

Inositol 1,4,5-trisphosphate receptor type 3 is involved in resistance to apoptosis and maintenance of human hepatocellular carcinoma

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Abstract. The expression of the inositol 1,4,5-trisphosphate receptor type 3 (ITPR3) in hepatocytes is a common event in the pathogenesis of hepatocellular carcinoma (HCC), regardless of the type of underlying liver disease. However, it is not

known whether ITPR3 expression in hepatocytes is involved in tumor maintenance. The aim of the present study was to determine whether there is an association between ITPR3 expression and clinical and morphological parameters using HCC samples obtained from liver explants from patients (n=53) with different etiologies of underlying chronic liver disease (CLD). ITPR3 expression, mitosis and apoptosis were analyzed in human liver samples by immunohistochemistry. Clinical and event-free survival data were combined to assess the relationship between ITPR3 and liver cancer growth in patients. RNA sequencing analysis was performed to identify apoptotic genes altered by ITPR3 expression in a liver tumor cell line. ITPR3 was highly expressed in HCC tumor cells relative to adjacent CLD tissue and healthy livers. There was an inverse correlation between ITPR3 expression and mitotic and apoptotic indices in HCC, suggesting that ITPR3 contributed to the maintenance of HCC by promoting resistance to apoptosis. This was confirmed by the upregulation of CTSB, CHOP and GADD45, genes involved in the apoptotic pathway in HCC. The expression of ITPR3 in the liver may be a promising prognostic marker of HCC

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Abbreviations: AFP, α fetoprotein; ALD, alcoholic liver disease; CAAE, Certificado de Apresentação de Apreciação Ética (Certificate of Presentation of Ethical Appreciation); CC, cryptogenic cirrhosis; CHILD, CHILD-Pugh scores; CHOP, C/EBP homologous protein; CLD, chronic liver disease; CTSB, cathepsin B; EFS, event-free survival; H&E, hematoxylin and eosin; HCC, hepatocellular carcinoma; HCV, hepatitis C virus; HepG2, immortalized liver cancer cell line; HPF, high-power fields; INR, international normalized ratio; ITPR, inositol 1,4,5-trisphosphate receptor; ITPR1, inositol 1,4,5-trisphosphate type 1 receptor; ITPR2, inositol 1,4,5-trisphosphate type 2 receptor; ITPR3, inositol 1,4,5-trisphosphate type 3 receptor; MELD, model for end-stage liver disease; OLT, orthotopic liver transplantation; PH, partial hepatectomy; RNA-seq, RNA sequencing; TBIL, total bilirubin

Key words: apoptosis, calcium signaling, chronic diseases, liver cancer, mitosis

Introduction

Hepatocellular carcinoma (HCC) is the commonest primary liver malignancy and its incidence is increasing worldwide (1). According to the GLOBOCAN database, in 2020, including intrahepatic bile ducts, there were 905,677 new cases of liver cancer around the world, accompanied by 830,180 new deaths (2). Liver cirrhosis is a major risk factor for HCC (3). Although liver transplantation is the primary long term curative treatment, HCC still has a recurrence rate of 10-15% in

transplant patients (4). Since biopsy is not mandatory for HCC diagnosis (5), the availability of tumor specimens for analysis is limited, which in turn has limited our understanding of the molecular mechanisms involved in carcinogenesis.

The calcium ion (Ca^{2+}) is an intracellular second messenger involved in energy metabolism, cell cycle control, gene expression, cell proliferation, cell migration, necrosis and apoptosis in nearly every cell type, including hepatocytes (6). The inositol 1,4,5-trisphosphate receptor (ITPR) is the only intracellular Ca^{2+} release channel expressed in the liver, with all three isoforms described in humans and rodents (7,8). The expression of each ITPR varies among cell type, having distinct subcellular distribution and biophysical properties (9). For instance, under physiological conditions type 1 (ITPR1) and 2 (ITPR2) are the predominant isoforms in hepatocytes (10,11), while isoform 3 (ITPR3) is absent or minimally expressed (12). However, ITPR3 becomes significantly expressed in hepatocytes in acute (13,14) and chronic liver diseases, as well as in HCC and in liver cancer cell lines (15,16). ITPR3 expression is also increased in other malignancies, including cholangiocarcinoma (17), colon cancer (18), melanoma (19), mesothelioma and prostate cancer (20).

Epigenetic events have been associated with the expression of ITPR3 in HCC, with hypomethylation of the promoter region of the ITPR3 gene upregulating its expression (16). In contrast, the promoter regions of ITPR1 and ITPR2 are already demethylated in normal hepatocytes (16), consistent with the observation that ITPR1 and ITPR2 are constitutively expressed in this cell type (10).

To understand the biological role of ITPR3 in HCC, the present study compared ITPR3 expression in HCC and adjacent cirrhotic parenchyma in patients with different types of underlying chronic liver disease. It also correlated ITPR3 expression with clinical and morphological parameters in order to investigate whether ITPR3 could serve as a prognostic marker.

Materials and methods

Sample characterization. A total of 53 liver explants from patients who underwent orthotopic liver transplantation (OLT) in Hospital das Clínicas, Universidade Federal de Minas Gerais (UFMG; Federal University of Minas Gerais), Brazil, between January 2002 and December 2017, were retrospectively reviewed after the study was approved by the local Ethics Committee, COEP-UFMG, in the city of Belo Horizonte, state of Minas Gerais, Brazil (CAAE 71206617.8.0000.5149) and written informed consent was obtained from the patients or their relatives. Histologically normal tissues were obtained from liver resections of patients with metastatic colon cancer between 2010 and 2017 at Hospital das Clínicas (UFMG), after rigorous examination by an experienced pathologist.

A representative slide of each case was selected and analyzed after staining with conventional hematoxylin and eosin (H&E) and immunohistochemistry. Formalin-fixed (concentration 10%; duration 12 h at room temperature) and paraffin-embedded samples were deparaffinized and antigen retrieval was performed in citrate buffer (10 mM) containing 0.6% peroxide of hydrogen (heating temperature, 65°C for 8 h

in the thermal oven), followed by four immersions in xylene (the first for 20 min and the others with 1 min duration) and three immersions in absolute alcohol (the first for 5 min and the others with 1 min duration). Afterwards, the slides were immersed in running water for 3 min. The Novolink Polymer Detection System (cat. no. RE7200-CE; Leica Microsystems, Inc.) was used in the subsequent steps as described previously (21). Primary antibodies against ITPR3 (anti-ITPR3; Sigma-Aldrich; Merck KGaA; cat. no. HPA003915) and ssDNA labeling (anti-Single Stranded DNA; cat. no. 18731; Immuno-Biological Laboratories, Co., Ltd.) were incubated overnight at room temperature in a 1:100 dilution, followed by incubation with detection polymer for 40 min at room temperature. DAB was used for signal detection.

Clinical pre-OLT data and histopathological parameters of the explanted liver were retrospectively collected from the medical records, including sex, age, etiology of the underlying chronic liver disease (CLD), model for end-stage liver disease (MELD) and CHILD-Pugh scores (CHILD); prognostic models that estimate the severity of underlying liver condition, pre-OLT serum levels of α fetoprotein (AFP), number of nodules, size of the greatest nodules, microvascular invasion and histological grade. The medians of MELD, AFP, total bilirubin, international normalized ratio, creatinine, alanine aminotransferase, aspartate aminotransferase and serum albumin were obtained from the sum of the values of each of these clinical data for each patient. Therefore, the medians reflect the mean of the total sum of the 53 patients in this study, between the minimum and maximum values.

Outcomes, such as death from any cause or tumor recurrence since OLT and event-free survival (EFS), defined as the time interval between the OLT date and the occurrence of the event or ending of the follow-up period (December 20, 2017), were analyzed.

Analysis of ITPR3 expression intensity and ssDNA labeling index. To evaluate ITPR3 expression, images of immunohistochemistry slides stained for ITPR3 from 10 different fields (magnification, x40) were captured in tumor and adjacent cirrhotic parenchyma through an optical microscope (Zeiss GmbH). The intensity of ITPR3 staining was evaluated by delimiting 10 cytoplasmic regions, excluding the nucleus, using ImageJ 1.50i software (National Institutes of Health). The software performed this analysis through a histogram, which expressed the intensity of the pixels in each selected region on a scale between 0-255. ITPR3 expression intensity was expressed as the mean of 10 analyzed cells in each of the 10 imaged fields.

Mitotic index evaluation. A total of two experienced liver pathologists identified areas of interest and counted the number of mitotic cells in 10 high-power fields (HPF) of H&E stained slides (magnification, x20) in order to identify areas containing mitotic figures, in metaphase or anaphase. Following Baak (22), the mitotic cells were counted only with a complete agreement between the two pathologists through a multi-head microscope. The delineation between low mitotic index in samples with >5 mitotic figures at 10 HPF and high mitotic index in samples with ≥ 5 mitotic figures at 10 HPF was based on previous studies (23-27).

Pre-processing of raw RNA sequencing (RNA-seq) data. Raw read sequences were pre-processed by filtering out read sequences with less than 36 bp, removing low quality or N bases from the read ends, trimming Illumina adapters, scanning the read with a 4-base wide sliding window and cutting when the average quality per base dropped below 15 (LEADING:3 TRAILING:3 SLIDINGWINDOW:4:15 MINLEN:36) by Trimmomatic v0.39 (28).

The raw RNA-seq data were obtained in our previous publication, through experiments with HepG2 (16), considered a cell line of liver cancer. A commercially available CRISPR/Cas9 system was used to eliminate ITPR3 in HepG2 liver cancer cells (Santa Cruz Biotechnology, Inc.). Cells were grown in Dulbecco's Modified Eagles Medium (cat. no. 11320033) supplemented with L-glutamine (1 mM; cat. no. 25030149), fetal bovine serum (10% v/v; cat. no. A4766801) and penicillin/streptomycin (100 units/ml and 100 mg/ml; cat. no. 15140122) at 37°C in a humid environment with an artificial atmosphere containing 5% CO₂.

Quantification and differential gene expression analysis. Quantification of the gene expression (-g) was performed by Salmon v1.2.1 (29). The transcriptome index was built using the cDNA sequences of the *Homo sapiens* assembly version GRCh38 downloaded from Ensembl release 104 (30). The estimated number of reads were used for the differential gene expression analysis. Genes with low expression levels were removed; genes with counts scaled by total mapped reads per million >1 in at ≥9 samples were retained for differential gene expression analysis. DESeq2 (31) was applied to evaluate the differential gene expression using DEBrowser v1.16.1 (32). Genes were considered differentially expressed when they had an adjusted P-value <0.05; fold-change ≤-1.5 (downregulated or more enriched in control condition).

Identification of differentially expressed genes associated with apoptosis. Genes participating in the human apoptosis route were extracted from the Kyoto Encyclopedia of Genes and Genomes (KEGG) Pathways (33) Apoptosis map (hsa04210) was obtained using the rest application programming interface of KEGG. Genes on the apoptosis pathway were mapped to their corresponding UniProt identifiers by UniProt's idmapping (The Uniprot Consortium 2021) (34). The same was done for the ensemble gene IDs of differentially expressed genes. An in-house custom python script was used to compare both lists and identify differentially expressed genes participating on the apoptosis pathway. A custom colorized version of the apoptosis pathway, containing upregulated genes in green, downregulated genes in red and non-differentially expressed genes in white was generated using KEGG Mapper (35).

Heatmap. Heatmaps of the top 50 differentially expressed genes sorted by the average Reads per Kilobase per Million of mapped reads were generated for each condition by Heatmapper (36) using average linkage clustering and Euclidean measurement methods.

Statistical analysis. Descriptive statistics summarized the data, which were expressed as numbers and percentages (categorical) and as median and interquartile range (continuous). A

normality test (Shapiro-Wilk) was performed for each continuous variable. For comparison between the etiological groups, χ^2 or the Fisher's exact test was applied in categorical variables and Mann Whitney or Kruskal-Wallis test in continuous data. In multiple comparisons, Dunn's test was applied after the Kruskal-Wallis test. Multivariate COX regression model with covariance structure was performed to determine the model for EFS. Moreover, to evaluate the correlation between the clinicopathological parameter and ITPR3 expression, Spearman's coefficient was used. SPSS software, v20 (IBM Corp.), was used for statistical analysis. Three technical replicates were used per sample in the analyses. Final results were expressed as the mean of the three values. P<0.05 was considered to indicate a statistically significant difference.

Results

The etiologies of the underlying chronic liver diseases among the explanted livers were distributed as follows: Hepatitis C virus (HCV) infection (n=31; 58.5%), alcoholic liver disease (ALD; n=16; 30.2%) and cryptogenic cirrhosis (CC; n=6; 11.3%). Multiple etiologies or coexistence of other liver pathologies were considered exclusion criteria.

Among all etiological groups, the median age of the patients was 57.09 years (range, 41.11-74.03). The median MELD score was 15.0 (range, 6.0-32.0) and the median AFP was 123.78 (range, 1.00-2883.00). The median total bilirubin was 2.65 (range, 0.50-8.70), the median international normalized ratio was 1.045 (range, 0.990-2.690) and the median creatinine was 1.02 (range, 0.14-3.00). The median alanine aminotransferase was 91.3 (range, 24.0-325.0), the median aspartate aminotransferase was 85.54 (range, 25.00-267.00) and the median serum albumin was 3.20 (range, 2.00-4.80). All clinical and laboratory data are summarized in Table I.

Morphological descriptions of the livers from HCC patients are presented in Table II. The number of nodules was categorized into two groups: ≤3 nodules and >3 nodules. Vascular invasion was categorized as present or absent. Differentiation grade, according to Edmondson and Steiner (37), was classified into I and II (Low grade; well differentiated) and III and IV (high grade; poorly differentiated). The non-categorical data (mitosis and nodule size) are expressed as median values.

As ITPR3 expression is a common event in HCC development (16), the presence and distribution of this Ca²⁺ channel was evaluated in HCC samples from different etiologies. ITPR3 staining of cholangiocytes was used as an internal positive control, as this liver cell type constitutively expresses ITPR3 (29). There was greater ITPR3 labeling, both in the non-tumor regions of the explants (Fig. 1A) and the tumor regions (Fig. 1B), relative to hepatocytes from healthy control livers. ITPR3 labeling in the tumor region of patients with HCC caused by HCV (Fig. 1C) or ALD (Fig. 1D) were higher in comparison with the adjacent non-tumor region, while no difference was observed between these two regions in CC patients (Fig. 1E). Although the level of ITPR3 staining was similar among the etiologic groups in both the cirrhotic parenchyma and the tumor regions (Fig. 1F), the pattern of labelling varied among groups. In the tumor regions and in the underlying cirrhotic parenchyma, homogeneous staining of ITPR3 was observed, with ITPR3 dispersed throughout the

Table I. Clinical and laboratory data of patients with hepatocellular carcinoma.

Parameter	Minimum	Median	Maximum
Age (years)	41.11	57.09	74.03
MELD	6.0	15.0	32.0
AFP (ng/ml)	1.00	123.78	2883.00
TBIL (mg/dl)	0.50	2.65	8.70
INR	0.990	1.045	2.690
Creatinine (mg/dl)	0.14	1.02	3.00
ALT (U/l)	24.0	91.3	325.0
AST (U/l)	25.00	85.54	267.00
ALB (g/dl)	2.00	3.20	4.80

MELD, model for end-stage liver disease; AFP α fetoprotein; TBIL, total bilirubin; INR, international normalized ratio; ALT, alanine aminotransferase; AST, aspartate transaminase; ALB, albumin.

Table II. Anatomopathological characteristics of liver from patients with hepatocellular carcinoma.

Clinical parameters	Value
Nodules number ^a	
≤ 3	26 (55.3%)
> 3	21 (44.7%)
Vascular invasion ^a	
Present	23 (54.8%)
Absent	19 (45.2%)
Differentiation degree ^a	
Low degree (I-II)	35 (72.9%)
High degree (III-IV)	13 (27.1%)
Histological type ^a	
Pseudo-acinar	6 (12.8%)
Solid	3 (6.4%)
Trabecular	22 (46.8%)
Pseudo-acinar/trabecular	16 (34.0%)
Mitosis (10 HPF) ^b	5.0 (0-22.0)
Nodule size (cm) ^b	3.0 (1-10.5)

Data are expressed as ^aabsolute numbers (percentage) or ^bmedian (interquartile range). HPF, high-power fields.

hepatocyte cytoplasm (Fig. 1A and B; blue arrows). In some cases, as observed in CC, a predominant perinuclear region of ITPR3 staining (Fig. 1A and B; black arrows) was present.

As in other malignancies, ITPR3 expression is an early event in HCC carcinogenesis (16,18) which may then relate to morphological aspects as well as clinical outcomes of patients affected by the disease. Thus, the correlation between ITPR3 expression and clinicopathological parameters was investigated. Table III showed that serum AST levels positively correlated with the intensity of the tumor ITPR3 labeling profile ($P < 0.02$), a correlation that was not observed in the

Table III. Correlation between ITPR3 labeling intensity with anatomopathological and clinical data parameters.

Clinical parameters	Cirrhosis P-value	Tumor P-value
Number nodules	0.120	0.296
Vascular invasion	0.637	0.746
Necrosis	0.114	0.461
Fibrosis	0.752	0.456
Tumor inflammation	0.383	0.552
Intracellular characteristics	0.927	0.825
Differentiation degree	0.677	0.668
Histological pattern	0.601	0.941
MELD	0.395	0.393
AFP (ng/ml)	0.703	0.781
TBIL (mg/dl)	0.496	0.694
INR	0.403	0.344
Creatinine (mg/dl)	0.216	0.248
ALT (U/l)	0.410	0.281
AST (U/l)	0.497	0.018 ^a
Mitosis (10 HPF)	-	0.0098 ^b

MELD, model for end-stage liver disease; AFP, α fetoprotein; TBIL, total bilirubin; INR, international normalized ratio; ALT, alanine aminotransferase (^a $P < 0.098$. Spearman post hoc test); AST, aspartate transaminase; ALB, albumin; HPF, high-power fields (^b $P < 0.098$; Student t-test).

adjacent parenchyma ($P > 0.4$; using Spearman's statistical test).

A high mitotic index ($\geq 5/10$ HPF) is a poor prognostic factor in HCC, as well as with intrahepatic metastasis, larger tumor size, higher AFP levels, advanced tumor stage (24,28), vascular invasion, poor histological differentiation and recurrence rate (26). Therefore, the mitotic index was correlated with the intensity of ITPR3 labeling. Mitotic index was categorized as either low (< 5) or high (> 5). The majority of cases (62.8%) had a low mitotic index ($n=22$), while 37.2% ($n=13$) exhibited high mitotic index. In addition, patients with greater ITPR3 staining exhibited a lower mitotic index, while patients with less ITPR3 labeling intensity exhibited a higher mitotic index (Fig. 2A). The mitotic index was inversely associated with ITPR3 expression in the tumor region ($P < 0.01$; using unpaired Student's t-test). These results suggest that cell proliferation is not the process by which ITPR3 regulates liver tumor maintenance. Therefore it was investigated whether Ca^{2+} may instead modulate apoptosis.

Apoptosis is involved in both cancer growth (19) and liver regeneration (30). Our group has previously demonstrated that ITPR3 expression can induce resistance to cell death in HCC (16). Thus, the correlation between the apoptotic marker ssDNA labelling and ITPR3 expression was investigated. In the HCC samples, 69.7% ($n=23$) had a low apoptotic index ($\leq 50\%$), whereas 30.3% ($n=10$) exhibited a high apoptotic percentage ($> 50\%$) (Fig. 2B). Patients with higher intensity of

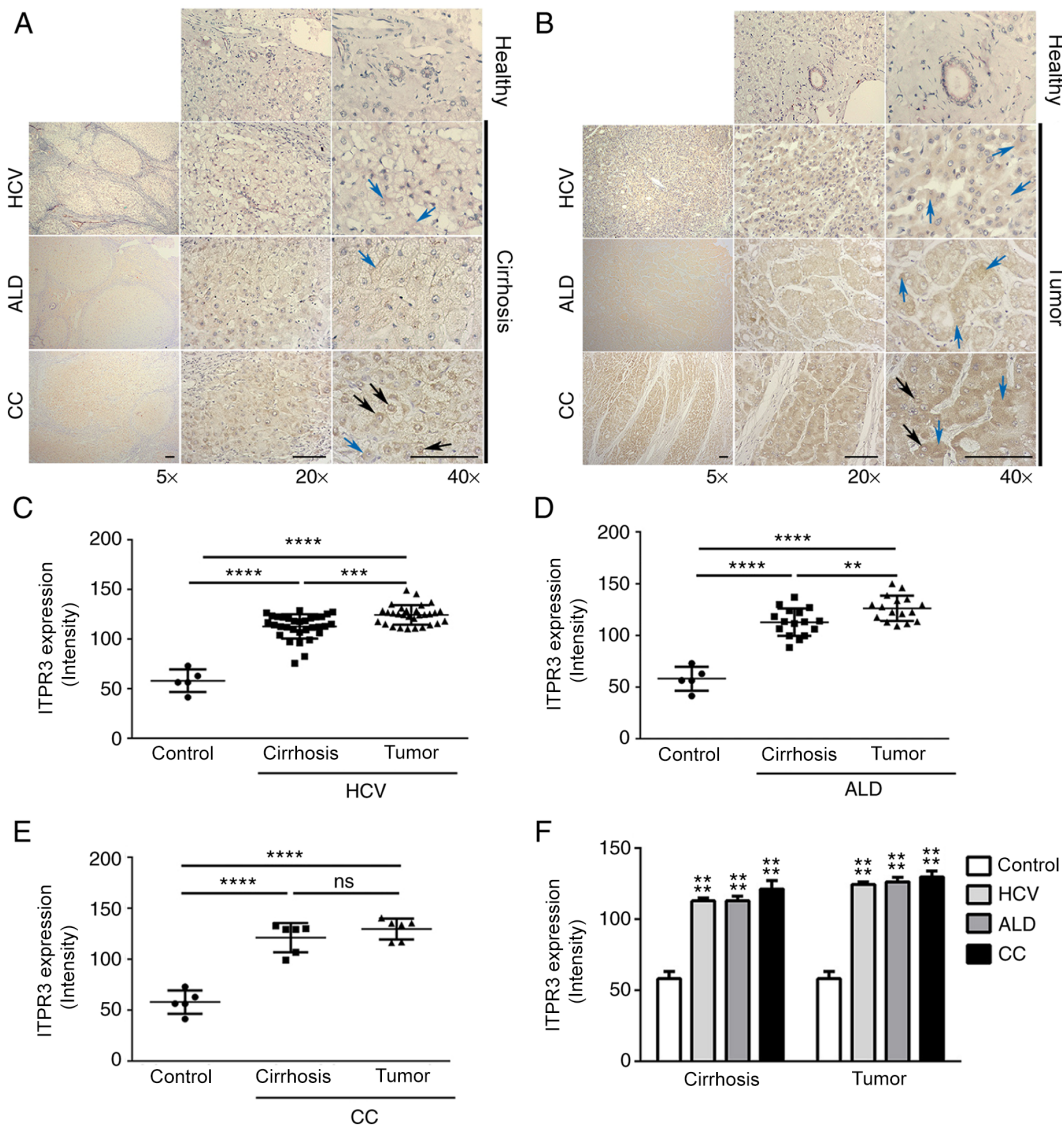


Figure 1. Representative images of ITPR3 labeling in HCC, underlying cirrhotic parenchyma and non-tumor livers (controls). (A) ITPR3 labeling in cirrhosis in patients diagnosed with HCV (n=31), ALD (n=16), CC (n=6) and healthy (control; n=2). Scale bar=200 μ m (magnification, x5), 50 μ m (magnification, x 20) and 25 μ m (magnification, x40). Black arrows indicate ITPR3 perinuclear localization in hepatocytes. Blue arrows indicate ITPR3 cytoplasmatic localization in hepatocytes. (B) ITPR3 labeling in tumor (HCC) region of patients diagnosed with HCV (n=31), ALD (n=16), CC (n=6) and healthy (control; n=3). Black arrows indicate ITPR3 perinuclear localization in hepatocytes. Blue arrows indicate ITPR3 cytoplasmatic localization in hepatocytes. (C) Representative graph of the intensity of ITPR3 marking in HCC regions and cirrhotic parenchyma of patients with HCV compared with non-tumor livers (control; n=31; ****P<0.0001; ***P<0.0001). (D) Representative graph the intensity of ITPR3 marking in HCC regions and cirrhotic parenchyma of patients with ALD compared with non-tumor livers (control; n=16; **P<0.0057; ***P<0.0001). (E) Representative graph the intensity of ITPR3 marking in HCC regions and cirrhotic parenchyma of patients with CC compared with non-tumor livers (control; n=06; ****P<0.0001). (F) Representative graph the ITPR3 labeling intensity in tumor samples and cirrhotic parenchyma comparing the three etiologies (****P<0.0001). ITPR3, inositol 1,4,5-trisphosphate type 3 receptor; HCC, hepatocellular carcinoma; HCV, hepatitis C virus; ALD, alcoholic liver disease; CC, cryptogenic cirrhosis; ALD, alcoholic liver disease.

ITPR3 staining exhibited a low apoptotic percentage, while patients with lower intensity of ITPR3 displayed a higher apoptotic percentage (P<0.0005; Fig. 2C). These findings suggested that upregulation of ITPR3 correlates with higher apoptosis resistance, which could contribute to tumor maintenance.

As the mitotic index and apoptotic percentage were both inversely associated with the intensity of tumor ITPR3 staining, whether these two parameters correlated with each other in HCC was evaluated. In the specimens of the present study, the degree of apoptosis and mitosis were directly related (P<0.05 by unpaired Student's t-test; Fig. 2D), indicating that these events were closely related in the context of the role of ITPR3 in the modulation of the cell cycle and tumoral maintenance in HCC.

To improved understanding of the mechanism by which ITPR3 expression in hepatocytes is protective, rather than permissive, of cell death by apoptosis, RNA-seq was analyzed comparing global gene expression patterns between wild-type and ITPR3 knockout HepG2 cells. The apoptotic genes CTSB, CHOP and GADD45 were found to be upregulated in the ITPR3 KO cells (Figs. 3 and S1). CTSB is a protease whose overexpression indicates poor prognosis of patients with HCC, facilitating tumor migration and invasion (38). CHOP is a potential pro-apoptotic oncogene in HCC, participating in the promotion of hepatic carcinogenesis through pathways involved in crosstalk with ATF6 (39). The GADD45 family of proteins responds to physiological and environmental cellular

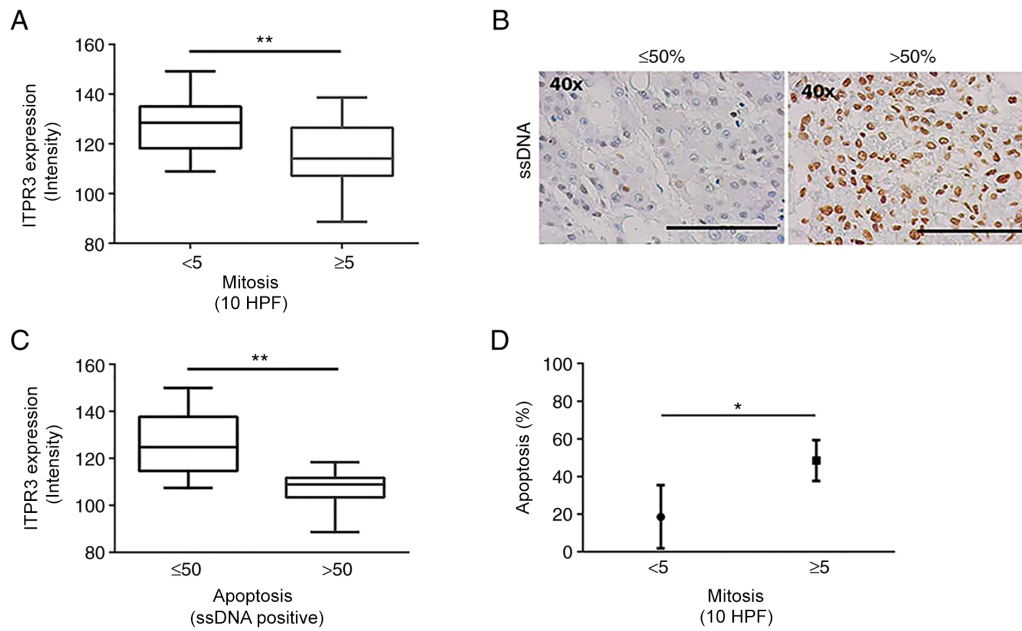


Figure 2. Correlation between ITPR3 labeling intensity and mitotic and apoptosis percentages (ssDNA). (A) Low mitotic (<math><5</math>), high mitotic (>math>\ge 5</math>); $n=35$. $^{**}P<0.0098$ Student's t-test. (B) Representative images of ssDNA labeling by immunohistochemistry, low nuclear labeling for ssDNA ($\le 50\%$) and high ssDNA labeling (>math>50\%</math>); Scale bar=25 μm (magnification, x 40). (C) Correlation graph between ITPR3 labeling intensity and apoptosis percentage. Low apoptosis percentage ($\le 50\%$), high apoptosis percentage (>math>50\%</math>); $n=33$. $^{**}P<0.0003$ Student's t-test. (D) Correlation graph between mitotic index and percentage of apoptosis ($n=15$; $^{*}P<0.0220$). ITPR3, inositol 1,4,5-trisphosphate type 3 receptor; ssDNA, single-stranded DNA; HPF, high-power fields.

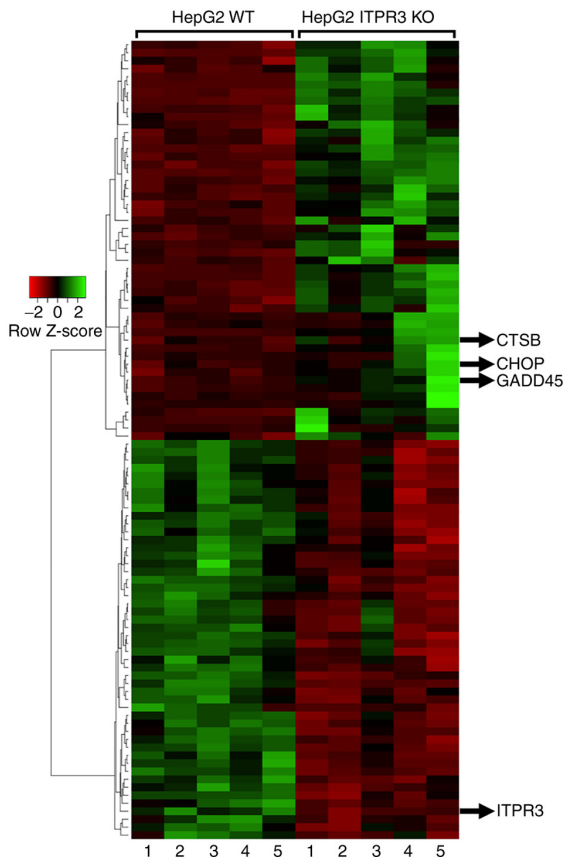


Figure 3. Heatmap of the most upregulated and downregulated genes in WT and ITPR3 KO HepG2 cells. Five different samples were analyzed by RNA-seq and several genes were altered when ITPR3 was not expressed. To confirm efficient depletion of ITPR3 in these cells, the ITPR3 gene was highlighted between the downregulated genes. CTSB, CHOP and GADD45 were the genes associated with apoptosis. WT, wild type; KO, knockout; ITPR3, inositol 1,4,5-trisphosphate type 3 receptor.

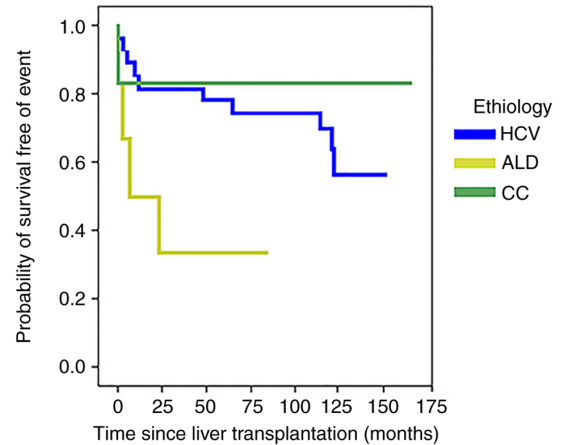


Figure 4. Analysis of the EFS among the three etiological groups. Kaplan-Meier curve, with a difference in the comparison of the EFS between the etiological groups (Renyi test=0.015). Blue line representing the EFS of patients with HCV-associated tumor; Green line representing the EFS of patients with CC-associated tumor and yellow line representing the EFS of patients with ALD-associated tumor. EFS, event-free survival; HCV, hepatitis C virus; CC, cryptogenic cirrhosis; ALD, alcoholic liver disease.

stress (40) through apoptosis (41), with a specific under-expression in HCC that correlates with worse prognosis (41). These results suggest that chronic ITPR3 expression may promote an anti-apoptotic response pathway in liver cancer through modulation of gene expression and proteasome activity.

In the survival analysis, the median event-free survival (EFS) was 113.2 months (range, 93.1-133.2), considering all etiologies together. The Kaplan-Meier curve showed a significant difference comparing the EFS among etiologic groups (Renyi test=0.015; Fig. 4). This difference was identified

between CC and HCV etiologies ($P=0.001$). However, neither ITPR3 expression in the tumor nor in the cirrhotic region was associated with EFS outcome. Together, the results suggested that ITPR3 expression in transformed hepatocytes is implicated in evasion of apoptosis, which contributes to the maintenance of HCC.

Discussion

Hepatocarcinogenesis is a process that involves multiple steps and in which several pathways can interact (42). Among them, regulators of cell proliferation and survival are prominent (43). Although different types of underlying CLD can lead to tumorigenesis through different mechanisms (44,45), *de novo* expression of the ITPR3 Ca^{2+} channel was noted as a common molecular signaling pathway that occurs in all HCC-related etiologies (16). The present study confirmed the involvement of ITPR3 in HCC of patients with HCV, ALD and CC.

Together, two main molecular classes of HCC can be identified (46). The proliferation class, characterized by a higher availability of signals involved in cell cycle proliferation and cell cycle progression, as well as progenitor cell markers, is associated with a more aggressive clinical course. Such tumors have higher AFP levels, less cell differentiation and more frequent vascular invasion (47,48). These characteristics are in turn associated with greater chances of recurrence after initial treatment and lower survival rates (48,49). HCC from hepatitis B virus infection is mostly in this class (50). The non-proliferative class of HCC possesses at least two central molecular characteristics: A predominance of Wnt signaling in $\leq 25\%$ of cases and an immune response in the remainder of cases; in the physiological context, the tumor transcriptome of this class is similar to normal liver physiology (50). From a clinical perspective, tumors in this class exhibit less aggressive characteristics, such as higher cell differentiation index and lower AFP levels. Regarding the underlying etiology, HCV and ALD-related HCC are more frequent in this class (48).

In the sample of the present study, the main CLD etiologies were HCV and ALD, with average AFP serum levels of 123.8 ng/ml, which is relatively low, as only levels above 400 ng/ml are considered as a threshold in the diagnosis of HCC (51). Regarding morphology, the tumor was well differentiated (grade I or II of Edmondson and Steiner score) in 72.9% of patients, the median nodule size was <3.0 cm and 62.8% of tumors had a low mitotic index. Taken together, these data indicated that the majority of the tumors in the present study would be classified as non-proliferative. Moreover, the tumor stage was not as advanced, which is compatible with some methods of treatment received; OLT is considered for patients fulfilling the Milan Criteria, as in our hospital: Hospital das Clínicas of the Federal University of Minas Gerais.

The prognostic importance of the mitotic index and tumor stage is appreciated in a number of malignancies (52,53). In addition, its value is similar to that of techniques such as nuclear antigen of cell proliferation (54) or DNA analysis (55,56). In the context of HCC, there is evidence that a high mitotic index is an unfavorable prognostic indicator (24,25,57,58), as well as an independent predictor of short-term survival in patients who undergo hepatectomy with curative intent (23). In addition, a

high mitotic index ($\geq 5/10$ HPF) is associated with intrahepatic metastasis, larger tumor size, higher AFP levels and advanced tumor stage (23,27), as well as vascular invasion (26), poor histological differentiation and high recurrence rate (23). In the samples in the present study, most patients (62.8%) had a low mitotic index ($<5/10$ HPF). This corroborated *in vivo* and *in vitro* studies which demonstrated that both HCV and ALD negatively affect cell cycle, promoting a G2/M phase arrest in HCC cell lines (59,60).

Demethylation of the promoter of the ITPR3 gene is necessary for its expression in hepatocytes and has been associated with cell proliferation in mouse livers after partial hepatectomy (PH) (16). The data of the present study showed that the higher the expression of ITPR3, the lower the mitotic index. These apparent discrepancies can be explained, in part, by the methodologies used in these studies (PCNA vs. mitotic index), as well as the type of samples analyzed (animal PH model vs. human HCC specimen). PCNA, used in the previous study, is known to serve an important role in the metabolism of nucleic acids and its main function is associated with DNA replication (61). However, PCNA may also be present in repair by DNA excision, cell cycle control, chromatin assembly and RNA transcription (61). Therefore, PCNA staining might also reflect cells in phases preceding mitosis. Another important factor is the type of sample used, either the PH model, in which hepatocytes are prompted to proliferate as a physiological process of the liver (62), or the liver with already established HCC. So, while previous work showed that ITPR3 stimulates hepatocyte proliferation and that its sustained expression in hepatocytes might lead to liver cancer (16), the findings of the current study supported a role for ITPR3 in the maintenance of the liver tumor by preventing cell death, rather than through enhanced hepatocyte proliferation.

It is known that liver growth involves apoptosis, both in physiological conditions, such as in regeneration (29) and in pathological situations, such as in HCC (16). Additionally, Ca^{2+} signaling through ITPR3 is part of apoptotic cell death (30) and the role of ITPR3 in carcinogenesis has been recently reviewed (63). ITPR3 can participate in both inhibiting and promoting apoptosis, depending on the type of tumor (16). The present study demonstrated an inverse correlation between apoptosis and liver tumor ITPR3 expression. This is consistent with a previous finding that ITPR3 triggers anti-apoptotic gene expression in HCC cell lines (16). Similarities occur in colon cancer cells, in which ITPR3 overexpression is associated with decreased apoptosis (18) and in hepatocytes infected with yellow fever virus, in which higher ITPR3 hepatocyte expression prevents cell death through apoptosis (14).

ITPR3 expression can induce resistance to cell death in liver cancer by increasing the expression of POU4F1 and decreasing SIAH2 in WT cells compared with HepG2 ITPR3KO (16). The present study extended these previous observations using RNA-seq and observed that more genes have their expression profile altered in relation to ITPR3 and, also, it is suggested that the signaling pathways associated with apoptosis resistance were altered in HepG2 ITPR3 KO cells (Fig. S1). In particular, a significant upregulation of CTSB, CHOP and GADD45 was observed, genes associated with either cell survival or apoptosis.

In the context of HCC, CHOP is pro-apoptotic (39), performing a pro-tumorigenic function, being an oncogene in hepatic carcinogenesis (39). CHOP is ubiquitously expressed at very low levels normally, but is strongly expressed in cells subjected to severe stress (64). However, activation of this gene is not universally observed in human HCC, which leads to the suggestion that there are other oncogenic pathways that promote hepatocarcinogenesis independently of CHOP. Thus, this gene may participate more in the progression of HCC rather than in its induction (39). The data from the present study suggested that the action of ITPR3 on CHOP expression was associated with the apoptotic signaling pathway by promoting maintenance of liver cancer development.

Growth arrest and damage to DNA 45 (GADD45) is a family composed of three homologous acidic proteins (α , β and γ) (65), which respond to cellular stresses of environmental and physiological origin (41). GADD45 proteins are also involved in hepatic tumorigenesis, in both carcinogenesis and established HCC (65). In the context of HCC, GADD45 β has a pro-apoptotic action and occurs at a low or absent expression level in neoplastic cells, a specific characteristic of this type of tumor (41). This is critically correlated with fundamental clinical-pathological characteristics of tumor development (41). The synergistic effect of aspirin and sorafenib treatment induces an increase in GADD45 β in HCC, promoting apoptosis and control of tumor growth (66), an opposite effect of growth in the non-neoplastic liver (65), reflecting the opposing effects of this gene depending on the biological context. GADD45 β responses reflect its dynamics in rapid adaptations (65), being a possible diagnostic and selection biomarker for the treatment of HCC (66). These data suggest that the appearance of ITPR3 in liver cancer may promote tumor maintenance through the suppressive effect of GADD45 expression, especially of the β isoform.

The present study demonstrated the upregulation of important genes involved in the apoptotic pathway in liver cancer, such as GADD45 and CHOP. However, high-throughput technology would add stronger evidence. Thus, the absence of microarray experiments is a limitation of the present study.

In other types of neoplasms or liver diseases, ITPR3 can assume several roles. In prostate cancer, the mechanisms that promote the degradation of ITPR3 and prevent apoptosis have been elucidated (19). In ischemia-reperfusion injury, ITPR3 serves a protective role in which expression of the protein serves to change the mechanism of hepatocyte death from necrosis to apoptosis (15). This demonstrates the versatility that ITPR3 assumes within the varied contexts of different diseases and organs, but all associated with apoptotic events. It has been demonstrated that ITPR3 serves a role in proliferative stimulation and apoptosis block in liver cell lines and normal mouse liver (16). The current study provided evidence in human HCC samples that ITPR3 serves a role in hepatocarcinogenesis by modulating apoptosis. Additional work will be needed to better understand the mechanism by which *de novo* expression of ITPR3 correlates with mitosis and apoptosis in the pathogenesis of HCC, as well as whether this represents a potential target for therapy.

In conclusion, ITPR3 was highly expressed in HCC tumor cells relative to its expression level in the underlying CLD and healthy livers. ITPR3 expression was inversely correlated with apoptotic and mitotic indices in HCC, suggesting that ITPR3 contributed to the maintenance of HCC, promoting resistance

to apoptosis. This resistance to apoptosis and alteration of tumor cell survival occurred, at least in part, through an intracellular signaling pathway in which ITPR3 demonstrated a negative effect on the expression of CHOP and GADD45, providing a reduction in apoptotic cell death. The expression of ITPR3 in the liver may be a promising prognostic marker of HCC and other types of liver cancer.

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Availability of data and materials

The datasets analyzed during the current study are available in the NCBI Sequence Read Archive under the Accession code PRJNA758563 (<https://www.ncbi.nlm.nih.gov/sra/?term=PRJNA758563>). All data generated or analyzed during this study are included in this published article.

Authors' contributions

The entire project was conducted, written and analyzed by MLDS, including carrying out the experiments. AF contributed in all aspects, mainly in the analysis and interpretation of data. RMF contributed with the assistance in carrying out the experiments, analysis and discussion of data and with the preparation of graphics and images. ACMLF contributed with the assistance in carrying out the experiments, analysis and discussion of data and with the preparation of graphics and images. PHD contributed to the statistical analysis and in the organization of the survival data. CAXG and VLC elaborated all the bioinformatics data, analysis and RNA-seq results. PVTV performed the mitotic cell count on the HCC samples in the tumor mitotic index experiment. In addition, PVTV helped to analyse and interpret the data in conjunction with the literature already published and was involved in the writing of the manuscript and in the critical review of important intellectual content. PVTV also helped with the construction of the data discussion. MFL participated in carrying out the experiments and articulating the data obtained from the works previously published by the group. FOL was involved in the writing of the manuscript and critical review of important intellectual content, and also contributed to the analysis of some of the data presented. MHN contributed to the analysis and discussion of the data in this study, in addition to approving the final version submitted for publication. GF participated in the orientation and design of the study, particularly in the contribution of immunohistochemical staining image analysis. CXL participated in the acquisition of clinical data from patients and in the interpretation of these data in conjunction with the results. MLDS and PVTV confirm the authenticity of all the raw data. All authors reviewed and approved the final manuscript.

Ethics approval and consent to participate

The present study was approved by the Ethics Committee of the Hospital das Clínicas, Federal University of Minas Gerais (approval number CAAE 71206617.8.0000.5149) and written informed consent was obtained from the patients or their relatives, including for use of their tissue in the research.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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