Liver fibrosis, an important health concern

HSCs were obtained from WT and [4,5]/C0 mice, and in combination with Axl inhibitor. Gas6 and Axl serum levels were measured in alcoholic liver disease (ALD) and hepatitis C virus (HCV) patients, inversely correlating with liver functionality.

Results: In primary mouse HSCs, Gas6 and Axl levels paralleled HSC activation. rGas6 phosphorylated Axl and AKT prior to HSC activation. Moreover, GBB324 blocked Axl/AKT phosphorylation and diminished HSC activation. In addition, Axd–/– mice displayed decreased HSC activation in vitro and liver fibrogenesis after chronic damage by CCl4 administration. Similarly, GBB324 reduced collagen deposition and CCl4-induced liver fibrosis in mice. Importantly, Gas6 and Axl serum levels increased in ALD and HCV patients, inversely correlating with liver functionality.

Conclusions: The Gas6/Axl axis is required for full HSC activation. Gas6 and Axl serum levels increase in parallel to chronic liver disease progression. Axl targeting may be a therapeutic strategy for liver fibrosis management.

Introduction

Activation of hepatic stellate cells (HSCs) is responsible for the liver fibrosis associated to chronic liver injury of any etiology, being HSCs the main collagen-producing cells in the damaged liver [1,2]. Liver fibrosis, critical pre-stage in the development of liver cirrhosis, may lead to hepatic transplantation or promote a favorable microenvironment for cancer development [3]. HSCs transform during chronic liver injury from a quiescent state into a myofibroblast-like phenotype, which proliferate and migrate towards areas of necrosis and regeneration [4,5]. Activated HSCs alter extracellular matrix (ECM) composition due to the upregulation of proteins such as α-smooth muscle actin (α-SMA), interstitial collagens such as Collagen 1A1 (COL1A1), and matrix metalloproteinases (MMPs) such as MMP9, as well as tissue inhibitor of metalloproteinases (TIMPs), and proteoglycans. Activated HSCs also generate hepatic cytokines such as TGF-β, PDGF, CTGF, FGF, HGF, and VEGF, and recruit inflammatory cells, mono- and polymorphonuclear leukocytes that produce chemokines, including MCP-1, RANTES, CCL21, CCR5. Although HSC critical role in liver fibrosis was proposed a decade ago [6], recent data demonstrates that irrespective of the underlying etiology of liver disease, the majority of myofibroblasts come from the liver-resident HSC population [7]. Moreover, after cessation of the fibrotic triggering insult, around 50% of the activated HSCs survive in an apparently quiescent state, being primed to quickly reactivate into myofibroblasts in response to fibrogenic stimuli [8,9]. Therefore, effective antifibrotic therapies aimed to inhibit activated HSCs, although positive to prevent extracellular matrix deposition, may be insufficient to definitely reverse fibrosis, probably requiring the elimination of activated...
HSCs for fibrosis resolution in the treatment of chronic liver disease.

Growth arrest-specific gene 6 (Gas6) product is a vitamin K-dependent protein that activates a family of receptor tyrosine kinases including Axl, MERTK and Tyro3, known as TAM receptors, whose immunologic and oncogenic properties have been described in detail [10,11]. Among them, Axl receptor signaling has been related to processes leading to cell differentiation and carcinogenesis. Gas6 possesses a high structural homology and sequence identity to the natural anticoagulant protein S (ProS). However, Gas6 and ProS have clearly different biological roles [12,13].

In liver pathologies, a hepatoprotective role for Gas6 has been reported in ischemia/reperfusion-induced damage [14], and in the wound healing response to liver injury [15,16]. In normal liver, Gas6 is mainly expressed in Kupffer cells, while Axl is found in macrophages and in quiescent HSC [17]. Moreover, after acute CCl4 administration increased Gas6 expression was observed in activated HSCs and macrophages, while Gas6 in vitro protection to HSCs was mediated by the Axl/Pi3-kinase/AKT pathway [17]. However, the role of Gas6/Axl signaling in chronic liver disease, the potential use of related proteins as serological markers of disease progression, and Gas6/Axl targeting in future liver therapies are aspects that merit further investigation.

To do so, we used both a genetic model of Axl deficiency (Axl−/−), and a pharmacologic approach, the Axl inhibitor BGB324 [18]. Our results revealed that Axl receptor is an interesting target to block HSC transformation in vitro and demonstrated the efficacy of both strategies, genetic and pharmacologic, to diminish experimental liver fibrosis after chronic administration of CCl4. Moreover, we analyzed data from patients at different stages of ALD and HCV infection providing evidence of the involvement of the Gas6/Axl axis in human liver fibrosis, and showing the correlation between Gas6/Axl serum levels and liver dysfunction.

In conclusion, our results underscore a critical role of the Gas6/Axl in fibrogenesis and in the progression of chronic liver diseases, suggesting that therapies aimed to inhibit Axl signaling deserve to be undertaken for the treatment of liver fibrosis, particularly now that small molecule inhibitors of Axl have been tested in clinical trials for cancer treatment [19].

Materials and methods

Animal procedures

All procedures were performed according to protocols approved by the Animal Experimentation Ethics Committee from the University of Barcelona. In vivo liver fibrogenesis was analyzed after chronic carbon tetrachloride (CCl4) administration. To this aim, WT or Axl−/− mice were treated with CCl4 at a dose of 5 μl (10% CCl4 in corn oil)/g of body weight, by intraperitoneal injection twice a week for five-six weeks. Control animals received corn oil alone. Treatment with Axl inhibitor (BGB324) or vehicle (saline solution) was performed daily for the last ten days of the study via oral gavage at a dose of 80 μg/g body weight. In previous experiments with rodents at similar doses, BGB324 reached serum concentration in the low micromolar range [18], being safe for animal treatment. Control animals received vehicle alone.

HSCs isolation and culture

Wild type and Axl knockout mice livers (male, 8–10 week-old littermates) (C57BL/6 strain) were perfused with collagenase and HSCs cultured as previously described [20,21]. Culture purity, assessed routinely by retinoid autofluorescence at 350 nm, was >95%. Lack of staining for F4/80 confirmed the absence of Kupffer cells. HSCs and LX2 human activated stellate cells [20,22] were cultured in DMEM supplemented with 10% FBS and antibiotics at 37 °C in a humidified atmosphere of 95% air and 5% CO2. Experiments to compare protein or mRNA content were always performed with cells extracted at the same time of culture, previously treated with recombinant Gas6 (R&D), Axl inhibitor (BGB324, BerGenBio), or siRNA silencing (Santa Cruz) after Lipofectamine 2000 exposure for the indicated periods of time.

SDS-PAGE and immunoblot analysis; RNA isolation and real-time RT-PCR; In Vitro Small Interfering RNA Transfection; Nuclear extract isolation; Immunohistochemical staining; and liver collagen determination

These methods were performed as previously indicated [20,21,23] with modifications as specified in Supplemental methods.

Determination of Gas6, and soluble Axl (sAxl) levels

Measurements of Gas6 and sAxl human levels were carried out using commercial antibodies (R&D Systems) to develop specific ELISAs that use the sandwich technique as described [24]. Serum Gas6 mouse levels were determined using a commercial kit (Duoset mGas6 ELISA, R&D). Serum sAxl mouse levels were determined by western blot.

Human samples

a) The ALD study group comprised serum samples from 40 individuals: ten healthy normal adult controls (C) and 30 alcoholic patients with different degrees of liver disease as diagnosed after hepatic biopsy and Fibroscan measurement: ten patients with initial fibrosis (Fibroscan score <7 KPa, mean = 5.2 ± 0.4) (F), ten patients with compensated cirrhosis (CH) and, 10 patients with decompensated cirrhosis (DCH), five of them due to ascitis, three due to spontaneous bacterial peritonitis (SBP) and two due to gastrointestinal bleeding by esophageal varices and portal hypertension. Relevant biochemical data are shown in Table 1. b) The HCV study group comprised serum samples from 51 individuals at different stages of liver fibrosis (8 F0, 15 F1, 17 F2, and 11 F3/F4), as stated in Table 1. c) The ALD study group comprised serum samples from 51 individuals at different stages of liver fibrosis (8 F0, 15 F1, 17 F2, and 11 F3/F4), as stated in Table 1. d) The HCV study group comprised serum samples from 51 individuals at different stages of liver fibrosis (8 F0, 15 F1, 17 F2, and 11 F3/F4), as stated in Table 1. e) The ALD study group comprised serum samples from 51 individuals at different stages of liver fibrosis (8 F0, 15 F1, 17 F2, and 11 F3/F4), as stated in Table 1. f) The HCV study group comprised serum samples from 51 individuals at different stages of liver fibrosis (8 F0, 15 F1, 17 F2, and 11 F3/F4), as stated in Table 1. g) The ALD study group comprised serum samples from 51 individuals at different stages of liver fibrosis (8 F0, 15 F1, 17 F2, and 11 F3/F4), as stated in Table 1. h) The HCV study group comprised serum samples from 51 individuals at different stages of liver fibrosis (8 F0, 15 F1, 17 F2, and 11 F3/F4), as stated in Table 1. i) The ALD study group comprised serum samples from 51 individuals at different stages of liver fibrosis (8 F0, 15 F1, 17 F2, and 11 F3/F4), as stated in Table 1. j) The HCV study group comprised serum samples from 51 individuals at different stages of liver fibrosis (8 F0, 15 F1, 17 F2, and 11 F3/F4), as stated in Table 1.

Statistical analyses

Results are expressed as mean ± standard deviation, unless indicated, with the number of individual experiments detailed in Figure legends. Statistical comparisons were performed using unpaired two-tailed Student’s t test or One-way ANOVA followed by Newman-Keuls Multiple Comparison Test (GraphPad Prism). A p value less than 0.05 was considered significant.

Results

TAM receptors and ligands levels during HSC activation

Gas6 and ProS are the ligands of the tyrosine kinase family of receptors named TAM (Tyro3, Axl, and MERTK), which have been involved in numerous processes related to cell transformation and cancer. Since TAM receptor participation in HSC activation has not been explored, we analyzed the presence of transcriptional changes during HSC transdifferentiation in mouse-derived primary cultures of HSCs. A significant increase in the mRNA levels of Gas6, but not of ProS, was detected (Fig. 1A). In parallel, strong upregulation of Axl was observed, with no significant changes in MERTK levels (Fig. 1B). Tyro3 mRNA levels were not detectable in these samples. Of note, increased secretion of Gas6 protein expression was confirmed in HSCs after different days in culture, as determined by ELISA in 24 h cell conditioned

JOURNAL OF HEPATOLOGY

Molecular and Cell Biology

Journal of Hepatology 2015 vol. 63 | 670–678

671
medium (Fig. 1C). Thus, the activation of WT HSCs is paralleled by an increase in expression of Axl, and the expression and secretion into the medium of Gas6.

Axl is required for full HSC activation and proliferation in vitro

To verify the repercussion of this correlation in HSC transformation, we analyzed the effect of reducing the expression of Axl in activated HSCs, by means of RNA silencing, and the consequences of Gas6 supplementation. Since primary HSCs are not easy to manipulate genetically, we used the LX2 human activated HSC cell line. LX2 cells with depleted Axl levels by siRNA transfection were obtained, cultured and analyzed in comparison to WT HSCs. A significant reduction in the mRNA levels of the markers -SMA and Col1a1 was observed in Axl deficient cells, showing a reduced proliferation and survival of HSCs via the AKT/NF-κB signaling pathways. However, since we also observed that rGas6 phosphorylated MERTK (Fig. 2C), at this point we cannot discard a potential role of MERTK in HSC signaling.

To better examine Axl contribution, HSCs from Axl−/− mice were obtained, cultured and analyzed in comparison to WT HSCs. A significant reduction in the mRNA levels of the markers for HSC activation α-SMA and Col1a1 was observed in Axl deficient mice, as well as in TIMP-1 mRNA, while no differences for TGF-β levels were detected (Fig. 2D). Indeed, Axl−/− HSCs showed decreased protein levels of α-SMA in vitro, showing a reduced activation state after 10 days and delayed proliferation, as manifested by MMP9 and PCNA levels, respectively (Fig. 2E). Although not all the variety of proteins induced during HSC activation are affected by Axl, as the absence of changes in TGF-β seem to indicate, taken as a whole, these results suggest that Axl signaling is required for full activation of HSCs in vitro.

Axl deficiency diminished liver fibrosis induced by carbon tetrachloride

To investigate if Gas6/Axl role in HSC activation in vitro may reflect a key participation of this system in the development of liver fibrosis, we used the chronic administration of CCl₄ as a model to generate liver damage and fibrogenesis in mice. First,
we wanted to analyze if Gas6 and Axl levels are modified in animals suffering liver fibrosis. After five weeks of CCl4 treatment mice exhibited increased Gas6 and sAxl serum levels compared to oil-treated animals (Fig. 3A), indicating that this pathway is upregulated during CCl4-induced liver fibrosis. Second, we analyzed in Axl−/− mice the effect of CCl4 administration. After five weeks, liver hydroxyproline levels, indicative of collagen deposition, were significantly lower in Axl-deficient animals treated with CCl4 (Fig. 3B), suggesting reduced liver fibrosis as confirmed in liver sections after Sirius Red staining and quantification (Fig. 3C and D). In accordance, liver homogenates exhibited an increase in α-SMA and MMP9 after CCl4 administration that was reduced in Axl−/− mice (Fig. 3E) indicative of HSC activation and changes in ECM composition. Analogously, α-SMA stained liver slides from CCl4-treated Axl−/− mice exhibited similar reduction (Supplementary Fig. 1A). Finally, we checked the degree of liver injury to verify that the lower liver fibrosis observed in Axl−/− mice is not a consequence of reduced hepatocellular damage induced by CCl4. Both WT and Axl deficient mice displayed similar ALT levels after CCl4 exposure, which allows discarding reduced liver fibrosis as a consequence of lesser hepatic damage.

It has been proposed that Gas6 deficiency could lead to a decline in liver fibrosis after CCl4 exposure due to reduced macrophage recruitment [25]. Therefore, we decided to analyze potential differences in liver inflammation and immune cell recruitment to the liver. When we quantified the mRNA levels of inflammatory cytokines (TNF), chemokines (MCP-1) or neutrophil infiltration (MPO) in WT and Axl−/− CCl4-treated mice they were similarly increased, as compared to their untreated controls (Supplementary Fig. 1B and D). In addition, no changes in MPO staining were observed between WT and Axl−/− CCl4-treated animals (Supplementary Fig. 1C). However, a minor level of macrophages (F4/80) and newly-recruited monocytes/-macrophages (CCR2) were detected in the livers of Axl deficient mice after CCl4 exposure (Supplementary Fig. 1D). These results support a role for the Gas6/Axl pathway in macrophage response to CCl4 exposure, as previously indicated [25], and are in line with other models of tissue damage such as advanced atherosclerotic plaques in Gas6 deficient animals [26].

**JOURNAL OF HEPATOLOGY**

**Fig. 1.** Gas6 and Axl levels are increased in WT HSCs during in vitro activation. (A and B) mRNA expression level of Gas6, ProS, Axl, and MERTK in HSCs at different times of in vitro activation, using β-actin as control. (n > 3). (C) Gas6 protein levels released to fresh culture medium during 24 h from HSCs at different time points of in vitro activation, detected by ELISA and corrected by cellular protein content. (n = 3). *p < 0.05, Student’s t test.

BGB324, small molecule inhibitor of Axl, blocks HSC activation in vitro and reduces CCl4-induced liver fibrosis in WT mice

Axl is an attractive target for the treatment of different human pathologies, particularly in cancer. Interestingly, a small molecule inhibitor of Axl (BGB324, BerGenBio) has entered clinical trials for cancer treatment. We tested the effect of BGB324 administration for 24 h on seven-day old WT HSC cells (Fig. 4A). Axl inhibitor administration was able to reduce the activation of primary HSCs, even inducing HSC elimination at higher doses in the micromolar range (Fig. 4A, upper image). This effect was specific for Axl inhibition since BGB324 administration did not affect MERTK phosphorylation (Fig. 4A, lower image), while effectively blocked AKT activation after short-term incubation with rGas6 (30 min., 200 ng/ml) (Fig. 4B).

To analyze whether inhibition of Axl, using BGB324, may play a role in the progression of liver fibrogenesis, mice were injected CCl4 twice weekly to stimulate HSC activation and promote liver fibrosis. After four weeks, animals started receiving BGB324 co-treatment via oral gavage on a daily basis for ten additional days. Determination of the hepatic hydroxyproline content showed a significant decrease in the accumulation of collagen fibers in animals treated with CCl4 that received Axl inhibitor compared to control animals (Fig. 4C). This result was confirmed after Sirius Red staining, showing less deposition of collagen fibers in animals treated with BGB324 and CCl4 compared to CCl4-vehicle treated mice (Fig. 4D), as denoted by the quantification of collagen content in different sections (Fig. 4E). Moreover, mice that received BGB324 exhibited reduced levels of MMP9 and α-SMA after CCl4 exposure (Supplementary Fig. 2A), and diminished α-SMA staining in liver slides compared to CCl4-treated mice without inhibitor administration (Supplementary Fig. 2B), confirming changes in ECM composition.
and HSC activation. Of note, serum levels of ALT after CCl₄ administration were similar in vehicle and Axl inhibitor-treated mice indicating that the antifibrotic effect of BGB324 is not a consequence of reduced hepatocellular damage after chemical exposure (Fig. 4F).

Interestingly, when we analyzed changes in liver inflammation and immune cell recruitment to the liver induced by BGB324, we obtained results in accordance with the data provided by the Axl⁻⁻ mice. While a comparable degree of TNF, MCP-1 or neutrophil infiltration was observed in all CCl₄-treated mice (Supplementary Fig. 2C and D), animals that received BGB324 exhibited decreased macrophage recruitment (Supplementary Fig. 2D), in line with previous results observed analyzing Gas6 KO mice [25].

Serum levels of Gas6 and sAxl correlate with liver dysfunction in human ALD, and increased during HCV-induced fibrosis progression (DCH), and compared them to control individuals (C). Both Gas6 and sAxl were found increased in serum levels of cirrhotic patients, showing close correlation with the severity of the disease, although behaving differently. Specifically, sAxl concentration was already augmented in individuals with compensated cirrhosis compared to initial fibrosis (Fig. 5B), while Gas6 levels were increased markedly in the DCH group (Fig. 5A).

To verify this observation, we examined the relationship between the serum levels of Gas6 and sAxl compared to the Model for End-Stage Liver Disease (MELD) score system, which assigns a value calculated from different biochemical parameters altered in chronic liver disease. The analysis revealed a remarkable correlation between the MELD score and both proteins (Fig. 5C and D), being better for Gas6 serum levels ($r^2 = 0.78$). Interestingly, we identified an algorithm containing sAxl and Gas6 that can achieve even stronger correlation ($r^2 = 0.86$) with the MELD score (Fig. 5D), suggesting that the measurement of both proteins provides a better evaluation of liver functionality.

However, since our ALD group contains only individuals with early F0/F1 fibrosis and with cirrhosis, compensated or decompenated, our measurements did not allow to verify an increase in the Gas6/sAxl system during the progression of fibrosis, or to validate Gas6/sAxl detection in other human hepatic pathologies.

To do so, we analyzed Gas6 and sAxl levels in the serum of HCV patients at different stages of liver fibrosis before starting treatments (Fig. 5E and F). Our data revealed that Gas6 levels were significantly different between individual with established fibrosis (F2) and patients with initial fibrosis (F0 and F1 groups).
MMP9 in liver extracts. (E) ALT serum levels.

with CCl4 or vehicle were measured: (B) Hydroxyproline levels in liver extracts. CCl4 (twice a week) for five weeks. (A) Gas6 and Axl levels in serum from WT mice treated with CCl4 or vehicle (corn oil). Additionally, in WT and Axl−/− mice treated with CCl4 or vehicle were measured: (B) Hydroxyproline levels in liver extracts. (C) Representative images of liver sections after Sirius Red staining (20 random sections from each animal. (E) Representative western blot of antifibrotic drug despite the experimental description of an array regression in chronic liver diseases is not accomplished by any ment [4]. Despite recent progress in understanding the biology ECM-remodeling phenotype that favors tumorigenic development. event that shifts HSCs from a normal vitamin A-storing to an HSC transdifferentiation represents a crucial cell reprogramming development in liver disease.

In addition, sAxl levels displayed significant changes between patients with F2 fibrosis and individuals with advanced fibrosis or cirrhosis (F3/F4 group). These findings underscore the relevance of the Gas6/Axl pathway during the development of ALD- and HCV-induced liver damage, supporting Gas6 and sAxl serum levels as indicative parameters of hepatic dysfunction and fibrosis development in liver disease.

Discussion

HSC transdifferentiation represents a crucial cell reprogramming event that shifts HSCs from a normal vitamin A-storing to an ECM-remodeling phenotype that favors tumorigenic development [4]. Despite recent progress in understanding the biology of HSCs, the mechanisms are not yet fully known. In fact, besides the treatment/withdrawal of the underlying cause, fibrosis regression in chronic liver diseases is not accomplished by any antifibrotic drug despite the experimental description of an array of pharmacological targets [1,5]. In this context, the characterization of the role of Gas6/Axl pathway in liver fibrosis, by participating in the activation of HSC may provide a new therapeutic target, not only for liver fibrosis, but also for different chronic liver diseases. Moreover, the existence of specific Axl inhibitors [30], already in clinical trials, may facilitate the biomedical translation of our results (Fig. 6).

Here, we used BGB324 (BerGenBio), an inhibitor of Axl ready to reach Phase Ib clinical trials for cancer treatment after showing good tolerability by healthy volunteers in doses up to 1.5 g/daily with a long plasma half-life [31]. BGB324 is highly specific for Axl inhibition, having exhibited >100-fold selectivity for Axl vs. Ab1 and 50- and >100-fold selectivity over TAM family kinases MERTK and Tyro3, respectively, in cells-based assays [18]. In fact, we observed that at doses effective to block AKT phosphorylation and HSC activation, BGB324 did not alter MERTK phosphorylation by rGas6. Although, based in our data, we cannot discard other off-target effects of BGB324 administration, the highly
similar results obtained between the Axl−/− mice and the BGB324-treated mice suggest that the anti-fibrotic action of BGB324 is mainly due to Axl inhibition.

High levels of Axl expression have been observed in many types of cancer correlating with poor survival; among them glioblastoma multiforme [32], acute myeloid leukemia [33], breast cancer [34], osteosarcoma [35] and renal cell carcinoma [36]. Moreover, Axl activation is a mechanism of drug resistance to therapies targeting EGFR mechanism in lung cancer [37]. However, Axl and MERTK are also expressed by macrophages and dendritic cells, where they limit excessive immune response [10,38]. This aspect has raised concerns and could limit the use of TAM receptors as targets in cancer, in special since blockage of Axl/ MERTK promote the development of tumor growth in an inflammatory environment such as colon cancer [39], a tumor that benefits of a pro-inflammatory milieu similarly as liver cancer does. Importantly, this deleterious effect seems to require the simultaneous absence of both kinases (Axl and MERTK) activities [40], and BGB324 is already effective at doses that provide serum concentration in the low micromolar range, with minimal affinity for MERTK [18]. This observation is supported by our results in mice, not having detected inflammation in treated animals, displaying similar levels of neutrophil infiltration after CCl4 administration. In fact, BGB324 has just received orphan-drug designation from FDA in the treatment of acute myeloid leukemia, particular for hepatocellular carcinoma, to act without favoring a pro-carcinogenic background. Of note, other approaches to block Axl signaling are already under study, for instance an Axl ‘decoy receptor’ has been recently engineered, showing capacity to inhibit metastasis and cancer progression in vivo [40]. Therefore, BGB324 and other molecules that antagonize the Gas6/Axl pathway deserve to be further analyzed in the context of advanced liver fibrosis, and most probably of liver cancer development.

Several reports have positioned Gas6 as a protective molecule against ischemia/reperfusion [14] and promoter of liver regeneration after acute liver damage [15,16], while its deficiency was associated to decreased liver fibrosis [25]. We considered using Gas6 KO mice to analyze Gas6/Axl system in HSC activation but previous results showing compensatory alterations such as Axl overexpression in the liver [25], refrained us to do it. In fact, we verified in murine Gas6−/− HSCs high expression of Axl, increased AKT phosphorylation, and elevated α-SMA levels, among other markers of HSC activation (Supplementary Fig. 3). Since Gas6/Axl axis is not blocked in Gas6−/− HSCs, we preferred to use the Axl−/− mouse model instead. In this sense, our data targeting directly Axl underscore the importance of Gas6/Axl pathway in liver disease, being supported by similar results obtained after administration of the small molecule inhibitor BGB324.

Although it has been reported an increase in Gas6/Axl proteins in patients with hepatocellular carcinoma from different etiologies [41], and sAxl levels are increased in end-stage heart failure patients undergoing heart transplantation [27], which
frequently suffer cardiac fibrosis, Gas6/Axl as serological markers of liver function have not been previously proposed. We have confirmed in serum samples from ALD and HCV patients an enhancement in Gas6 and sAxl serum levels. Although we can not discard that the prominent increase in Gas6/Axl observed in these ALD cirrhotic patients may be partially due to a deficient liver protein clearance, the observation that a similar increase is also detected in fibrotic HCV patients, even in patients without cirrhosis, suggests otherwise. Moreover, an algorithm combination of Gas6 and sAxl levels display an excellent correlation with the degree of liver dysfunction in ALD patients determined by the MELD index ($r^2 = 0.86$ for sAG index) suggesting that the measurement of both proteins provides additional information to analyze liver functionality. In this sense, we consider that preclinical research may also benefit from Gas6/Axl levels to measure fibrosis progression, since numerous targets for antifibrotic agents have problems to be analyzed or to enter early-phase clinical studies due to the lack of sensitive markers to follow the effects [42]. These results point to Gas6/sAxl determination as a significant diagnostic tool for chronic liver disease, and help us to position Gas6/Axl signaling pathway as a relevant in human liver pathology.

In conclusion, Gas6/Axl is a profibrogenic route that is activated in patients with chronic liver disease. Therefore, small molecule inhibitors against Axl, that effectively eliminate HSC activation and reduce experimental fibrosis progression, may be an interesting therapeutic tool for future clinical trials.

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Conflict of interest

P.G.F. is inventor on a patent filed for use of sAxl for diagnosis/prognosis of heart failure syndrome (U.E. patent number EP 13703603.4). Other authors declare no competing interests.

Authors’ contributions

C.B., M.S., A.T., L.J., L.M., and A.M. performed the experiments; C.G.R., P.S.-B., and J.C. analyzed clinical data and discussed the results; C.B. drafted the manuscript; J.F.C., M.M., C.V.R., P.G.F., and A.M. designed experiments and revised the results; P.G.F. and A.M. were primarily responsible for writing the manuscript. All authors contributed to manuscript editing and approval.

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Supplementary data

Supplementary data associated with this article can be found, in the online version, at http://dx.doi.org/10.1016/j.jhep.2015.04.013.

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Authors names in bold designate shared co-first authorship.

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