β-Lapachone Selectively Kills Hepatocellular Carcinoma Cells by Targeting NQO1 to Induce Extensive DNA Damage and PARP1 Hyperactivation

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Hepatocellular carcinoma (HCC) is the second leading cause of cancer-related death globally. Currently there is a lack of tumor-selective and efficacious therapies for hepatocellular carcinoma. β-Lapachone (ARQ761 in clinical form) selectively kills NADPH: quinone oxidoreductase 1 (NQO1)-overexpressing cancer cells. However, the effect of β-Lapachone on HCC is virtually unknown. In this study, we found that relatively high NQO1 and low catalase levels were observed in both clinical specimens collected from HCC patients and HCC tumors from the TCGA database. β-Lapachone treatment induced NQO1-selective killing of HCC cells and caused ROS formation and PARP1 hyperactivation, resulting in a significant decrease in NAD+ and ATP levels and a dramatic increase in double-strand break (DSB) lesions over time in vitro. Administration of β-Lapachone significantly inhibited tumor growth and prolonged survival in a mouse xenograft model in vivo. Our data suggest that NQO1 is an ideal potential biomarker, and relatively high NQO1:CAT ratios in HCC tumors but low ratios in normal tissues offer an optimal therapeutic window to use β-Lapachone. This study provides novel preclinical evidence for β-Lapachone as a new promising chemotherapeutic agent for use in NQO1-positive HCC patients.

Keywords: beta-lapachone, NQO1, hepatocellular carcinoma, DNA damage/reactive oxygen species, NAD+/ATP depletion
INTRODUCTION

As the most common primary liver cancer and the second leading cause of cancer-related deaths worldwide (1), hepatocellular carcinoma (HCC) patients are usually diagnosed with advanced disease, resulting in only 15% of HCC patients being eligible for surgical resection or liver transplantation and the median survival time for HCC patients in intermediate to advanced stages being only 1-2 years (2). Moreover, chemotherapy against HCC has limited benefits because of the high resistance to currently available chemotherapeutic agents. Sorafenib, the first-line drug used for patients with advanced HCC, has been used for over 10 years, but the overall outcomes are unsatisfactory (3). These have led to more extensive research focusing on personalized medicine with increased selectivity and efficacy.

A growing body of data demonstrates that NADPH:quinone oxidoreductase 1 (NQO1), a phase II two-electron reductase that can bioactivate certain quinone molecules and shows a protective effect against natural and exogenous quinones, is abnormally upregulated in many solid cancers, such as lung, pancreatic, breast, prostate, and colon cancers (4-11). In liver cancer, it has been reported that NQO1 was increased 18-fold in HCC versus normal livers (12). Recently, NQO1 overexpression was reported to be a potent independent biomarker for prognostic evaluation of HCC (13) and enhanced apoptosis inhibition of liver cancer cells via the SIRT6/AKT/XIAP signaling pathway (14, 15). NQO1 overexpression in tumors has the advantage of preferentially killing cancer cells and sparing normal cells when anticancer drugs that are bioreductively activated by NQO1, such as β-Lapachone (β-lap), are used (5).

β-lap has gained increasing attention for its tumor-selective and antitumor effects in many cancers, including lung cancer, breast cancer, prostate cancer, pancreatic cancer, and leukemia (4-7, 9-11, 16, 17). Its toxicity is closely correlated with NQO1 expression and activity. Our studies suggest that NQO1 metabolizes β-lap through a futile redox cycle in which β-lap is converted into a highly unstable hydroquinone form and then spontaneously reacts with oxygen to revert back to the parent compound, causing rapid NAD(P)H oxidation. This process generates high levels of reactive oxygen species (ROS) (e.g., H$_2$O$_2$), resulting in genomic instability and DNA damage (6, 11). In addition, catalase (CAT) can bypass β-lap toxicity by neutralizing hydrogen peroxide produced by β-lap (18). We have previously reported that β-lap alone or combined with other inhibitors had profound toxicity in pancreatic cancer and nonsmall-cell lung cancer (NSCLC) (11, 19-21). However, the effect of β-lap on HCC is virtually unknown.

Here, we demonstrate that HCC patient samples have significantly elevated levels of NQO1 but concomitantly low catalase levels compared with associated normal tissues. HCC cells were efficiently killed by β-lap, along with increases in ROS production and PARP1 hyperactivation, severe NAD$^+/$/ATP depletion and DNA damage. NQO1-dependent HCC killing was confirmed in HCC cells with stable NQO1 overexpression and knockout. Furthermore, a human HCC subcutaneous xenograft mouse model exhibited efficient β-lap-induced control of tumor growth and prolonged mouse survival.

MATERIALS AND METHODS

Human HCC Cell Lines and Clinical Samples

Human HCC cell lines (SNU-182, PLC/PRF/5, Huh7, HepG2, and Li7) were purchased from Guangzhou Cellcook Biotechnology Co., Ltd. (Guangzhou, China), and PLC/PRF/5 and SK-HEP1 cells were purchased from ATCC. The authentication of these cell lines was performed via comparisons with the STR database. Cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal bovine serum (FBS; HyClone), 100 U/ml penicillin, and 100 U/ml streptomycin at 37°C and 5% CO$_2$.

Reagents and Chemicals

β-lap, synthesized by Dr. Bill Bornmann (M.D. Anderson, Houston, TX) was dissolved in DMSO for in vitro experiments or 20% HPβCD for in vivo experiments, and the concentrations were verified by spectrophotometry. Hydrogen peroxide (H$_2$O$_2$) and dicoumarol (DIC) were purchased from Sigma-Aldrich. HPβCD (>98% purity) was purchased from Cydodextrin Technologies Development, Inc. The ROS-Glo™ H$_2$O$_2$ assay kit, NAD/NADH-Glo kit, and CellTiter-Glo® 2.0 kit for the ATP assay were obtained from Promega Corporation. An alkaline comet assay kit was purchased from Trevigen, Inc.

NQO1 Knockout/Knockin Cells

CRISP/Cas9 NQO1 knockout PLC/PRF/5 cells and NQO1-overexpressing SK-HEP1 cells were generated by our lab. Vectors of guide RNA sensing human NQO1 or nontarget control (LV04) and Cas9 expression (CAS9NEO) were provided by Sigma-Aldrich, and the guide RNA targeting sequences were AGGATACTGAAAGTTCGCAAGG and CAACATATCTGGGAGTCAGATGG. Vectors with NQO1 or empty control (EX-Z0563-Lv205, EX-NEG-Lv205) were purchased from Sigma-Aldrich, and transfection of SK-HEP1 with these vectors was performed with Lipofectamine 3000 reagent (Thermo Fisher) according to the manufacturer’s protocol.

Cell Survival Assay

A total of 10,000 cells/well were seeded on 48-well plates 24 h prior to treatment. Varying doses of β-lap dissolved in DMSO ± DIC were added and incubated for 2 h at 37°C and 5% CO$_2$. After treatment, the media were replaced with fresh complete media and allowed to grow for 7 days. After 7 days, the cells were washed with 1x PBS, 200 µl of H$_2$O was added, and the cells were frozen at -80°C for at least 2 h. After thawing, 200 µl/well TNE
buffer (50 mM Tris–HCl (pH 7.4), 100 mM NaCl, 0.1 mM EDTA) with 1 μg/mL Hoechst 33258 was added and incubated for 1 h at room temperature in the dark. Cell growth was determined by absorbance at 560 nM with a multilabel plate reader (PerkinElmer). Percentage of cell growth = (100× (cell experimental – blank)): (cell control).

**Antibodies**

Antibodies for immunofluorescence and western blotting included NQO1 (A180, Santa Cruz, La Jolla, CA), PARP1 (F-2, Santa Cruz), β-actin (C4, Santa Cruz), α-tubulin (B-7, Santa Cruz), PAR (Trevigen, Gaithersburg, MD), γH2AX (JBW301, Millipore, Temecula, CA), H2AX (938CT5.1.1, Cell Signaling, Danvers, MA), and catalase (12980S, Cell Signaling, MA).

**Western Blotting Analysis**

Cells were seeded on plates at approximately 70% confluence 24 h in advance and then treated with/without β-lap for 2 h. Next, the cells were lysed in lysis buffer. Approximately 40 μg of protein was resolved by SDS-PAGE, transferred onto PVDF membranes, and probed with antibodies. The protein-antibody complexes were detected using Super Signal West Femto Substrate (Thermo Fisher, Waltham, MA) and exposure to film.

**Real-Time PCR**

Assays were performed as previously described (22). The primer sequences were as follows: GAPDH-sense: 5'-CTGCTGTAGTCCCCATGTTC-3'; GAPDH-antisense: 5'-CATCCACAGTCTTCTGGGTGG-3'; NQO1-sense: 5'-GCCATGTATGCAAAAGGA CCC-3'; NQO1-antisense: 5'-ACTTGGAGGCCACGAAATAGC-3'; CAT-sense: 5'-CTTCTGACCAAGCAACATGC-3'; CAT-antisense: 5'-GGGTGAGTGTCAGGATA GG-3'. 2-ΔΔCt was used to calculate the fold change of mRNA expression.

**NQO1 and Catalase Activity Assays**

Extracts were obtained from different HCC cell lines. Then NQO1 and catalase enzyme activities were assayed by NQO1 Activity Assay Kit (Abbkine), respectively, according to the manufacturer's manual.

**ATP, NAD+ and Hydrogen Peroxide (H2O2) Assays**

Cells were cultured (1 × 10^5 cells/well) 24 h in advance in 96-well white-walled clear-bottom tissue culture plates (Sigma) and treated with β-lap with or without DIC for 2 h. Then, ATP (CellTiter-Glo), hydrogen peroxide (H2O2) (ROS-Glo), and NAD/NADH (NAD/NADH-Glo) were assayed at the indicated time points after treatments using specific assays (Promega).

**Comet Assay**

Total DNA damage was measured by the alkaline comet assay (Trevigen) according to the manufacturer’s manual. Slides were stained with SYBR green, and images were acquired with a Leica DM5500 fluorescence microscope. Comet tail lengths were quantified by NIH ImageJ.

**Immunofluorescence Staining of γH2AX**

The treated cells were fixed with 4% paraformaldehyde for 30 min, permeabilized with 0.2% Triton-X 100 for 10 min at 4°C, blocked with 3% bovine serum albumin for 30 min at room temperature, and incubated overnight at 4°C with γH2AX antibody (diluted at 1:1000). Cells were washed 3 times for 5 min in PBS and then incubated for 2 h with AlexaFluor secondary antibody (diluted 1:1000 in blocking buffer). DAPI was used to stain nuclei. γH2AX foci were visualized with a laser scanning confocal microscope (LSM 510 Meta), and the number of H2AX foci per nucleus was quantified.

**In Vivo Antitumor Study**

All animal procedures were approved by the IU IACUC committee. For the in vivo xenograft model, PLC/PRF/5 cells (5×10^6) were subcutaneously inoculated into the right flank of nonobese diabetic/severe combined immunodeficiency (NOD/SCID) male mice (6~8 weeks old). Tumor volumes were measured with a caliper and calculated by the formula 0.5×length×width^2. When tumor volumes reached ~150 mm³, mice were randomly divided into vehicle (n=7) and treatment (n=8) groups with no significant differences in tumor sizes. Then, the mice were treated with HPβCD or HPβCD-β-lap (12.5 mg/kg) by intratumor injection every other day for a total of five injections. When the tumor volume reached ~150 mm³, the mice were sacrificed, and a survival curve was plotted.

**Bioinformatic Analysis**

The liver hepatocellular carcinoma (LIHC) dataset was downloaded from the Broad Institute TCGA Genome Data Analysis Center, https://doi.org/10.7908/C11G0KMK9. Only samples with both tumor and matched normal samples were selected for further analysis of NQO1 and CAT levels. The differences in gene expression levels for individual genes or fold changes of two genes between normal and tumor tissues were identified by paired t test.

**Statistical Analysis**

All the experimental results were analyzed using two-tailed Student’s t tests for independent measures with Holm-Sidak correction for multiple comparisons if >1 comparison was performed. The minimum replicate size for experiment was n=3. Statistical analysis were performed in GraphPad Prism 8 (GraphPad Software, Inc. CA, USA). Images are representative of the results of experiments or staining repeated 3 times. Data are presented as the mean ± SD. The survival rate was analyzed by Kaplan-Meier survival curves. A p value of < 0.05 was considered statistically significant between the compared groups. *p < 0.05; **p < 0.01 and ***p < 0.001.

**RESULTS**

**High Expression of NQO1 in Hepatocellular Carcinoma Patients**

Our previous studies and other reports have shown that NQO1 enzyme levels were elevated, whereas catalase (gene: CAT) levels
were lower in NSCLC and pancreatic cancers than in associated normal tissues, suggesting that the NQO1/CAT ratios in tumor tissue versus associated normal tissue are an important and highly exploitable therapeutic window (8, 11, 23). To investigate whether the NQO1:CAT ratios are a potential therapeutic window in liver cancer, we analyzed NQO1 and CAT expression in liver hepatocellular carcinoma (LIHC) from The Cancer Genome Atlas (TCGA). Our analysis revealed that the mRNA levels of NQO1 were significantly elevated (Figure 1A, \( p = 1.5 \times 10^{-9} \)), while CAT levels were notably lower (Figure 1B, \( p = 7.8 \times 10^{-6} \)) in tumor tissues than in matched normal tissues. Consistently, markedly higher NQO1:CAT ratios were observed in these tumor samples than in normal samples (Figure 1C, \( p = 1.5 \times 10^{-9} \)). Moreover, we also found that patients with high NQO1 expression had a significantly lower overall survival rate than those with low NQO1 expression (Figure 1D, \( p = 0.0075 \)). Together, these results indicate that the NQO1:CAT ratio is an ideal therapeutic window in liver cancer for NQO1 bioactivatable drugs.

Elevation of NQO1 Expression and Enzyme Activity in Hepatocellular Carcinoma Patients and Cell Lines

To further confirm the above observations, we collected 62 pairs of clinical HCC patient samples and associated normal tissues to detect the mRNA levels of NQO1 and CAT. Indeed, 69.4% (43/62) of HCC patient samples showed relatively higher NQO1 mRNA levels than associated normal tissues (\( p = 0.0005 \), Figure 2A). In contrast, notably lower CAT mRNA levels (54/62 = 87.0%, \( p < 0.0001 \), Figure 2B) were observed in tumor tissues than in associated normal tissues. Concomitant high NQO1 and low CAT mRNA levels (high NQO1:CAT ratios (Figure 2C, \( p < 0.0001 \)) in HCC tumor tissue offer an ideal target for NQO1 bioactivatable drugs. Consistently, our western blotting analysis revealed that NQO1 protein levels were obviously elevated in 43.8% (28/64) of HCC tumor tissues, while catalase levels were repressed significantly in most HCC tumor tissues (Figure 2D and Supplementary Figures S1A, B). These results confirm our above observations that the relatively high NQO1:CAT ratios in HCC patients might be an exploitable therapeutic target in liver cancer. On the other hand, similar to the analysis in HCC patient samples, we observed that the liver cancer cell lines HepG2, Huh7 and Li7 showed high NQO1 levels; PLC/PRF/5 cells exhibited moderate NQO1 expression; and Hep3B, SNU-182, and SK-HEP1 cells expressed low or undetectable NQO1 (Figure 2E). Consistently, the NQO1 enzyme activity assay exhibited similar NQO1 activity in these cell lines (Figure 2F). Meanwhile, relatively high catalase protein levels accompanied by relative high catalase enzyme activities were observed in HepG2, Huh7 and Hep3B liver cancer cells (Figures 2E, G).

Selectively and Effective Killing of Hepatocellular Carcinoma Cells by β-Lapachone

It has been reported that NQO1 is a promising therapeutic target in multiple solid tumors, and the anticancer efficacy of β-lap is mainly mediated and promoted by NQO1 (7, 11, 17). Based on our above findings, we hypothesized that β-lap could effectively control cell growth in NQO1+ HCC cells. To this end, we treated...
Accumulating evidence suggests that exposure to β-lap causes DNA lesions in NQO1⁺ NSCLC, breast cancer, and pancreatic cancer cells, resulting in PARP hyperactivation in terms of the accumulation of poly(ADP-ribose)-PARP (PAR-PARP) posttranslational protein modification (11, 18, 24). To investigate whether PARP1 is involved in β-lap-induced cell death in HCC cells, we examined the effect of β-lap on poly(ADP-ribose)ylated protein (PAR) accumulation, which is an indicator of PARP1 hyperactivation. First, β-lap-treated PLC/PRF/5 and Hep7 cells were analyzed for PAR formation using western blotting analysis. As shown, a lethal dose of β-lap (10 µM for PLC/PRF/5, 4 µM for Hep7 cells) significantly induced PARP1 hyperactivation, as indicated by a rapid rise in PAR formation and then an increase in DNA damage, as indicated by γH2AX expression over time (Figure 4A and Supplementary Figure S2A). To confirm that the β-lap-induced hyperactivation of PARP1 is NQO1-dependent, we next examined PAR formation in NQO1 knockout PLC/PRF/5 and NQO1-expressing SK-HEP1 cells. As expected, PAR formation was significantly inhibited by β-lap after NQO1 was stably knocked out in PLC/PRF/5 cells, accompanied by no detectable γH2AX expression (Figure 4B). A rapid increase and continuous level of PAR was detected after NQO1 re-expression in these SK-HEP1 cells. In addition, no significant change of catalase levels were noted in these β-lap-treated cells (Figures 4A–C). Taken together, these data indicate that β-lap induces PARP1 hyperactivation and DNA damage in NQO1⁺ HCC cells.

Intracellular ROS production is crucial for cancer cell death (25), and our previous studies show that NQO1 metabolizes β-lap in a futile redox cycle manner to generate ROS in other solid cancer cells (6, 26). Therefore, we investigated whether ROS are involved in β-lap-induced HCC cell death. We examined the levels of hydrogen peroxide (H2O2) as an indicator of intracellular ROS. After 2 h of exposure to a sublethal dose (4 or 8 µM) of β-lap, PLC/PRF/5 cells exhibited a significantly higher level of H2O2 (p < 0.001) than the untreated group (Figure 4D), while NQO1 KO cells showed no
FIGURE 3 | The cytotoxicity of β-Lapachone in hepatocellular carcinoma cells is NQO1-dependent. (A–F) HCC cells (HepG2 (A), Huh7 (B), Li7 (C), PLC/PRF/5 (D), SK-HEP1 (E), SNU-182 (F), and Hep3B (G)) were exposed to β-lap (0–10 µM) ± dicoumarol (DIC, 50 µM) for 2 h, and then relative survival was assessed. (H, I) NQO1-overexpressing SK-HEP1 cells (H) and stable NQO1 knockout PLC/PRF/5 cells (I) were treated as in (A), and then cell viability was assessed. All error bars are means ± SDs. ***p < 0.001 (t tests). No addition, DMSO alone; DIC, dicoumarol; β-lap, β-Lapachone. Inset, NQO1 expression in NQO1-overexpressing SK-HEP1 and NQO1 knockout PLC/PRF/5 cells was confirmed by western blotting. %T/C, the mean of %Treated/Control.

FIGURE 4 | β-Lapachone induces NQO1-dependent PARP1 hyperactivation, ROS formation and NAD+/ATP loss. (A–C) PLC/PRF/5 (A), NQO1 knockout PLC/PRF/5 (B) and NQO1-overexpressing SK-HEP1 (SK-HEP1 NQO1+) (C) cells were exposed to 10 µM β-lap for the indicated times. Cells were then harvested, and western blotting analysis was performed to detect the levels of PAR (PARP1 hyperactivation), PARP1, NQO1, catalase and γH2AX. The protein levels of PAR and γH2AX were quantified by Image J and normalized to α-tubulin, which was used as a loading control. (D–F) PLC/PRF/5 cells were treated with or without β-lap (4 or 8 µM) ± DIC (50 µM) for 2 h. Then, cells were subjected to measurement of H2O2 levels (D), NAD+ levels (E) and ATP levels (F). Data represent at least three independent sets of experiments. Error bars are means ± SDs. ***p < 0.001, **p < 0.01 (t tests). T/C, the mean of Treated/Control. %T/C, the mean of %Treated/Control. NS, Not Statistically Significant.
changes in H$_2$O$_2$ levels. Moreover, no obviously increase of H$_2$O$_2$ levels was observed in SK-HEP1 cells that have undetected NQO1 expression, while a significant increase of H$_2$O$_2$ levels was noted after reconstitution with NQO1 (Supplementary Figure S2B). Similarly, a significant increase in H$_2$O$_2$ levels in the β-lap-treated group was observed in NQO1$^+$ Huh7 cells, and DIC blocked this increase (Supplementary Figure S2C). As reported previously by us and others (11, 27), β-lap-induced PAR-PARP1 formation consumes NAD$^+$ and ATP, and together with our above results that PAR is induced rapidly and then decreases over time. Therefore, we examined NAD$^+$ and ATP levels in β-lap-treated HCC cells. As shown in Figures 4E, F and Supplementary Figures S2D, E, dramatic NAD$^+$ and ATP depletion was observed after NQO1$^+$ cells were exposed to β-lap for 2 h. All these depletions were blocked in the NQO1 KO cells or DIC-treated group. When NQO1 expression was restored in SK-HEP1 cells, markedly increase of NAD$^+$ and ATP levels were observed after treatment with β-lap (Supplementary Figures S2F, G). Together, these results demonstrate that β-lap induces cell stress in HCC cells by generating ROS, and disrupts essential metabolic nucleotides.

β-Lapachone Causes Dramatic NQO1-dependent Total DNA Damage and Double-Strand Breaks in Hepatocellular Carcinoma Cells

According to previous reports, ROS release can mediate and promote DNA damage (28, 29), and our above results show that β-lap generates ROS in HCC cells. Therefore, we investigated total DNA damage under β-lap treatment via an alkaline comet assay in PLC/PRF/5 cells. In NQO1-expressing PLC/PRF/5 wild-type cells, a sublethal dose of β-lap (4 µM) caused total DNA damage, as indicated by the comet tail, as early as 30 min, and a lethal dose of β-lap (10 µM) markedly increased DNA damage (Figure 5A). In contrast, when NQO1 was knocked out, no obvious DNA tails were observed even at a lethal dose of β-lap (10 µM) (Figure 5A). Consistently, quantification of the comet tail length confirmed these observations. On the other hand, our previous study suggested that when PARP hyperactivity is exhausted, cells attempt to replicate despite the damage to AP sites or the presence of SSBs, and the damage becomes hypersensitive to the oxidative stress caused by β-lap and then induces DSBs (21). In fact, we observed γH2AX expression after β-lap treatment in NQO1$^+$ HCC cells, especially when PAR levels were exhausted (Figure 4). To further confirm whether β-lap induces DSBs in HCC cells, we examined γH2AX expression via immunofluorescence staining. As shown in Figure 5B, dramatic increases in DNA DSB formation were noted in the PLC/PRF/5 cells that were treated with a sublethal or lethal dose of β-lap compared to the untreated cells as early as 30 min. Even at a lethal dose of β-lap (10 µM), NQO1 knockout cells showed no obvious increase in γH2AX foci. The observations were confirmed by the quantification of γH2AX foci formation (Figure 5B). Taken together, these data suggest that exposure of NQO1$^+$ HCC cells to β-lap results in cell death due to significant DNA DSBs.

FIGURE 5 | β-Lapachone induces NQO1-dependent DSB formation in hepatocellular carcinoma cells. Wild-type or NQO1 knockout PLC/PRF/5 cells were treated with a sublethal (4 µM) or lethal (10 µM) dose of β-lap for the indicated time (min), and then total DNA lesions were assessed using the alkaline comet assay. (A) Comet tail lengths were imaged under an immunofluorescence microscope. (B) DSBs were quantified by γH2AX foci/nuclei. Data represent the means ± SDs. Student’s t tests were performed. Scale bar = 10 µm, ***p < 0.001, *p < 0.05.
**β-Lapachone Significantly Suppresses Tumor Growth and Prolongs Survival in HCC Mouse Models**

Relatively high NQO1 and low catalase expression in various HCC patients and cell lines indicates a potent therapeutic window in liver carcinoma for β-lap, and our above data confirmed that β-lap efficiently represses tumor cell growth in vitro. To address the antitumor efficacy of β-lap in vivo, we established xenograft liver cancer models. First, 5×10⁶ PLC/PRF/5 cells/mouse were implanted subcutaneously into NOD/SCID mice. When the tumor size reached 150 mm³, the mice were grouped and treated with vehicle (HPβCD, intratumor (i.t.)) or HPβCD-β-lap (hereafter referred to as β-lap, at 12.5 mg/kg, i.t.) every other day for a total of five injections. Tumor volume and survival were scored to monitor tumor growth and response (Figures 6A–C). Exposure to β-lap resulted in a dramatic decrease in tumor growth compared with vehicle group mice, confirmed by the quantification of tumor volume (Figures 6A, B). Moreover, overall survival also showed significant antitumor efficacy of β-lap (Figure 6C). This model suggests that β-lap is a good candidate agent to kill NQO1⁺ HCC in vivo.

**DISCUSSION**

Here, we show a potential antitumor effect of β-lap in HCC and reveal that β-lap efficiently killed NQO1-overexpressing HCC cells without affecting NQO1-low-expressing cells or tissues. As the most common type of primary liver cancer, HCC often occurs in people with long-term liver diseases and is generally diagnosed with advanced disease, leading to systemic therapy (30, 31). At present, there are only a few efficacious drugs for HCC treatment or the drugs cause severe side effects (32, 33). Thus, exploiting new drugs and identifying differences between carcinomas and healthy tissue are critical for HCC treatment.

Previous reports suggest that β-lap is a competent tumor-selective agent against many NQO1⁺ solid cancers, such as NSCLC, pancreatic and breast cancers (6, 11, 34–36), while the antitumor effect of this drug in liver cancer is still unknown. Except the expression and activity of NQO1, the cytotoxicity of β-lap is also driven by ROS-metabolizing enzymes catalase and SOD1 expression and activity. Catalase is an important resistance factor in β-lap-induced cytotoxicity and this resistance could be enhanced by superoxide dismutase (SOD) (37). The NQO1:CAT ratios are suggested to be a therapeutic window in NSCLC, pancreatic and breast cancers (11, 18). In our study, analysis of 64 clinical patient samples of HCC revealed that 43.8% of patients exhibited NQO1 overexpression, and the majority of patients showed low CAT levels in tumors compared with adjacent normal tissues. Moreover, LIHC data from TCGA demonstrated that patient overall survival significantly correlated with NQO1 expression. These results suggest that relatively high NQO1:CAT ratios in tumor tissues could provide a therapeutic window for using NQO1 bioactivatable...
drugs such as β-lap to kill HCC. In fact, cell viability showed that β-lap efficiently killed PLC/PRF/5, Huh7, and Li7 cells, which have high NQO1 expression, but did not affect NOQ1 cell growth (Figure 2).

β-lap was reported to induce cancer cell death via an NQO1-dependent programmed necrotic pathway, which caused robust ROS elevation and PARP1 hyperactivation (6, 38). In HCC cells, we observed β-lap-induced NQO1-selective elevated H2O2 and a rapid and transient increase in PAR formation, followed by DNA damage over time. On the other hand, PARP1 uses NAD+ as a substrate to perform PAR posttranslational modification of proteins, resulting in NAD+ and ATP losses (27, 39). Our β-lap-treated HCC cells indeed exhibited NQO1-dependent NAD+ and ATP depletion. Collectively, our investigation of clinical patient samples of HCC and in vitro results offer a therapeutic window and potential use of β-lap in HCC.

β-lap has an apparently broader NQO1-dependent therapeutic window in HCC. NQO1 is overexpressed in numerous human cancers, and our previous studies together with others demonstrate that β-lap has efficacy in tumor-selective cell growth control and produced promising preclinical results (6, 11, 19, 34, 36). Our preclinical model exhibited an efficient antitumor effect of β-lap in HCC in which β-lap-treated mice had dramatically decreased tumor growth and prolonged survival compared to control mice. Together with our in vitro results, we anticipate that β-lap would be an extremely efficacious tumor-selective therapy against HCC and other kinds of liver cancer. Furthermore, the data presented in this study reveal that NQO1 and the NQO1:CAT ratio could be used as biomarkers to examine the efficacy of NQO1 bioactivatable drugs in HCC or other kinds of liver cancers. In addition, our in vitro and in vivo data could translate our findings regarding β-lap in HCC to the clinic. Moreover, because β-lap alone induces tumor programmed necrotic cell death that could induce many cytokines or other side effects in vivo, our recent report revealed that low-dose β-lap combined with a PARP inhibitor switched the pathway to apoptosis (11), which implies that combination therapy between β-lap and other clinical drugs would be worth exploring. Finally, immunotherapy with immune checkpoint inhibitors such as PD-1 inhibitors has shown promise in HCC (40). Clinical data showed that only approximately 15–20% of HCC patients exhibited a response, and a fraction of HCC patients could benefit from this therapy (41, 42). We recently revealed that β-lap not only directly kills tumor cells but also increases tumor immunogenicity by triggering immunogenic cell death and overcoming immunotherapy resistance (43). Thus, we propose that β-lap could exert a synergistic effect with immune checkpoint inhibitors and enhance the antitumor immune response in HCC.

Taken together, our study demonstrated that β-lap, a novel NQO1 bioactivatable drug, selectively kills HCC cells expressing NQO1 through inducing ROS and PAR formation, NAD+ and ATP depletion and lethal DNA damage. High NQO1:CAT ratios in HCC tumors but low ratios in normal tissues offer an optimal therapeutic window and an ideal therapeutic target for β-lap.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/Supplementary Material. Further inquiries can be directed to the corresponding author.

ETHICS STATEMENT

The studies involving human participants were reviewed and approved by Zhongshan Hospital of Xiamen University. The patients/participants provided their written informed consent to participate in this study. The animal study was reviewed and approved by the IU IACUC committee.

AUTHOR CONTRIBUTIONS

LJ, WZ, and XH designed the experiments, analyzed the data, and wrote the manuscript. XH supervised the project. SL, YY, HZ, and JW. analyzed the bioinformatic data. TF, FF, YL, YZ, XS, and JWV provided help with the experiments. LJ, WZ, YC, JW, and XH reviewed and edited the manuscript. All authors contributed to the article and approved the submitted version.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fonc.2021.747282/full#supplementary-material

Supplementary Figure S1 | NQO1 and catalase expression in hepatocellular carcinoma patients. (A, B) Western blotting analysis of NQO1 and catalase protein expressions in 54 pairs of HCC patient tumor samples and adjacent normal tissues. N, Normal; T, Tumor.

Supplementary Figure S2 | β-Lapachone induces NQO1-dependent PARP1 hyperactivation, ROS formation and NAD+/ATP loss in Huh7 cells. (A) Huh7 cells were exposed to 4 µM β-lap at the indicated time, then cells were harvested and western blotting analysis was to detect the levels of PAR (PARP1 hyperactivation) and γH2AX. (B-G) SK-HEP1, SK-HEP1 NQO1+, and Huh7 cells were treated with or without β-lap + DIC (50 µM) for 2 h. Then cells were measured for H2O2 levels (B, C). NAD+ levels (D, F), and ATP levels (E, G). Data represent at least three independent sets of experiments. Error bars are means ± SD. ***p < 0.001, **p < 0.01, *p < 0.05 (t tests).
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