**BASIC SCIENCE FOR THE CHEST PHYSICIAN**

**Vitamin D and lung disease**

Paul E Pfeffer, Catherine M Hawrylowicz

**ABSTRACT**

The classic role of vitamin D in the regulation of calcium and phosphate metabolism, and its effects on bone health, are well established. More recently a critical role in immunity and respiratory health has been proposed. This arises from evidence for the capacity to generate the active metabolite, 1α,25-dihydroxyvitamin D3 (1,25(OH)2D3), locally in many tissues beyond the kidney; expression of the vitamin D receptor (VDR) in immune and structural cells not involved in calcium-phosphate homeostasis; and control by 1,25(OH)2D3 of the transcription of genes associated with numerous different biological processes through its nuclear VDR. Abnormalities in the vitamin D axis, including a high prevalence of vitamin D insufficiency worldwide, now appear important in a wide range of pulmonary diseases including viral and bacterial respiratory infection, asthma, chronic obstructive pulmonary disease, and cancer. Actions of vitamin D on innate immune responses, for example, production of antimicrobial peptides and autophagy, and on adaptive immune responses, for example, promoting regulatory lymphocytes, are believed to underpin these associations.

The principal source of vitamin D is sunlight; solar ultraviolet B radiation converts 7-dehydrocholesterol in the skin to pre-vitamin D3—our environment, lifestyle and our genes greatly influence the efficacy of this pathway. Dietary intake of vitamin D is limited (figure 1). Although rickets and osteomalacia occur with vitamin D deficiency (circulating levels of 25-hydroxyvitamin D <25 nmol/litre, although there is no consensus for the absolute cut-off value), evidence is accumulating of disease associations that occur in vitamin D insufficiency (<75 nmol/litre). Importantly, vitamin D insufficiency is surprisingly common, with reports that the majority of adults in the UK are vitamin D insufficient year round.2

Asthma, chronic obstructive pulmonary disease (COPD), interstitial lung disease and cystic fibrosis show correlations between severity and low vitamin D status, as does incidence of pulmonary infections and lung cancer.4 For example, paediatric studies in Costa Rica, North America and the UK have found associations between asthma exacerbations and vitamin D insufficiency. Although issues of confounding and reverse causation are problematic to these studies, gene studies and mechanistic science strongly support the role of vitamin D in pulmonary health. For example, polymorphisms in the vitamin D receptor (VDR) gene are associated with lower respiratory tract infections and severe respiratory syncytial virus bronchiolitis in children.4 In addition, levels and polymorphisms of vitamin D binding protein are associated with COPD.5
regulatory T-cell (T-reg) responses and T-cell production of IL-10. Translational research has shown that vitamin D treatment is able to restore ex vivo steroid sensitivity in T cells from people with steroid-refractory asthma, and these data compliment increasing evidence for associations of low vitamin D status with poor asthma control.\(^\text{10}\) Interactions between lymphocytes and other cells, necessary for coordinated balanced immune responses, are also regulated by vitamin D—for example lymphocyte expression of CD200.\(^\text{11}\)

This scientific evidence supports a profound role of vitamin D in orchestrating immune responses. Rising levels of local 1,25(OH)\(_2\)D\(_3\) during the immune response to a bronchial infection will boost innate immune responses and then regulate adaptive responses to prevent an excessive response. Mechanistically vitamin D insufficiency is thought to lead to more severe exacerbations in asthma and COPD as a result of an impaired innate response to the pathogen followed by an excessive adaptive immune response with increased production of inflammatory cytokines and increased bronchial inflammation.

**INTERVENTION AND THE FUTURE**

Research is now moving into large intervention studies examining the role of patient supplementation of vitamin D to treat diseases and also maternal/early life supplementation to prevent development of pulmonary diseases, for example the Vitamin D Antenatal Asthma Reduction Trial (VDAART) study investigating whether maternal supplementation in pregnancy may reduce the incidence of asthma and allergic illness. However, several issues need to be considered in these studies. It is unknown whether there may be continual improvement in disease activity with any increase in vitamin D levels upon patient supplementation (continual effect), or whether there will only be improvement in patients whose low vitamin D levels are elevated by supplementation over a threshold level (threshold effect). Different VDR polymorphisms strongly affect vitamin D action and may affect whether supplementation is beneficial. For example, Martineau \textit{et al} showed that vitamin D supplementation of patients with tuberculosis reduced the time for sputum conversion, although this was only significant in patients with the \textit{tt} TaqI polymorphism of the VDR gene.\(^\text{12}\) Bolus versus steady-state vitamin D supplementation, and the form of vitamin D supplement may also be important. Importantly, the low dose of vitamin D in over-the-counter supplements and in calcium supplements for preventing osteoporosis is much lower than the dose used in current trials.

Future vitamin D therapies for cancer, infection and asthma show great promise, although the forms of treatment and specific subgroups of patients who will respond are still being clarified through ongoing research.

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