

Vitamin D supplementation for the prevention or depletion of side effects of therapy with alemtuzumab in multiple sclerosis

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Purpose of review: Not only the multiple sclerosis specialist but also the general neurologist and primary care practitioner are increasingly aware of possible adverse events (AEs) by treatment with alemtuzumab (over 47% risk of secondary autoimmune-mediated diseases). Vitamin D supplementation's effect (VDS) to reduce these autoimmune AEs is poorly performed in routine practice. This article seeks to justify why this simple, inexpensive, patient-friendly therapy should be seriously discussed.

Recent findings: Patients who have developed autoimmunity also show a high basal level of IL-21, a cytokine which increases the growth of auto-reactive T-cells. For side effects such as thyroid dysfunction, autoimmune thrombocytopenia, autoimmune hemolytic anemia, autoimmune hepatitis, diabetes mellitus type 1, and alopecia areata/alopecia totalis, VDS may have an impact on the immunological mechanism, in particular lowering levels of IL-17 and IL-21.

Summary: The potential role of vitamin D in influencing autoimmune diseases is evident. If a life-threatening side-effect can be prevented by high-dose VDS, it is ethical to initiate this add-on therapy despite contradictory results in studies on the effectiveness of VDS.

Keywords: alemtuzumab, adverse events, hemolytic and endocrine diseases, autoimmune hepatitis, vitamin D supplementation

Introduction

Alemtuzumab (AI) is a humanized monoclonal antibody anti-CD52 that effectively depletes lymphocytes and has been approved for the treatment of adult patients with relapsing-remitting multiple sclerosis (RRMS) with active disease and is approved in more than 70 countries. AI acts by targeting CD52, an antigen primarily expressed on T and B lymphocytes, resulting in their depletion and subsequent repopulation. MS is an inflammatory condition, which is believed to be autoimmune in nature. Treatment with AI has been shown to increase the risk of secondary autoimmune-mediated conditions, particularly thyroid disorders (mainly AI-induced Graves' disease [GD]), including AI-associated (secondary) autoimmune thrombocytopenia (AsITP), or rarely autoimmune hemolytic anemia (AIHA), autoimmune neutropenia, red cell aplasia and single cases of Goodpasture syndrome (antiglomerular basement membrane disease) as well as membranous nephropathy. An extraordinary observation in the use of AI in MS is the occurrence of acquired autoimmune disease, which is not observed in other conditions treated with this drug – this indicates a disease-specific phenomenon.¹

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The AI therapy of RRMS requires at least 4 years of surveillance. So far, a drug prophylaxis of these serious side effects is not established in practice.

Baker et al have made a hypothesis after analyzing the T- and B-cell repopulation after AI treatment, regarding the induction of secondary autoimmune diseases. The B-cell repopulation occurs much faster than that of the T-cells, especially the regulatory T-cells. Reconstitution of B-cells without adequate control (hyperrepopulation) by regulation of T-cells could be considered as the triggering cause of GD, Hashimoto's thyroiditis, and secondary immune thrombocytopenia (ITP).²

It is evident that vitamin D supplementation (VDS) yields multiple beneficial immunological effects. These positive effects could be used to prevent or mitigate secondary autoimmune diseases.

AI-associated secondary autoimmune thrombocytopenia (AsITP)

AsITP usually develops in 1–3.5% of patients with a latency period of 14–36 months (delayed ITP) after the first AI infusion; there are also reports for the period of 4–51 months. Mild transient decrease in platelet count during AI administration (5-day therapy cycle) is not associated with clinically relevant events and is likely related to the cytokine-release syndrome. Normalization of platelets occurring within 8 weeks without specific therapy has been observed. However, not always self-limiting, fully reversible or predictable thrombocytopenias have been registered.^{3–5} Monitoring in the following months after AI-infusion should be considered good clinical practice.

AI-induced AIHA

AI can be associated with rare but severe AIHA during MS treatment, also in combination with glomerular nephropathies.^{6,7} AIHA is defined as the increased destruction of red blood cells in the presence of anti-red blood cells autoantibodies with or without complement activation. Barcellini investigated the pathogenesis of AIHA and favored a reduced Th1 and a predominant Th2-like profile in the immunopathogenesis of AIHA, in contrast to the prevalent role of CD8+ interferon (IFN)- γ -secreting cytotoxic T-cells in organ-specific autoimmunity, such as MS.⁸

Several immunologic mechanisms are involved in the pathogenesis of AIHA, autoantibodies, antibody-dependent cell-mediated cytotoxicity, phagocytes, B and T lymphocytes,

specific CD4+/CD25+ regulatory T-cells (Tregs), cytokines, and the complement system. Overview by⁸ in the occurrence of two or more autoimmune diseases and MS, the pathophysiology is extremely complex and so far unclear in all details.

Polyautoimmunity in patients with MS – an immunological labyrinth

Numerous publications have shown persons with multiple sclerosis (PwMS) to be susceptible to developing chronic autoimmune diseases alone without AI therapy, suggesting an autoimmune predisposition. Overview by⁹ studies demonstrated a pivotal role of homeostatic proliferation driven by IL-21, arising from mature T-cells having escaped AI treatment and leading to a predominance of oligoclonal, highly proliferative, and chronically activated effector memory T-cells.^{10,11} The tendency to polyautoimmunity is demonstrated by the simultaneous occurrence of ITP and autoimmune thyroid disease following AI treatment in RRMS and simultaneous early-onset severe AIHA and nephropathy.^{7,11} AsITP is associated with a unique form of ITP, is characterized by delayed onset, responsiveness to conventional therapies, and prolonged remission following treatment.¹²

Vitamin D (VD) deficiency – a remarkable pathophysiological problem

VD deficiency has been implicated in the pathophysiology of various inflammatory diseases including MS and hematologic autoimmune diseases.

Mean serum 25 (OH) D3 (VD) levels have been determined to be significant lower in hematological disorders, such as ITP, AIHA, Evan's syndrome, and chronic idiopathic neutropenia (CIN), than in controls.¹³ VD deficiency is very common in children with newly diagnosed or chronic ITP form.¹⁴

Which immunological mechanisms are known in hematological diseases?

Several autoimmune (eg, MS) and hematological disorders share a similar cytokine profile of higher IFN γ -, IL-6-, and IL-17-levels, as was observed at ITP, AIHA, Evan's syndrome, and CIN. IL-21, a cytokine promoter in differentiation of T helper cells (Th) 17, follicular helper T-cell and B-cell may thereby play an important role in controlling the autoimmune process in ITP. Circulating IL-21 levels are significantly higher in ITP-patients than that in healthy

controls.¹⁵ IL-17A and IL-21 induce Th17-cells and inhibit T regulatory (Treg)-cells re-differentiation via Th17-associated signaling pathway in ITP patients in vitro.¹⁶

More recently, Th17-cells, which are characterized by secretion of IL-17, have been identified as key effectors in the development of many autoimmune diseases (eg, MS), including AIHA. Elevated frequency of Th17-cells and increased IL-17 secretion were found closely correlated with the disease activity in AIHA patients.⁸ Higher levels of IL-21 have been shown to be associated with developing autoimmune disease post-AI therapy.¹⁷

Which immunological mechanisms justify a VD supplementation?

It is biologically plausible that sufficient VD levels may be important in modulating inflammatory processes. VD has numerous effects on cells within the immune system. It inhibits B-cell proliferation and blocks B-cell differentiation and immunoglobulin secretion. VD additionally suppresses T-cell proliferation and results in a shift from a Th1 to a Th2 phenotype. Furthermore, it affects T-cell maturation with a skewing away from the inflammatory Th17 phenotype and facilitates the induction of T regulatory cells. This leads to a decline in the production of inflammatory cytokines, such as IL-12, IL-17, and increased production of anti-inflammatory cytokines such as IL-10.^{13,18}

VD also has effects on monocytes, macrophages and dendritic cells. It inhibits production of inflammatory cytokines, such as IL-1, IL-6, IL-8, IL-12, and TNF α .¹⁹ Especially, 1,25-dihydroxyvitamin D3 (1,25(OH)2D3) has potent immunomodulatory properties that have promoted its potential in the prevention and treatment of autoimmune disease.²⁰ Jeffery et al observed that stimulation of CD4+CD25-T cells in the presence of 1,25(OH)2D3 inhibited production of pro-inflammatory cytokines and stimulated expression of high levels of cytotoxic T-lymphocyte antigen 4 as well as FoxP3. It is now established that FoxP3+ regulatory T-cells are critical to the prevention of autoimmunity. The maintenance of a significant population of such cells is therefore required for lifelong health.²⁰

Overall, the majority of cells involved in hematological diseases express a high level of vitamin D receptor (VDR). This suggests that these cells may be responsive to VD treatment.²¹ In ITP, AIHA and CIN VD receptor(R) expression levels were higher in the disease state compared to normal donors.¹³

From a prophylactic point of view, to avoid AsITP, VDS should be carried out to an optimal 25(OH)D

serum level. Benefits were demonstrated by VD administration in ITP.^{13,22,23} ITP also showed a lower number of platelets in cases with very low VD levels, and Liu et al also observed lower 1,25(OH)2D3 values with active ITP.²⁴

1,25(OH)2D3 has two important immunomodulatory functions: Lower regulation of CD4+ cell overproduction combined with up-regulation of T-regulatory cells (Treg) could therefore reduce the autoantibody response and restore normal platelet levels. 1,25(OH)2D3 as a potent immunomodulator inhibits inter alia the production of the proinflammatory cytokines IL-17 and IL-21, which could be a potential marker for increased autoimmunity in MS, as well as increased production of anti-inflammatory cytokine IL-10. IL-10 plays an essential and highly complex role in the modulation of adaptive immune responses. IL-10 inhibits production of a number of proinflammatory cytokines, including IL-1 β , IL-6, IL-12, IL-18, granulocyte-macrophage colony-stimulating factor, and TNF α . Overview by²⁵ regarding the regulatory role of B-cells and IL-10 production in the resolving of autoimmune diseases, there is currently a significant increase in knowledge. The interpretation of elevated levels of IL-10 is also complicated by the reports of IL-10-independent mechanisms in autoimmune diseases.²⁶ The pleiotropic nature of IL-10 makes it difficult to introduce the benefit of IL-10-modulating therapies into the clinic.

Practical recommendations – AsITP

All therapeutic and diagnostic options must be exploited to prevent or detect timely and rare adverse effects. However, rare and late adverse events (AEs) are often identified only during post-marketing observation.²⁷

VD may very well be a significant factor in preventing the loss of tolerance to self and resultant polyautoimmunity. The optimal VD supplementation results in multiple beneficial immunological effects in multiple sclerosis.²⁸ The demonstrated immunological mechanisms in hematological diseases – ITP, AIHA-demand VD supplementation, especially since the VD application has few side effects. Although VD therapy is initially hypothetical for preventing side effects from AI, its effectiveness should be confirmed to increase patient safety.

If platelet counts of $\geq 50,000/\mu\text{L}$ but $< 100,000/\mu\text{L}$ (two short-term controls after a few days) are registered, immediate initiation of high-dose VD supplementation may be initiated prior to the side-effect rich first-line therapy. These platelet counts could be the putative “window of

therapeutic opportunity” with VD being a meaningful “supportive first line option”. If cutaneous signs of thrombocytopenia occur (petechiae, ecchymoses) and the platelet counts are $<50,000/\mu\text{L}$, equally high-dose VD would be indicated as “first-line co-medication”. The differential diagnosis of thrombocytopenia is facilitated if AI-treated patients refrain from the consumption of quinine-containing drinks and medicines (tonic water, bitter lemon, etc.), as this substance may cause secondary (drug-related) sudden, severe thrombocytopenia.²⁹

Furthermore, there is a correlation of VD deficiency and thyroid autoimmune diseases. Serious VD deficiency ($<10\text{ ng/mL}$) corresponds with Hashimoto’s thyroiditis, GD, and hypothyroidism.

Considerations for clinical practice in hematological AEs

The current state of knowledge gives no reference of a predisposition to PwMS that could lead to developing an autoimmune disease. There are no biomarkers that signal polyautoimmunity which could occur with AI therapy. The results of the studies allow a VDS without restriction. 25 (OH)D3-serum levels are typically lower in PwMS and hematological disease compared to healthy individuals. Diseased cells typically express a high level of VDR. In abnormal hematological cells, VDS reduced the production of pro-inflammatory cytokines.²¹ In patients with autoimmune cytopenias, it was possible to verify these theses: 1) VD is reduced and its receptor increased in autoimmune cytopenias; 2) VD deficiency is associated with ITP and AIHA severity at onset; 3) low VD levels are associated with increased relapse rates in AIHA; and 4) VD inhibits anti-erythrocyte autoantibody production in vitro.¹³ For the prevention of side effects (autoimmune and hematological diseases) under therapy with AI, VDS should be used in particular for AsITP and AIHA. It could affect the severity of the disease, as it has protective and immunomodulatory effects.¹³ The recent case reports on AIHA in RRMS treatment with AI³⁰ could expand the surveillance program by determining the Coomb’s test.^{6,7} Due to the possible life-threatening comorbidities due to an autoimmune side effect, a physician applying this treatment should be cautious. The successful implementation of an automated electronic support system for safety monitoring of multiple sclerosis patients demonstrates its importance. In one case of AsITP, while the treating neurologist was on leave, the automated analysis of the pathology results provided rapid treatment.³¹

Quick recognition of unexpected adverse effects as prevention of life-threatening situations

Secondary hemophagocytic lymphohistiocytosis

Rare and late AEs are often identified only during post-marketing observation. In the monthly AI monitoring program (blood tests), the determination of C-reactive protein (CRP) in the differential diagnosis may be helpful. In the presence of anemia, thrombocytopenia, pancytopenia and elevated CRP, a rare secondary hemophagocytic lymphohistiocytosis may be the cause that has been previously reported in two cases.³² This hyperinflammatory syndrome has occurred among others in autoimmune disease and this syndrome is characterized mainly by fever, pancytopenia, hyperferritinemia, pathological liver counts, raised soluble IL-2 receptor (CD 25), lymphadenopathy, and hemophagocytosis.

Acute acalculous cholecystitis (AAC)

If acute abdominal pain occurs in the upper right quadrant shortly after AI infusion, a differential diagnosis of very acute AAC must be considered. An AAC represents a new and potential life-threatening adverse event. Croteau et al described eight cases with AAC, four cases were assessed as probable while four were possible. The clinical symptoms occurred during or shortly after AI infusion. Although the pathophysiological mechanism has not been fully elucidated, an “acute cytokine release syndrome” may be the cause. One patient experiences an AAC about 45 days after completion of the first cycle of AI infusion.³³

VDS as a further prevention measure of rare side effects of AI therapy

Alopecia areata (AA)/alopecia totalis (AT)

Rare side effects of AI therapy are the occurrence of AA or AT,^{34,35} which comes with a substantial health care burden. Zimmermann et al and Leussink et al provide evidence that AA and AT is an additional, but barely recognized, secondary autoimmune disease after AI application.^{34,35} The predisposition of patients with MS for polyautoimmunity is corroborated by these case histories. AA is a common, recurrent, autoimmune hair disorder.³⁶ Severe forms of AA, such as AT or alopecia universalis are strongly linked to thyroid autoimmunity ranging from 25% to 40% of all cases.^{37,38} The complexity of pathophysiology is demonstrated by the fact that thyroid

disorders predominantly show B-cell but also a T cell mediated autoimmunity. AA is an autoimmune disease resulting from T-cell mediated (Th1 category) damage to the hair follicle.³⁹ CD8+NKG2D+T cells promote AA pathogenesis, acting as cytolytic effectors responsible for autoimmune attack of the hair follicle. IFN- γ produced by CD8 T-cells leads to the collapse of immune privilege in the hair follicle.³⁹ However, the lesional skin in AA also shows a mixed profile – including signs of a classic Th2 disease.⁴⁰ Serum 25(OH)D levels are lower in patients with AA than healthy control samples.^{41,42} These levels were negatively correlated with disease severity and pattern of hair loss. This may provide evidence about the role of VD deficiency in AA pathogenesis. Severe AA showed by far the lowest VD levels compared with cases with mild or moderate disease.⁴³ This result was suggested to be caused by its effects on the hair and the immune system. Expression of VD-receptors in keratinocytes is necessary for maintenance of the normal hair cycle. VD inhibits the secretion of proinflammatory cytokines, including IL-2 and IFN- γ . IL-2 and its receptors play a key role in the proliferation of autoreactive T-cells, but also in the loss of immune tolerance in MS. Because serum 25(OH)D levels were significantly lower in AA, VD supplementation with serum levels in the upper physiological range is also justified here.^{41,42}

Chan et al report on another case of AA and they could register a great therapeutic success by a “therapeutic” coincidence.⁴⁴ Local therapy with triamcinolone acetonide with intralesional injections was not very successful. Because of a relapse, the female MS patient intravenous received 1,000 mg methylprednisolon/day over 5 days. This “therapeutic coincidence” led to a distinct improvement in AA.

AI-induced thyroid dysfunction (AITD)

AI-induced thyroid disease (thyroid AEs combined with abnormal thyroid function test) occurs in up to 40% of patients. AITD can significantly affect the quality of life of MS patients if this side effect is not recognized in time and adequately responded to by all specialist disciplines. Before AI therapy, the TSH-(thyroid-stimulating hormone) level and the thyroid gland should be examined. In the detection of TPO-(thyroid peroxidase)-AB before the start of treatment, the risk of developing an AITD was 69% compared to TPO-negative patients with only 31% risk. Eighty-five percent of patients who developed AITD later

were TPO-AB-negative before therapy, meaning baseline AB-negativity is no protection from AITD.¹

Daniels et al have observed 102 episodes of thyroid dysfunction in 216 AI-treated patients. Seventy-three patients (34%) developed AITD, which manifested in 65.8% of the cohort as GD, in 20.5% as hypothyroidism (HYPOT) and in 12.3% as subacute thyroiditis (SLT).¹ Of 86 patients, Tuchy et al diagnosed 41% with AITD, of which 63% developed GD, 34% developed HYPOT (with positive TPO AB), and one patient with transient thyroiditis.⁴⁵ Cossburn et al reported a high AITD rate of 77%, with 71% developing GD, 17% HYPOT, and 12% developing transient thyroiditis.⁴⁶

Several studies reaffirm that the GD is the most common occurring condition.⁴⁷ For the practical procedure, the reports about the variants of the clinical course and the laboratory parameters of the AITD are helpful. Overt Graves’ hyperthyroidism spontaneously resolved itself in 36.7% of patients, 20.4% became euthyroid, whereas 16.3% became hypothyroidic.¹ The SLT with hyper, euthyroid, and hypothyroid phases develops in about 50% of patients to definitive hypothyroidism.⁴⁸

Between GD and thyroiditis, there are transitional forms and one disease can transition into the other. Unexpected clinical courses that turn the GD into HYPOT both after long antithyroid treatment as well as without antithyroid drugs transformed must be accounted for within the long-term care plan. In the review by Rotondi et al, hyperthyroid patients have an unusually high rate of spontaneous shifts to a hypothyroidism.⁴⁹ In a recent study, thyroid dysfunction was reported in 42% of patients, of which 72% developed GD, 12% HYPO, 6% Hashimoto's thyroiditis, and 5% thyroiditis (defined as thyrotoxicosis followed by spontaneous euthyroidism or hypothyroidism, with negative TSH receptor autoantibodies (TRAbs) and/or absent tracer uptake on technetium scan).⁵⁰ In contrast to previous reports, Pariani et al noted a more frequent incidence of GD requiring definitive or prolonged antithyroid drug treatment.⁵⁰ The cause of unusual clinical observations and changing biochemical markers could be explained by an imbalance of TSH receptor stimulating AK (TSAbs) and TSH-blocking AK (TBAb) is coming.¹

In general, spontaneous remissions of Graves’ hyperthyroidism occur when TRAbs disappear, when TSAbs are balanced by antibodies that block thyrotropin action, or when persistent TSAbs are thwarted by associated autoimmune thyroiditis.^{1,50}

Numerous case-based reports of an AITD document the colorful image with light forms and heavier gradients (especially in smokers) of different responses to the therapy and variants in the course of the disease.⁵¹ In isolated cases of thyroid AB without evidence of subclinical HYPOT or HYPERT and a lack of pathological TSH levels, one check is sufficient.

Relationship between VD serum levels and autoimmune thyroid dysfunction

There is an evident correlation between VD deficiency and thyroid autoimmune disease.⁵²

Although the results from studies are controversial and unresolved questions remain, there is evidence that a correlation exists between severe VD deficiency (<10 ng/mL) and Hashimoto's thyroiditis and GD,^{53–55} as well as hypothyroidism.⁵⁶ Low 25(OH)D3-serum levels (20–30 ng/mL=50–75 nmol/L) were found particularly in young women with elevated TG-AK⁵⁷ and moderately elevated TPO-AK.⁵⁸ On the other hand, low TSH levels were seen in individuals with high VD levels.⁵⁹

If AITD develops in more than 40% of patients after AI therapy, using the current state of our knowledge, it should be discussed whether it would be better to initiate an add-on treatment with high-dose oral VD (eg, 4–10,000 IU/day) before the series of infusions. In addition, for the argumentation of a VD supplementation (14,000 IU/day), the success rate indicates that the number of active lesions in MRI was about 1/3 lower and the volume increase in the T2 lesions was halved. In young MS patients (under 30 years), no new T1 lesions were registered in 86% of cases.⁶⁰

Where it is demonstrable that over 40% of patients develop an AITD after AI therapy and VD deficiency may encourage the development of autoimmune thyroid disease, VDS should take priority in the interests of patient safety.

Considerations for clinical practice in thyroid dysfunction

Hyperthyroid patients (AI-induced GD [reconstitutions GD]) have an unusually high rate of spontaneous shift to hypothyroidism. The remission rate of Graves' hyperthyroidism, both spontaneous and after antithyroid drugs is unexpectedly high.⁴⁹ However, long-term observations also show that GD is more likely to require definitive or prolonged antithyroid drug treatment. For reasons of safety, because of clinically atypical forms, it may be

advisable to carry out the monitoring with annual TSH level determinations over a longer period of time.⁵⁰

Early neutropenia after AI infusion cycles – non-immunologically induced

Increased attention must be given to AI-induced (early) neutropenia. The polymorphonuclear neutrophils also express CD52 and may be depleted by AI, thus potentially contributing to the infections that develop post-AI treatment. The degree of neutropenia was generally mild. Grade 3–4 neutropenia (<1.0×10⁹/L) occurred in <1.5% of PwMS in each AI treatment cycle. Two PwMS developed severe neutropenia-related AEs.⁶¹ Gaitán et al report two cases with early neutropenia. In one case, 4 weeks after standard AI induction, a severe neutropenia was developed with an absolute neutrophil count (ANC) of 470/μL. In the second case, a neutropenia with values of 300/μL was observed in the first AI infusion series. In the second series of AI infusions, a transient neutropenia occurred after 5 weeks (ANC 398/μL) for 6 days.⁶² Galgani et al reported a case of asymptomatic leukopenia with neutropenia detected approximately 1 month after the first AI course with spontaneous resolution.⁶³ A case of early neutropenia with fatal outcomes in a 47-year-old PwMS patient 23 days after AI infusion was described. Weekly blood tests for the first 2 months after the first infusion could prevent major infections.⁶⁴ This “early neutropenia after alemtuzumab infusion” is to be distinguished from a “neutropenia as a result of alemtuzumab-induced secondary autoimmunity” which occurs much later in the context of immune reconstitution.⁶⁵

Diabetes mellitus type I after AI therapy

The tendency to polyautoimmunity in PwMS is confirmed by the occurrence of autoimmune diabetes mellitus type 1 (T1DM) and autoimmune thyroid disease (GD) after AI therapy in three cases.^{66,67} The observation of three autoimmune diseases in PwMS is associated with hyperactivity of the adaptive immune system (T- and B-cells lymphocytes). An adequate VD status has been observed with a decreased risk for T1DM.⁶⁸ VD represents a candidate protective factor for T1DM as it regulates the immune system and autoimmunity. The chance to prevent the T1DM lies in the administration of VD before the onset of AI cycles. The therapeutic window for VD supplementation lies in the period of “prevention”, since the destruction of β-cells cannot be reversed.

Before diagnosis, patients with T1DM had lower 25(OH) D levels than controls.⁶⁹ Patients with various autoimmune

diseases showed a low VD level.⁷⁰ The benefit of adequate VD supplementation was also demonstrated by administration of 2,000 IU/day from the first to the 31st year of age. There, children had a 78% lower risk of developing T1DM compared to this person who did not receive this supplementation.^{71,72} Adults with lower serum 25(OH)D levels were at higher risk for insulin-dependent diabetes than those at higher levels. A 3.5-fold lower risk was associated with a serum of 25(OH) \geq 60 nmol/L. Individuals with a serum level of 25(OH)D > 100 nmol/L had a 70% lower risk of developing insulin-dependent diabetes than subjects below 43 nmol/L.⁷³ There is a significant association between inadequate levels of 25(OH)D and higher levels of cytokine (eg, IFN γ , TNF- α , IL-6, IL-1 β , etc.), high titers of ICA (islet cell antibodies), anti-GAD-AK (glutamic acid decarboxylase antibody) and IAA (anti-insulin antibodies).⁷⁴

VD supplementation up to a dose of 10,000 IE/day must be given to achieve an individual, effective 25(OH)D level, checking VD serum levels and serum Ca levels. Uniform data on the VD daily dose for all PwMS are not possible because various specific genetic errors in the VD metabolism determine the serum VD level (eg, abnormalities of the gene of enzyme 1 α -hydroxylase CYP27B1 [cytochrome P 450 family 27 subfamily B member], SNPs [single-nucleotide polymorphisms] rs 703,842 or rs10877,013).²⁸ Another pathogenesis factor in the development of autoimmune disease is a VDR polymorphism. Genetic VD receptor mutations may result in an alteration of the effects produced by the binding of 1,25(OH)₂ D in the promoter regions of response to VD. The association of numerous SNPs and the risk of islet autoimmunity is increasingly confirmed.⁷⁰ The determination of specific diabetes autoantibodies before the first AI infusion has the advantage of prematurely detecting high risk.

“Surprises” in the repertoire of unwanted secondary autoimmune diseases

Lambert–Eaton myasthenic syndrome (LEMS)/myasthenia gravis

New symptoms in PwMS can easily be misinterpreted when clinical signs are incorrectly attributed to the course of MS. Hoffman et al diagnosed LEMS after AI therapy.⁷⁵ LEMS is caused by autoantibodies direct against calcium channels (anti-VGCC-AK) on the neuromuscular junction. IL-21 drives secondary autoimmunity in PwMS, following therapeutic lymphocyte depletion with AI.¹⁷ Jones et al suggest that serum IL-21 may, therefore, serve as a biomarker for

the risk of developing autoimmunity month to years after AI treatment. Before treatment, PwMS who went on to develop secondary autoimmunity had more than twofold greater levels of serum IL-21 than the nonautoimmune group.¹⁷ Patients with myasthenia gravis had lower 25(OH)-plasma levels⁷⁶ and by a “high-dose vitamin D treatment” in one case a complete remission could be achieved. However, the VD serum levels were in the extremely high range (400–700 ng/mL).⁷⁷ VDS (1,25(OH)D) can attenuate the potential increase of IL-21 levels. Neurological autoimmune diseases as a side effect are a diagnostic pitfall.

Autoimmune hepatitis

The tendency to polyautoimmunity is complemented by publications on autoimmune hepatitis (AIH) in two cases after the second AI cycles and simultaneous manifestation of a Grave’s disease.^{78,79} Canham et al diagnosed an autoimmune hepatitis in a 43-year-old Caucasian female after the second infusion-AI cycle 1 year later.⁸⁰ There is an increase in the prevalence of AIH in PwMS compared to the general population (0.17–1% vs 0.02%).^{81,82} In terms of differential diagnosis, AIH must be distinguished from a drug-induced liver injury caused by AI, which can be confirmed by re-exposure.⁸³ As a supplement, liver injury after pulsed methylprednisolone therapy in PwMS with RRMS has been observed both as acute drug-induced liver injury and as AIH.^{84,85} AIH was observed in 40 reported cases in PwMS receiving disease-modifying drug therapy.⁸⁶ However, patients with AIH have a high prevalence of VD deficiency.⁸⁷ Low serum VD levels are associated with severe histological features and poor response to therapy in patients with autoimmune hepatitis.⁸⁸ 1,25-dihydroxyvitamin D can inhibit immune cell proliferation, promote an anti-inflammatory cytokine profile, expand regulatory T-cells, enhance glucocorticoid actions, increase glutathione production, and inhibit hepatic stellate cells. Loss of VD-dependent homeostatic mechanism may promote disease progression.⁸⁹ Severe VD deficiency is associated with treatment non-response, progression to cirrhosis, and liver-related death or need for liver transplant. Severe VD deficiency is a prognostic biomarker in AIH.^{90,90} VD supplementation initially requires a higher VD dose (6,000–10,000 IU/day) to rapidly achieve an effective serum level (at least 30 ng/mL), followed by a maintenance dose of 3,000–6,000 IU/day). These high doses are necessary because the AIH has excessive systemic inflammation at the time of diagnosis, supports obligatory

glucocorticoid therapy and may have a “glucocorticoid saving effect”.^{90,91}

The general importance of the daily dose of VD is pointed out by the study of O’Connell et al⁹². The CISAVID trial (“Dose-related effects of vitamin D3 on immune responses in patients with clinically isolated syndrome” [NCT01728922; interventional, double-blind randomized placebo-controlled trial]) which also aimed to examine the immunologic effects of VD supplementation at two doses (5,000 IU or 10,000 IU/day) in patients with CIS over a 24-week treatment period. This study from Ireland has been completed, but no results of the study have been reported yet.⁹³ The dose–response effect is decisive for the success of the therapy.

Rare side effects – often a diagnostic puzzle?

The importance of polyautoimmunity is demonstrated by the case report of the simultaneous occurrence of GD and acquired hemophilia A (AHA). AHA is extremely rare in patients treated with AI (0.2% in clinical trials).⁹⁴ For bleeding, both the platelet count and the routine coagulations test (prothrombin time) and active partial thromboplastin time) should be included in the laboratory serological monitoring^{95,96} madeley of patients treated with AI.

In the presence of thyroid antibodies and manifestation of new neurological symptoms (eg, paresthesia, epilepsy) autoimmune encephalitis may rare develop. Giarola et al report a case of autoimmune encephalitis (“Hashimoto’s Encephalopathy”) manifesting as a polymorphic epilepsy partialis continua/status epilepticus 7 months after the second course of AI in a patient with previous autoimmune hypothyroidism and immune thrombocytopenic purpura.⁹⁷ Metz et al report a case of disseminated necrotizing leukoencephalopathy and severe AIHA 8 months after AI treatment.⁹⁸

Vitiligo

Ruck et al describe three cases of AI-treated patients with RRMS developing vitiligo 14, 18, 52 months after AI initiation (T-cell-mediated secondary autoimmune disease with increased IL-21).⁹⁹ Eichau et al diagnosed another case of vitiligo in a 28-year-old female patient 5 months after the first AI cycle.¹⁰⁰ Vitiligo and alopecia are both autoimmune diseases. VD represents a potential player in the pathogenesis of vitiligo. Low 25-hydroxyvitamin D levels are associated with vitiligo.¹⁰¹ IL-17 was significantly higher whereas VD was found to be lower among the patients. A

significant positive correlation was noted between VD levels and disease duration.¹⁰² Zhou et al showed increased circulating Th17-cell frequencies and elevated serum IL-17A, TGF- β 1, and IL-21 levels in patients with non-segmental vitiligo.¹⁰³ In a pilot study, 16 patients with vitiligo received 35,000 IU VD/day for 6 months in combination with a low calcium diet. Avoidance of milk products and calcium-enriched foods as well as consumption of oats, rice, and soy milk is necessary. A fluid intake of at least 2.5 L per day had to be maintained. Fourteen of 16 vitiligo-patients showed a repigmentation of 25–75%. All patients presented low VD status (serum 25(OH)D \leq 30 ng/mL).¹⁰⁴

Myalgia as an indication of secondary autoimmune myositis

Aouad et al report on a first case of secondary myositis after AI therapy. Six months after the induction therapy with AI a 44-year-old patient with RRMS myalgia affecting both arms and legs in a predominantly proximal distribution developed. The blood test showed an extremely high level of creatine phosphokinase (CK) (>50-fold). The complete regression of the symptoms and the increased CK after oral glucocorticoid therapy speak for an autoimmune genesis.¹⁰⁵ Azali et al could prove low serum levels of VD in idiopathic inflammatory myopathies (polymyositis, dermatomyositis, inclusion body myositis).¹⁰⁶

Serum IL-17A level was high in patients with dermatomyositis and polymyositis and just as serum IL-6 and IFN γ but also associated with disease activity.¹⁰⁷ VDS is being considered as an adjuvant therapy. The supplementation of VD can also reduce the synthesis of inflammatory cytokines (IL-21, IL-17) and increase the synthesis of anti-inflammatory cytokines.

The knowledge of the tendency toward polyautoimmunity is the key to the exact interpretation of symptoms caused by unwanted side effects

Sarcoidosis

Tyshkov et al confirm the trend toward polyautoimmunity through the report of comorbidity of sarcoidosis in 10 PwMS.¹⁰⁸ Systematic sarcoidosis can either follow or precede the diagnosis of MS or occur simultaneously.^{109,110} In addition to these publications, Willis et al report on three cases of sarcoidosis after two AI cycles over a period of 2–4 years. These also showed comorbidity with hypothyroidism, GD, and ITP.¹¹¹ Graf et al observed acute sarcoidosis

(Löfgren's syndrome) 1.5 years after the second AI cycle and thus demonstrated the invasive effects of AI on the immune system.¹¹² IFN- β -induced sarcoidosis has also been reported repeatedly.¹¹³ However, VD administration is not unproblematic. Up to 10% of cases of sarcoidosis develop hypercalcemia.^{114,115} In patients with sarcoidosis, hypercalcemia occurs as a result of the activity of ectopic 25(OH)D-1-hydroxylase (CYP27B1) expressed in macrophages.^{114,115} The constellation of elevated 1,25-dihydroxyvitamin D and low 25(OH) D serum values may be possible, the latter being used to control overdose.¹¹⁶ In 40–60%, there is a hypercalciuria.¹¹⁷ There are controversial opinions on VD supplementation as add-on therapy for sarcoidosis.¹¹⁵ Miedema et al demonstrate a key role of Th17-cell plasticity in granuloma formation and maintenance.¹¹⁸ VD supplementation would affect the disease by suppressing the activity of Th-17cells and Kamphuis et al recommend supplementation because there was a significant negative correlation between 25(OH)D serum levels and disease activity.¹¹⁹ VD increases the therapeutic effects of glucocorticoids via an mTORc1-dependent upregulation of the glucocorticoid receptor.¹²⁰ The risk of osteoporosis in the primary therapy of sarcoidosis with a glucocorticoid could be prevented by VD supplementation.

It is challenging to assign an absolute serum VD concentration over which toxicity is always present.¹⁰³ However, serum total 25(OH)D3 concentrations >80 ng/mL (200 nmol/L) should be a warning mark on monitoring, only concentrations typically severalfold higher than 80 ng/mL showed hypercalcemia.¹⁰³ Monitoring by determination of serum calcium and phosphate is required.

A study on VD homeostasis in sarcoidosis is currently being conducted. This study evaluates the relationship between vitamin-D status and severity of sarcoidosis, and the effects of vitamin-D repletion in vitamin-D insufficient patients with sarcoidosis.¹²¹

Conclusion

Vitamin D, as an immune regulator, can modulate the immune system via endocrine, paracrine, and intracrine mechanisms. Extensive epidemiological research supports the causality of the link between low VD values and the occurrence and development of MS and autoimmune diseases. PwMS who developed autoimmune secondary diseases after AI treatment as side effects showed high basal levels of IL-21, a cytokine that increases the growth of autoreactive T- cells. Serum IL-21 levels may serve as biomarkers for the risk of developing autoimmunity months to

years after treatment with AI.¹⁷ VD is one of the factors that can regulate the function of Treg cells. Therapy with VD is inexpensive and without significant side effects. The mechanisms of polyautoimmunity with its “immunological labyrinth” are currently unclear, as are the exact physiological influences of VDS. However, randomized clinical trials testing VDS on polyautoimmunity are not possible due to virtually insurmountable practical difficulties.

Escasany's demands can only be supported by the fact that new side effects that have occurred in the meantime must be communicated to the doctors in good time, especially as not all side effects are published. Although life-threatening adverse events are rare in long-term monitoring and there are no markers prior to alemtuzumab therapy to detect the possibility of secondary autoimmunity,¹²² interdisciplinary collaboration is the key to detecting rare adverse events. In the case of polyautoimmunity with rare combinations, e.g. MS and myasthenia gravis, a differential diagnostic distinction must be made between “natural occurrence of such coexistence” and secondary autoimmunity in the context of alemtuzumab treatment.

It is very likely that for this problem of side effects of AI therapy, no randomized clinical trials with VDS testing are possible in the short term. For side effects, such as thyroid dysfunction, autoimmune thrombocytopenia, AIHA, acquired hemophilia A, diabetes mellitus type 1, autoimmune hepatitis, Lambert–Eaton myasthenic syndrome, sarcoidosis and AA/AT, vitiligo, or autoimmune myositis, VDS may have an impact on the immunological mechanism, in particular lowering levels of IL-17 and IL-21. The daily intake of VD and the attained 25(OH)-serum level are decisive for the success of the therapy (dose–response effect). In daily practice, it should be noted that PwMS with oral VD supplementation has a lower increase in 25(OH)D serum levels (genetically caused hypovitaminosis D) than healthy individuals. The therapeutic potential of VD supplementation can only be used if regular monitoring of VD serum levels is performed to achieve individual optimal values (30 ng/mL [75 nmol/L] or higher). This control protects against rare intoxications observed at values >150 ng/mL [375 nmol/L].¹²³ A prudent upper limit of 100 ng/mL (250 nmol/L) for toxicity has a wide safety margin.¹²⁴ The differential diagnosis of unexpected “atypical” symptoms in PwMS can be facilitated if doctors are sensitized to possible polyautoimmunity. Older age, females, higher education, obesity, and familial autoimmunity were significantly associated with polyautoimmunity. Ever smokers were marginally more likely to be

polyautoimmune. Family autoimmunity may lead to earlier onset of MS.¹²⁵

AEs under AI therapy as well as the tendency toward polyautoimmunity in PwMS require intensive interdisciplinary cooperation among physicians.

Human and animal rights and informed consent

This article does not contain any studies with human or animal subjects.

Disclosure

The author declares no conflicts of interest in this work.

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Supplementary material

Patient safety requires a rapid transfer of new findings into practice, which can currently take more than 15 years. Louis Pasteur already regretted more than 100 years ago that this delayed translation into practice is to the detriment of patients.

The famous French Scientist (1822–1895) once said: “To him who devotes his life to science, nothing can

give more happiness than increasing the number of discoveries, but his cup of joy is full when the results of his studies immediately find practical applications”.¹

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