The effect of vitamin D in the evolution of experimental autoimmune encephalomyelitis

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Abstract
Vitamin D3 deficiency is associated with an increased risk of multiple sclerosis (MS) and unfavorable MS disease progression. Here we demonstrated immunomodulatory effect of vitamin D in Experimental autoimmune encephalomyelitis a experimental model for studying MS.

Key words: multiple sclerosis, vitamin D, EAE.

Introduction
EAE is an antigen-driven autoimmune model in which immunization against myelin autoantigens elicits strong CD4 T lymphocyte responses, which initiate its pathology with central nervous system myelin destruction. Material and Methods: EAE was induced by immunization with MOG35-55 peptide emulsified in complete Freund adjuvant. The clinical expression of the disease was graded on a clinical scale 0-5 according to the severity of the disease. Vitamin D3 (cholecalciferol D3) Sigma Aldr. Mo, USA) was diluted in polyethylene glycol and give orally (5µg/Kg/day) during 2 weeks. The control group was fed with vehicle alone (15 animal/group were studied in three independent experiments. The expression of cytokine mRNA was evaluated by quantitative RT-PCR.

Results and Discussion
We demonstrated that the oral administration of vitamin D reduced significantly the severity of EAE. The reduction of the severity of the disease was accompanied by the reduction of proinflammatory cytokines such as IFNγ and increase of antinflammatory cytokines such as IL-10 and IL-27.

Conclusions
We conclude that vitamin D3 presents immunomodulatory effect in the EAE model. The immunosuppressive mechanism may inhibit the migration of autoreactive T lymphocytes into central nervous system. Moreover, the study suggest that supplementation with vitamin D3 may be beneficial to patients with multiple sclerosis, that present a functional vitamin D receptor.

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¹Reference: ²Vitamin D3 Induces IDO+ Tolerogenic DCs and Enhances Treg, Reducing the Severity of EAE

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