Molecular characterization of autophagic and apoptotic signaling induced by sorafenib in liver cancer cells

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Sorafenib is the unique accepted molecular targeted drug for the treatment of patients in advanced stage of hepatocellular carcinoma. The current study evaluated cell signaling regulation of endoplasmic reticulum (ER) stress, c-Jun-N-terminal kinase (JNK), Akt, and 5′ AMP-activated protein kinase (AMPK) leading to autophagy and apoptosis induced by sorafenib. Sorafenib induced early (3–12 hr) ER stress characterized by an increase of Ser51P-eIF2α/eIF2α, C/EBP homologous protein (CHOP), IRE1α, and sXBP1, but a decrease of activating transcription factor 6 (ATF6) expression, overall temporally associated with the increase of Thr183,Tyr185P-JNK1/2, Thr172P-AMPKα, Ser413P-Foxo3a, Thr308P-Akt/Akt and Thr32P-Foxo3a/Foxo3a ratios, and reduction of Ser2481P-mammalian target of rapamycin (mTOR)/mTOR and protein translation. This pattern was related to a transient increase of tBid, BimEL, Beclin-1, Bcl-xL, Bcl-2, autophagy markers, and reduction of myeloid cell leukemia-1 (Mcl-1) expression. The progressive increase of CHOP expression, and reduction of Thr308P-Akt/Akt and Ser473P-Akt/Akt ratios were associated with the reduction of autophagic flux and an additional upregulation of BimEL expression and caspase-3 activity (24 hr). Small interfering-RNA (si-RNA) assays showed that Bim, but not Bak and Bax, was involved in the induction of caspase-3 in sorafenib-treated HepG2 cells.

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1 | INTRODUCTION

Hepatocellular carcinoma (HCC) is the most common type of liver cancer and the second most frequent cause of cancer-related death worldwide (Llovet et al., 2016; Torre et al., 2015). HCC develops within a cirrhotic liver in 80%-90% of cases that influences therapeutic decisions (Fattovich, Stroffolini, Zagni, & Donato, 2004; Sanyal, Yoon, & Lencioni, 2010). The difficulty in forming an early diagnosis, at the first stage of the disease, contributes to the poor response of patients to current clinical treatments. Currently, only one-third of newly diagnosed patients with HCC are an at early stage (0 or A), according to the Barcelona Clinic Liver Cancer (BCLC) staging system, and are eligible for potential curative therapies such as local ablation, resection, or orthotopic liver transplantation with a 5-year survival rate of 50%-80% (Bruix, Reig, & Sherman, 2016). The recommended first-line treatment for patients with locally advanced/metastatic disease (BLCL C) is sorafenib with a mean overall survival of 11 months (Cheng et al., 2009; Llovet et al., 2008).

Sorafenib inhibits tyrosine kinase receptors such as vascular endothelial growth factor receptor (VEGFR) 2, VEGFR 3, platelet-derived growth factor receptor-β (PDGFR-β), Flt3 and c-KIT, as well as molecular components of the Raf/MEK/extracellular signal-regulated kinase (ERK) signaling pathway (Cervello et al., 2012). Different studies have suggested that the increased susceptibility to cell death by sorafenib is associated with down-regulation of cell survival pathways in hepatoma cells (Prieto-Dominguez et al., 2016). In particular, Chen et al. (2010) showed that sorafenib enhanced TRAIL-induced cell death through SH2 domain-containing tyrosine phosphatase (SHIP-1)-dependent reduction of signal transducers and activators of transcription type 3 (STAT3) phosphorylation (Ty705STAT3) and related proteins members myeloid cell leukemia-1 (Mcl-1), survivin, and cyclin D1 in hepatoma cells. Sorafenib-induced cell death is also related to upregulation of the proapoptotic "BH3-domain only" PUMA and BIM, as well as a decrease of Mcl-1 and survivin (Fernando et al., 2012). It has also been shown that c-Jun-N-terminal kinase (JNK)-dependent GADD45β (Ou et al., 2010) and PUMA (Fernando et al., 2012) expressions are also involved in the proapoptotic properties of sorafenib in HCC cells. Shimizu et al. (2012) showed that sorafenib enhanced autophagy, pharmacological inhibition due to which increased apoptosis and decreased cell viability in sorafenib-treated Huh7, HLF, and PLC/PRF/5 cells. However, other studies did not find a relationship between autophagy induction and resistance to treatment in sorafenib-treated HCC cells (Fischer, Wang, Vlada, Kim, & Behrns, 2014). The current study showed that the induction of autophagy in sorafenib-treated HepG2 cells is related to IRE-1 and eIF2α-dependent ER stress, JNK and 5’ AMP-activated protein kinase (AMPK)-dependent signaling. The shift from autophagy to apoptosis is related to a decline of Beclin-1, Bcl-2, Mcl-1, Bcl-xL, Bcl-2, and tBid and the increase of proapoptotic BimEL, Bak, and Bax.

2 | MATERIAL AND METHODS

2.1 | Chemical and reagents

Sorafenib (Ref FS10808; Carbosynth Limited, Berkshire, UK) was dissolved in dimethyl sulfoxide as stock solution (100 mM). 3-MethylAdenine (3MA; Ref M9281), chloroquine (CQ; Ref C6628), SP600125 (Ref S5567), and dorsomorphin (Ref P5499) were obtained from Sigma-Aldrich (Sant Louis, MO).

2.2 | Primary human hepatocytes, cell lines, and culture conditions

Human hepatocytes were prepared from liver biopsies obtained from three patients (two females and one male, aged 66 ± 3.0 years) submitted to surgical resection for liver tumors after obtaining the patients’ written consent. Hepatocytes isolation was based on a two-step collagenase procedure (Pichard et al., 2006). HepG2 and Hep3B cell lines were obtained from the American Type Culture Collection (LGC Standards, S.L., Barcelona, Spain). Huh7 cell line was purchased from Apath, LLC (Brooklyn, NY). Cell lines were selected according to their p53 content: HepG2 (WT p53 expression), Huh7 (p53 mutated isofrom, codon 220), and Hep3b (nonsense p53 mutation). All lines were negative for mycoplasma contamination. The cells were cultured in minimum essential medium (MEM) with Earle’s salts with L-glutamine (Ref E15–825, PAA) with 10% fetal bovine serum (FBS; F7524, Sigma-Aldrich, Lot No: 085686).
022M3395, endotoxin < 0.2 EU/ml), sodium pyruvate (1 mM; Ref S11-003, PAA), nonessential amino acids (Ref M11-003, PAA), penicillin-streptomycin solution (100 U/ml-100 µg/ml; P11-010, PAA), at 37°C in a humidified incubator with 5% CO₂. The cells were cultured at a cell density of 100,000 cells/cm², treatment added 24 after plating, and lysates were obtained at different time points after treatments. Sorafenib was administrated at different concentrations (1 nM–100 µM). Additional experimental interventions, such as 3MA (5 µM) and CQ (50 µM), were preadministered (2 hr), and dorsomorphin (5 µM) was coadministered with sorafenib. The Biobank of the Institution approved the use of primary biopsies of patients for the present research (Ref. no 33140031, April 1, 2014).

2.3 | Effect of sorafenib administration on tumor growth in a xenograft mouse model

Sixty male Hsd:Athymic Nude-Foxn1nu mice (Harlan Laboratories, Barcelona, Spain), aged 4 weeks (weighing 18–22 g), were randomly distributed in three groups according to the subcutaneous implantation of HepG2, Hep3b, and Huh7. Cells (10 × 10⁶) were diluted in matrigel (Ref 356231, Lot 36821, Becton Dickinson, NJ) and administered subcutaneously into the right flank of the mice. Each experimental group was subdivided into the group receiving sorafenib (200 mg/kg) and the group receiving a solvent, which was constituted by ethanol:Koliphor EL (C5135, Sigma) receiving a solvent, which was constituted by ethanol:Koliphor EL 1:1 (v/v). Sorafenib or solvent was administered daily by oral gavage when the tumor diameter grew to 5 mm (11 days) and ended at the sacrifice of animals when tumor diameter was 15 mm (24 days). The tumor volume was calculated using the formula: 
\[ V = \frac{\text{length} \times \text{width} \times \text{depth}}{6} \]
All animal care and experimentation conformed to the Guide for the Care and Use of Laboratory Animals published by the National Academy of Sciences. The Ethical Committee of the Institution approved the in vivo experiments (Ref. no CEP2104 PI/016, January 30, 2014).

2.4 | Immunohistochemistry

Tumor sections (5 µm) were obtained to analyze the expression of α-smooth muscle actin (α-sm), Ki67, CD31, and CD34 by immunohistochemistry. Sections were deparaffinized, hydrated through graded ethanol steps, briefly rinsed in water, and blocked at room temperature using TBSA-BSAT (10 mM Tris, 0.9% NaCl, 0.02% sodium azide, 2% bovine serum albumin and 0.1% Triton-X100). Slices were incubated overnight at room temperature with primary antibodies against: 1:250 α-sm (ab5694; Abcam, Cambridge, UK), 1:500 (Ki67, IR626; DAKO, Glostrup, Denmark), 1:50 CD31 (ab28364; Abcam), and 1:200 CD34 (ab64480; Abcam), followed by incubation with the corresponding secondary antibodies, either Alexa 488 anti-rabbit IgG (A11008; Invitrogen, Carlsbad, CA, Spain) or Alexa 488 anti-mouse IgG (A11001; Invitrogen), for 5 hr diluted in TBSA-BSAT (1:500). Nuclear staining was performed using DRAQ-5th (red fluorescent cell-permeable DNA probe; Biostatus Limited, Loughborough, UK). Immunofluorescence analysis was performed using an Olympus BX61 microscope (Tokyo, Japan). Fluorescence quantification was performed using Leica Application Suite Advanced Fluorescence software and ImageJ software (Wetzlar, Germany).

2.5 | Ultrastructural analysis by transmission electronic microscopy

Treated and control cells were isolated and prefixed in 1.5% formaldehyde in 1% cacodylate buffer, pH 7.4, for 2 hr at 4°C. After three washes in cacodylate buffer, the extracts were fixed in 1% osmium tetroxide for 60 min at 0–4°C. The samples were dehydrated in graded ethanol and embedded in Epon resin. After overnight incubation at 65°C, ultrathin sections (70 nm) were cut with a diamond knife using an Ultracut S ultramicrotome and placed on 200-mesh copper grids. All sections were stained with uranyl acetate, counterstained with lead citrate, and viewed using a Carl Zeiss (Oberkochen, Germany) EM10C electron microscope at 4,646×, 10,000×, and 46,460× magnifications in the Biomedical Research Center at the University of Granada’s Health Technology Park.

2.6 | Measurement of cell death

Caspase-3-associated activities were determined using Caspase-Glo® 3 Assay Systems (G8091; Promega, Madison). Cells were treated with Caspase-Glo® 3 in an "add-mix-measure" format resulting in cell lysis, caspase-3-dependent cleavage of the substrate and generation of a "glow-type" luminescent signal. The signal generated was proportional to the amount of caspase activity. The values are extrapolated into a calibration curve included in the assay. Cell death in tissue sections was determined using the DeadEnd™ fluorometric terminal deoxynucleotidyl transferase mediated dUTP nick-end labeling (TUNEL) System (G3250, Promega) that measures the fragmented DNA of apoptotic cells by catalytically incorporating fluorescein-12-dUTP at 3'-OH DNA ends using the enzyme terminal deoxynucleotidyl transferase (TdT), which forms a polymeric tail using the principle of the TUNEL assay. The fluorescein-12-dUTP-labeled DNA was visualized using an Olympus BX61 microscope. Fluorescence quantification was performed using Leica Application Suite Advanced Fluorescence software and the ImageJ software.

2.7 | Cell proliferation

The measurement of bromodeoxyuridine (BrdU) incorporation was used as marker of cell proliferation (Ref. 1164729001; Roche Diagnostics, Indianapolis, IN). Cells were seeded in 96-well plates at low density (12,500 cells/cm²). Two hours before sampling, 20 µl of BrdU (10 µM) was added to the cultures. DNA was denaturalized with 200 µl FixDenat solution included in the commercial assay for 30 min at room temperature. After removal, the cells were incubated with 100 µl monoclonal anti-BrdU antibody HRP conjugated for 90 min at room temperature. Afterwards, the cells were washed with PBS and incubated with 100 µl reveling solution including hydrogen peroxide, luminol and...
4-iodophenol for 15 min at room temperature. Absorbance was measured at 370 nm using an Infinite 200 PRO Microplate Reader (TECAN, Männedorf, Switzerland).

2.8 Assessment of protein expression of ER stress, autophagy, and apoptosis markers

Cells or tumor tissues were treated with lysis solution including 50 mM HEPES pH 7.5, 5 mM ethylenediaminetetraacetic acid (EDTA), 150 mM NaCl, 1% NP-40, a commercial proteases inhibitor cocktail (P8340, Sigma-Aldrich), 1 mM phenylmethanesulfonyl fluoride (PMSF), 1 mM NaF, and 1 mM Na₂VO₄. Protein expression were determined by SDS-PAGE coupled to western blot analysis. Proteins (50–100 µg) were separated by Any kD™ Criterion™ TGX Stain-Free™ Protein Gel, 18 well, 30 µl (#5678124; BioRad, Hercules, CA) and transferred to polyvinylidene difluoride (PVDF) membranes. The system uses stain-free technology, which is a method that appears to be more reliable as a protein loading control than measurement of housekeeping proteins (Colella et al., 2012). The membranes were incubated with the corresponding commercial primary and secondary antibodies coupled to horseradish peroxidase revealing protein content by Clarity™ Western ECL substrate (Ref 170-5061; BioRad) and analyzed in a ChemiDoc™ Touch Imaging System. Antibodies for western-blot, tBid (#2002), elf2α (#5324), inositol-requiring enzyme 1α (IRE1α, #3294), p62 (#5114), Ser51P-elf2α (#9721), Ser2481P-幅 phosphorylated ribosomes in 5 min (mTOR) (#2974), AMPK (#5832), Tβr177P-AMPK (#2525), Bcl-xL (#27625), Bcl-2 (#2872), Bax (#5023), Bak (#12105), Bin (#2933), FoxO3a (#2497), Ser413P-FoxO3α (#8174), Thr32P-FoxO3a (#9464), and cleaved PARP (#9542) were commercially obtained from Cell Signaling Technology (Danvers, MA); activating transcription factor 6 (ATF6, ab122897) was purchased from Abcam; LC3 (PM036) was purchased from MBL International (Woburn, MA); Beclin-1 (sc-11427), GADD153/CHOP (sc-575) and Akt (sc-1618) were obtained from Santa Cruz Biotechnology Inc. (Texas); Tβr303P-AKT (SAB4504740), Ser473P-AKT (SAB430042), JNK1/2 (SAB420176), Thr183/Thr186P-JNK1/2 (J644) and Mcl-1 (SAB4501843) were obtained from Sigma-Aldrich. The corresponding secondary antibodies anti-rabbit (sc-2004), anti-mouse (sc-2005) and anti-goat (sc-2768) IgG-HRP labeled were purchased from Santa Cruz. Densitometric analysis was performed using the software Image Lab 6.0 of BioRad.

2.9 XBP1s/XBP1 mRNA expression ratio by qRT-PCR

RNA was isolated using TRIzol reagent (BIO-38033; Bioline Reagents Ltd., London, UK), and submitted to retrotranscription (iScript™ cDNA Synthesis Kit #1708891; BioRad) and amplification by real-time quantitative reverse transcription polymerase chain reaction (RT-qPCR) (iTaq™ Universal SYBR® Green Supermix #1725142; BioRad). RT-PCR was performed in Viia™ 7 Real-Time PCR System (Applied Biosystems, Thermo Fisher Scientific, Waltham, MA) with a QuantumStudio™ Real Time PCR Software. Reactions were performed in 96-well plates with an optical sealing tape (Thermo Fisher Scientific) in 25 µl total volume containing SyberGreen Mix and the corresponding cDNA. Histone 3 (H3B3) was used as an internal control in parallel for each run. The conditions for amplification were as follows: polymerase activation and a DNA denaturation step of 95°C for 40 s; 30 cycles of amplification composed of a denaturation step of 95°C for 20 s and annealing/extension;plate read at the optimized temperature of each couple of primers (54–60°C) for 60 s; and a final melt curve analysis of 95°C 15 s, 60°C 60 s and 95°C 15 s. All primers were designed using online Primer3 software Primer3input and purchased from Eurofins (Eurofins Genomics):

<table>
<thead>
<tr>
<th>Primer</th>
<th>Sequence 5’-3’</th>
<th>Tm, °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>XBP1s Fw</td>
<td>CGG GTT TGC TGA TGC CGC AGC AG</td>
<td>69.6</td>
</tr>
<tr>
<td>XBP1 Fw</td>
<td>GCA GGT GCA GGC CCA GTT GTC AC</td>
<td>67.8</td>
</tr>
<tr>
<td>XBP1s Rv</td>
<td>CCC CAC TGA CAG AGA AAG GGA G</td>
<td>66</td>
</tr>
<tr>
<td>XBP1 Rv</td>
<td>CCC CAC TGA CAG AGA AAG GGA G</td>
<td>66</td>
</tr>
</tbody>
</table>

2.10 si-RNA interference

HepG2 cells were seeded in 6-well plates the day before transfection to reach a confluency of 60% in 24 hr. Transfection was conducted with the siTran 1.0 Transfection Reagent (siTranTM siRNA transfection; Origene, Rockville, MD) according to recommendations of the manufacturer. The si-RNA of IRE1α, pancreatic ER kinase (PKR)-like ER kinase (PERK), and ATF6 were synthesized by Origene. Their targeting sequences can be accessed from the Origene webpage with the references SR301457 (IRE1α, NM_001433), SR306267 (PERK, NM_004836), and SR307883 (ATF6, NM_007348). A nontargeting si-RNA (SR30004, Triencer-27 Universal Scrambled Negative Control siRNA Duplex) also purchased from Origene was used as negative control. Soralenib was added 48 hr after transfection.

si-RNA for knock-down Bax (L-003308-01-0005), Bak (L-003305-00-0005), Bim (L-004835-00-0005), FoxO3a (L-003007-00-0005), and scramble (D-001810-10-05) were obtained from Dharmacon (Lafayette, CO) and used in HepG2 cells cultured at 20,000 cells/cm². si-RNA (25 nM) were mixed with the transfection reagent DharmaFECT1 (Ref T-001-02) for 20 min at room temperature and transferred to culture medium without antibiotic/antimycotic solution and kept for 24 hr prior to treatment. Afterwards, transfected cells were maintained until the end of the experiment following the manufacturer’s recommendations.

2.11 pDest-LC3-GFP-mCherry transient transfection and detection of punctuate LC3-positive structures

Expression of pH-sensitive sensor constructed as a tandem fusion of mCherry and GFP to LC3B (pDest-LC3-GFP-mCherry) was used to monitor autophagy in HepG2 cells (kindly provided by Dr. Terje Johansen). The GFP tag is acid-sensitive, whereas the mCherry tag is acid-insensitive. Yellow merge punctate structures indicate autophagosomes which neutral pH allows both tags emit
fluorescent light resulting in a yellow staining. Subsequent fusion of autophagosomes with lysosomes results in red-positive acidic vacuoles, where the green fluorescence from GFP is lost. HepG2 cells were seeded in 24-well plates with coverslips at 2 × 10^5 cells/ml, and 500 µl of antibiotic free medium per well. After 24 hr, the cells were transiently transfected with pDest-LC3-GFP-mCherry construct with Lipofectamine 2000 (Invitrogen), according to the manufacturer’s protocol (2 µl of Lipofectamine and 3 µg of plasmid per well). The following day, the cells were treated with Earle’s balanced salt solution (Sigma, E2888) and CQ (50 µM) as negative and positive control, respectively. Sorafenib was administered at 10 µM and 100 µM. All these treatments lasted for 6, 12, or 24 hr. Once the treatments were finished, the medium was recovered, and the cells were washed with phosphate saline buffer (PBS1X). Nuclei were stained with Hoechst 33342 and immediately fixed by using a 4% (w/v) paraformaldehyde solution in PBS1X for 30 min. The cells were transferred to the second dish. After scraping the cells corresponding to the second dish, the whole juice was transferred to a 1.5 ml Eppendorf tube. The tubes were incubated at 4°C with gentle end-over-end rotation for 10 min and then centrifuged at 16,000 g for 8 min at 4°C in a refrigerated microfuge. The corresponding supernatants were recovered, and the A260 was measured using a NanoDrop ND-1000 Spectrophotometer (Thermo Fisher Scientific). About 10 absorption units of A260 were layered on top of 7%–50% (w/v) sucrose gradients prepared in 50 mM Tris-acetate, pH 7.5, 50 mM NaCl, 12 mM MgCl2, 200 µg/ml cycloheximide, 200 µg/ml heparin, 2 mM dithiothreitol (DTT), 0.5% NP40) was added to one dish, cells were scraped, and transferred to the second dish. After scraping the cells corresponding to the second dish, the whole juice was transferred to a 1.5 ml- Eppendorf tube. The tubes were incubated at 4°C with gentle end-over-end rotation for 10 min and then centrifuged at 16,000 g for 8 min at 4°C in a refrigerated microfuge. The corresponding supernatants were recovered, and the A260 was measured using a NanoDrop ND-1000 Spectrophotometer (Thermo Fisher Scientific). About 10 absorption units of A260 were layered on top of 7%–50% (w/v) sucrose gradients prepared in 50 mM Tris-acetate, pH 7.5, 50 mM NaCl, 12 mM MgCl2, 1 mM DTT. The gradients were centrifuged at 260,110 g (39,000 rpm) in a Beckman SW41 rotor at 4°C for 2 hr 45 min. To dissociate the vacant 80S ribosomes, high-salt conditions were achieved by adding NaCl to a final concentration of 0.25 M in the gradients. Gradient analysis was performed with an ISCO UA-6 system (Isco, Inc. Lincoln, NE) equipped to continuously monitor the A260.  

### 2.12 | Assessment of protein translation

The protocol adapted for polysome preparation of HepG2 cells was the one we routinely employ for yeast cells (Kressler, de la Cruz, Rojo, & Linder, 1997). Briefly, HepG2 cells were grown to 80% confluency in 175 cm² dishes as described above. Normally, two dishes were used per condition assayed. Before harvesting the cells, 200 µg/ml cycloheximide was added and incubated for 5 min at 37°C. Each dish was then placed on ice, the medium was collected, and the cultures washed twice with PBS without Ca²⁺ and Mg²⁺ containing 200 µg/ml cycloheximide. Then, 600 µl of lysis buffer (10 mM Tris–HCl, pH 7.4, 150 mM NaCl, 10 mM MgCl2, 200 µg/ml cycloheximide, 200 µg/ml heparin, 2 mM dithiothreitol (DTT), 0.5% NP40) was added to one dish, cells were scraped, and transferred to the second dish. After scraping the cells corresponding to the second dish, the whole juice was transferred to a 1.5 ml-Eppendorf tube. The tubes were incubated at 4°C with gentle end-over-end rotation for 10 min and then centrifuged at 16,000 g for 8 min at 4°C in a refrigerated microfuge. The corresponding supernatants were recovered, and the A260 was measured using a NanoDrop ND-1000 Spectrophotometer (Thermo Fisher Scientific). About 10 absorption units of A260 were layered on top of 7%–50% (w/v) sucrose gradients prepared in 50 mM Tris-acetate, pH 7.5, 50 mM NaCl, 12 mM MgCl2, 1 mM DTT. The gradients were centrifuged at 260,110 g (39,000 rpm) in a Beckman SW41 rotor at 4°C for 2 hr 45 min. To dissociate the vacant 80S ribosomes, high-salt conditions were achieved by adding NaCl to a final concentration of 0.25 M in the gradients. Gradient analysis was performed with an ISCO UA-6 system (Isco, Inc. Lincoln, NE) equipped to continuously monitor the A254.

### 2.13 | Expression of Bim, Bak, and Bax in Beclin-1 immunoprecipitated fraction

A volume of the cell lysate (500 µg protein) was mixed with immunoprecipitation buffer (20 mM Tris–HCl pH 7.5, 1% Triton X-100, 150 mM NaCl, 10% glycerol, 1 mM Na2VO3, 50 mM NaF, 2 mM EDTA, 1 mM PMSF, commercial proteases inhibitor cocktail (P8340, Sigma-Aldrich), and 2 µg of anti-Beclin-1 antibodies (sc-48341, Santa Cruz Biotechnology Inc.) at 4°C overnight. Afterward, protein G Sepharose (35 ll; GE Healthcare) was added, agitated for 3 hr, and the washed resin was loaded onto a 10% SDS-PAGE electrophoresis gel. The expression of Bim, Bak, and Bax was assessed by western-blot analysis as described above. The immunoprecipitation using unspecific mouse IgG (sc-2025; Santa Cruz Biotechnology) and anti-Beclin-1 antibodies were run in parallel. The expression of Bim, Bak, and Bax has also been measured in cell lysate (50 µg proteins).

### 2.14 | Statistical analysis

All results are expressed as the mean ± standard error of the mean of independent experiments (n = 4–10). Data were compared using the analysis of variance with the least significant difference’s test as post-hoc multiple comparison analysis (homogeneity of variances) or Games–Howell analysis (nonhomogeneity of variances). If Shapiro–Wilk’s test showed nonnormal distribution of data, nonparametric Kruskal–Wallis coupled to U Man–Whitney posthoc analysis with Finner’s correction was done. The level of significance was set at *p* ≤ 0.05, **p** ≤ 0.01, ***p** ≤ 0.001. The groups significantly different (p ≤ 0.05) were indicated with different letters.

### 3 | RESULTS

#### 3.1 | Sorafenib administration reduced tumor cell growth in xenograft mouse model

The administration of sorafenib tosylate (30–100 mg/kg) orally once daily reduced growth and angiogenesis while promoting apoptosis in tumors developed by PLC/PRF/5 (Liu et al., 2006) and patient-derived cell lines (Huynh et al., 2009) subcutaneously implanted in severe combined immunodeficient mice. Sorafenib tosylate (200 mg/kg) was administered to nude mice once daily when the diameter of the tumors reached 5 mm. The weight of animals did not change with the treatment (Supporting Information Figure S1a). The administration of sorafenib reduced tumor volume (Figure 1a and Supporting Information Figure S1b) in parallel to an increase of caspase-3 activity (Figure 1b), cleaved PARP (Figure 1c) and autophagic markers (Beclin-1 and LC3II/LC3I; Figure 1d) in tissue homogenate from tumors derived after implantation of HepG2 cells in nude mice. The expression of p62 in tumors was not reduced in tumors from sorafenib-treated
animals (Figure 1d). Sorafenib also reduced the volume of tumors derived after implantation of Huh7 and Hep3B in nude mice (Supporting Information Figure S1b). Sorafenib increased apoptosis (TUNEL staining; Supporting Information Figure S1c) and reduced cell proliferation (Ki67; Supporting Information Figure S1d), fibrogenesis (α-SMA; Supporting Information Figure S1e), angiogenesis (CD31; Supplementary Figure S1f) and stemness (CD34; Supporting Information Figure S1g).

3.2 Sorafenib reduced cell proliferation and increased apoptosis

Primary human hepatocytes were resistant to the proapoptotic properties of sorafenib (Figure 2a). However, the administration of sorafenib (10 µM) increased caspase-3 in the three HCC cell lines tested at 24 hr, with HepG2 being more susceptible than p53-deficient HCC cell lines (Hep3B and Huh7) to the treatment (Figure 2a). Differently, sorafenib moderately reduced cell proliferation in primary hepatocytes, whereas it drastically blocked cell proliferation in HCC cell lines with a half maximal inhibitory concentration (IC50) close to 10 µM (Figure 2b; 24 hr).

3.3 Induction of endoplasmic reticulum stress by sorafenib

The ER is responsible for the regulation of protein folding and cellular calcium concentrations through three ER transmembrane proteins: PERK, ATF6, and IRE1 (Fribley, Zhang, & Kaufman, 2009; Szegedi, Logue, Gorman, & Samali, 2006). Shi et al. (2011) have reported that sorafenib (20 µM) induces ER stress independently of MEK/ERK
through PERK and IRE1 pathways that are related to increased apoptosis and autophagy markers respectively in HepG2 and PLC/PRF/5. In agreement with these studies, our data showed that sorafenib (≥10 µM) not only increased the ratio of Ser51P-eIF2α/eIF2α protein expression (Figure 3a) and the protein expression of the C/EBP homologous protein (GADD153 or CHOP; Figure 3b) at 6 hr, but also that of IRE1α (Figure 3c), and the mRNA expression of XBP1s/XBP1 ratio (Supporting Information Figure S2) at 12 hr. Interestingly, we have observed that sorafenib reduced ATF6 protein expression (Figure 3d) at the same range of concentration and time period as increased PERK and IRE1 pathways (Figure 3a and 3c, respectively).

3.4 | Sorafenib decreased Ser2481P-mTOR phosphorylation, impaired initiation of protein synthesis and increased transient autophagy

The induction of apoptosis by sorafenib has been related to an alteration of protein translation characterized by a reduction of eIF4E phosphorylation and Mcl-1 translation, but unassociated with MEK/ERK down-regulation in HCC cells (Liu et al., 2006). The mTOR integrates nutrient and mitogen signals to regulate cell growth and cell proliferation. Different pathways have been shown to regulate mTOR signaling by antitumoral agents. The ATF4-mediated increase in the expression of sestrin-2 caused by nelfinavir- and bortezomib-induced AMPK activity

![Figure 3](https://example.com/figure3.png)

**FIGURE 3** Expression of markers from all three branches of ER stress altered by sorafenib in HepG2 cells. A kinetic analysis of the protein expression of Ser51P-eIF2α/ eIF2α (a), CHOP (b), IRE1α (c) and ATF6 (d) were determined at different sorafenib concentrations (0, 10 nM, 10 µM, and 100 µM) by western-blot analysis as described in Material and Methods. Results are expressed as mean ± standard error of the mean, and with the blots are representative of five independent experiments. *p ≤ 0.05, **p ≤ 0.01, and ***p ≤ 0.001 between control and sorafenib treated cells. The groups with different letters (a, b, c, d, or e) were significantly different (p ≤ 0.05). ATF6, activating transcription factor 6; CHOP, C/EBP homologous protein; ER, endoplasmic reticulum.
inhibits mTOR and triggers autophagy in breast, ovarian, and cervical cancer cell lines (Brüning, Rahmeh, & Friese, 2013). Sorafenib promoted an early perturbation of the mitochondrial function, ATP depletion and AMPK-dependent inhibition of the mTORC1 pathway in breast cancer cells (Fumarola et al., 2013). In addition, the sustained activation of phosphatidylinositol 3-Kinase (PI3K)/AKT upregulating mTOR has also been related to sorafenib resistance in HCC cells (Chen et al., 2011). In the current study, we observed that sorafenib (10 µM) caused an early and progressive reduction (3–24 hr) of Ser2481P-mTOR phosphorylation in HepG2 (Figure 4a), as well as in Hep3B and Huh7 cells (Supporting Information Figure S3). Sorafenib (10 µM) induced an early upregulation of Thr308P-AKT phosphorylation (6–12 hr) followed by downregulation of Thr308P-Akt and Ser473P-Akt phosphorylation (24–48 hr; Figure 4b). Sorafenib also induced a rapid and transient increase of Thr172P-AMPK phosphorylation (6–12 hr; Figure 4c). Data suggest that mTOR inhibition may be related to the combined signaling of AKT inhibition and AMPK activation.

The inhibition of mTOR by sorafenib has a great impact on protein translation. The polysome profiles in control cells were obtained by observing free 40S and 60S ribosomal subunits, 80S ribosomes, and actively translating polysomes. However, sorafenib (10 µM) resulted in abnormal polysome profiles, with a dramatic increase of the 80S peak and a reduction of polysomes (Figure 4d). Salt treatment dissociated most of the 80S peak into free 40S and 60S ribosomal subunits, indicating that the large 80S ribosome population was mostly nontranslating vacant ribosomes rather than monosomes engaged with mRNA (Supporting Information Figure S4).

The nutrient deprivation, growth factor withdrawal or hypoxia are related to autophagy induction (He & Klionsky, 2009). The most important pathways involved in autophagy modulation are mTORC1 and AMPK (Prieto-Dominguez et al., 2016). We used the double fluorescent-tagged LC3 (pDest-LC3-GFP-mCherry) plasmid to distinguish autophagosomes from autolysosomes, transmission electron microscopy (TEM), and biochemical markers to follow

**FIGURE 4** Kinetic study of the phosphorylation of mTOR (a), AKT (b), AMPK (c), as well as the alteration in the polysome profile (d) in HepG2 cells treated with sorafenib (10 µM). The protein expression was determined by western-blot. Polysome profiles were generated following the procedure described in Material and Methods after 12 hr of sorafenib (10 µM) treatment. Results are expressed as mean ± standard error of the mean, and with the blots are representative of four independent experiments. *p ≤ 0.05, **p ≤ 0.01 and ***p ≤ 0.001 between control and sorafenib-treated cells. The groups with different letters (a, b, c, or d) were significantly different (p ≤ 0.05). AMPK, 5′AMP-activated protein kinase; mTOR, mammalian target of rapamycin.
the induction and progression of autophagy flux. We found that sorafenib induced autophagolysosome formation in mRFP-EGFP-LC3-transfected cells (Figure 5a), a fact confirmed by TEM imaging (Figure 5b). The quantification of the phagolysosomes (black line) and phagosomes (grey line; Supporting Information Figure S5) and biochemical markers (p62, Beclin-1, and LC3II/I) in biohecha markers (p62, Beclin-1, and LC3II/I) in HepG2 (Figure 5c) showed that autophagy induced by sorafenib (10 µM) was only transiently active (6–12 hr).

3.5 | The inhibition of the ER stress and AMPK downregulated autophagy and increased apoptosis in sorafenib-treated HepG2 cells

We have evaluated the specific role of ER stress arms and AMPK on autophagy and apoptosis using si-RNA technology and dorsomorphin, respectively. Data showed that IRE1α and PERK were deeply involved in sorafenib-induced autophagy, whereas a minor effect of ATF6 was also observed (6 hr; Figure 6a). Thus, the reduction of autophagy by si-IRE1α and si-PERK increased sorafenib (10 µM)-induced caspase-3 activity (Figure 6b) and PARP cleavage (Figure 6c) in HepG2.

Moreover, the AMPK inhibition by dorsomorphin reduced Thr172P-AMPK/AMPK, Ser413P-Foxo3a/Foxo3a, and Beclin-1 expression induced by sorafenib (Figure 6d; 12 hr). However, the sustained reduction of the activating phosphorylation of Thr172P-AMPK/AMPK by dorsomorphin was related to a drastic upregulation of Beclin-1 expression (Figure 6d), and downregulation of caspase-3 activity (Figure 6e) and PARP cleavage (Figure 6f) at 24 hr in sorafenib-treated HepG2 cells. In addition, the antiapoptotic properties of autophagy were further confirmed using autophagic inhibitors 3MA and CQ by the upregulation of sorafenib (10 µM)-induced caspase-3 activity (Figure 6g) and PARP cleavage (Figure 6h) in HepG2 (24 hr).

3.6 | Sorafenib regulated the activation of JNK and Foxo3a, and the expression of proapoptotic, and antiapoptotic Bcl-2 family members

The Bcl-2 family is a well characterized protein family including antiapoptotic members (Bcl-2, Mcl-1, and Bcl-xL), proapoptotic effectors (Bax and Bak) and proapoptotic BH3-only members, which include the BH3-only direct activators Bim, Bid, and Puma (Kim et al., 2006) and the sensitizers/repressors such as Noxa, Bad, Bik, Bmf, and HRK (Willis et al., 2007). The interactions among them, as well as with the autophagic component Beclin-1, result in a potential switch between autophagic and apoptotic cell death pathways. The induction of apoptosis by sorafenib has been related to activation of Bax and Bak, and increased expression of proapoptotic "BH3-domain only" PUMA and BIM, as well as decreased protein levels of the antiapoptotic Mcl-1 (Fernando et al., 2012). The expression of Bim, Bak, Bax, tbid, Bcl-2, Bcl-xL, Beclin-1, and Mcl-1 in sorafenib (10 nM and 10 µM)-treated HepG2 cells is shown in Figure 7a and Supporting Information S6a and S6b. The line chart in Figure 7b shows the profile throughout time in the expression of mentioned proteins. Our data suggest that sorafenib (10 µM) induced a rapid drop of Mcl-1 protein expression, as well as transient and moderate increase of tbid, BimIL, Bcl-2 and Bcl-xL (3–6 hr) overlapping with the increase of Beclin-1 (3–12 hr). The expression of tbid drastically dropped below the control value at 6 hr, and Bcl-2, Bcl-xL and Beclin-1 expressions were reduced at 12 hr after sorafenib treatment (Figure 7b). Among the proapoptotic members, only BimIL showed a sustained increase throughout the kinetic study, reaching the highest value at 24 hr (Figure 7). Bak also showed a moderate rise throughout the study, while Bax expression increased transiently (3–6 hr) returning expression to the control level (Figure 7).

The activation of JNK has been related either to ER-related autophagy (Ogata et al., 2006) or apoptosis (Dhanasekaran & Reddy, 2008) according to different experimental conditions. The stimulation of ER-induced autophagy by sorafenib showed above was associated with the transient activation of JNK as measured by Thr183, Tyr185P-JNK1/2/JNK1/2 ratio (3–12 hr; Figure 7c and 7d). The inhibition of the Thr183, Tyr185P-JNK1/2/JNK1/2 ratio was assessed using the specific inhibitor SP600125 (Figure 7d). The treatment reduced cell proliferation in control and sorafenib-treated HepG2 cells (33% and 41% at 12 hr, and 53% and 22% at 24 hr, respectively; Figure 7e). In contrast, SP600125 increased caspase-3 in control and sorafenib-treated HepG2 cells (Figure 7f).

The activation of Foxo and its proapoptotic properties is under regulation of multiple cell signaling (Farhan et al., 2017). The inhibitory phosphorylation Thr32P-Foxo3a was transiently upregulated at 6–12 hr, while dropped afterwards in sorafenib (10 µM)-treated HepG2 cells (Figure 7g).

The role of Bim, Bak, and Bax has also been assessed using si-RNA technology. The reduction of Bim, but not Bak and Bax, by si-RNA recovered caspase-3 control levels in sorafenib (10 µM)-treated HepG2 (Figure 8a; 24 hr). The reduction of apoptosis by si-RNA Bim was associated with upregulation of p62 expression and LC3II/LC3I ratio, but without alteration of Beclin-1 expression in sorafenib-treated HepG2 cells (Figure 8b). The immunoprecipitation experiments showed that Bim, but not Bak and Bax, was able to bind Beclin-1 (Figure 8c).

4 | DISCUSSION

Sorafenib is, to date, the only approved first-line systemic therapy for patients in advanced stages of HCC. This chemotherapeutic drug has demonstrated a moderate but significant increase of mean overall survival in patients with good liver function based on the results in SHARP and Asia-Pacific clinical trials (7.9 to 10.7 months in Llovet et al., 2008; and from 4.2 to 6.5 months in Cheng et al., 2009; respectively). Beneficial effects have also been reported in patients with thyroid cancer, acute myeloid leukemia, advanced renal cell carcinoma, or prostate cancer (Antar, Kharfan-Dabaja, Mahfouz, & Bazarbachi, 2015; Chi et al., 2008; Escudier et al., 2007; Luo et al., 2014). Although sorafenib has been effective in extending the overall
FIGURE 5 Evaluation of autophagic flux (a), presence of autophagolysosomes by electronic transmission microscopy (b) and expression of autophagy markers p62, Beclin-1, and LC3II/I (c) in HepG2 cells treated with sorafenib (10 nM and 10 µM). The autophagic flux was assessed in HepG2 transfected with double-tagged pDest-LC3-GFP-mCherry plasmid, which is a pH-sensitive sensor to monitor autophagy in live cells (6 hr). The GFP tag is acid-sensitive while the mCherry tag is acid-insensitive. Yellow merge punctate structures indicate autophagosomes which neutral pH allows both tags emit fluorescent light resulting in a yellow staining. Subsequent fusion of autophagosomes with lysosomes results in red-positive acidic vacuoles where the green fluorescence from GFP is lost. Arrows indicate a typical example of colocalized particles of GFP and mCherry signal, while the arrowhead indicates a typical example of a particle with mCherry signal but without GFP signal due to its degradation. Sorafenib was used at a concentration of 10 µM, while EBSS was used as positive control treatment and chloroquine (50 µM) was used to inhibit autophagic flux. The presence of autophagolysosomes in HepG2 cells treated with sorafenib (10 µM) was also detected by transmission electronic microscopy (12 hr) following the procedure described in Material and Methods. The protein expression of autophagy markers (p62, Beclin-1, and LC3 II/I) was assessed at 6, 12 and 24 hr in sorafenib (10 nM and 10 µM)-treated HepG2 cells by western-blot procedure as described in Material and Methods. Results are expressed as mean ± standard error of the mean, and with the blots and images are representative of four independent experiments. *p ≤ 0.05, **p ≤ 0.01, and ***p ≤ 0.001 between control and sorafenib-treated cells. The groups with different letters (a, b, c, or d) were significantly different (p ≤ 0.05) [Color figure can be viewed at wileyonlinelibrary.com]
FIGURE 6  Effect of knock-down of ER stress branches, and inhibition of AMPK and autophagy in sorafenib-induced autophagy and apoptosis in HepG2. HepG2 cells were first transfected with IRE1α, PERK, and ATF6 si-RNA duplexes for 48 hr to assess their role in autophagy (a) and apoptosis (b,c). The expression of IRE1α, Ser51P-eIF2α/eIF2α and ATF6 was determined to validate si-IRE1α, si-PERK, and si-ATF6 treatments (a). Afterwards, cells were treated for 6 hr with sorafenib (10 nM and 10 µM) and harvested for analysis. The inhibition of AMPK was carried out by dorsomorphin (5 µM) with or without coadministration with sorafenib (10 µM) being measured Thr172P-AMPK/AMPK, Ser413FOXO3a/FOXO3a, and Beclin-1 (d) as well as caspase-3 activity (e) and cleaved PARP (f) at 12 and 24 hr. The inhibition of autophagy by 3-methyl adenine (3MA; 5 µM) or chloroquine (50 µM) administered 2 hr before sorafenib (10 µM) induced caspase-3 activity (g) and cleaved PARP protein expression (h) at 24 hr. Autophagy was biochemically followed by the measurement of p62, Beclin-1 and LC3II/I protein expression by western-blot analysis as described in Material and Methods. Apoptosis was analyzed by caspase-3 activity and cleaved PARP protein expression as described in Material and Methods. Results are expressed as mean ± standard error of the mean, and with the blots are representative of four independent experiments. *p ≤ 0.05, **p ≤ 0.01 and ***p ≤ 0.001 between Control and sorafenib-treated cells. The groups with different letters (a, b, c, d, e, or f) were significantly different (p ≤ 0.05). AMPK, 5′AMP-activated protein kinase; ER, endoplasmic reticulum; PERK, pancreatic ER kinase (PKR)-like ER kinase.
survival of patients, only a minor percentage of patients (2%) displayed partial response to therapy based on response evaluation criteria in solid tumors (RECIST) criteria (Llovet et al., 2008), the low response rate being mainly attributed to intrinsic resistance of HCC cells to sorafenib (Villanueva & Llovet, 2012). Several studies have provided mutational profile and identified an average of 30–40 mutations per tumor; among which 5–8 might be driver mutations, as well as high-level amplifications of oncogenes in specific chromosome..

**FIGURE 7** Kinetic study of the expression of Bcl-2 family members and Beclin-1 (a,b), JNK phosphorylation (c), and its inhibition on JNK phosphorylation (d), cell proliferation (e) and caspase-3 (f), as well as Foxo3a phosphorylation (g) in sorafenib-treated HepG2 cells. The effect of sorafenib (10 nM and 10 µM) on the expression of Bcl-2 family members and Beclin-1 were followed at 3, 6, 12, and 24 hr by western-blot analysis. The effect of sorafenib (10 nM and 10 µM) on the Thr183,Tyr185P-JNK1/2/JNK1/2 and Thr32P-Foxo3a/Foxo3a protein expression ratio was determined at different time points according to the experiment (3–48 hr). The inhibition of JNK was assessed by the administration of SP600125 (20 µM) added to cell culture 12 hr before sorafenib (10 µM) treatment. Cell proliferation was determined by BrdU incorporation as described in Material and Methods. Caspase-3 activity was measured by commercial chemiluminescence-based assay as described in Material and Methods. Results are expressed as mean ± standard error of the mean, and with the blots are representative of four independent experiments. *p ≤ 0.05 and **p ≤ 0.01 between control and sorafenib-treated cells. The groups with different letters (a, b, c, d, e, or f) were significantly different (p ≤ 0.05). JNK, c-Jun-N-terminal kinase.
regions. All influence high resistance and reduced extended survival rate during sorafenib treatment in HCC (Llovet, Villanueva, Lachmann, & Finn, 2015).

Sorafenib triggers antiproliferative and antiangiogenic effects through inhibition of the activity of several tyrosine kinase receptors including VEGFR-2/3, PDGFR-β, Flt3, and c-Kit, nonreceptor tyrosine kinase receptors such as Flt3, as well as Raf serine/threonine kinase (Cervello et al., 2012; Wilhelm et al., 2004). Antiproliferative and antiangiogenic effects induced by sorafenib, both in vitro and in vivo conditions, are driven through different mechanisms and promote apoptosis via both the extrinsic and the intrinsic pathway in HCC cells (Fernando et al., 2012). The in vivo administration of sorafenib (200 mg/kg) reduced tumor growth characterized by reduction of cell proliferation (ki67) and promoted apoptosis (caspase-3, cleaved PARP, and TUNEL) in tumors developed from subcutaneously HCC implanted cells in nude mice (Figure 1 and Supporting Information Figure S1). Sorafenib also increased Beclin-1 and LC3II/LC3I, but did not reduce the p62 expression in tissue homogenate, which may reflect high autophagic turnover in coexistence with apoptotic processes (Figure 1d). The effectiveness of sorafenib is probably a result of a balance between targeting cancer cells and the microenvironment (Llovet et al., 2015). In this sense, it is interesting to notice that the administration of sorafenib reduced fibrogenesis (α-SMA), angiogenesis (CD31), and the presence of hematopoietic stem cells (CD34) in tumors (Supporting Information Figure S1). These results, as well as those related to the antiproliferative
properties of the drug, are mostly dependent on the inhibition of tyrosine kinase receptors and the Raf/MEK/ERK.

The mechanism by which sorafenib induces apoptosis is not fully elucidated, and the effectiveness and adverse events are related to the dose used. The mean trough sorafenib concentration is 4.3 ± 2.5 mg/L in cancer patients under sorafenib monotherapy with the recommended dose (400 mg, 12 hr), which are increased up to 7.7 ± 3.6 mg/L in patients who experienced grade 3 drug-related adverse events (Blanchet et al., 2009). The corresponding therapeutic concentration of sorafenib used in the study is 10 µM. It has been associated with the inhibition of phosphorylation of the initiation factor eIF4E and loss of the antiapoptotic Bcl-2 family members Mcl-1 which appear to occur through a MEK/ERK independent mechanism in different cancer cell lines treated with sorafenib (Rahmani, Davis, Bauer, Dent, & Grant, 2005; Yu et al., 2005). Mcl-1, an antiapoptotic member of the Bcl-2 family, has been shown to be an important factor for apoptosis resistance in HCC (Fleischer et al., 2006). Sorafenib (10 µM) reduced the level of P-eIF4E and Mcl-1 at 2 and 16 hr after treatment in HepG2 and PLC/PRF/5 cells (Liu et al., 2006). However, we found that there is no temporal relationship between Mcl-1 downregulation and induction of apoptosis by sorafenib in HCC cells. In our conditions, caspase-3 activity was significantly increased 24 hr after the administration of sorafenib (10 µM) in HepG2 cells. HCC cells with mutated p53 (Huh7) and nonsense p53 mutation (Hep3B) required high sorafenib concentration (100 µM) to induce caspase-3 activation (Figure 2a). Data showed that the antiproliferative and proapoptotic kinetics induced by the drug were more potent in HCC cell lines than in primary human hepatocytes (Figure 2). In this sense, primary human hepatocytes were resistant to the proapoptotic properties of sorafenib treatment (Figure 2a). This aspect is relevant for the specificity of sorafenib for liver cancer cells instead of healthy hepatocytes. We undertook experiments to identify the differential pattern of miRNA expression upon sorafenib treatment among primary human hepatocytes and cancer cells.

One well-characterized function of mTORC1 is to maintain protein synthesis through phosphorylation of at least two direct targets, eukaryotic initiation factor (eIF) 4E-binding proteins (4E-BPs) and ribosomal protein S6 kinases (S6ks) (Hay & Sonenberg, 2004). mTOR is regulated by different cell signaling involving ER stress, Akt, and AMPK (Rashid, Yadav, Kim, & Chae, 2015). The induction of PERK signaling by antitumoral drugs inhibits mTOR and triggers autophagy in different cancer cell lines (Bruning et al., 2013). Sorafenib (10 µM) significantly increased PERK- and IRE1α-related signaling at 6 hr in HepG2 cells (Figure 3). The significant reduction of ATF6 (Figure 3d) may be related to the alteration of trafficking between the ER and the Golgi apparatus, as well as the overall secretory pathway induced by sorafenib (Yi et al., 2012). The activation of PERK and IRE1α signaling was associated with alteration of Thr172P-AMPK (Figure 4c) and Ser2448P-mTOR (Figure 4a). In addition, the early downregulation of mTOR activation appeared not to be related to reduced Akt phosphorylation. In fact, Thr385P-Akt increased 6–12 hr after sorafenib administration in HepG2 (Figure 4b). The activation of survival pathways on mTOR inhibition has been associated with upregulation of IGF-1 receptor in breast (O’Reilly et al., 2006) and rhabdomyosarcoma (Wan, Harkavy, Shen, Grohar, & Helman, 2007) cell lines. The inhibition of the mTOR pathway is in agreement with the drastic alteration of polyome profile we observed in sorafenib-treated HepG2 cells. In agreement with Sauzay et al. (2018), sorafenib induced a blockage of translation at initial stage (Figure 4d and Supporting Information Figure 4).

The most important pathways involved in autophagy modulation are mTORC1 and AMPK (Prieto-Dominguez et al., 2016). The inhibition of mTOR by sorafenib appeared to be stronger in HepG2 (Figure 4a) than in Hep3B and Huh7 (Supporting Information Figure S3). However, the activation of autophagic process using the tandem pDest-LC3-GFP-mCherry plasmid transfected to sorafenib-treated cells (Figure 5a), TEM (Figure 5b) and biochemical markers (Figure 5c) were more closely correlated to the activation of ER stress (Figure 3) and AMPK (Figure 4c) than mTOR inhibition (Figure 4a). The respective use of si-RNA and inhibitory compounds showed that ER stress (Figure 6a) and AMPK activation (Figure 6d) played an essential role during sorafenib-induced autophagy in HepG2. The inhibition of autophagy by si-IRE1α and si-PERK, as well as 3MA, and CQ increased caspase-3 (Figure 6b and 6g, respectively) and PARP cleavage (Figure 6c and 6h, respectively). It was also interesting to notice that the downregulation of autophagy by reduction of Thr172P-AMPK/AMPK ratio using dorsomorphin observed at 12 hr was followed by a drastic upregulation of Beclin-1 (Figure 6d) and reduction of caspase-3 activity (Figure 6e) and PARP cleavage (Figure 6f) in sorafenib-treated HepG2 cells at 24 hr. The experiments confirmed Thr172P-AMPK/AMPK ratio was positively correlated to the Ser413P-Foxo3a/Foxo3a ratio (Figure 6d). Our results support the survival properties of autophagy, and it may suggest an autophagic alternative activation pathway in dorsomorphin and sorafenib cotreated HepG2 cells.

Urano et al. (2000) reported that ER stress and, in particular the IRE1 kinase activity, increases JNK activity. The autophagic process, as apoptosis, appeared to be tightly regulated by anti- and proapoptotic members of the Bcl-2 family. The activation of the autophagic process (6–12 hr) by sorafenib was associated with a drastic increase of Beclin-1 expression, which it was preceded by rapid downregulation of Mcl-1 and upregulation of tBid and BimEL, while Bcl-2 and Bcl-xL remained at moderate levels during this period after drug administration (Figure 7a and 7b, and Supporting Information Figure S6). Interestingly, Bcl-xL and BimEL have been shown to promote autophagy by displacing Beclin-1 from its interaction with Bcl-2 (Erlich et al., 2007; Marquez & Xu, 2012). In fact, Beclin-1 has been identified as a BH3-only protein (Maiuri et al., 2007) that allows its interaction with Bcl2 and its homologue Bcl-xL (Oberstein, Jeffrey, & Shi, 2007). The expression of Beclin-1 (Park et al., 2009) and Bcl-2 phosphorylation that disrupts Beclin-1-Bcl-2 complex-1 (Wei, Pattingre, Sinha, Bassik, & Levine, 2008) have been related to the activation of JNK1. It was interesting to observe that Thr183 Tyr185P-JNK1/2/JNK1/2 was induced by sorafenib (10 µM) during the period in which autophagy was activated (3–12 hr; Figure 7c). The inhibition of Thr183, Tyr185P-JNK1/2/JNK1/2 (Figure 7d) by SP600125 was related to a reduction in cell proliferation (Figure 7e) that resulted in an increased apoptosis (Figure 7f), suggesting that JNK played a survival role during
sorafenib-induced apoptosis. In addition, a timeframe relationship existed between JNK and AMPK activation, supporting the existence of a complex signaling module involving LKB1, AMPK and JNK, in which early phase of JNK activation was also promoting cell survival by inducing AMPK activity, whereas the late phase of JNK activation induced apoptosis (Yun et al., 2009).

The activation of apoptosis by sorafenib was coincident with a sustained increase in BimEL expression, remaining Bak and Bax at a moderate expression level (Figure 7a and 7b, and Supporting Information Figure S6). JNK has also been observed to induce phosphorylation of Bim, promoting its release from its binding to dynein and myosin V motor complexes (Lei & Davis, 2003). Sorafenib (10 µM) induced a sustained and progressive increase of CHOP expression (6–24 hr; Figure 3b). The main effector of PERK-mediated apoptosis is the proapoptotic transcription factor CHOP which can be induced by ATF4, ATF6, as well as XBP1s (Verfaillie, Salazar, Velasco, & Agostinis, 2010). Interestingly, CHOP suppresses the expression of the antiapoptotic Bcl-2 (McCullough, Martinlde, Klotz, Aw, & Holbrook, 2001) and promotes the transcription of the proapoptotic BH3-only protein Bim (Puthalakath et al., 2007). The induction of thapsigargin-induced ER has also been related to a stabilization of Bim by PP2A-related dephosphorylation that prevents its ubiquitination and degradation, as well as by transcriptional regulation though CHOP-ATF4 in breast cancer cells (Puthalakath et al., 2007). The expression of Bim is also under positive transcriptional regulation of Foxo3a (Nakamura & Sakamoto, 2008) whose activity is regulated by a plethora of kinases such as GSK3β, MST1, JNK, CS1, AMPK, and AKT (Farhan et al., 2017). As described above, a positive correlation appeared between Thr^{172}p-AMPK/AMPK ratio and related activation phosphorylation of Ser^{413}p-Foxo3a/Foxo3a ratio (Figure 6d). In addition, PI3K/Akt survival signaling increases the inhibitory phosphorylation Thr^{32}p-Foxo3a, preventing its nuclear translocation and transcriptional activity (Brunet et al., 1999). Our study also showed that the expression of BimEL raised shortly after sorafenib administration which was maintained at moderate expression levels during autophagic-related phase (6–12 hr). This expression level might be the result of the interaction between the upregulation of Thr^{308}p-Akt (Figure 4b)-dependent Thr^{32}p-Foxo3a (Figure 7g), and Thr^{172}p-AMPK (Figure 4c)-dependent Ser^{413}p-Foxo3a (Figure 6d) in sorafenib-treated HepG2. Additional studies should be undertaken to identify the relevance of each pathway in the expression of BimEL. The sustained increase of PERK and CHOP expressions (Figure 3a and 3b, respectively) and reduction of Akt phosphorylation (24 hr; Figure 4b) supports BimEL upregulation during sorafenib-induced apoptotic phase (24 hr) in HepG2. The administration of Bim si-RNA, but not Bak or Bax, prevented sorafenib-induced caspase-3 activity in HepG2 cells (Figure 8a). The reduction of apoptosis by Bim si-RNA was associated with upregulation of p62 expression and LC3II/LC3I ratio, but without alteration of Beclin-1 expression in sorafenib-treated HepG2 cells (Figure 8b). The proapoptotic BH3-only members such as BimEL, and others like BNIP3, Bad, Bik, Puma, and Noxa, induce autophagy by disrupting the Beclin-1-Bcl2 complex (Marquez & Xu, 2012). The immunoprecipitation studies demonstrated the formation of Beclin-1 and Bim complex (Figure 8c). In this sense, our study may also suggest that the absence of Bim by si-RNA promotes the induction of early autophagic structures by sorafenib (10 µM) independently of Beclin-1 expression which remain at low level at 24 hr (Figure 8b).

In conclusion, our study showed that the induction of ER stress by sorafenib was the driving mechanism involved in the sequential induction of autophagy and apoptosis in HepG2 cells (Figure 9). The early sorafenib-induced ER stress and regulation of JNK and AMPK-dependent signaling were related to the induction of survival autophagic process (3–12 hr). The sustained treatment with sorafenib induced a progressive increase of ER stress and PERK-CHOP-dependent rise of BimEL, which induced the shift from autophagy to apoptosis (24 hr). However, the kinetic of BimEL expression profile might also be related to the tight balance between Akt- and AMPK-related signaling in Foxo3a-dependent BimEL upregulation.

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**CONFLICTS OF INTEREST**

The authors have no commercial associations that might create a conflict of interest in connection with the present submitted
manuscripts. The authors declare there are any competing financial interests in relation to the work described.

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**REFERENCES**


**SUPPORTING INFORMATION**

Additional supporting information may be found online in the Supporting Information section at the end of the article.