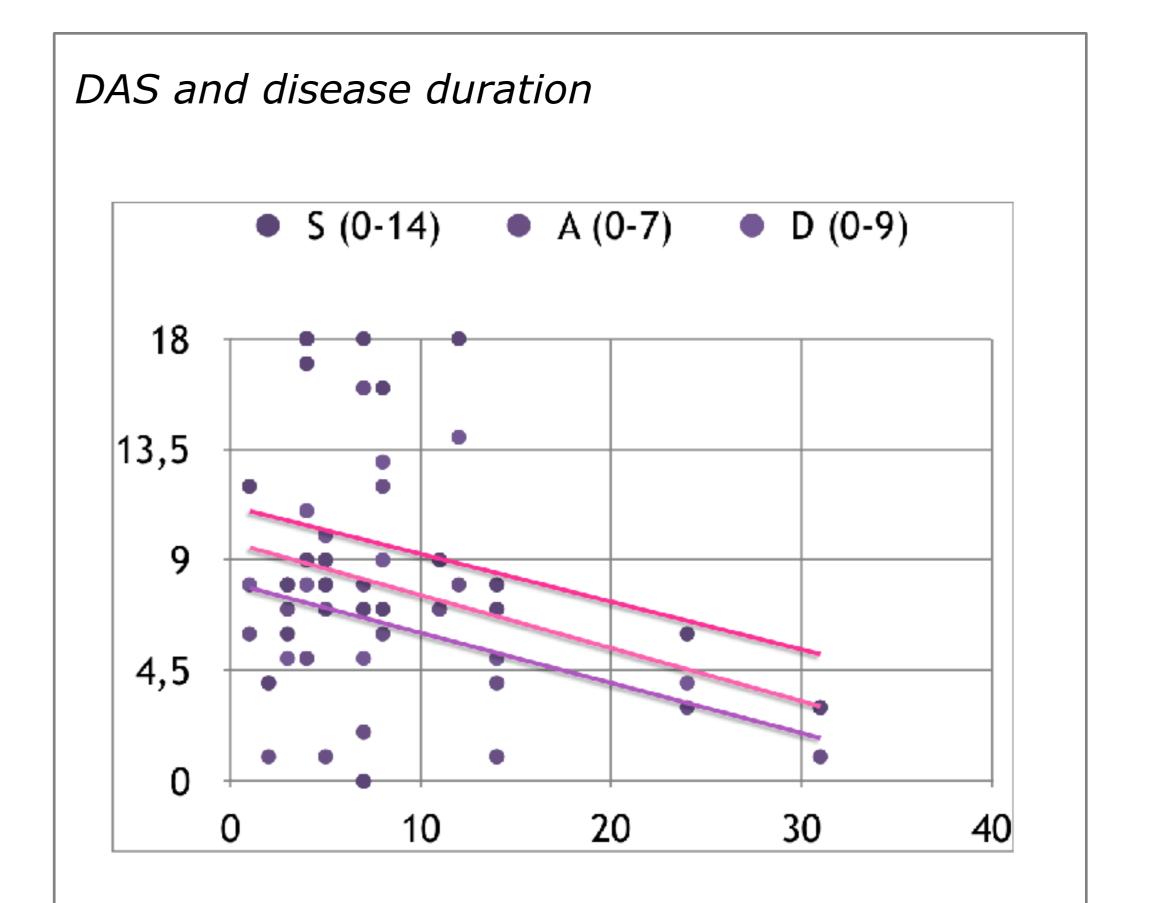
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Introduction

We screened a consecutive population of outpatients of the MS service for cognitive decline and emotional disturbances using a single prospective protocol. Purpose of the study was to assess factors associated with abnormal MoCA scores.

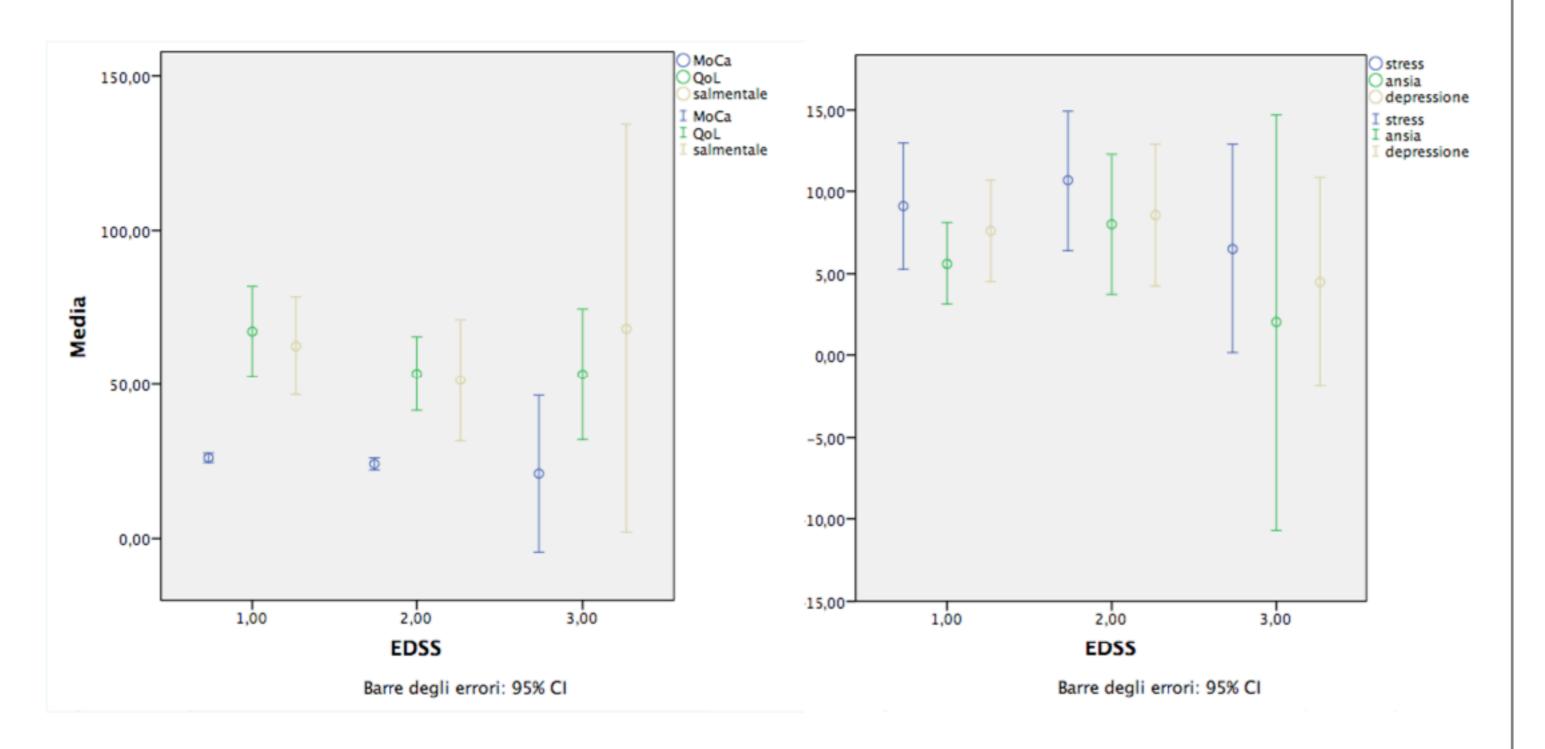
Methods

Materials: We interviewed in 3 months 25 patients of the 235 followed up, 3 of them were excluded because of language barriers or refusal. A formal interview was done using MoCA, DASS-21, MSQOL-54, EDSS, 9-HPT, T25FWT. Methods: Statistical analysis was done using Mann Whitney U test for qualitative MoCA scores using a cut off score of 26, while one-way ANOVA was used to test potential association for quantitative variables , Pearson's r was used as correlations screening. The study population consisted of 20 MSRR outpatients with median EDSS 3. Median age 42.5 yrs. Scholarity 13 yrs. Seven years of median disease duration.

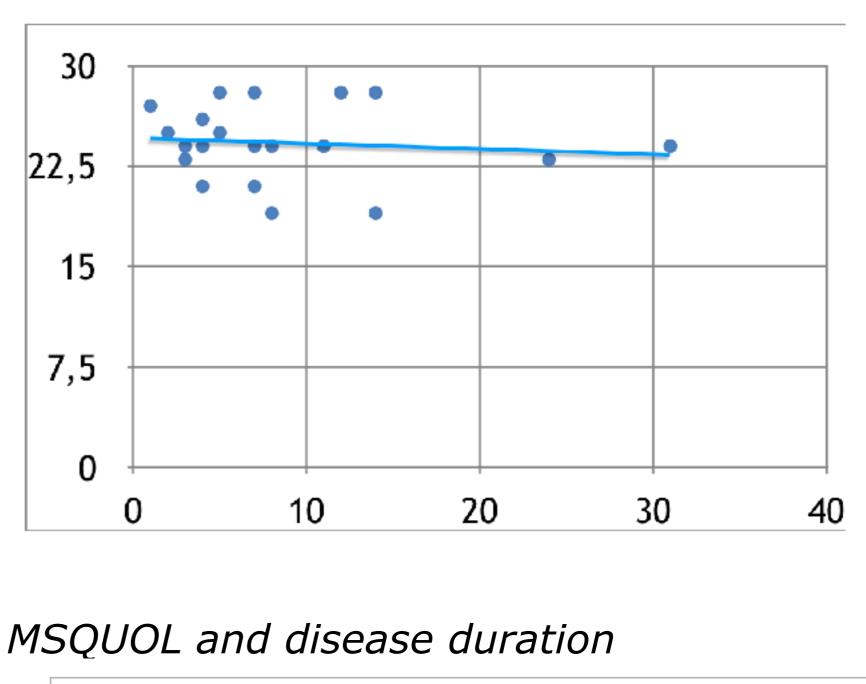


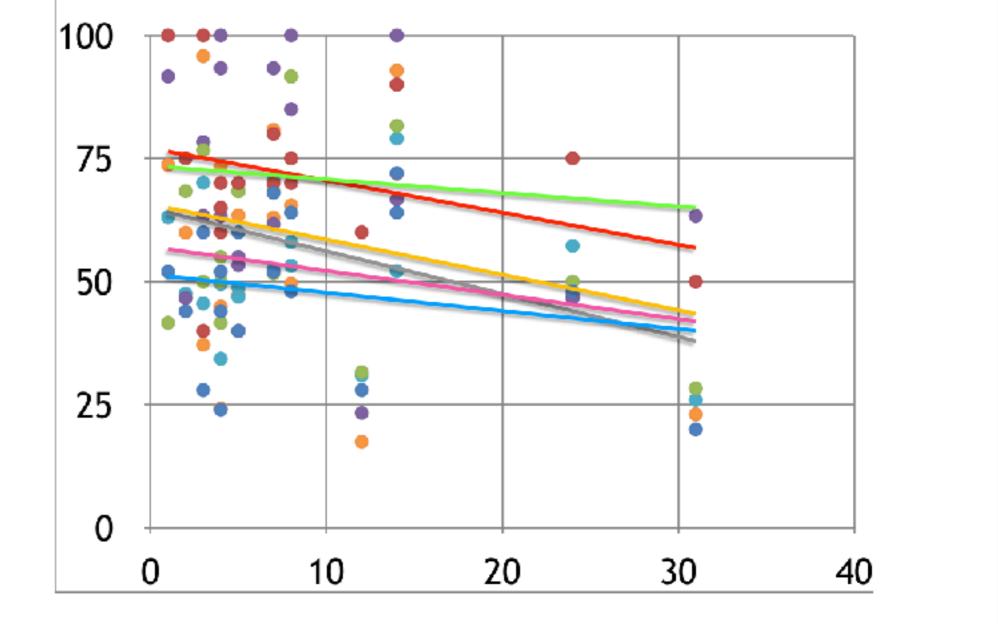
Results

The mean values of corrected MoCA scores were 24.5 (SD 2.84); DASS-21 Stress: 9.52, Anxiety: 6.29, Depression: 7.71; MSQOL -54: Mental health: 59.29, Physical health 54.3. We found a significant correlation according to Pearson's R (p<0.01) between MoCA scores and QoL scores. We found no correlation between MoCA scores and perceived cognitive functioning according to MSQOL-54. One-way ANOVA demonstrated a protective effect on MoCA scores in subjects with ongoing Natalizumab infusional therapy.



MOCA and disease duration





Conclusions

Cognitive decline is often underdiagnosed in many cases of multiple sclerosis. The 65% of our outpatients suffers of a misdiagnosed cognitive impairment. The patient perception of his personal cognitive functioning is seriously compromised. Conclusions: specific screening campaigns should be provided in order to minimize the misdiagnosis of cognitive decline. We found a protective effect of specific disease modifying therapies as already reported in literature.

Bibliografia

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