

Activation of the β -TrCP/I κ B α /inflammation axis limits the sensitivity of liver cancer cells to neddylation inhibition

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Abstract. Upregulation of protein neddylation occurs in numerous types of human cancer, including liver cancer. MLN4924, a potent neddylation-inhibiting pharmacological agent, demonstrates anticancer ability in numerous cancers. However, the sensitivity of MLN4924 in liver cancer remains unsatisfactory due to factors causing resistance. RT-qPCR and western blotting were utilized to assess the mRNA and protein levels of genes, respectively. Cell Counting Kit-8 assay and colony formation assays were employed to assess cell viability and proliferation. The pathway of protein degradation and stability were determined by western blotting after treatment with MG132 and cycloheximide. An immunoprecipitation assay was utilized to detect the ubiquitination of protein. An *in vitro* ubiquitination assay was used to determine the ubiquitin linkage. To the best of our knowledge, the present study was the first to demonstrate that NF- κ B inhibitor α (I κ B α) downregulation and subsequent inflammation in response to MLN4924 limited the antitumor potential of MLN4924. Ectopic expression of I κ B α enhanced the antitumor potential of MLN4924 in liver cancer cells. Moreover, the results of the present study demonstrated that MLN4924 decreased I κ B α via promoting the K48 linkage of ubiquitin to I κ B α . Mechanistic studies demonstrated that MLN4924 enhanced the protein stability of β -transducin repeat-containing protein (β -TrCP), promoting the ubiquitination of I κ B α , which led to the

ubiquitin-mediated degradation of I κ B α . In addition, the results of the present study also demonstrated that β -TrCP knockdown markedly inhibited MLN4924 from suppressing the growth of liver cancer cells, via attenuating MLN4924-mediated I κ B α downregulation and inflammation. Collectively, these results indicated that the β -TrCP/I κ B α /inflammation pathway may act as a novel resistance factor of MLN4924, and targeting β -TrCP may be beneficial for the treatment of liver cancer.

Introduction

Hepatocellular carcinoma (HCC) accounts for ~80% of primary liver cancers and is the third leading cause of cancer-related mortality worldwide (1,2). In patients with early-stage liver cancer, surgery resection and thermal ablation are the most common available treatment options (3,4). However, the majority of patients with liver cancer are diagnosed at an advanced stage, and are therefore unlikely to respond to these treatment options due to the rapid development and distant metastases of HCC (5). Over the past decade, treatment options for HCC have progressed and include multi-kinase inhibitors (sorafenib and lenvatinib) and immune checkpoint inhibitors. However, treatment of advanced HCC remains unsatisfactory, as the overall survival rates of patients with liver cancer are marginally extended following treatment (6). Therefore, further investigations are required for the development of novel potential therapies for liver cancer including HCC.

Protein neddylation is an important post-translational modification that adds the ubiquitin-like molecule NEDD8 to substrate proteins (7). This is carried out via a series of processes that are successively catalyzed by E1 NEDD8-activating enzyme (NAE1), E2 neddylation conjugation enzymes and E3 neddylation ligases. Previous studies have demonstrated that neddylation regulates protein function and stabilization. Well-established substrates of the neddylation pathway are the Cullin family of proteins that are the scaffold components of cullin-RING ligases (CRLs). Neddylation of Cullin is indispensable for the activation of CRLs and the subsequent ubiquitination and degradation of CRL substrates (8). Results of a previous study revealed that protein neddylation is upregulated in numerous human cancers, including liver cancer, and inhibition of neddylation exhibits anticancer potential (9).

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MLN4924, also known as pevonedistat, is a specific inhibitor of NAE. MLN4924 suppresses the entire protein neddylation modification (10). Results of a previous study demonstrated that MLN4924 exhibits suppressive activity against a variety of human cancer cells. Moreover, MLN4924 has been advanced into several Phase II/III clinical trials against solid tumors and hematologic malignancies (11-14). Mechanistic studies that focus on the role of MLN4924 in cancer suppression revealed that MLN4924 effectively induced the DNA damage response, nonhomologous end-joining repair, DNA re-replication stress and oxidative stress at the biochemical level, and induced cell cycle arrest, apoptosis, autophagy and senescence at the cellular level (15). However, results of a previous study demonstrated that the anticancer effects of MLN4924, particularly in solid tumors, remain unsatisfactory due to drug resistance (16). This factor limits the clinical use of MLN4924; thus, exploring the mechanisms of drug resistance of cancer cells to MLN4924 may help overcome chemoresistance and improve the associated clinical outcomes.

Chemoresistance is a major obstacle to curing cancer. Previous research has demonstrated that inflammation is an important factor that attenuates the chemosensitivity of cancer cells to anticancer agents. For example, doxorubicin activates the NF- κ B signaling pathway, resulting in transactivation of inflammatory factors and potent anti-apoptosis genes, which leads to the chemoresistance of cancer cells (17). In addition, multiple previous studies have demonstrated that the NF- κ B signaling pathway is aberrantly activated in multiple cancer types, such as HCC (18), cervical cancer (19) and lung cancer (20), providing a rationale for overcoming chemoresistance by targeting the NF- κ B pathway. Notably, NF- κ B inhibitor α (I κ B α), the upstream molecule of NF- κ B, plays an important role by inhibiting NF- κ B activity (21). In the resting state, I κ B α combines with the NF- κ B dimer to retain its existence in the cytoplasm. On the other hand, I κ B α is triggered by diverse extracellular signals in the activating state, such as lipopolysaccharide, tumor necrosis factor (TNF) and growth factors. Subsequently, I κ B α is phosphorylated by the I κ B kinase complex and recognized by an E3 ubiquitin ligase for ubiquitination-dependent degradation (21). Moreover, I κ B α is considered a tumor-suppressing factor in different types of cancer, such as breast (22), ovarian (23), gastric (24), colorectal (25) and liver cancer (26), mainly due to its inhibition of NF- κ B activity. However, it remains unclear whether the levels of I κ B α can be regulated by MLN4924 in liver cancer cells.

Results of the present study demonstrated that downregulation of I κ B α and the subsequent inflammation in response to MLN4924 limits the antitumor potential of MLN4924 in liver cancer cells. Mechanistic studies revealed that MLN4924 stabilized β -transducin repeat-containing protein (β -TrCP), promoting the K48 linkage of ubiquitin to the I κ B α protein, leading to the degradation of I κ B α via proteasomes. In addition, β -TrCP knockdown sensitized liver cancer cells to MLN4924. Collectively, the results of the present study revealed that the β -TrCP/I κ B α /inflammation axis limited the sensitivity of liver cancer cells to neddylation inhibition, suggesting that interference of this pathway may act as a novel target for developing new sensitizing strategies of MLN4924 to liver cancer treatment.

Materials and methods

Reagents. MLN4924 and MG132 (a proteasome inhibitor) were purchased from Selleck Chemicals. Opti-MEM was purchased from Invitrogen (Thermo Fisher Scientific, Inc.). Cycloheximide (CHX) was purchased from Chengdu XiYa Chemical Technology Co., Ltd. RNAiso Plus reagent, SYBR Green Mix and Reverse Transcription kit were purchased from Takara Bio, Inc. X-tremeGENE HP DNA Transfection Reagent, X-tremeGENE siRNA Transfection Reagent and Cocktail were purchased from Roche Diagnostics, GmbH. Cell Counting Kit-8 (CCK-8) was purchased from Dojindo Laboratories, Inc. Small interfering (si)RNAs targeting β -TrCP and control siRNA were synthesized by Shanghai GenePharma Co., Ltd.

Cell lines and culture. The human liver cancer cell line LM3 was purchased from BeNa Culture Collection (cat. no. BNCC342335), the HepG2 cell line (cat. no. HB-8065) was obtained from the American Type Culture Collection and the Huh7 cell line (cat. no. SCSP-526) was purchased from the National Collection of Authenticated Cell Cultures of the Chinese Academy of Sciences (<https://www.cellbank.org.cn/>). All cells were cultured in DMEM containing 10% fetal bovine serum at 37°C in a 5% CO₂ incubator.

Plasmid construction. The DNA fragment encoding I κ B α (NM_020529) was synthesized by Shanghai GeneChem Co., Ltd., and inserted into the vector pcDNA-3.1. The resulting plasmid was named pcDNA-I κ B α or overexpression (OE)-I κ B α . The His-ub (His-ubiquitin) plasmids and the corresponding mutant plasmids were constructed by Tsingke Biotechnology Co., Ltd.

Reverse transcription-quantitative (RT-q)PCR. Total RNA was isolated from cultured cells using RNAiso Plus reagent, and the first-strand cDNA was synthesized using the PrimeScript RT Reagent Kit with gDNA Eraser (Takara Bio, Inc.) according to the manufacturer's instructions. qPCR was performed using a One-Step RT-PCR kit (ComWin) and SYBR Green Mix. The PCR cycle parameters were as follows: 95°C for 30 sec, and 40 cycles at 95°C for 5 sec and 60°C for 1 min. Results were calculated based on the quantification cycle (C_q), and the relative fold change was determined using the 2^{- $\Delta\Delta$ C_q} method (27). Primers used for RT-qPCR are listed as follows: IL-6-forward (F), 5'-TACCCAGGAGAAGATTCC-3' and IL-6-reverse (R), 5'-TTTTCTGCCAGTGCCTCTTT-3'; IL-8-F, 5'-CGGAAGGAACCATCTCACTGTG-3' and IL-8-R, 5'-AGAAATCAGGAAGGCTGCCAAG-3'; TNF- α -F, 5'-AACCTCTCTGCCATCAA-3' and TNF- α -R, 5'-CCAAGTAGCCTGCCAGA-3'; β -TrCP-F, 5'-CAGTCTCTGCCTGCGTTTC-3' and β -TrCP-R, 5'-CTCACTACCAGCTGTCCCT-3'; I κ B α -F, 5'-AAGTGTATCCGCCAGGTGAG-3' and I κ B α -R, 5'-CTGCTCACAGGCAAGGTGTA-3'; and β -actin-F, 5'-GTGAAGGTGACAGCAGTCGGTT-3' and β -actin-R, 5'-GAAGTGGGGTGGCTTTTAGGA-3'.

Transient transfection. Liver cancer cells were seeded and cultured in 6-well plates overnight at 37°C in an incubator, and small interfering RNAs (siRNAs, 100 pmol) or plasmids (2 μ g)

were transfected into liver cancer cells using X-tremeGENE siRNA Transfection Reagent or X-tremeGENE HP DNA Transfection Reagent for 48 h in 37°C incubator, respectively, according to the manufacturer's instructions. The sequences of siRNAs are listed as follows: si- β -TrCP (100 pmol), 5'-AAG UGGAAUUUGUGGAACAUC-3' and si-negative control (si-NC; 100 pmol), 5'-UUCUCCGAACGUGUCACGUTT-3'.

MG132 treatment protocol. Following treatment with MLN4924 for 48 h, LM3 and Huh7 cells were incubated with or without 20 μ M MG132 for 5 h at 37°C. Subsequently, western blotting was employed to determine the protein levels of I κ B α .

Western blot analysis. Cells were lysed with RIPA lysis buffer (Beyotime Institute of Biotechnology) supplemented with a protease inhibitor mixture (Roche Diagnostics), and then the lysates were centrifuged at 16,000 \times g for 15 min at 4°C. The protein concentrations were then detected using a BCA kit (Beyotime Institute of Biotechnology), according to the manufacturer's instructions. Following denaturing, samples (100 μ g/lane) were separated by 10% SDS-PAGE (Beyotime Institute of Biotechnology) and then transferred to NC membranes (Cytiva). The NC membranes were then blocked in 5% nonfat milk in Tris-buffered saline containing 0.1% Tween-20 (TBST) for 1 h at room temperature and then incubated with the indicated primary antibodies. Primary antibodies against ubiquitin (cat. no. sc-271289; 1:1,000 dilution), β -TrCP (cat. no. sc-390629; 1:1,000 dilution) and I κ B α (cat. no. sc-373893; 1:1,000 dilution) were purchased from Santa Cruz Biotechnology, Inc. Anti-cleaved caspase-3 was purchased from Cell Signaling Technology (product no. 9664; 1:1,000 dilution). Primary antibodies against GAPDH (cat. no. AG019; 1:5,000 dilution) and tubulin (cat. no. AT819; 1:5,000 dilution) were purchased from Beyotime Institute of Biotechnology. All primary antibodies were incubated at 4°C overnight. Secondary antibodies, HRP-labeled goat anti-rabbit (cat. no. 7074S) and anti-mouse (cat. no. 7076S) immunoglobulin G, were purchased from Cell Signaling Technology, Inc. The secondary antibodies were diluted in 5% skim milk at a ratio of 1:5,000 and incubated at room temperature for 1 h, and then the protein signals were measured using chemiluminescence detection reagent (cat. no. WBKLS0500; Millipore; Merck KGaA) and the Bio-Rad Imaging System (Image Lab 4.1; Bio-Rad Laboratories, Inc.).

Immunoprecipitation (IP). Liver cancer cells were lysed using IP lysis buffer purchased from Beyotime Institute of Biotechnology, and then the lysates were centrifuged at 10,000 \times g for 10 min at 4°C. Subsequently, the supernatant was collected into another EP tube. The anti-I κ B α antibody (cat. no. sc-373893) was then added to the lysate in a ratio of 2:1 mg total protein. Following overnight incubation at 4°C, 40 ml protein A/G agarose beads (Santa Cruz Biotechnology, Inc.) were added to each sample and incubated at room temperature for 2 h. The immunoprecipitates were washed five times with ice-cold PBS and isolated by centrifugation (600 \times g, 3 min, 4°C). Finally, the precipitates were mixed with 1X SDS-PAGE loading buffer and boiled at 100°C for 10 min, and then assessed by western blot analysis.

In vitro ubiquitination assay. To examine the ubiquitination of I κ B α by MLN4924, liver cancer cells were co-transfected with His-ub (His-ubiquitin) and HA-I κ B α , and subsequently incubated with MLN4924 (0.4 μ M) for 48 h at 37°C. MG132 (20 μ M) was added to the medium for 5 h, cells were harvested and subsequently split into two aliquots; one for direct western blotting and one for *in vitro* ubiquitination. Briefly, cells were lysed in buffer A [6 mol/l guanidinium-HCl, 0.1 mol/l Na₂HPO₄/NaH₂PO₄, 10 mmol/l Tris-HCl (pH 8.0), 5 mmol/l imidazole and 10 mmol/l β -mercaptoethanol] on ice, sonicated to reduce viscosity and mixed with 50 μ l Ni²⁺-NTA-agarose beads (Qiagen, Inc.) overnight. Beads were successively washed with buffer A with 10 mM β -mercaptoethanol, buffer B [8 mM urea, 0.1 M Na₂HPO₄/NaH₂PO₄, 10 mM Tris/HCl (pH, 8.0), and 10 mM β -mercaptoethanol], buffer C [8 mM urea, 0.1 M Na₂HPO₄/NaH₂PO₄, 10 mM Tris/HCl (pH, 6.3), 10 mM β -mercaptoethanol and 0.2% Triton X-100], and buffer C with 10 mM β -mercaptoethanol and 0.1% Triton X-100. Subsequently, His6-tagged ubiquitinated proteins were eluted with buffer D [200 mM imidazole, 0.15 M Tris-HCl (pH, 6.7), 30% glycerol, 0.72 M β -mercaptoethanol and 5% SDS]. The polyubiquitination of I κ B α was detected using western blot analysis, using an anti-HA antibody (cat. no. 11867423001, 1:5,000; Roche Diagnostics).

Cell proliferation assays. Liver cancer cells were seeded in 96-well plates at 3,000 cells per well overnight at 37°C, and transiently transfected with the indicated plasmids or siRNAs, and then incubated with MLN4924 for 48 h. Subsequently, the CCK-8 agent was added to each well at a volume of 10 μ l per well, and then incubated at 37°C for 30-60 min, followed by detection of the absorption value at 450 nm with a microplate reader (Thermo Scientific, Inc.).

Colony formation assays. Liver cancer cells were seeded in 6-well plates at 500 cells per well overnight at 37°C, and transiently transfected with the indicated plasmids or siRNAs or incubated with MLN4924 (0.4 μ M) for 10 days. Subsequently, the cells were fixed with 4% formaldehyde (15 min at room temperature) and stained with 0.1% crystal violet at room temperature for 10 min. During colony growth, the culture medium was replaced every 3 days. Colonies with >50 cells were counted manually, 10-14 days after plating.

Bioinformatics analysis. UALCAN and GEPIA online databases were utilized to determine the mRNA level of I κ B α (NFKBIA). Briefly, NFKBIA was selected after logging in the website (UALCAN, <http://ualcan.path.uab.edu/analysis.html>). Subsequently, liver hepatocellular carcinoma (LIHC) was selected as the subject of analysis. Thereafter, the button 'Explore' was selected and the result for the expression of NFKBIA in LIHC based on sample tapes was obtained. For the usage of GEPIA online database (Log2FCI Cutoff: 1; P-value Cutoff: 0.01), NFKBIA was typed into the box under 'Enter gene/isoform name' after logging in the website (GEPIA, <http://gepia2.cancer-pku.cn/#index>), and then the 'Boxplots' button was selected followed by LIHC. Thereafter, the button 'Plot' was selected and the result of the mRNA level of NFKBIA in 160 normal liver tissues and 369 liver cancer tissues was obtained. For the analysis of the protein level of

I κ B α in different human cancers and corresponding normal tissues, Clinical Proteomic Tumor Analysis Consortium (CPTAC, <http://ualcan.path.uab.edu/analysis-prot.html>) online database was employed. NFKBIA was typed into the box under 'Enter gene names' column after logging in the website, and then LIHC was selected, followed by the analysis of the protein level of NFKBIA between normal and liver cancer tissues. For the analysis of normal and primary cancer of pan-cancer, the 'Pan-cancer view' link was selected. Z-values represent standard deviations from the median across samples for the given cancer type in analyzing the protein level of I κ B α .

Statistical analysis. GraphPad Prism 8.0 (GraphPad Software, Inc.) was used to analyze data. Data are presented as the mean \pm standard deviation. Comparisons between two groups were performed using a two-tailed unpaired Student's t-test. Comparisons between multiple groups were performed using one-way ANOVA, followed by Tukey's post hoc test. The data are presented as the mean \pm SD of at least three independent experiments. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

MLN4924 suppresses the survival of liver cancer cells and downregulates I κ B α . A previous study demonstrated that MLN4924 exhibits antitumor potential (28). To confirm this phenomenon, the effects of MLN4924 on the growth of liver cancer cells were investigated in the present study. As revealed in Fig. 1A, 0.8 μ M MLN4924 increased the natant cells compared with the control group, while 0.4 μ M MLN4924 exhibited no notable effect on cell morphology (Fig. 1A). Therefore, this concentration (0.4 μ M) of MLN4924 was selected to investigate the chemoresistance of liver cancer cells in the subsequent experiments. Moreover, the results of the CCK-8 assay demonstrated that MLN4924 suppressed the growth of liver cancer cells in a dose-dependent manner. However, the half maximal inhibitory concentration (IC_{50}) values of MLN4924 in LM3 and Huh7 cells were 13.01 and 12.36 μ M, respectively (Fig. 1B and C), indicating that MLN4924 markedly suppressed the growth of liver cancer cells at high concentrations. As I κ B α is considered a tumor-suppressing factor in various cancers due to its inhibition of NF- κ B activity, the expression levels of I κ B α in cancer tissues were further investigated using bioinformatics analysis. The results obtained from the UALCAN database demonstrated that the I κ B α mRNA expression levels exhibited no significant difference between 50 healthy liver tissues and 371 LIHC tissues (Fig. S1A). These results were further confirmed using an alternative database, GEPIA, containing 160 healthy liver tissues and 369 HCC tissues (Fig. S1B). Notably, protein expression analysis using data from the Clinical Proteomic Tumor Analysis Consortium (CPTAC) revealed that the I κ B α protein expression levels were markedly reduced in HCC (Fig. S1C), ovarian cancer, uterine corpus endometrial carcinoma and lung adenocarcinoma (Fig. S1D). These results suggested that the regulation of I κ B α during liver carcinogenesis and development may occur at a post-translational level. Subsequently, the effects of MLN4924 on I κ B α expression were explored. As shown in Fig. 1D and E, MLN4924 significantly decreased I κ B α protein

expression levels in a dose-dependent manner; however, it exerted no effect on the mRNA expression. Furthermore, it was determined whether MLN4924 inhibited the protein level of I κ B α in a time-dependent manner. As revealed in Fig. S2, the protein level of I κ B α was markedly decreased following treatment with MLN4924 for 48 and 72 h, and thus 48 h was selected in the subsequent experiments. As I κ B α plays a key role in anti-inflammation (29), mRNA expression levels of interleukin (IL)-6, IL-8 and TNF- α , downstream molecules of I κ B α , were explored in the present study. As shown in Fig. 1F and G, MLN4924 significantly increased the mRNA expression levels of IL-6, IL-8 and TNF- α . Collectively, these findings indicated that MLN4924 possesses an antitumor role in liver cancer cells; however, decreased I κ B α expression may impact its antitumor potential.

I κ B α overexpression sensitizes MLN4924 to inhibit liver cancer cells. To clarify whether MLN4924-induced I κ B α downregulation is involved in the insensitivity of liver cancer cells to MLN4924 treatment, LM3 and Huh7 cells were treated with MLN4924 in the presence or absence of OE-I κ B α . Firstly, the efficacy of the plasmid transfection (OE-I κ B α) was determined using qPCR and it was determined that the mRNA level of I κ B α was significantly increased after transfection with OE-I κ B α plasmid, suggesting that transfection with OE-I κ B α successfully overexpressed I κ B α in both LM3 and Huh7 cells (Fig. S3). As revealed in Fig. 2A and B, cotreatment with MLN4924 and OE-I κ B α significantly suppressed the growth of liver cancer cells compared with MLN4924 or OE-I κ B α treatment alone. Moreover, the results of the colony formation assay revealed that cotreatment with MLN4924 and OE-I κ B α significantly inhibited the number of colonies, compared with MLN4924 or OE-I κ B α treatment alone (Fig. 2C and D). These results indicated that downregulation of I κ B α is a novel resistance factor of MLN4924, and overexpression of I κ B α sensitizes MLN4924 to inhibit liver cancer cells.

MLN4924 decreases the protein stability of I κ B α . The mechanism by which MLN4924 decreases the expression of I κ B α was explored in the present study. The results of the RT-qPCR analysis revealed that I κ B α mRNA expression levels exhibited no significant change following MLN4924 treatment (Fig. 1E), suggesting that MLN4924 downregulated I κ B α at a post-translational level. Thus, I κ B α protein stability was determined following the addition of CHX in the presence or absence of MLN4924. As demonstrated in Fig. 3A-C, MLN4924 treatment markedly decreased the protein stability of I κ B α and reduced the half-life of I κ B α protein in LM3, Huh7 and HepG2 cells. The results of previous research demonstrated that proteasomes play an important role in protein degradation (30). To verify whether the proteasome is involved in MLN4924-mediated I κ B α degradation, the proteasome inhibitor MG132 was utilized. A previous study demonstrated that MG132 (20 μ M) always exerts its function to inhibit protein degradation at 4-6 h (31), thus, MG132 (20 μ M) was added and incubated with LM3 and Huh7 cells at 37°C for 5 h before harvesting. As shown in Fig. 3D and E, MG132 treatment markedly attenuated MLN4924-induced I κ B α downregulation. These results indicated that MLN4924 promoted I κ B α protein degradation via a proteasome-associated pathway.

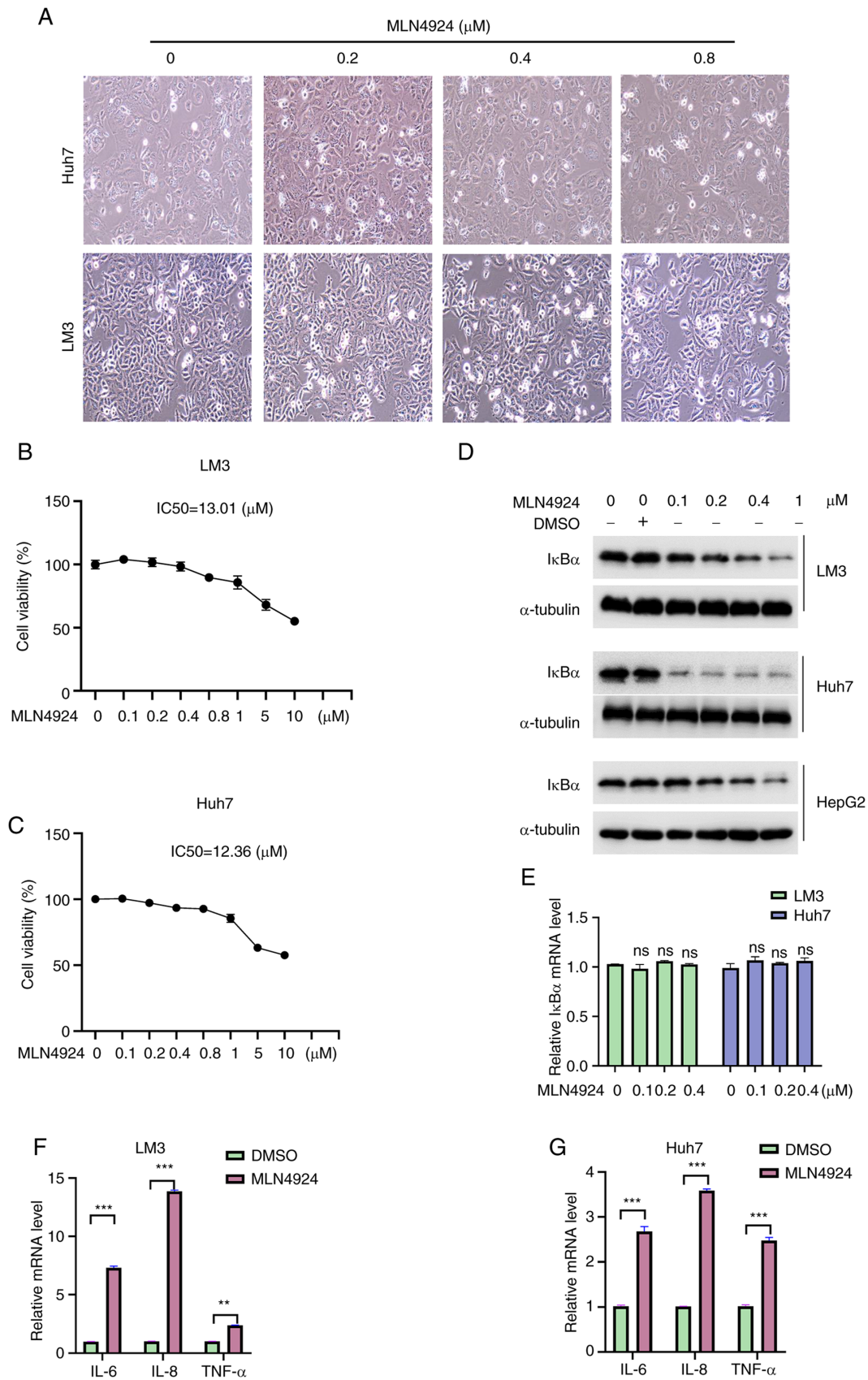


Figure 1. MLN4924 suppresses the survival of liver cancer cells and downregulates I κ B α expression. (A) LM3 and Huh7 cells were treated with the indicated concentrations of MLN4924 for 48 h, and cells were imaged under a phase-contrast microscope (magnification, x200). (B and C) LM3 and Huh7 cells were treated with the indicated concentrations of MLN4924 for 48 h, and a Cell Counting Kit-8 assay was utilized to evaluate the cytotoxicity of MLN4924. (D) Following treatment with the indicated concentrations of MLN4924 for 48 h, the protein expression levels of I κ B α were determined in LM3, Huh7 and HepG2 cells using western blot analysis. α -Tubulin was used as a loading control. (E) LM3 and Huh7 cells were treated as previously described, and the mRNA expression levels of I κ B α were determined using RT-qPCR. (F and G) LM3 and Huh7 cells were treated with 0.4 μM MLN4924 for 24 h, and RT-qPCR was used to assess the mRNA expression levels of IL-6, IL-8 and TNF- α . β -Actin was used as a loading control. ** P <0.01 and *** P <0.001. I κ B α , NF- κ B inhibitor α ; RT-qPCR, reverse transcription-quantitative PCR; IL, interleukin; TNF, tumor necrosis factor; DMSO, dimethyl sulfoxide.

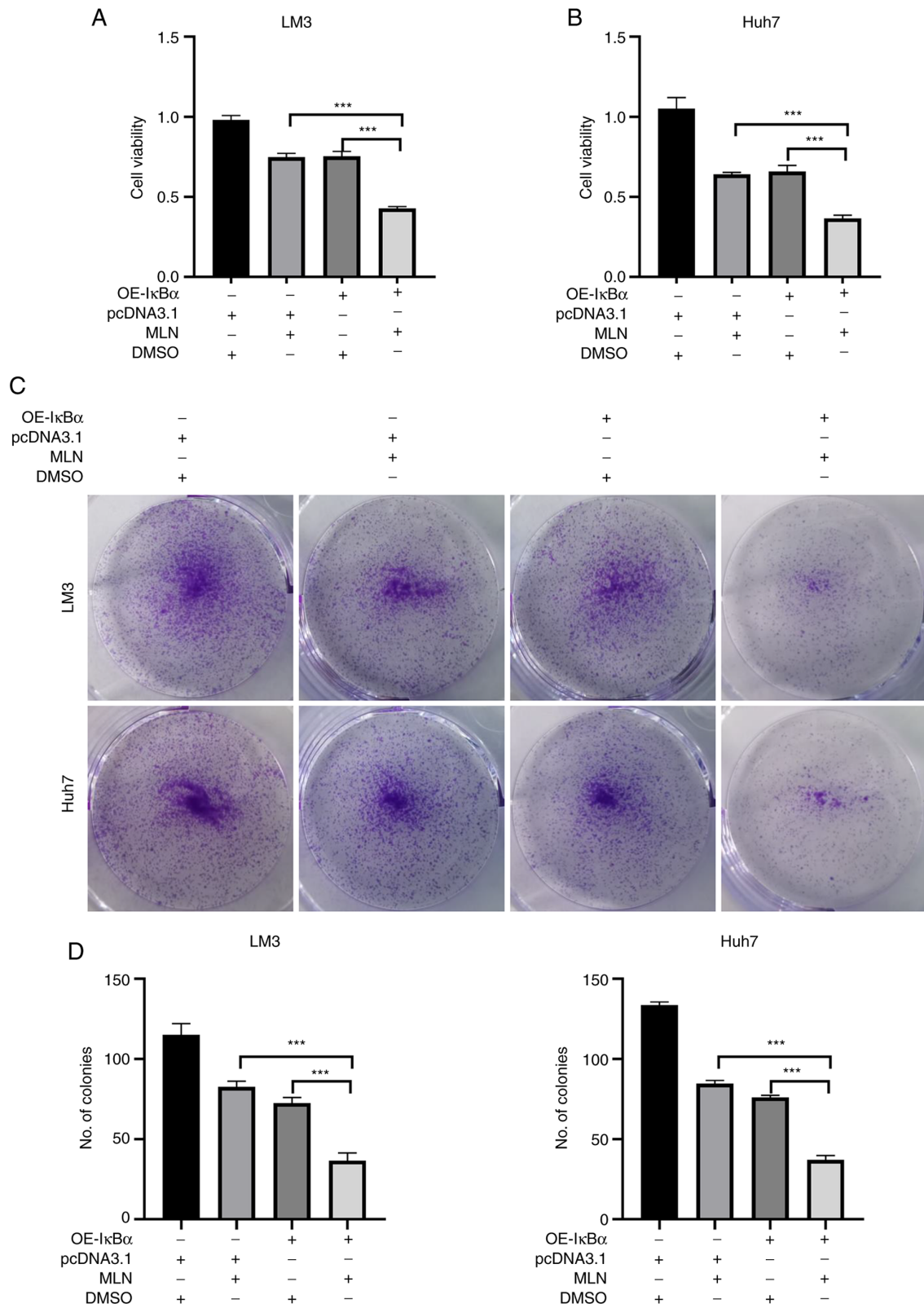


Figure 2. Overexpression of I κ B α sensitizes MLN4924 in the treatment of liver cancer cells. (A and B) Following transfection with OE-I κ B α or pcDNA3.1 for 24 h, LM3 and Huh7 cells were treated with 0.4 μ M MLN4924 or DMSO for 48 h. A Cell Counting Kit-8 assay was utilized to determine the growth of liver cancer cells. (C and D) Following transfection with OE-I κ B α or pcDNA3.1 for 24 h, LM3 and Huh7 cells were treated with 0.4 μ M MLN4924 or DMSO for 10 days. Colonies of LM3 and Huh7 cells were stained using crystal violet and counted. ***P<0.001. I κ B α , NF- κ B inhibitor α ; OE, overexpression; DMSO, dimethyl sulfoxide; MLN, MLN4924.

MLN4924 promotes the K48 linkage of ubiquitin to I κ B α protein. A previous study demonstrated that the ubiquitin-proteasome pathway is key in controlling the degradation of

proteins in eukaryocytes (30). Thus, the role of MLN4924 in I κ B α protein degradation via the ubiquitin-proteasome pathway was explored. As revealed in Fig. 4A and B,

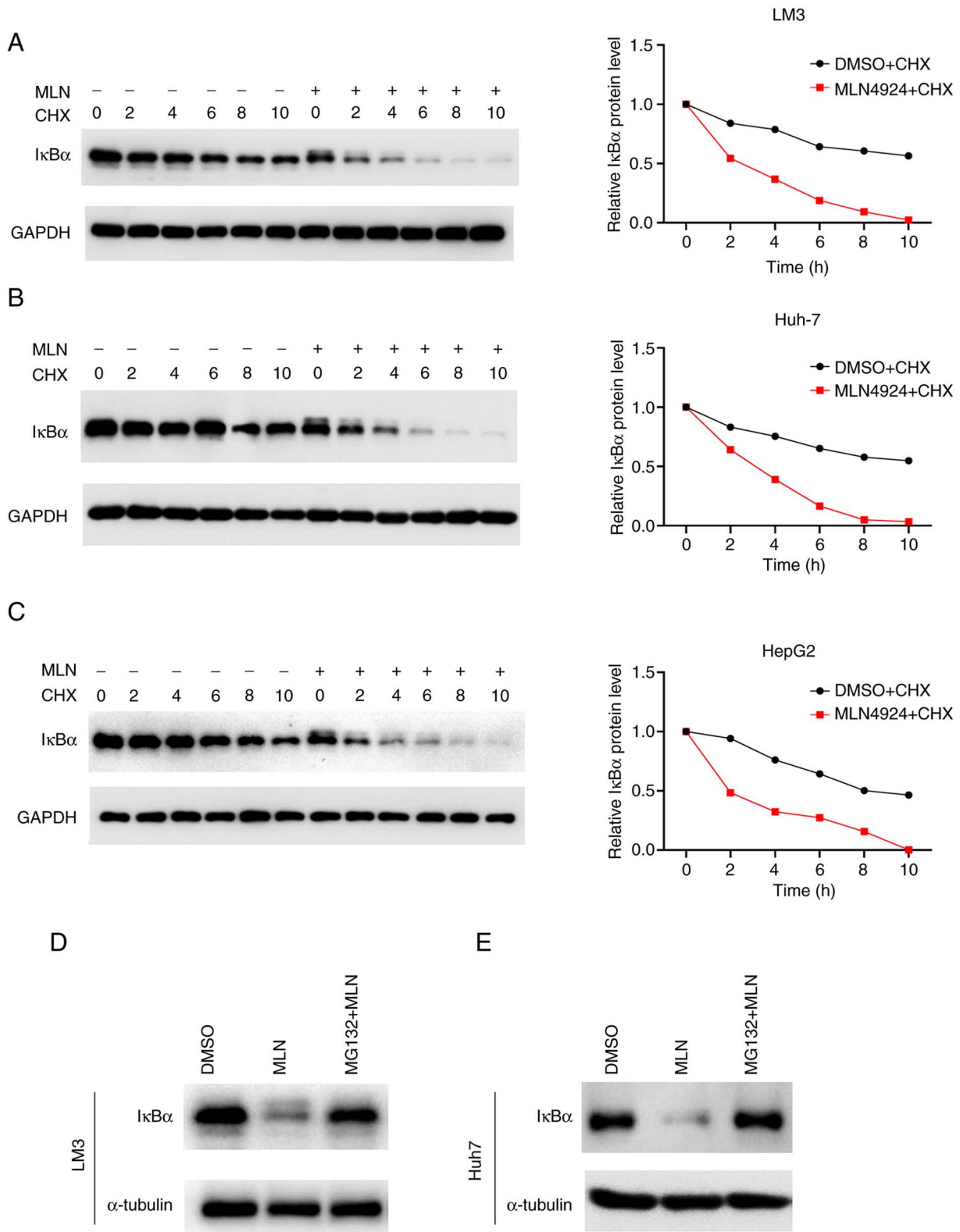


Figure 3. MLN4924 decreases the protein stability of IκBα. (A-C) Following treatment with 0.4 μM MLN4924 or DMSO for 48 h, LM3, Huh7 and HepG2 cells were incubated with 10 μg/ml cycloheximide for the indicated time-points. Protein expression levels of IκBα were determined using western blot analysis and quantified using Image Lab. (D and E) Following treatment with 0.4 μM MLN4924 or DMSO for 48 h, LM3 and Huh7 cells were incubated with or without 20 μM MG132 for 5 h at 37°C. Western blotting was carried out to determine the protein expression levels of IκBα. IκBα, NF-κB inhibitor α; DMSO, dimethyl sulfoxide; CHX, cycloheximide; MLN, MLN4924.

MLN4924 markedly enhanced the ubiquitination of endogenous IκBα, and this was further confirmed using exogenous overexpression of IκBα (Fig. 4C and D). These results

indicated that MLN4924 promoted the polyubiquitination of IκBα protein. To determine the type of ubiquitin chains involved in MLN4924-mediated IκBα polyubiquitination, six

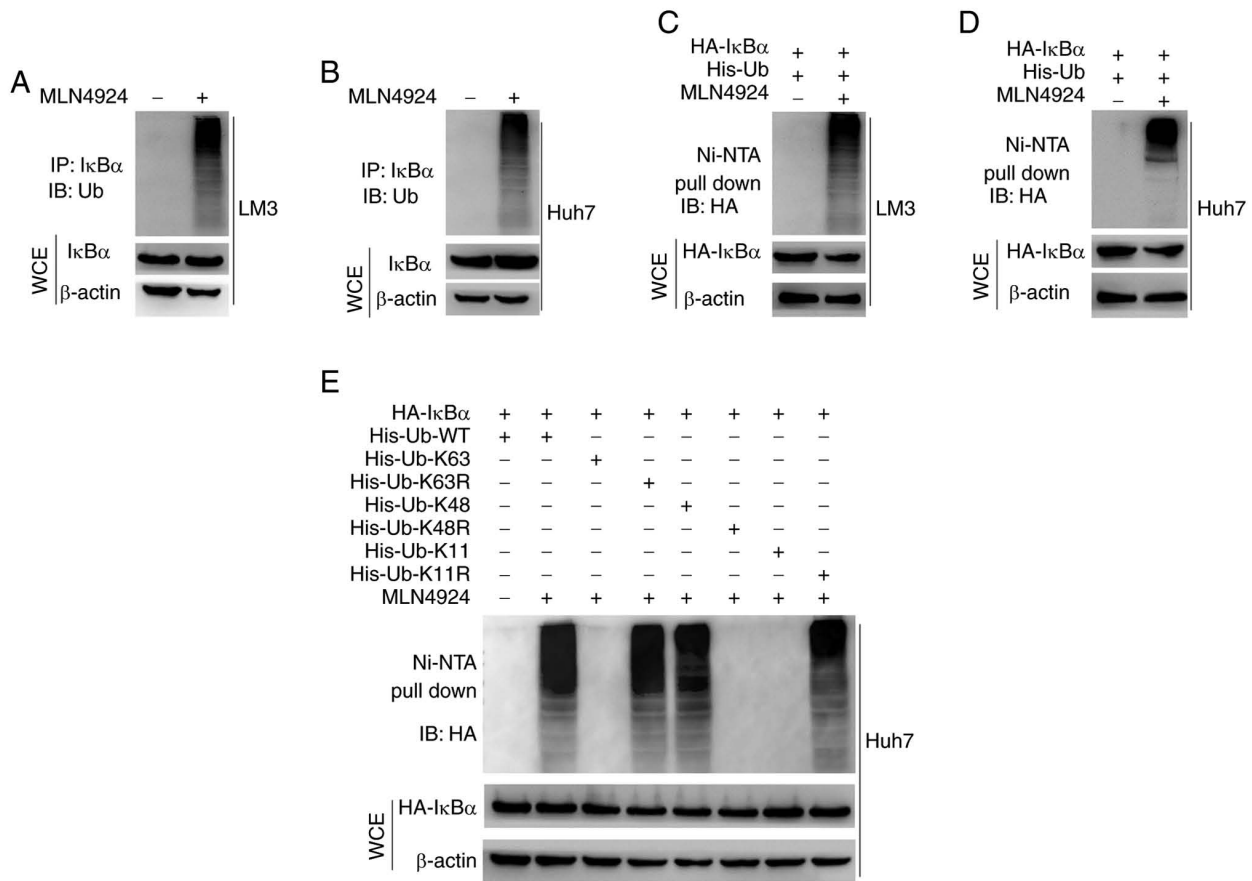


Figure 4. MLN4924 promotes the K48 linkage of ubiquitin to IκBα protein. (A and B) LM3 and Huh7 cells were treated with 0.4 μM MLN4924 or DMSO for 48 h, and 20 μM MG132 was subsequently added for 5 h at 37°C. IκBα protein was immunoprecipitated using anti-IκBα. Moreover, ubiquitination of IκBα in the immunoprecipitated fraction was assessed by western blot analysis, using anti-ubiquitin. (C and D) Following co-transfection with HA-IκBα and His-ub plasmids for 24 h, LM3 and Huh7 cells were treated with 0.4 μM MLN4924 or DMSO for 24 h, and 20 μM MG132 was subsequently added for 5 h at 37°C. Ni-NTA beads were used to pull down the His-ub tagged proteins. Ubiquitination of HA-IκBα in the pull-down fraction was assessed by western blot analysis, using anti-HA. Whole cell lysates were subjected to western blot analysis. (E) Huh7 cells were transfected with the indicated plasmids for 48 h, incubated with 20 μM MG132 for 5 h at 37°C, lysed with 6 M guanidine solution, pulled down with Ni-NTA beads and western blot analysis was carried out to determine the levels of HA-IκBα. The whole cell lysates were subjected to western blot analysis. IκBα, NF-κB inhibitor α; DMSO, dimethyl sulfoxide; His-ubiquitin, His-ub; WCE, whole cell extract.

ubiquitin mutants (K63, K63R, K48, K48R, K11 and K11R) were used along with the wild-type (WT) ubiquitin. Results of the *in vitro* ubiquitination assay indicated that K63 and K11 ubiquitin mutants attenuated the formation of the polyubiquitin chain to IκBα, compared with the K48 ubiquitin mutant and WT ubiquitin. Moreover, the results of the present study demonstrated that the K48R ubiquitin mutant significantly attenuated the formation of the polyubiquitin chain to IκBα, compared with WT, and the K63R and K11R mutant forms of ubiquitin (Fig. 4E). Collectively, these results indicated that MLN4924 promoted the K48-linked ubiquitination of IκBα.

MLN4924 promotes the degradation of IκBα protein via stabilizing β-TrCP. The mechanism underlying MLN4924-induced degradation of IκBα was further explored in the present study. The results of a previous study demonstrated that β-TrCP, also named FBXW1, is an important E3 ligase for the ubiquitin-proteasome pathway-dependent degradation of IκBα (32). Thus, the role of MLN4924 in IκBα degradation via β-TrCP was explored. The results of the western blot analysis demonstrated that MLN4924 markedly enhanced the protein expression levels of β-TrCP in LM3,

Huh7 and HepG2 cells compared with the controls (Fig. 5A). In addition, LM3 and Huh7 cells were transfected with si-β-TrCP or si-NC in the presence or absence of MLN4924. As shown in Fig. 5B and C, β-TrCP knockdown markedly reduced MLN4924-mediated IκBα downregulation. These results indicated that MLN4924 promoted the degradation of IκBα via upregulation of β-TrCP. Moreover, the protein stability of β-TrCP was evaluated by adding translation inhibitor CHX in the presence or absence of MLN4924. The results of the present study demonstrated that MLN4924 significantly enhanced the protein stability of β-TrCP and prolonged the half-life of β-TrCP in LM3, Huh7 and HepG2 cells (Fig. 5D-F). Collectively, these results indicated that MLN4924 induces the degradation of IκBα via stabilizing β-TrCP.

β-TrCP knockdown enhances the antitumor potential of MLN4924 in liver cancer cells via inhibiting inflammation. To clarify whether knockdown of β-TrCP enhanced the antitumor potential of MLN4924 in liver cancer cells via inhibiting inflammation, LM3 and Huh7 cells were treated with MLN4924 in the presence or absence of si-β-TrCP. As revealed in Fig. 6A and B, transfection with si-β-TrCP significantly reduced

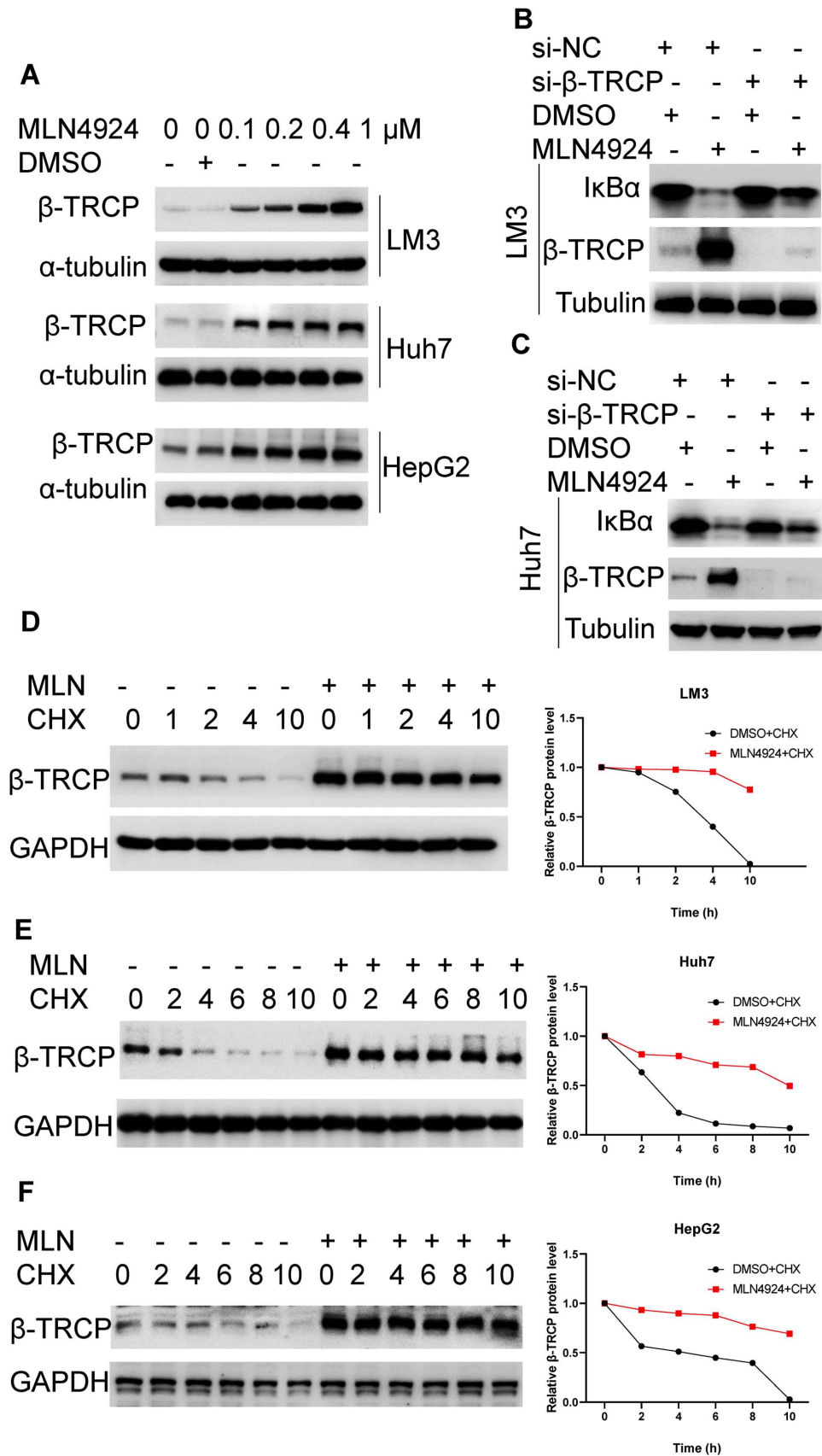


Figure 5. MLN4924 promotes the degradation of $\text{I}\kappa\text{B}\alpha$ protein via stabilizing β -TrCP. (A) Following treatment with the indicated concentrations of MLN4924 for 48 h, the protein expression levels of β -TrCP were determined in LM3, Huh7 and HepG2 cells using western blot analysis. (B and C) Following transfection with si- β -TrCP or si-NC for 24 h, LM3 and Huh7 cells were treated with 0.4 μ M MLN4924 or DMSO for 24 h. Subsequently, the protein expression levels of $\text{I}\kappa\text{B}\alpha$ and β -TrCP were determined using western blot analysis. (D-F) Following treatment with 0.4 μ M MLN4924 or DMSO for 48 h, LM3, Huh7 and HepG2 cells were incubated with 10 μ g/ml cycloheximide for the indicated time-points. Subsequently, the protein expression levels of β -TrCP were determined using western blot analysis and quantified using Image Lab. $\text{I}\kappa\text{B}\alpha$, NF- κ B inhibitor α ; β -TrCP, β -transducin repeat-containing protein; siRNA, small interfering RNA; si-NC, negative control siRNA; DMSO, dimethyl sulfoxide; CHX, cycloheximide; MLN, MLN4924.

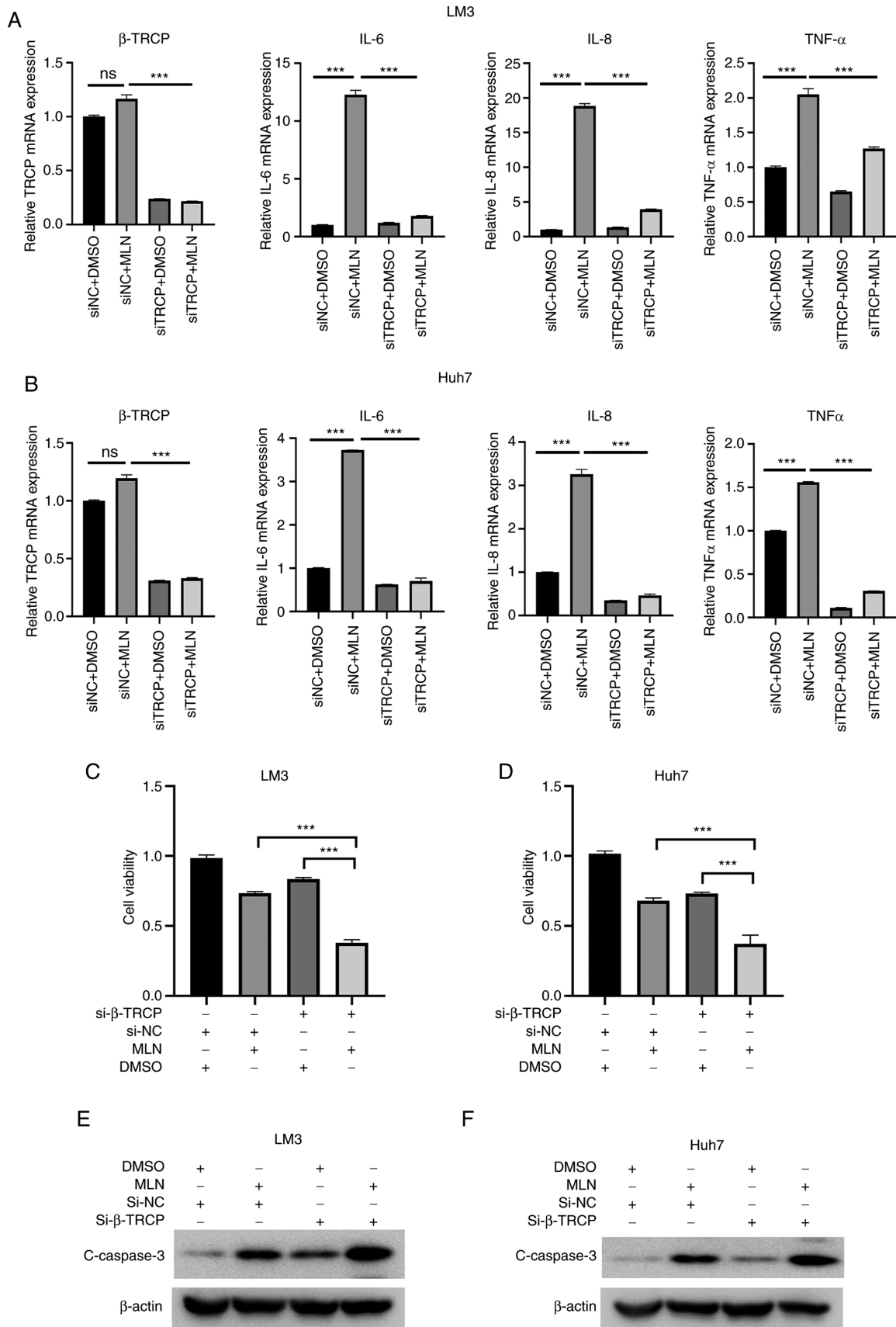


Figure 6. β -TrCP knockdown enhances the antitumor potential of MLN4924 in liver cancer cells via inhibiting inflammation. (A and B) Following transfection with si- β -TrCP or si-NC for 24 h, LM3 and Huh7 cells were treated with 0.4 μ M MLN4924 or DMSO for 24 h. Subsequently, the mRNA expression levels of IL-6, β -TrCP, IL-8 and TNF- α were assessed using reverse transcription-quantitative PCR. (C and D) Following transfection with si- β -TrCP or si-NC for 24 h, LM3 and Huh7 cells were treated with 0.4 μ M MLN4924 or DMSO for 48 h. A Cell Counting Kit-8 assay was used to determine the growth of liver cancer cells. (E and F) LM3 and Huh7 cells were treated as previously described, and the protein expression levels of caspase-3 were determined using western blot analysis. *** P <0.001. β -TrCP, β -transducin repeat-containing protein; siRNA, small interfering RNA; si-NC, negative control siRNA; DMSO, dimethyl sulfoxide; IL, interleukin; TNF, tumor necrosis factor; ns, not significant.

the mRNA expression of β -TrCP. Moreover, MLN4924 treatment notably enhanced the mRNA expression levels of IL-6, IL-8 and TNF- α , which was significantly attenuated following β -TrCP knockdown in LM3 and Huh7 cells. Moreover, the results of the CCK-8 assay demonstrated that cotreatment with si- β -TrCP and MLN4924 significantly suppressed the growth of liver cancer cells, compared with MLN4924 or si- β -TrCP treatment alone (Fig. 6C and D). In addition, the results of the western blot analysis demonstrated that β -TrCP knockdown markedly enhanced MLN4924-induced caspase-3 upregulation compared with the controls (Fig. 6E and F). Collectively, these results demonstrated that β -TrCP knockdown enhanced the antitumor potential of MLN4924 in liver cancer cells via inhibiting inflammation.

Discussion

The results of a previous study (28) demonstrated that MLN4924 exhibits high suppressive activity against a variety of human cancer cells. However, further investigations into the specific mechanisms underlying MLN4924 as an anticancer treatment are required. To the best of our knowledge, the results of the present study were the first to demonstrate that downregulation of I κ B α and subsequent inflammation is key for attenuating the antitumor potential of MLN4924 in liver cancer cells. The mechanistic study of MLN4924-induced I κ B α downregulation revealed that MLN4924 stabilized β -TrCP, promoting the ubiquitination and degradation of I κ B α ; thus, enhancing the degradation of I κ B α and subsequent inflammation. This therefore attenuated the antitumor potential of MLN4924 in liver cancer cells. Moreover, the results of the present study demonstrated that β -TrCP knockdown enhanced the antitumor potential of MLN4924 in liver cancer cells via inhibiting inflammation. Collectively, the afore mentioned results may provide a novel potential strategy for sensitizing MLN4924 in the treatment of liver cancer.

The results of previous studies have demonstrated that inflammation could lead to chemoresistance of cancer cells (17,33). Zhong *et al* revealed that the NF- κ B-IL6-STAT3 axis activated by intratumoral LPS promoted the expression of cyclin D1, c-Myc, Bcl-2 and survivin, facilitating prostate cancer proliferation and docetaxel chemoresistance (33). Neddylation-inhibitor MLN4924 exhibits broad anticancer potential and may be used as a chemotherapeutic agent in the future. However, it remains to be fully elucidated whether inflammation is involved in the chemoresistance of liver cancer cells in response to MLN4924 treatment. The results of the present study demonstrated that MLN4924 promoted I κ B α degradation and increased the expression of inflammatory factors (IL-6, IL-8 and TNF- α), leading to the chemoresistance of liver cancer cells to MLN4924. Notably, the results of a previous study demonstrated that MLN4924 inhibited activation of the NF- κ B pathway via inhibiting CRL-mediated I κ B α degradation. However, the results of this study did not indicate that treatment with MLN4924 decreased the protein expression levels of I κ B α (34). Moreover, I κ B α knockdown significantly increased the MLN4924-mediated proliferation inhibition in esophageal cancer cells (34). Therefore, the contrasting results of the two studies may be ascribed to the different cancer cell lines, as

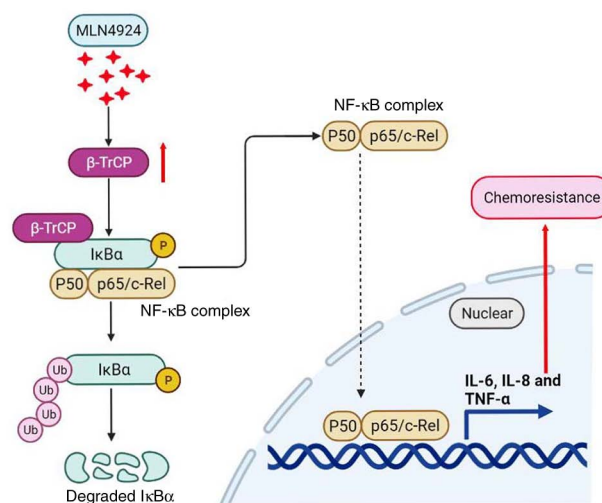


Figure 7. Schematic model of the mechanism by which activation of the β -TrCP/I κ B α /inflammation axis limits the sensitivity of liver cancer cells to neddylation inhibition. MLN4924 treatment leads to the upregulation of β -TrCP stability, promoting the ubiquitination and degradation of I κ B α protein, derepressing the NF- κ B complex and promoting its nuclear location, enhancing the transcription of inflammatory cytokines including IL-6, IL-8 and TNF- α , which attenuates the sensitivity of liver cancer cells to neddylation inhibition. β -TrCP, β -transducin repeat-containing protein; I κ B α , NF- κ B inhibitor α ; IL, interleukin; TNF, tumor necrosis factor.

liver cancer is one of several cancers that is often associated with chronic inflammation.

MLN4924, a neddylation inhibitor, suppresses CRLs via inhibiting NAE-mediated cullin neddylation (28). Notably, the results of the present study demonstrated that MLN4924 promoted the proteasome-dependent degradation of I κ B α and shortened the half-life of I κ B α , following treatment with MG132 and translation inhibitor CHX. Moreover, the K48-linkage of ubiquitin to I κ B α was significantly increased following treatment with MLN4924, which reflects that some E3 ligases may be involved in MLN4924-mediated I κ B α degradation. The expression of β -TrCP, a well-established E3 ligase for I κ B α degradation (35), was markedly increased following MLN4924 treatment in liver cancer cells. However, the molecular mechanism by which MLN4924 stabilizes β -TrCP remains unclear. Further studies are warranted to clarify which protein is targeted by MLN4924 directly and participates in the regulation of β -TrCP stability. Moreover, in the present study it was determined that MLN4924 promoted I κ B α degradation via stabilizing β -TrCP, despite a previous study which reported that β -TrCP is inhibited by MLN4924 (36). Therefore, it was hypothesized that increase of β -TrCP may be a resistance response of liver cancer cells when treated with neddylation inhibitor MLN4924.

β -TrCP acts as a substrate receptor and constitutes an active SCF $^{\beta$ -TrCP ligase with a scaffold protein cullin 1 (CUL1), a RING protein RBX1 and an adaptor protein SKP1 (37). β -TrCP plays a critical role in the regulation of various physiological and pathological processes, including signal transduction, cell cycle progression, cell migration, DNA damage response and tumorigenesis, by governing large amounts of key regulators for ubiquitination and proteasomal degradation (38). The results of previous studies have demonstrated that β -TrCP plays a vital role in carcinogenesis via promoting the degradation of its

substrates, including oncoproteins and tumor suppressors (38). Specifically, the results of previous studies demonstrated that β -TrCP is upregulated in liver cancer tissues (39,40). Previous research has also demonstrated that β -TrCP promotes HCC progression and metastasis via degrading leucine zipper tumor suppressor 2 (LZTS2) in HCC cells (41). These results indicated that β -TrCP acts as an oncoprotein in liver cancer, which provides clear rationale for targeting β -TrCP to overcome chemoresistance. The results of the present study demonstrated that β -TrCP was increased following MLN4924 treatment, and β -TrCP knockdown enhanced the antitumor potential of MLN4924 in liver cancer cells. Moreover, MLN4924 enhanced the protein expression levels of β -TrCP via inhibiting its degradation and prolonging its half-life. However, the mechanism by which MLN4924 stabilizes β -TrCP remains to be fully elucidated. Previous investigations demonstrated that β -TrCP1 is ubiquitinated and degraded by SCF^{SKP2} ubiquitin ligase, SAG-CUL5 ubiquitin ligase with UBC10 and UBE2S E2s, as well as the SMURF2:UBCH5 complex (38). Moreover, members of the de-ubiquitinating enzyme (DUB) family, such as USP24 (42) and USP47 (43,44), enhance the stability of β -TrCP. Further investigations into the role of E3 ligases and DUBs in MLN4924-mediated β -TrCP upregulation are required.

In conclusion, the results of the present study demonstrated that downregulation of $\text{I}\kappa\text{B}\alpha$ and the subsequent inflammation attenuate the antitumor potential of MLN4924 in liver cancer cells. Mechanistic studies demonstrated that MLN4924 enhanced the protein stability of β -TrCP, promoting the ubiquitination of $\text{I}\kappa\text{B}\alpha$ protein, leading to the ubiquitin-mediated degradation of the $\text{I}\kappa\text{B}\alpha$ protein, which is summarized in a working diagram (Fig. 7). In addition, the results of the present study demonstrated that β -TrCP knockdown markedly sensitized MLN4924 in suppressing the growth of liver cancer cells via attenuating MLN4924-mediated $\text{I}\kappa\text{B}\alpha$ downregulation and inflammation. The results of the present study not only provide novel insights into the molecular mechanisms by which MLN4924 promotes the degradation of $\text{I}\kappa\text{B}\alpha$ and subsequent inflammation, but also offer a potential therapeutic strategy for sensitizing MLN4924 in the treatment of liver cancer.

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Availability of data and materials

The datasets used and/or analyzed during the present study are available from the corresponding author on reasonable request.

Authors' contributions

ZhixiangY and ZhenzhouY supervised the present study. HX designed the research, performed the experiments and drafted

the manuscript. HX, DZ, YL and LM_a contributed to the acquisition of data. HX, DZ and YL analyzed and interpreted the data. HX, YL, LM_a and LM_{eng} performed the statistical analysis. HX and ZhixiangY confirm the authenticity of all the raw data. All authors have read and approved the final manuscript.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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