

Vitamin D deficiency in patients with liver cirrhosis

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Abstract

There is ongoing evidence that vitamin D is related to the pathophysiology of cirrhosis. Although the incidence of vitamin D deficiency in chronic liver diseases and cirrhosis is strongly documented, its pathogenic association with advanced liver fibrosis remains controversial. There is evidence of a significant relation of 25(OH)D levels with the degree of liver dysfunction, considering that an inverse correlation of 25(OH)D levels with both Child-Pugh score and Model for End-Stage Liver Disease has been reported. In addition, vitamin D deficiency has been shown to increase the risk for overall mortality and infections in patients with cirrhosis. Vitamin D deficiency has been also associated with advanced stages of hepatocellular carcinoma and poor prognosis. Finally, there are studies suggesting that patients with chronic hepatitis C and normal vitamin D levels have higher virological response to treatment. However, there are not enough studies conducted in cirrhotic-only populations. The association between vitamin D and cirrhosis demonstrates a great potential for clinical application. The relation between vitamin D deficiency and the degree of liver function, degree of fibrosis and infectious complications could support its use as a prognostic index and a diagnostic tool.

Keywords Vitamin D, vitamin D deficiency, vitamin D insufficiency, liver cirrhosis, prognosis

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Introduction

Vitamin D is a secosteroid hormone, which is mostly known as a regulator of calcium and bone metabolism. However, vitamin D has pleiotropic effects including cellular proliferation, differentiation and immunomodulation [1]. These extra-skeletal effects have been related to the pathogenesis and treatment of infections, cardiovascular, autoimmune and degenerative diseases and several types of cancer [2].

The role of vitamin D in chronic liver diseases is not well known, but there are reports suggesting that this hormone has anti-inflammatory and anti-fibrotic effects and, consequently, it has a significant role in the natural history of chronic liver diseases, such as chronic hepatitis C and non-alcoholic fatty liver disease (NAFLD) [3]. The aim of this review was to assess vitamin D deficiency in patients with advanced liver fibrosis

and cirrhosis. We searched (CK, PT) Medline, Scopus and Google Scholar databases from November 1950 to May 2015 using the textwords: "vitamin D", "cirrhosis", "liver disease" and "deficiency" and abstracts from major Gastroenterology and Liver meetings.

Vitamin D metabolism

The basic metabolism of vitamin D has been extensively studied. Dietary sources [such as fatty fish, eggs and artificially fortified foods (oral supplements)] provide vitamin D₂ and vitamin D₃, absorbed in the intestine by biliary acids and then transported via chylomicrons to the circulation [4-6]. The main source of vitamin D is biosynthesis from epidermal cells as a result of exposure to sunlight. Initially, 7-dehydrocholesterol, a metabolite of cholesterol, is converted into pre-vitamin D₃ by ultraviolet-B radiation in the lower epidermis. Then, pre-vitamin D₃ is transformed into vitamin D₃ (cholecalciferol) in a heat-dependent process. A significant percentage of vitamin D₃ is metabolized by sunlight and so the equilibrium between pre-vitamin D₃ and vitamin D₃ is maintained. Vitamin D that comes from dietary sources or skin synthesis can be stored in the adipocytes or it may undergo hepatic 25-hydroxylation [5]. Vitamin D₃ is bound to vitamin D-binding protein (DBP) or albumin and it is transferred to the liver, where 25-hydroxylation takes place [4] (Fig. 1).

Eighty eight per cent of 25(OH)D is bound to DBP, a protein synthesized by the liver and a member of the albumin

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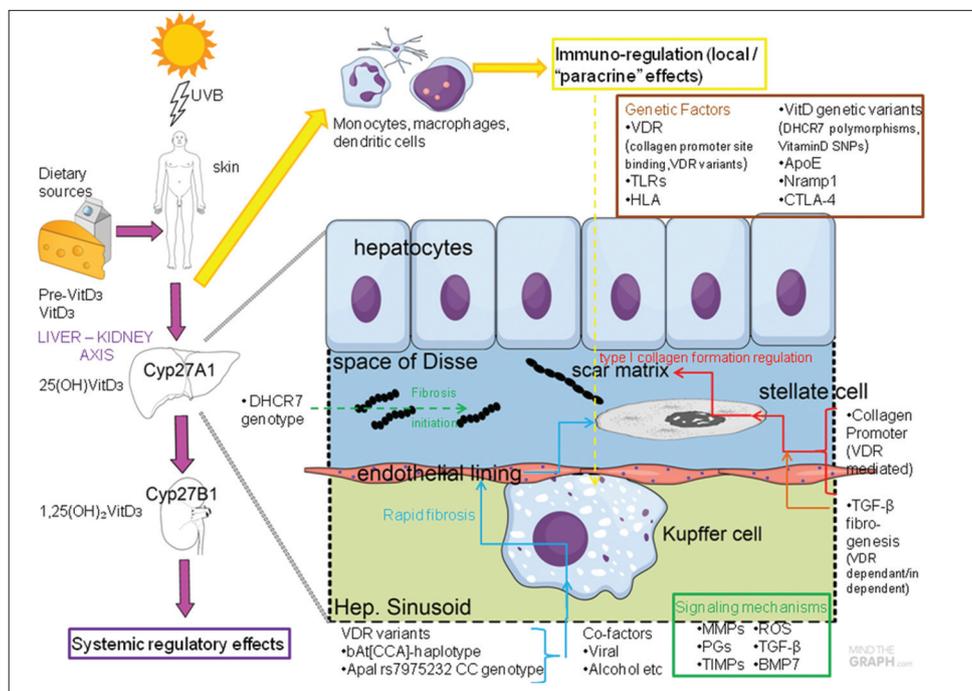


Figure 1 Vitamin D and liver fibrosis. Dietary sources provide vitamin D₂ and vitamin D₃, which are absorbed in the intestine. The main source of vitamin D is biosynthesis from epidermal cells as a result of exposure to sunlight. Vitamin D₃ is transferred to the liver, where 25-hydroxylation takes place. 25(OH)D is transported to the kidney where it is again enzymatically hydroxylated to form 1,25(OH)₂D₃ or calcitriol (the active form of vitamin D₃). Calcitriol through its receptor, VDR, exerts its systemic (endocrine – purple axis) regulatory effects on many different target organs. A second 1,25(OH)₂D₃ pool is proposed (paracrine - yellow axis), that involves the immune system and the local production of calcitriol by immune cells, which could contribute to immune regulation

The complex interplay between environmental, genetic, cell signaling and other host factors that drive the inflammatory process and fibrogenesis remains a topic of research. “Key players” of this process are presented in this figure

- *DHCR7* genotype / polymorphisms = The *DHCR7* gene has been proposed to be able to affect vitamin D serum levels. A link with the severity of fibrosis has been suggested, namely with the (GG) homozygote of the *DHCR7* gene. A stronger association of the susceptible *DHCR7* SNPs (single nucleotide polymorphisms) with fibrosis initiation rather than progression has been suggested
- Collagen promoter = There is evidence that 1,25-(OH)₂D₃ inhibits type I collagen formation in stellate cells, mainly by binding to specific sites. At least two sites have been identified, with the proximal Sp 1-1 site being the most recognizable
- *VDR* variants = *VDR* polymorphisms have been associated with liver fibrosis
- Signaling mechanisms = A host of enzymes, growth factors and other physiologically active compounds are in complex interaction, resulting in the regulation of liver inflammation and fibrosis

UVB, ultraviolet B; VitD, vitamin D; Cyp27A1, cytochrome P450, family 27, subfamily A, polypeptide 1; Cyp27B1, cytochrome P450, family 27, subfamily B, polypeptide 1; VDR, vitamin D receptor; TLRs, toll-like receptors; HLA, human leukocyte antigen system; ApoE, apolipoprotein E; *Nramp1*, natural resistance-associated macrophage protein one; *CTLA-4*, cytotoxic T lymphocyte antigen-4; *DHCR7*, 7-dehydrocholesterol reductase; MMPs, matrix metalloproteinases; ROS, reactive oxygen species; PGs, prostaglandins; BMP7, bone morphogenetic protein 7; TGF-β, transforming growth factor-β; TIMPs, tissue inhibitor metalloproteinases

gene family, homologous to albumin and α-fetoprotein [5]. 25(OH)D is transported to the kidney where it is again enzymatically hydroxylated to form 1,25(OH)₂D₃ or calcitriol, the active form of vitamin D₃. CYP27B1 is the enzyme responsible for 1^α-hydroxylation in the proximal tubule of the kidney. The production of calcitriol is regulated by rising serum calcium and phosphorus levels [4]. 1,25(OH)₂D is also highly bound to DBP and this complex activates the vitamin D receptor (VDR). VDR is expressed in many human tissues, such as liver, gastrointestinal tract, pancreas and immune cells (e.g. T lymphocytes, B lymphocytes, natural killer cells). VDR regulates the expression of more than 200 genes and, thus, influences cell proliferation, differentiation, apoptosis, immunomodulation and angiogenesis [7] (Fig. 1).

Vitamin D deficiency

Vitamin D insufficiency and deficiency are considered to be common in the general population and more frequent among elderly people and individuals with chronic diseases. It has been reported that 1 billion people have inadequate serum levels of 25(OH)D levels [8]. However, the normal range of vitamin D levels has been debated [9]. In general, optimal vitamin D status ranges from 30 to 50 ng/mL (i.e. 75-125 nmol/L) [10,11]. Vitamin D deficiency has been defined as serum 25(OH)D levels lower than 20 ng/mL (i.e. 50 nmol/L) and vitamin D insufficiency has been defined as serum levels between 20 and 30 ng/mL (i.e. 50-75nmol/L) [5,12-14]. According to the Institute of Medicine (IOM) of the National Academies

in the United States, vitamin D concentration of 20 ng/mL is adequate [15]. However, the Endocrine Society (Maryland, USA) [16] recommends levels of at least 30 ng/mL (i.e. 75 nmol/L) as adequate and concentrations between 40 and 60 ng/mL (i.e. 100-150 nmol/L) as optimal. There is still no definition regarding the optimal vitamin D levels for patients with chronic liver diseases.

Vitamin D and chronic liver disease

Vitamin D has an important role in various chronic diseases, such as infectious and cardiovascular diseases, diabetes mellitus and some types of cancer [17]. In addition, vitamin D has been associated with chronic liver diseases and it has been reported that low vitamin D status is a common feature in different types of liver diseases [18,19].

According to recent studies, the prevalence of vitamin D insufficiency and deficiency is higher in patients with chronic liver disease than in general population ranging between 64 and 92% [20,21]. It has been also reported that the incidence of vitamin D deficiency increases as the liver disease progresses [20,21]. In a study by Fisher *et al*, vitamin D deficiency was higher in cirrhotic patients in Child-Pugh class C than in patients in Child-Pugh class A [21]. Similar results were demonstrated from studies that evaluated vitamin D levels in patients with NAFLD and non-alcoholic steatohepatitis (NASH) [22,23]. According to Barchetta *et al* [23], patients with NAFLD had lower 25(OH)D than controls (14.8±9.2 versus 20.5±9.7 ng/mL).

As summarized by Stokes *et al*, a variety of mechanisms contribute to vitamin D deficiency [9]. In chronic liver diseases, the decreased vitamin D levels are associated with both malnutrition and low exposure to sunlight. Moreover, liver disease is characterized by low intestinal absorption of vitamin D and low levels of binding proteins (DBP and albumin), which can transfer the hormone to the liver and kidney, in order to be activated. In addition, hepatic hydroxylation of vitamin D is impaired leading to low production of the active hormone, whereas the catabolism of the vitamin is increased [9].

Vitamin D and hepatitis C virus (HCV) infection

The majority of patients with chronic hepatitis C have vitamin D deficiency compared to controls (serum 25(OH)D, 25.07 mg/L versus 43.06 mg/L) [24]. According to a cohort study including 468 patients, low serum vitamin D levels at baseline were associated with failure to achieve sustained virological response (SVR) in HCV genotypes 1, 2, and 3 following treatment with pegylated interferon and ribavirin [25]. Other studies showed that patients with chronic HCV infection (genotypes 1, 2/3) using oral vitamin D supplements had higher response to therapy and lower relapse rates [26,27]. Data from *in vitro* studies supported the significance of vitamin D in chronic hepatitis C, suggesting 25(OH)D as a

suppressive factor of HCV replication [28,29]. Vitamin D ameliorates the necroinflammatory process and inhibits liver fibrosis, and subsequently, its deficiency could contribute to the progression of chronic hepatitis [24,25].

NAFLD

NAFLD is the most common chronic liver disorder in economically developed countries, and its prevalence is strongly linked to current lifestyle. Approximately 30% of patients with NAFLD have NASH in liver biopsy which might progress to liver cirrhosis [30]. Insulin resistance, a component of metabolic syndrome, is implicated in the progression to NASH [31]. Vitamin D has a great impact on NASH, considering that an optimal vitamin D status reduces the incidence of metabolic syndrome [32]. 25(OH)D improves insulin resistance by accelerating the metabolism of proinsulin to insulin [33,34]. According to Targher *et al*, vitamin D deficiency is implicated in steatosis, necroinflammation and liver fibrosis, regardless of other aspects of metabolic syndrome [22]. Nakano *et al*, using a rat model, examined the role of vitamin D produced by phototherapy on the progression of NASH. Phototherapy reduced inflammation and fibrosis of hepatic cells compared to controls. It has been also suggested that phototherapy improves insulin resistance and inhibits the expression of profibrotic factors, such as transforming growth factor (TGF)- β [35].

Vitamin D and liver transplantation

Vitamin D has immunomodulatory effects with direct actions at dendritic cells, monocytes, macrophages, B-cell, and T-cell functions. Considering that vitamin D and VDR are expressed by several cellular populations of the immune system, such as Th1 and Th2, it has been suggested that calcitriol has a significant role in liver transplantation [2]. Trautwein *et al* studied 193 patients before and after liver transplantation and concluded that liver transplantation improved serum vitamin D levels and enhanced bone metabolism [36]. Another study suggested that patients with pre-transplant low serum 25(OH)D levels (<12.5 ng/mL), were more likely to have organ rejection [37].

Vitamin D and primary biliary cholangitis (PBC)

PBC is an autoimmune liver disease of unknown etiology. It is characterized by a T-lymphocyte-mediated slowly progressive destruction of small intralobular bile ducts, which results in cholestasis and, eventually, in cirrhosis and liver failure. Vitamin D has been implicated into the pathogenesis of PBC [38] through several cell signaling mechanisms namely matrix metalloproteinases, reactive oxygen species,

prostaglandins, and TGF- β in which vitamin D normally plays a regulatory role (of varying degree) [38]. Genetic studies have yielded interesting results on identifying genes that confer susceptibility to PBC. A firm association between PBC and several genes within the major histocompatibility class II region (MHC) has now been reported [39,40].

VDR has been a topic of research in the study of non-MHC genes that link vitamin D to PBC pathogenesis. Three recent meta-analyses on VDR polymorphisms have provided slightly diverse results [41-43] (Table 1). A number of other genetic factors, namely Toll-like receptors (TLRs), apolipoprotein E (ApoE), Nramp1, and Cytotoxic T lymphocyte antigen-4 (CTLA-4) have also been associated with vitamin D and PBC [38].

Low serum levels of vitamin D have been reported in PBC patients [38,44] especially compared to controls (significantly lower mean values) [38,45,46]. According to Agmon-Levin *et al* [45], vitamin D deficiency is highly prevalent in the PBC group with almost one third of these patients exhibiting serum vitamin D levels below 10 ng/mL. Treatment with ursodeoxycholic acid (UDCA) resulted in increased levels of vitamin D. An inverse correlation between vitamin D levels and advanced liver disease/histological stage [45,47] has been reported; the authors have commented on the potential use of this finding as a prognostic marker (disease severity, mortality, need for liver transplantation). The same authors also suggest an inverse association between vitamin D levels and both the markers of PBC disease activity, (namely alkaline phosphatase and bilirubin) [38,47], and the presence of an additional autoimmune disease [38]. PBC patients who failed to respond to UDCA therapy (according to Paris-I/Barcelona criteria) had significantly lower baseline vitamin D levels [47]. Thus, Guo *et al* suggest a link between pre-treatment vitamin D levels and the response to treatment, irrespectively of the stage of the disease [47]. In addition, Agmon-Levin *et al* suggest that low vitamin D levels also correlate with the absence of UDCA therapy (among other factors like advanced liver disease and autoimmune comorbidity) [45].

Vitamin D and autoimmune hepatitis (AIH)/primary sclerosing cholangitis (PSC)

AIH is a chronic inflammatory disease of the liver of unknown etiology. Liver cirrhosis/failure is a possible outcome if left untreated. AIH diagnosis is based on serological, immunological and histopathological findings, namely hyperglobulinemia, circulating autoantibodies and evidence of interface hepatitis

on liver biopsy [48]. Patients with AIH have significantly lower vitamin D levels compared with controls [49,50]. Several studies have reported a link between genetic factors and pathogenetic mechanisms of AIH. It has been suggested that a number of genes inside the MHC region confer susceptibility to AIH. Other (non-MHC) vitamin D-related genetic factors include: VDR polymorphisms, CTLA-4, TLRs, cytochrome P450 Cyp2D6, regulatory T cells, and the forkhead/winged helix transcription factor 3, but data on these factors is limited [50]. Efe *et al* link low 25(OH)D levels in AIH to advanced fibrosis and severe inflammation [49], whereas at the same time Luong *et al* suggest that vitamin D may have a beneficial role on AIH [50], and predict response to treatment [49].

The literature on the connection of vitamin D with PSC is scarce. Vitamin D deficiency has been reported in PSC patients which is far exaggerated in the pre-transplantation setting [51]. Very recently it has been reported that CD28⁺ T cells accumulate near the bile ducts of PSC patients, thus promoting inflammation in a tumor necrosis factor- α -rich microenvironment. The authors suggest that vitamin D reverses this condition *in vitro* [52].

Vitamin D and liver cirrhosis

The deranged metabolism of vitamin D in liver cirrhosis was first reported in the late '70s [53-55] and it was mainly attributed to impaired 25(OH)-vitamin D hydroxylation of the precursor vitamin D due to insufficient liver function [53,55]. Before the year 2000, the majority of the studies on vitamin D in cirrhosis [56-58] focused on the association of hepatic insufficiency with bone demineralization, osteomalacia, osteoporosis, minerals metabolism/equilibrium (calcium, phosphorus), possible endocrine disturbances (parathormone – secondary hyperparathyroidism) and, in general, with the homeostasis involving the liver-kidney-gut-calcium axis. In the past two decades, there have been considerable advances in the understanding of the pathophysiology of vitamin D and its possible clinical implications in chronic liver diseases [59-65].

Serum levels of vitamin D in patients with liver cirrhosis

Vitamin D deficiency in cirrhosis is related to liver dysfunction rather than etiology and it is no longer considered

Table 1 Recent meta-analysis studies on the association of the most well studied vitamin D receptor polymorphisms (BsmI, ApaI, and TaqI) and the risk of primary biliary cholangitis. Overall refers to overall analysis, whereas subgroup refers to subgroup analysis based on ethnicity. Negative reveals no association while positive confirms an association; the degree (significant/non-significant) of association is commented in brackets

	BsmI overall	BsmI subgroup	ApaI overall	ApaI subgroup	TaqI overall	TaqI subgroup
Fang <i>et al</i> , 2015 [43]	Negative	Negative	Positive (significant)	Asian (significant)	Negative	Caucasian
Mo <i>et al</i> , 2014 [42]	Negative	Negative	Negative	Negative	Negative	Negative
Li <i>et al</i> , 2014 [41]	Negative		Negative		Positive	

prevalent only in cholestatic disorders [19,66]. Malham *et al* [19] compared vitamin D status between patients with alcoholic cirrhosis (ALC) and PBC. ALC patients had lower vitamin D levels compared to PBC patients. A number of studies have supported the prevalence of hypovitaminosis D in chronic liver disease and cirrhosis [67-76] with one study [77] reporting a low prevalence of 25(OH)D deficiency in a cohort of patients with genotype 1 chronic HCV infection and compensated liver disease (15% cirrhotic patients): 48% and 16% of the cohort had vitamin levels of <75 nmol/L and <50 nmol/L, respectively (Table 2).

Relationship between vitamin D and liver fibrosis

An association of low 25(OH)D levels with advanced fibrosis has been reported in both HCV mono-infected [24] and HCV-HIV co-infected cohorts [67]. In the latter study [67], a significant correlation of 25(OH)D levels with the histological METAVIR fibrosis score was observed. Two recent genetic studies have further supported the relation between vitamin D and fibrosis. Baur *et al* [78] studied both 25(OH)D serum

levels and *VDR* gene (NR1H1) polymorphisms in a cohort of patients with chronic hepatitis C. The authors concluded that both deficient 25OHD levels and presence of an unfavorable bAt[CCA]-haplotype, ApaI rs7975232 CC genotype) increased the risk for fibrosis progression. Likewise, Grunhage *et al* [79] conducted a large-scale (712 patients) study of genetic variants affecting serum 25(OH)D levels in chronic liver disease (6.6% F3 and 57% F4 patients). Serum levels of 25(OH)D inversely correlated with transient elastography and fibrosis stages. Homozygous carriers of the rare *DHCR7* allele or the common *CYP2R1* allele had reduced 25(OH)-vitamin D levels. The variant rs12785878 in the *DHCR7* locus was correlated with liver stiffness in transient elastography. Their results imply that vitamin D has a greater impact on the initiation than on the progression of liver fibrosis. Petta *et al* [80] studied the association between liver fibrosis and certain genetic variants affecting 25(OH)D serum levels, in a cohort of 260 biopsy-proven genotype 1 chronic HCV-infected patients (17.3% F3, 11.2% F4 fibrosis). *DHCR7* GG genotype was identified as an independent risk factor for severe fibrosis and was associated with lower vitamin D serum levels. In a recent meta-analysis by Garcia - Alvarez *et al* [76], a significant association between vitamin D status and liver fibrosis [cut-

Table 2 Studies evaluating Vitamin D levels in patients with chronic liver disease and/or cirrhosis

Author [ref]	Year (publication)	Etiology of liver disease	Study population (n)	Proportion of cirrhotics/ALF patients included	Study in favor of hypovitaminosis	Vitamin D levels
Terrier <i>et al</i> [67]	2011	HIV-HCV	189	25% F3-F4 0% F0	Yes	F3/F4: 16.2±10.0 ng/mL; F2: 18.9±8.5 ng/mL; F1: 20.9±11.1 ng/mL
Putz-Bankuti <i>et al</i> [68]	2011	Various, mainly alcohol (61%)	75	100% cirrhotic (33 CP-A, 32 CP-B, 10 CP-C)	Yes	Baseline: 16.0±9.2 ng/mL
Bitetto <i>et al</i> [69]	2011	HCV	211	Decompensated disease excluded, baseline staging: 2 (0-6 Ishak stage)	Yes	Median: 20.7 ng/mL (2.1-59.6); 46.4% <20 ng/mL; 16.1% <10 ng/mL
El-Maouche <i>et al</i> [70]	2013	HIV-HCV	116 African-American	13 cirrhotics	Yes	41% <15 ng/mL
Venu <i>et al</i> [71]	2013	Various, mainly alcohol and HCV	63	Liver transplant candidates	Yes	75% <20 ng/mL; 6.3% <10 ng/mL
Stokes <i>et al</i> [72]	2014	Various, mainly alcohol (66%)	65	100% cirrhosis, 82% with advanced disease (CP: B,C stage)	Yes	Median: 8.2 ng/mL (4.0-95.8)
Anty <i>et al</i> [73]	2014	Various, mainly alcohol (71%)	88	100% cirrhosis, with active infection	Yes	Median: 8.8 (5.3-14.1) ng/mL
Savic <i>et al</i> [74]	2014	Alcohol	30	100% cirrhosis	Yes	66.7% <50 ng/mL
Finkelmeier <i>et al</i> [75]	2014	Various, mainly alcohol and HCV	200	HCC	Yes	Mean: 17±13 ng/mL (1-72)
Kitson <i>et al</i> [77]	2013	HCV genotype 1	274	15% cirrhotics with compensated disease	No	Mean: 79.6 nmol/L; 48% <75 nmol/L; 16% <50 nmol/L

ALF, advanced liver fibrosis; CP, Child-Pugh stage; HCC, hepatocellular carcinoma

offs of 10 ng/mL (odds ratio, OR: 2.37, 95% CI: 1.20-4.72) and 30 ng/mL (OR: 2.22, 95% CI: 1.24-3.97)] was observed.

However, two studies [69,70], similar to the study by Kitson *et al* [77], challenged the results of the earlier studies regarding the relation of vitamin D with the stage of liver fibrosis. The authors found no association between the baseline 25(OH)D levels and either the SVR rate or the fibrosis stage. They reported that season, race and geographic latitude were independent predictors of 25(OH)D status. Mean 25(OH)D levels did not significantly vary between fibrosis stages (F0-F4). This discrepancy compared to other studies might be partly related to the lack of certain clinical and laboratory baseline factors (namely season of baseline blood sampling) and to the use of non-standardized commercially available automated assays to measure 25(OH)D levels. Fig. 1 summarizes main environmental/genetic factors that implicate vitamin D in the inflammatory/fibrogenetic process.

Prognosis/mortality/infectious complications

Bankuti *et al* [68] reported a significant association of 25(OH)D with the degree of liver dysfunction. They enrolled 75 patients with liver cirrhosis (various etiologies and severity; approximately 14% of the patients had Child-Pugh C cirrhosis). All patients were followed-up until hepatic decompensation or death. 25(OH)D levels were inversely correlated with MELD score and Child-Pugh score. Patients at the first (6.9 ± 1.9 ng/mL) vs. the third/reference (27.1 ± 6.3 ng/mL) 25(OH)D tertile had a relative risk for hepatic decompensation and mortality of 6.37 (95%CI: 1.75-23.2; $P=0.005$) and 4.31 (95%CI: 1.38-13.5; $P=0.012$), respectively, after adjustment for age and sex.

Three more studies, two in cirrhotic cohorts [72,73] and another one in patients with hepatocellular carcinoma [75], reached similar conclusions. In the first study, the authors studied the association of vitamin D deficiency with survival in a cohort of 92 patients with advanced liver cirrhosis (82% Child-Pugh stages B and C) [72]. They reported a considerable incidence of vitamin D deficiency (86%). In the univariate analysis, low vitamin D levels were associated with mortality; levels ≤ 6 ng/mL were associated with an unfavorable outcome, whereas vitamin D levels < 20 ng/mL were not found significant. Multivariate analysis confirmed that both MELD score and serum vitamin D ≤ 6 ng/mL were independent predictors for mortality. 25(OH)D levels ≤ 6 ng/mL had a positive predictive value of 50% mortality within 24 months. Moreover, mortality from infectious complications was more frequent in this subgroup. The second study was conducted in a cohort of hospitalized cirrhotics of various etiologies [73]. Almost 60% of the patients had severe vitamin D deficiency (< 10 ng/mL). The authors found an inverse correlation of 25(OH)D levels with the Child-Pugh score, but no correlation with the MELD score. Infections were more frequent in patients with severe vitamin D deficiency compared to non-deficient patients (54 vs. 29%). In addition, severe vitamin D deficiency was independently associated with infections together with the Child-Pugh score and the C-reactive protein

levels. In the multivariate Cox regression analysis including Child-Pugh score, infections and severe vitamin D deficiency, only the presence of infection was associated with mortality. Lastly, Finkelmeier *et al* prospectively evaluated the role of serum vitamin D status as a prognostic marker in 200 patients with hepatocellular carcinoma [75]. The authors reported that 25(OH)D3 levels negatively correlated with both the stage of cirrhosis and the stage of hepatocellular carcinoma. Patients with severe 25(OH)D3 deficiency had the highest mortality risk. In the multivariate analysis, very low 25(OH)D3 levels were independently associated with mortality together with MELD score and high α -fetoprotein levels (> 400 ng/mL). The authors concluded that 25(OH)D3 deficiency was associated with advanced stages of hepatocellular carcinoma and was also a prognostic indicator of poor outcome.

SVR in patients with cirrhosis and chronic hepatitis C

The data in the literature regarding the virological response of antiviral treatment combined with Vitamin D supplementation in patients with liver cirrhosis are scarce. Most of the published data concern patients with chronic hepatitis C treated with interferon-ribavirin regimens with a small number of cirrhotic patients (usually with compensated disease, Child-Pugh class A) [24,67]. As mentioned before, Petta *et al* [24] reported evidence supporting that low 25(OH)D levels are an independent risk factor for SVR (interferon-based therapeutic regimens). On the contrary, data from both the HIV-HCV co-infected cohort of Terrier *et al* [67] and the HCV mono-infected cohort of Kitson *et al* [77], showed no association between the baseline 25(OH)D status and SVR.

There are two recent meta-analyses in this field [76,81]. The first one reported a high prevalence of vitamin D deficiency and higher likelihood of SVR in the individuals with higher serum vitamin D levels and the individuals on vitamin D supplementation [81]. The second meta-analysis [76] reported a significant association with SVR only for a vitamin D cut-off of 20 ng/mL (OR: 0.53, 95% CI: 0.31-0.91). When the outliers were excluded, significant pooled ORs were observed for all patients included [10 ng/mL (OR: 0.48, 95% CI: 0.34-0.67), 20 ng/mL (OR: 0.58, 95% CI: 0.45-0.76)] and specifically for genotype 1/4 patients [10 ng/mL (OR: 0.53, 95% CI: 0.34-0.81), 20 ng/mL (OR: 0.54, 95% CI: 0.39-0.74)].

Clinical implications of vitamin D in the cirrhotic setting

Several clinical applications of 25(OH)D levels have been suggested, including its use as a non-invasive marker of liver fibrosis in chronic hepatitis C [24], as a prognostic predictive factor for mortality and infections in patients with liver cirrhosis [73] and as a marker of unfavorable outcome and advanced disease stage in patients with hepatocellular carcinoma [75]. Malham *et al* [19] emphasized the importance

of monitoring vitamin D in all cirrhotic populations, especially those with alcoholic liver cirrhosis, and commented on the efficacy of treatment in liver insufficiency-associated bone disease, on the possible extraskelatal benefits (muscle function, cancer risk, and immune impairment) and on the probable benefit of higher than standard doses of vitamin D supplementation for repletion. Garcia-Alvarez *et al* [76] also recommended vitamin D screening in HCV patients. Finkelmeier *et al* [75] showed that, in patients with hepatocellular carcinoma and vitamin D supplementation, the mean serum 25(OH)D levels did not differ significantly from those that did not receive supplementation. However, because of the small number of treated patients, these results should be interpreted with caution.

The possible benefit of vitamin D substitution, as a preventive measure for the development of liver fibrosis in patients with chronic hepatitis C, has also been suggested [78]. The same authors comment on expanding the indications for vitamin D supplementation to all patients with chronic liver diseases irrespective of the presence of bone disease. A recent meta-analysis suggested the cutoff of 30 ng/mL as an appropriate threshold to prevent both fibrosis and treatment failure in patients with chronic hepatitis C [76]. Another study suggested vitamin D supplementation as a preventive, early treatment, strategy against fibrosis in the cohort of HCV-infected patients [82]. Lastly, Lim *et al* [66] suggested periodic monitoring of 25(OH)D in patients with chronic liver disease and cirrhosis, and substitution therapy in those with levels <30 ng/mL, which includes administration of 5000 IU of vitamin D3 daily or 50000 IU of vitamin D2 or D3 weekly for 3 months, followed by 1000 IU/day indefinitely.

The Endocrine Society Clinical Practice Guideline (ESCPG) [16] recommend screening for vitamin D deficiency (cutoff <20 ng/mL) in individuals at high risk for deficiency, including those with hepatic failure, and recommends vitamin D supplementation in cases of deficiency. They also suggest that vitamin D requirements may be greater for sick patients than for healthy individuals and that serum vitamin D levels above 30 ng/mL may have additional benefits in reducing the risk for various disease conditions. They recommend that all adults who are vitamin D deficient to be treated with 50000 IU of vitamin D2 or vitamin D3 once a week for 8 weeks or its equivalent of 6000 IU of vitamin D2 or vitamin D3 daily to achieve a serum level of 25(OH)D above 30 ng/mL, followed by maintenance therapy of 1500–2000 IU/day. Furthermore, they have set minimal daily dietary recommendations of vitamin D intake for patients at high risk for vitamin D deficiency, depending on their age group. The extent of the recommended screening indications is limited to hepatic failure and does not cover the full spectrum of liver diseases.

International liver study associations (both EASL [83] and AASLD [84]) recommend fat-soluble vitamin substitution for the management of all patients with cholestatic liver diseases. These recommendations focus mostly on the prevention of osteoporosis. More specifically, they propose clinical assessment of the risk of osteoporosis for all cholestatic patients and emphasize the importance of both identifying reversible risk factors and applying appropriate lifestyle changes. It

is highlighted that the risk of osteoporosis is increased in decompensated disease and in high degree of cholestasis and it is suggested at least annual screening intervals following diagnosis. Finally, it is recommended that calcium and vitamin D supplementation should be considered in all cholestatic disease patients [83].

Discussion

Vitamin D has an emerging role on immunity, cancer, infectious diseases, fibrosis and chronic liver diseases [85]. The pleiotropic effects of this hormone including the regulation of transcription of over 200 genes involved in cell proliferation and differentiation, immunomodulation, inflammation and fibrogenesis and their effects on liver disease have been extensively reviewed in the literature [3,24,85]. Han *et al* [85] suggested two separate pools of 1,25(OH)2D3 with distinct purposes. The first pool, consisting of the traditional liver-kidney loop, facilitates intestinal absorption of calcium by mediating active calcium transport (calbindin) across the intestinal mucosa, which maintains calcium homeostasis in blood and allows for bone calcium deposition (Fig. 1). The second pool involves the immune system and the local production of calcitriol by immune cells (monocytes, macrophages, dendritic, B and T cells and lymphocytes), which could contribute to immune regulation (with a possible protective role against infections). These separate pools, although not well defined, could contribute to two separate, in essence endocrine and paracrine, homeostasis functions. Our current understanding of vitamin D pathophysiology with regard to liver diseases/cirrhosis is probably best summarized by Petta *et al*: “a complex interplay between liver damage, vitamin D and genetic determinants of vitamin D deficiency” [80].

The majority of the studies on patients with cirrhosis [19,68,71-74] confirm the prevalence of vitamin D deficiency in this setting. The question of whether liver damage precipitates the disturbance in vitamin D homeostasis or the other way around, still remains, leading to a type of “chicken or the egg” causality dilemma. However, whatever its role in liver disease (epiphenomenon of cirrhosis or co-factor of liver fibrosis and necroinflammation), vitamin D must still be considered as a diagnostic tool and prognostic index.

There is ongoing evidence supporting the importance of vitamin D status on mortality not only in the general population but also in patients with chronic liver diseases, cirrhosis and hepatocellular carcinoma [72-75]. Furthermore, the association of low vitamin D levels with liver insufficiency and infections [72,73] supports the use of vitamin D as a prognostic marker in the population of cirrhotics.

The role of vitamin D supplementation on the achievement of SVR in patients with chronic hepatitis C treated with interferon-based treatment is debatable [76,77]. This needs to be elucidated as most published studies included patients with well-preserved liver function and minor fibrosis. However, in the era of direct acting antiviral therapy, the interest in

vitamin D supplementation for higher SVR rates is expected to decrease together with the declining interferon use.

Current clinical guidelines cover the issue of vitamin D supplementation for bone disease in liver cirrhosis and cholestatic disorders. However, some argue that the established definitions of vitamin D deficiency and insufficiency might not apply in patients with cirrhosis. It is of major importance to define better the details of vitamin D supplementation, such as the threshold for commencing supplementation, optimal duration, other liver-related extra-skeletal indications, dosage customization (maximized dosages in cirrhotics)/optimal level of supplementation), way of delivery and bioavailability, pre-treatment screening intervals, in-treatment efficacy/monitoring intervals, post-treatment surveillance intervals. Moreover, there are concerns regarding the accuracy of the different 25(OH)D assays that need to be addressed [77,86].

In conclusion, the association of vitamin D with liver cirrhosis shows great potential for clinical application. In the near future, we expect a variety of extra-skeletal indications to be explored. The relation between vitamin D deficiency and the degree of liver function, degree of fibrosis and infectious complications could support its use as a prognostic index and diagnostic tool. The role of vitamin D in liver cirrhosis needs to be further evaluated and validated in large prospective cohort studies and randomized trials.

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