







RESEARCH PAPER

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The human vault RNA enhances tumorigenesis and chemoresistance through the lysosome in hepatocellular carcinoma

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ABSTRACT

The small non-coding *VTRNA1-1* (vault RNA 1–1) is known to confer resistance to apoptosis in several malignant cell lines and to also modulate the macroautophagic/autophagic flux in hepatocytes, thus highlighting its pro-survival role. Here we describe a new function of *VTRNA1-1* in regulating *in vitro* and *in vivo* tumor cell proliferation, tumorigenesis and chemoresistance. Knockout (KO) of *VTRNA1-1* in human hepatocellular carcinoma cells reduced nuclear localization of TFEB (transcription factor EB), leading to a downregulation of the coordinated lysosomal expression and regulation (CLEAR) network genes and lysosomal compartment dysfunction. We demonstrate further that impaired lysosome function due to loss of *VTRNA1-1* potentiates the anticancer effect of conventional chemotherapeutic drugs. Finally, loss of *VTRNA1-1* reduced drug lysosomotropism allowing higher intracellular compound availability and thereby significantly reducing tumor cell proliferation *in vitro* and *in vivo*. These findings reveal a so far unknown role of *VTRNA1-1* in the intracellular catabolic compartment and describe its contribution to lysosome-mediated chemotherapy resistance.

Abbreviations: ATP6V0D2: ATPase H⁺ transporting V0 subunit d2; BafA: bafilomycin A1; CLEAR: coordinated lysosomal expression and regulation; CQ: chloroquine; DMSO: dimethyl sulfoxide; GST-BHMT: glutathione S-transferase N-terminal to betaine–homocysteine S-methyltransferase; HCC: hepatocellular carcinoma; LAMP1: lysosomal associated membrane protein 1; LLOMe: L-leucyl-L-leucine methyl ester; MAP1LC3B/LC3: microtubule associated protein 1 light chain 3 beta; MAPK: mitogen-activated protein kinase; MITF: melanocyte inducing transcription factor; MTT: 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide; ncRNA: non-coding RNA; RNP: ribonucleoprotein; SF: sorafenib; SQSTM1/p62: sequestosome 1; STS: staurosporine; tDRs: tRNA-derived RNAs; TFE3: transcription factor binding to IGHM enhancer 3; TFEB: transcription factor EB; vtRNA: vault RNA transcript.

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Introduction


A new class of small non-coding RNA (ncRNA), named vault RNA (vtRNA), was discovered in 1986 in the Rome lab [1]. Vault RNAs are integral components of the large 13-MDa vault particle, a hollow barrel-shaped ribonucleoprotein (RNP) complex, consisting of multiple copies of three proteins: MVP (major vault protein), PARP4/vPARP (poly(ADP-ribose) polymerase family member 4) and TEP1 (telomerase associated protein 1) [2]. Notably, only a minor fraction of their cellular transcript is associated with the vault particles [3,4] hinting to complex-independent functions. Vault RNAs are short POLR3 (RNA polymerase III) transcripts with a length varying between about 80 and 150 nucleotides (nt). In humans, four vtRNA paralogs have been identified, *VTRNA1-1*, *VTRNA1-2*, *VTRNA1-3* and *VTRNA2-1* [4,5]. vtRNAs have been suggested to be involved in a multitude of functions including cell proliferation [6,7], apoptosis [8], autophagy [9], serving as microRNA precursors [10] and have also been linked to chemotherapy

resistance [11,12]. Many of these findings are compatible with the hypothesis that altered vault RNA expression is associated with tumorigenesis. Recently, we demonstrated that the *VTRNA1-1* paralog protects various human cancer cell lines from undergoing apoptosis in a vault particle-independent manner [8,13]. Yet, the exact function and mechanism of action of vault RNAs remains undeciphered in molecular terms.

Other classes of small ncRNAs, such as microRNAs (miRNA) and tRNA-derived RNAs (tDRs) have also been linked to tumorigenesis. miRNA levels were found to be altered in all cancer types studied [14] and have been suggested to function as tumor suppressors or oncogenes (oncomiRs) [15]. miRNAs seem to exert multifaceted functions on tumor progression, modulating tumor growth, metastatic potential, chemoresistance and regulation of metabolism [16,17]. Similarly, tDRs have been identified to be significantly upregulated under various stress conditions and therefore have garnered much attention in the recent years [18–21]. This newly discovered class of ncRNAs has been shown

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to influence cancer development by modulating global translation via regulating the expression of genes coding for ribosomal components [22], interacting with RNA binding proteins [23], by regulating kinase activity [24] and promoting cell proliferation [25]. It is evident that small ncRNAs are important regulatory molecules orchestrating gene expression that can either drive or prevent oncogenic processes and therefore possess potential as prognostic biomarkers or putative drug targets for cancer treatment.

Autophagy is among the biological processes that have been recently connected to vault RNA levels [9]. Autophagy is an evolutionarily conserved degradation pathway activated by cellular stress [26]. Basal autophagy is essential for the degradation and the recycling of proteins and organelles. Especially during stress conditions an effective autophagy machinery is beneficial for cell survival [26]. Damaged organelles and proteins are taken up by autophagosomes which fuse with lysosomes for degradation [27]. Lysosomes are a key entity of the cellular digestive system and their function requires the concerted action of hydrolases, the acidification machinery and luminal protective membrane proteins [28]. Lysosome biogenesis and function are closely coordinated by TFEB (transcription factor EB). TFEB orchestrates the coordinated lysosomal expression and regulation (CLEAR) network genes, involved in lysosomal biogenesis, lysosome-to-nucleus signaling and lipid catabolism [28]. The acidic environment of the lysosomes is a double-edged sword for chemotherapy. Numerous anti-tumor drugs, due to their formulation as a weak-base and hydrophobicity, can be sequestered into the lysosomal lumen, a phenomenon known as lysosomotropism, thus reducing drug-sensitivity and inducing chemoresistance [29].

Here, we describe a novel pro-survival function of *VTRNA1-1/vtRNA1-1* in regulating *in vitro* and *in vivo* tumor cell proliferation, tumorigenesis and chemoresistance, via its newly disclosed role in supporting the function of lysosomes. We showed that *VTRNA1-1* knockout (KO) in human hepatocellular carcinoma (HCC) cells, leads to lysosomal compartment dysfunction by inhibiting TFEB nuclear translocation thus resulting in a downregulation of the CLEAR network genes. We observed that *VTRNA1-1* depletion resulted in an increased activation of the MAPK (mitogen-activated protein kinase) cascade, MAPK1/ERK2-MAPK3/ERK1, responsible for TFEB inactivation and cytoplasmic retention. Finally, we demonstrated that lack of *VTRNA1-1* reduces drug lysosomotropism and significantly inhibits tumor cell proliferation *in vitro* and in an *in vivo* mouse model. These findings reveal a role of *VTRNA1-1* in supporting the function of the intracellular catabolic compartment in human hepatocellular carcinoma cells and highlight its importance in lysosome-mediated chemotherapy resistance.

Results

VTRNA1-1 plays a crucial role in tumor cell proliferation and tumorigenesis

The anti-apoptotic effect of *VTRNA1-1/vtRNA1-1* has been the subject of intensive studies in our laboratory [8,13] and lead us to hypothesize about its involvement in tumorigenesis

and cancer progression. To test the influence of *VTRNA1-1* expression on tumor cell proliferation and colony formation *in vitro*, we used the human hepatocellular carcinoma cell lines Huh-7 (WT), and Huh-7 *VTRNA1-1* KO (Figure 1A). We found that proliferation and colony formation were significantly reduced in Huh-7 *VTRNA1-1* KO cells (Figure 1B-D). Re-introducing the *VTRNA1-1* gene could fully rescue cell proliferation in these complementation cells (Figure 1A, C). Likewise, the colony formation assay showed a similar trend upon *VTRNA1-1* complementation (Figure 1D). The lack of *VTRNA1-1* seems to suggest an important role of this ncRNA in tumor cell viability and proliferation. To further assess the effects of vtRNA1-1 on the tumor growth using an *in vivo* subcutaneous xenograft mouse model, we injected Huh-7 cells into mice and allowed to develop measurable tumors. Consistent with our *in vitro* results, the growth was significantly reduced in the *VTRNA1-1* KO-transplanted tumors compared to the WT (Figure 1E, F).

Interestingly, *VTRNA1-1* expression seems to be also modulated in human derived primary hepatocytes (Figure S1A). In culture, primary hepatocytes are unable to proliferate and progressively de-differentiate from their cell-specific functions [30,31]. Interestingly, we observed a significant increase in *VTRNA1-1* expression in hepatocyte cultures for up to 14 days (Figure S1A) while hepatocytes still maintained their differentiated state (Figure S1B). Hepatocytes in culture are known for spontaneously activating apoptosis pathways [32,33]. Consistently, we detected PARP1 (poly(ADP-ribose) polymerase 1) cleavage already at day 0 (before seeding) together with an increased expression of BCL2 (BCL2 apoptosis regulator) (Figure S1C), hinting at the already described activation of an anti-apoptotic cellular response [34]. Thus, increased *VTRNA1-1* expression might be similarly involved in pro-survival pathways.

These observations reveal that the cellular levels of *VTRNA1-1/vtRNA1-1* contribute to proliferation in immortalized cell lines and are modulated in response to stresses in primary hepatocytes. All together our data indicate that vtRNA1-1 plays a key role in tumor cells proliferation and tumorigenesis.

Defective lysosome function impairs autophagy-mediated clearance in VTRNA1-1 KO cells

Because *VTRNA1-1* levels have been reported previously to affect apoptosis [8,13] and autophagy [9], we next asked the question how and if these cellular processes are linked via *VTRNA1-1* expression in hepatocytes. Huh-7 sensitivity toward apoptosis was not affected by *VTRNA1-1* depletion, since upon staurosporine (STS) treatment, a broad-spectrum protein kinase inhibitor inducing apoptosis, the IC₅₀ values were similar for both WT and *VTRNA1-1* KO cell lines (Figure 2A). In agreement cleavage of pro-apoptotic PARP1 was not observed or was negligible in these cells (Figure 2B). These results fit with the observation that Huh-7 are remarkably resistant to apoptosis even after treatment with STS [35]. Similar results were obtained using the pro-apoptotic DNA-damaging anticancer drug doxorubicin (Figure S2A) as well

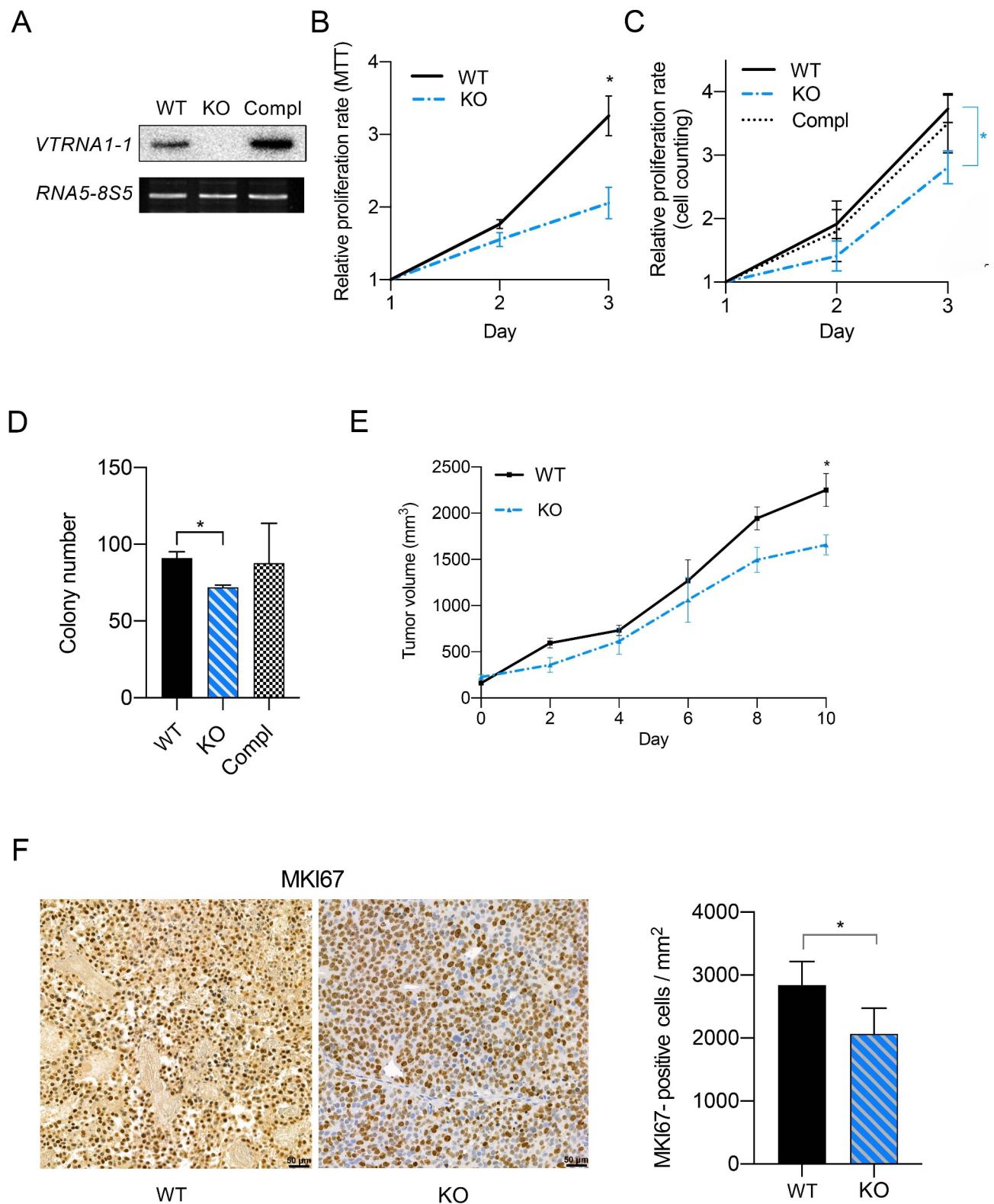


Figure 1. *VTRNA1-1/vtRNA1-1* plays a crucial role in tumor cell proliferation and tumorigenesis. (A) Total RNA was extracted from Huh-7 cells and analyzed by northern blotting to confirm successful *VTRNA1-1* complementation in Huh-7 KO cells. *RNA5-8S5* (RNA, 5.8S ribosomal 5) serves as internal loading control. (B) Mean \pm SD relative proliferation of Huh-7 WT, KO and complementation cells was measured with the MTT assay. Values were normalized to day 1, $n = 3$. (C) Three-day cell proliferation (cells/ml) by automated cell counter. Values were normalized to day 1, $n = 5 \pm$ SD. (D) Clonogenic assay: data are expressed as the mean values of three independent experiments \pm SD. (E) Tumor volume \pm SEM of Huh-7 subcutaneous xenografts mouse model (WT $n = 4$ and KO $n = 5$). (F) Huh-7 subcutaneous xenografts immunohistochemistry for MKI67. Counterstain: hematoxylin. Scale bar: 50 μ m. The graphs show the mean percentage \pm SD of MKI67-positive cell/area per tumor/mouse (five random fields were used for the analysis per tumor/mouse). P values < 0.05 were considered statistically significant and are indicated as follows: * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$; **** $P < 0.0001$; ns, not significant.

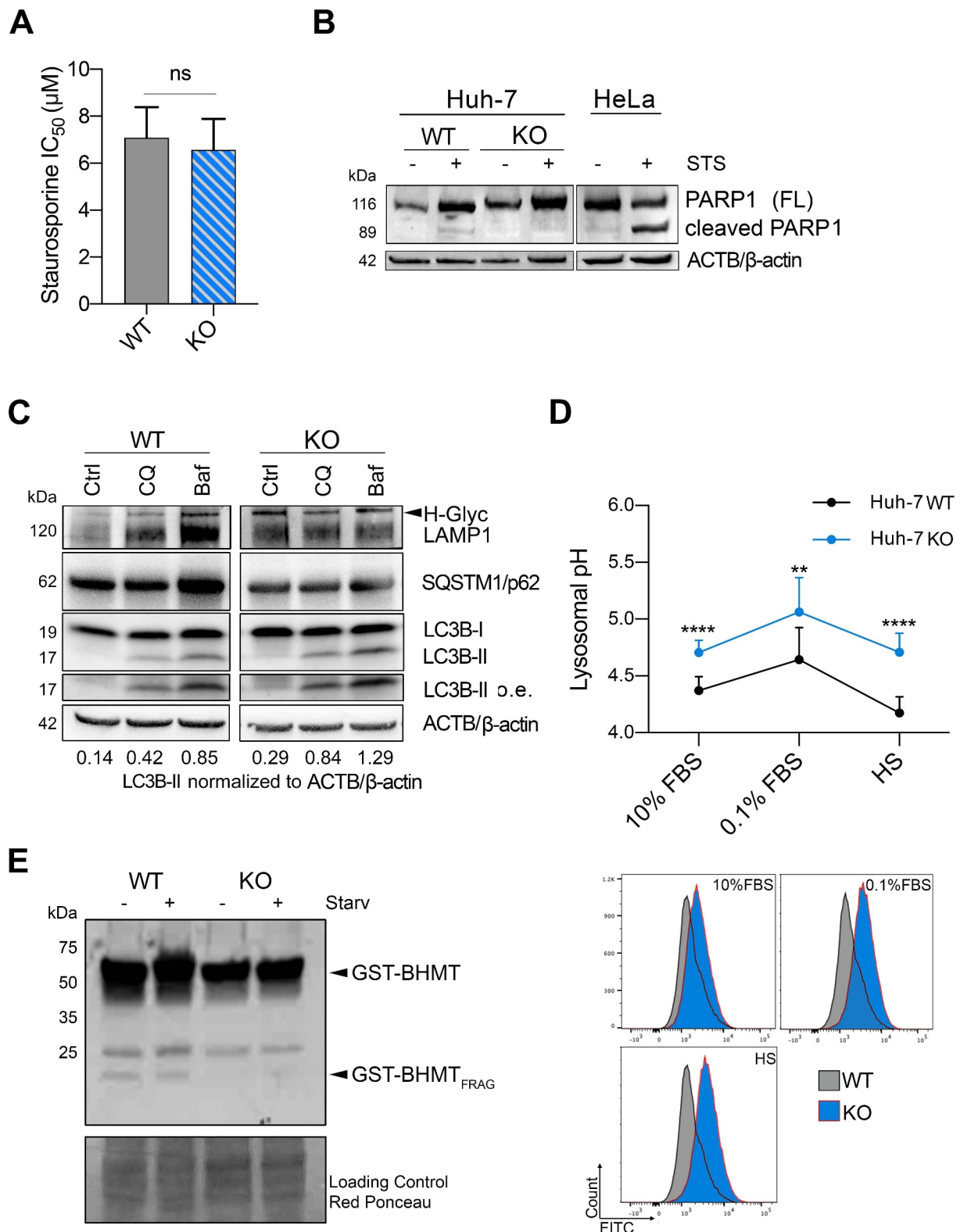


Figure 2. Defective lysosome function impairs autophagy-mediated clearance in *VTRNA1-1* KO cells. (A) IC₅₀ values of Huh-7 WT and KO cells after 24 h of staurosporine treatment. (B) Representative immunoblots for PARP1 (FL, full-length) and ACTB/β-actin in Huh-7 and HeLa cells after Staurosporine IC₃₀ treatment (24 h, 5 µM). HeLa cells were used as control for apoptosis induction. (C) Representative immunoblots for LAMP1, SQSTM1/p62, LC3B-I, LC3B-II (o. e., overexposed) and ACTB/β-actin in Huh-7 cells under complete culture medium (10% FBS – Ctrl) and treatment with chloroquine (CQ – 20 µM, 4 h) or bafilomycin A₁ (BafA, 100 nM, 4 h). (D) Top panel, mean ± SD of lysosomal pH values measured by flow cytometry in Huh-7 pre-incubated in culture medium supplemented with FITC-dextran (0.1 mg/mL, 72 h) followed starvation (0.1% FBS) for 24 h and high starvation (HS, with low glucose, without amino acids and FBS) for 6 h. n = 3. Bottom panel, illustrative histograms from flow cytometry showing the shift of FITC-dextran emission wavelength. (E) Representative immunoblots for GST-BHMT of total lysate obtained from Huh-7 cells transiently transfected with the GST-BHMT construct and either maintained in complete culture media or starving media supplemented with leupeptin and E64d and without essential amino acids and FBS for 6 h. Red Ponceau staining served as loading control. P values < 0.05 were considered statistically significant and are indicated as follows: *P < 0.05; **P < 0.01; ***P < 0.001; ****P < 0.0001; ns, not significant.

as during high stress culture conditions (serum-deprived culture medium) where KO of *VTRNA1-1* actually slightly enhances cell survival compared to WT (**Figure S2B**). This is in stark contrast to the effects seen in HeLa cells, where staurosporine treatment and starvation increase apoptotic rates and induces PARP1 cleavage (**Figure 2B** and [13]).

A recent study suggested that *VTRNA1-1* acts as a regulator of autophagy by interacting with SQSTM1/p62 (sequestosome 1), which is involved in targeting substrates of the autophagy machinery, preventing its self-oligomerization in Huh-7 cells [9]. As already shown by Horos et al., *VTRNA1-1*-dependent impairment of the autophagic flux can be detected by treating Huh-7 cells with the two late-stage autophagy inhibitors, chloroquine (CQ) and bafilomycin A₁ (BafA). Western blot analysis revealed an increased amount of MAP1LC3B/LC3 (microtubule-associated protein 1 light chain 3 beta) expression in KO cell lines compared to WT (**Figure 2C, S2C**). Conversion of LC3B-I to LC3B-II is the hallmark for autophagosome formation. Interestingly, we detected a pronounced LC3B-II protein accumulation in cells lacking *VTRNA1-1* already at basal level, under normal culture conditions, suggesting that loss of this ncRNA induces autophagy impairment (**Figure 2C**). Of note, no significant changes in SQSTM1/p62 levels were apparent under control conditions in the absence of *VTRNA1-1* (**Figure 2C**).

Autophagosomes are intermediate structures in a very dynamic process, where their intracellular localization results from a balance between their generation and their conversion rate into autolysosomes [36]. Thus, autophagosome accumulation may derive from either autophagy induction or impaired lysosomal clearance [28]

To identify the phenomenon causing autophagosome accumulation in *VTRNA1-1* KO cells, we first aimed to examine lysosomal compartment structural proteins focusing on the expression levels of LAMP1 (lysosomal associated membrane protein 1). Interestingly, LAMP1 levels were increased, together with an enhanced expression of its hyperglycosylated (H-Glyc) form only in *VTRNA1-1* KO cells (**Figure 2C**). Further, in variance to WT cells we found that CQ- and BafA-induced upregulation of LAMP1, a clear sign of a larger accumulation of undigested autolysosomes, was not detectable in the absence of *VTRNA1-1* (**Figure 2C**). An increased expression of H-Glyc LAMP1 has been reported to be linked with intrinsic lysosomal compartment instability [37]. Moreover, in addition to lysosomal compartment stability, lysosome-mediated cellular clearance processes require the concerted action of hydrolases and the intraluminal acidification machinery [28]. Interestingly, in *VTRNA1-1* KO cells we detected a significant increase in lysosomal pH toward more alkaline values (**Figure 2D**). Taking advantage of the GST-BHMT assay, previously developed for measuring autophagic degradation capabilities [38], we next assessed the lysosomal proteolytic activity in the cells. In line with our previous observations, the ability of lysosomes to degrade GST-BHMT was markedly reduced in absence of *VTRNA1-1*, indicating an impaired

lysosomal proteolytic activity (**Figure 2E**). Conversely, in HeLa cells, where *VTRNA1-1* was shown to confer apoptosis resistance [8,13], neither changes in lysosomal pH nor in lysosomal proteolytic activity were apparent when *VTRNA1-1* was depleted (**Figure S2D and E**).

Collectively, our data demonstrate that *VTRNA1-1* has a crucial role in Huh-7 in modulating lysosomal function, especially for regulating their intraluminal pH and proteolytic activity.

***VTRNA1-1* levels influence TFEB nuclear translocation and affect TFEB-driven CLEAR network gene expression**

Lysosomal function is closely monitored to respond and adapt to environmental stimuli by its master TFEB regulator. Under basal conditions, TFEB is located in the cytoplasm and under specific stimuli, such as starvation, it is rapidly translocated into the nucleus. The cellular localization and activity of TFEB are mainly controlled by phosphorylation, which depends predominantly on MTORC1 and MAPK1/ERK2 [39,40]. To gain further mechanistic insight into the molecular events regulating TFEB activation, we tested whether its upstream regulators might be influenced by *VTRNA1-1* expression. While we did not observe any difference in MTOR phosphorylation (**Figure 3A**), *VTRNA1-1* depletion resulted in a clear basal increase of MAPK pathway activation (**Figure 3B and S3A**). We observed a higher phosphorylation level of MAPK1/3, its upstream kinases MAP2K1/MEK1-MAP2K/MEK2 as well as RPS6KA1/90RSK (ribosomal protein S6 kinase A1), its downstream target (**Figure 3B and S3A**). Importantly, the elevated markers of phosphorylation returned to WT-levels in the complementation cells (**Figure 3B and S3A**) thus suggesting that re-introducing *VTRNA1-1* restores signaling. Surprisingly, the MAPK pathway was also regulated in HeLa *VTRNA1-1* KO cells, but in an opposite manner (**Figure S3B**). Our data suggests that the loss of *VTRNA1-1* leads to a hyperactivation of MAPK1/3, which might in turn prevent TFEB nuclear translocation. We indeed found the amount of nuclear TFEB accumulation to be significantly lower in KO cells compared to the WT (**Figure 3C and S3C**). Other transcription factors of the same family known to regulate starvation-induced lysosomal biogenesis and autophagy [41,42] were either lowly expressed in Huh-7 and thus below the western blot detection limit (MITF), or did not show changes in protein expression or altered localization upon *VTRNA1-1* removal (TFE3) (**Figure 3D**). These findings are in accordance with a previous study demonstrating no nuclear translocation of TFE3 or MITF in Huh-7 cells under the applied starvation conditions [43]. Therefore, our results reveal *VTRNA1-1* levels to specifically influence TFEB as the major transcription factor for lysosome biogenesis.

TFEB orchestrates lysosomal expression and regulation (CLEAR) network genes, involved in lysosomal biogenesis, lysosome-to-nucleus signaling and lipid catabolism [28]. Therefore, we analyzed the expression of known TFEB target genes in KO cells compared to WT at basal level and under serum-deprived culture conditions, where especially tumor

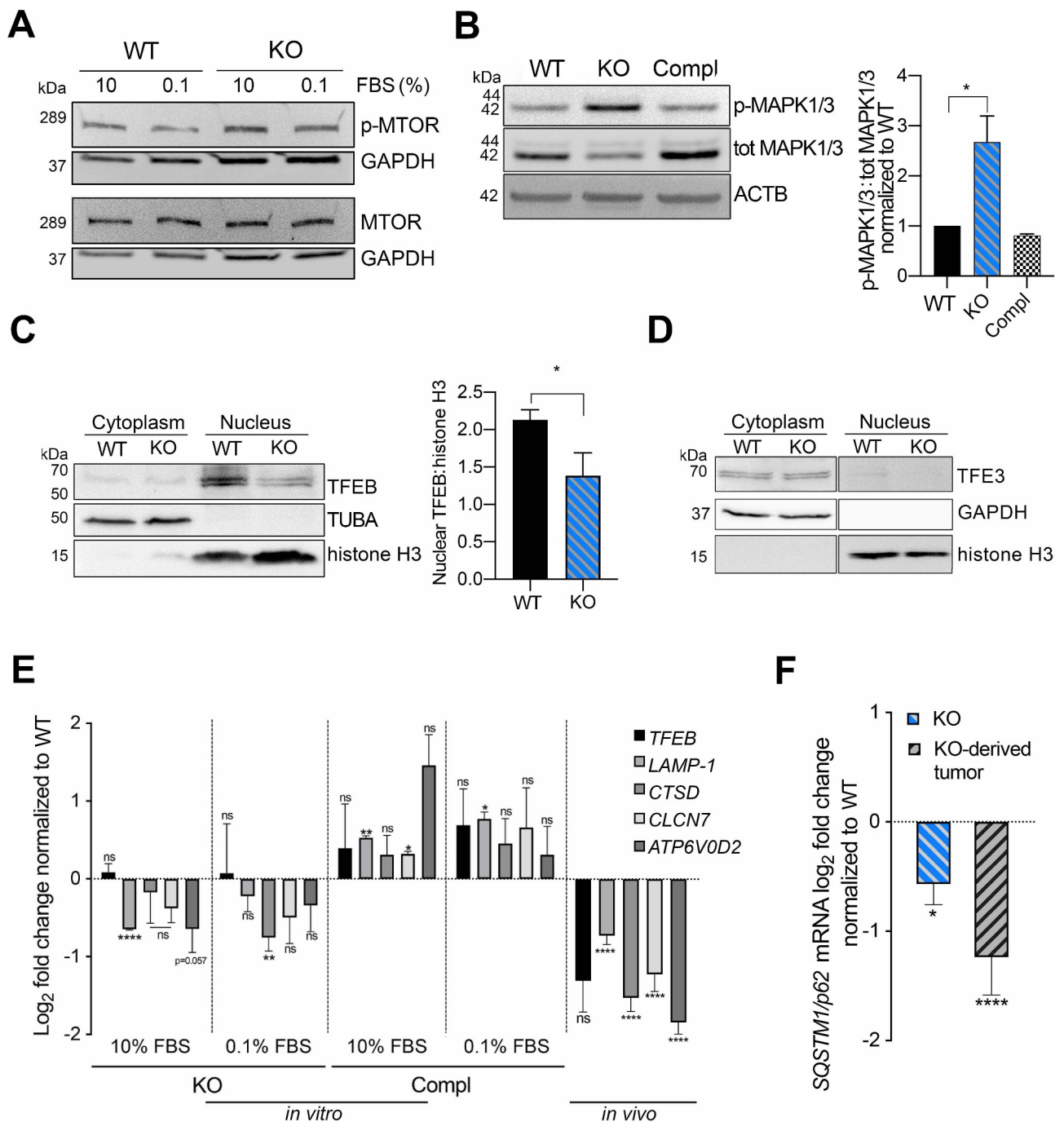


Figure 3. *VTRNA1-1/vtrna1-1* plays a key role in TFEB nuclear translocation and TFEB-driven CLEAR network genes expression. (A) Representative immunoblots for p-MTOR, total MTOR and GAPDH in Huh-7 WT and *VTRNA1-1* KO cells under normal culture conditions (10% FBS). (B) Left, representative immunoblots for p-MAPK1/ERK2-MAPK3/ERK1, total MAPK1/3 and ACTB/ β -actin in Huh-7 WT, *VTRNA1-1* KO and complementation cells under normal culture conditions (10% FBS). Right, quantification of p-MAPK1/3 obtained by normalizing the p-MAPK1/3 and total MAPK1/3 levels by ACTB/ β -actin followed by the ratio of p-MAPK1/3 over total MAPK1/3. Values are expressed as the mean values of three (for WT and KO) or two (for complementation) independent experiments \pm SD normalized to WT level. (C) Left, representative nuclear/cytoplasm fractionation immunoblots for TFEB, TUBA/ α -tubulin, histone H3 in Huh-7 WT and *VTRNA1-1* KO cells under high starvation for 3 h. Right, quantification of TFEB nuclear levels normalized to histone H3. Values are expressed as the mean values of three independent experiments \pm SD. (D) Representative nuclear-cytoplasm fractionation immunoblots for TFE3, GAPDH, histone H3 in Huh-7 WT and *VTRNA1-1* KO cells under high starvation for 3 h. (E) Real-time qPCR data (mean \pm SEM) of *TFEB*, *LAMP1*, *CTSD*, *CLCN7* and *ATP6V0D2* mRNA levels in Huh-7 *VTRNA1-1* KO cells grown under complete medium (10% FBS - Ctrl) and starving culture conditions (0.1% FBS) for 24 h, normalized to Huh-7 WT (n = 3) and *VTRNA1-1* KO-derived tumors normalized to WT-transplanted ones (WT n = 4, KO n = 9). (F) Real-time qPCR mean \pm SEM data of *SQSTM1/p62* in Huh-7 KO cells and derived tumors normalized to Huh-7 WT and WT-transplanted tumors respectively. P values < 0.05 were considered statistically significant and are indicated as follows: *P < 0.05; **P < 0.01; ***P < 0.001; ****P < 0.0001; ns, not significant.

cells more heavily rely on their intracellular catabolic machinery. In fact, the recycling of intracellular constituents provides alternative energy sources during periods of metabolic stress to maintain homeostasis and viability [44]. While the mRNA

levels of *TFEB* remained unchanged, its target genes, such as *LAMP1*, *CLCN7* (chloride voltage-gated channel 7), *CTSD* (cathepsin D) and *ATP6V0D2* (ATPase H⁺ transporting V0 subunit d2) were downregulated in the absence of *VTRNA1-1*

(Figure 3E). Upon complementation, Huh-7 cells restored *VTRNA1-1* levels (Figure 1A), concomitantly with an increased expression of CLEAR network genes (Figure 3E). Overexpressing a TFEB gain-of-function mutant with pronounced nuclear localization characteristics ([45] and Figure S3D) in the *VTRNA1-1* KO cells rescued expression of CLEAR network genes (Figure S3E). These observations substantiate the interdependence of *VTRNA1-1* expression, TFEB nuclear localization and lysosome activity. Consistently, we also confirmed that the CLEAR network genes were downregulated in tumors in *VTRNA1-1* KO-injected mice compared to the WT control (Figure 3E), indicating that the previously observed altered lysosome function in the *VTRNA1-1* KO cells (Figure 2C) can be attributed to the downregulation of CLEAR gene expression. Of note, SQSTM1/p62 expression is also under the transcriptional control of TFEB [46] and in fact, in line with our previous findings, its mRNA levels were significantly downregulated in *VTRNA1-1* KO cells compared to WT in cell culture as well as in the *in vivo* mouse model (Figure 3F).

In summary these data provided evidence that *VTRNA1-1* plays a role in establishing lysosome function,

where its absence lead to a marked loss of autolysosome proteolytic activity, decreased TFEB nuclear translocation and subsequent reduced expression of CLEAR network genes.

Removal of *VTRNA1-1* potentiates the cytotoxicity of sorafenib *in vitro* and *in vivo*

Our data provide evidence that loss of *VTRNA1-1/vtRNA1-1* interferes with lysosome function, thus leading to a deficiency in the autophagic machinery. Weak base anticancer compounds can be sequestered in the acidic lumen of lysosomes by cation trapping and consequently are no longer able to reach their target, thereby reducing their effectiveness [47]. It has been reported that the cytotoxic potential of sorafenib (SF), the first-line treatment option for advanced hepatocellular carcinoma patients and a known tyrosine kinase inhibitor of the MAPK pathway [48], is significantly decreased due to its passive lysosomal compartment trapping [37]. Therefore, we next investigated if loss of *VTRNA1-1* leads to lysosomal malfunction and thus consequently to increased drug

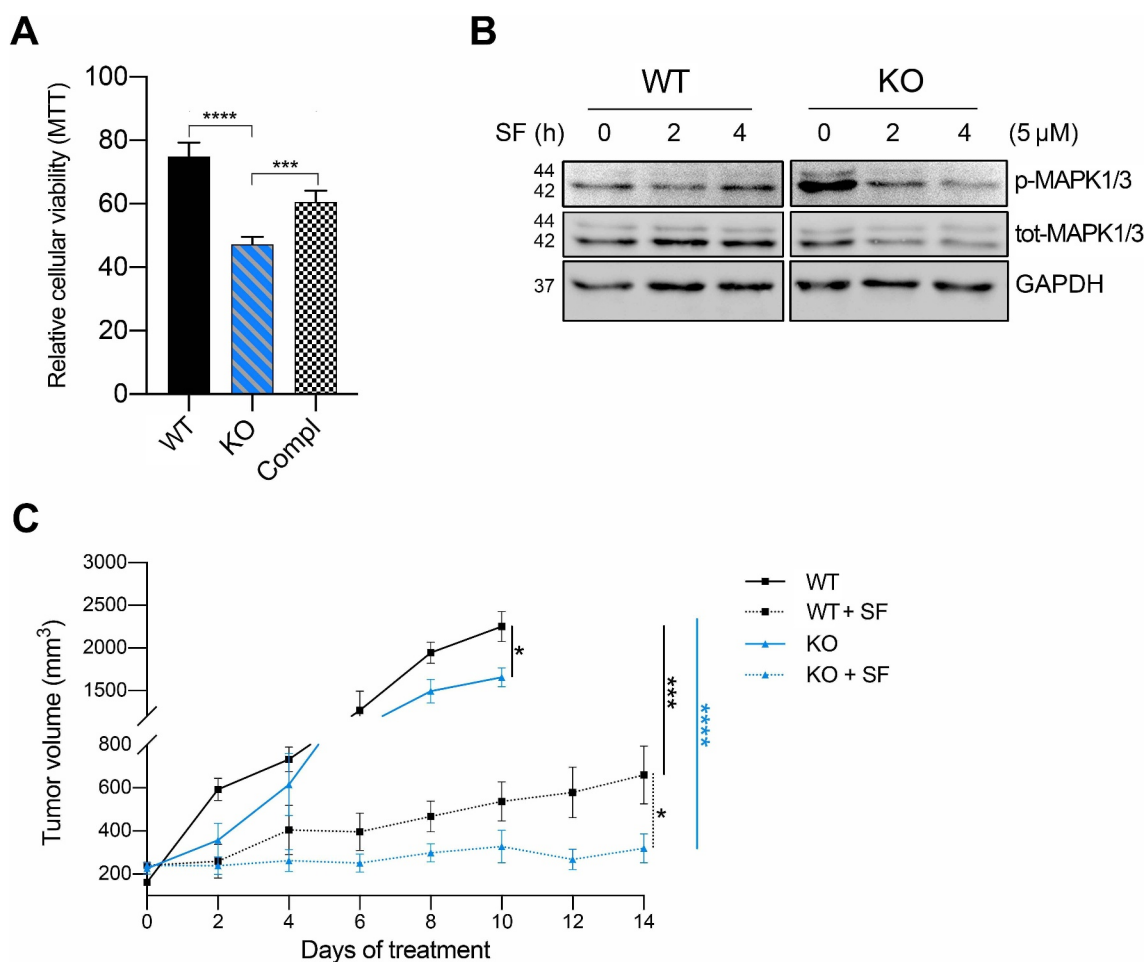


Figure 4. Removal of *VTRNA1-1* potentiates the cytotoxicity of sorafenib *in vitro* and *in vivo*. (A) Mean \pm SD relative viability of Huh-7 WT, KO and complementation cells, after sorafenib IC_{30} treatment (24 h, 14 μ M) measured with the MTT assay. Values were normalized to untreated cells, $n = 3$. (B) Representative immunoblots for p-MAPK1/3, total MAPK1/3 and GAPDH in Huh-7 WT and *VTRNA1-1* KO cells in presence or absence of sorafenib (5 μ M) as indicated. (C) Tumor volume of Huh-7-transplanted xenografts mice treated with vehicle and sorafenib (SF – 100 mg/kg – daily oral gavage) for 14 days (SF $n = 6$ /group; vehicle WT $n = 4$, KO $n = 5$). P values < 0.05 were considered statistically significant and are indicated as follows: * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$; **** $P < 0.0001$; ns, not significant.

sensitivity. When treated with the IC₃₀ dose of SF, Huh-7 KO cells showed a significantly decreased viability compared to WT cells after treatment (Figure 4A). This increased SF cytotoxicity indeed depends on *VTRNA1-1* levels since the rescued cell line behaved similarly to the WT cells (Figure 4A). Interestingly, we found that SF markedly downregulates MAPK1/3 phosphorylation in a time-dependent manner in Huh-7 KO cells compared to the WT (Figure 4B), suggesting a higher amount of drug being able to reach its intracellular molecular target due to a decreased lysosomotropism. On the other hand, the lysosomotropic compound LLOMe (L-leucyl-L-leucine methyl ester), which is an acidic pH-driven lysosome-specific membrane damage inducer, showed a markedly reduced cytotoxicity in *VTRNA1-1* KO cells (Figure S4A). Huh-7 WT cells viability was more strongly decreased compared to *VTRNA1-1* KO cells after LLOMe administration (Figure S4A). We observed a significantly lower LLOMe-induced lysosomal pH alkalinization in Huh-7 *VTRNA1-1* KO cells compared to the WT (Figure S4B) further confirming a weaker LLOMe lysosomal compartment tropism and trapping of LLOMe, mainly due to the previously shown elevated alkaline intraluminal pH observed for the KO cells (Figure 2D). Corroborating our *in vitro* data, SF treatment significantly impaired

tumor growth and progression in the Huh-7 KO-transplanted mice as compared to animals transplanted with the WT control cells (Figure 4C).

These data exhibit a role of *VTRNA1-1* in lysosomal compartment functionality, which seems to involve in the positive modulation of the pro-tumorigenic MAPK cascade activation (Figure 5). Cumulatively we found that *VTRNA1-1* plays a significant role in determining the intracellular fate and potency of lysosomotropic compounds and thus affects their molecular target site distribution and antitumor effectiveness.

Discussion

The human genome encodes on chromosome 5 four vault RNA paralogs, *VTRNA1-1*, *VTRNA1-2*, *VTRNA1-3* and *VTRNA2-1* [5]. These ncRNAs are around 100 residues long and they received their name based on an early observation that they are integral to the vault particle, a gigantic hollow RNP with a molecular mass of 13 MDa [1]. Despite decades of dedicated research, the biological function of the vault particle or the vault RNAs remained enigmatic [2,49]. More recently, it was demonstrated that only a minor fraction of vault RNA is actually associated with the vault complex [4,50] thus suggesting cellular functions of these ncRNAs beyond the vault

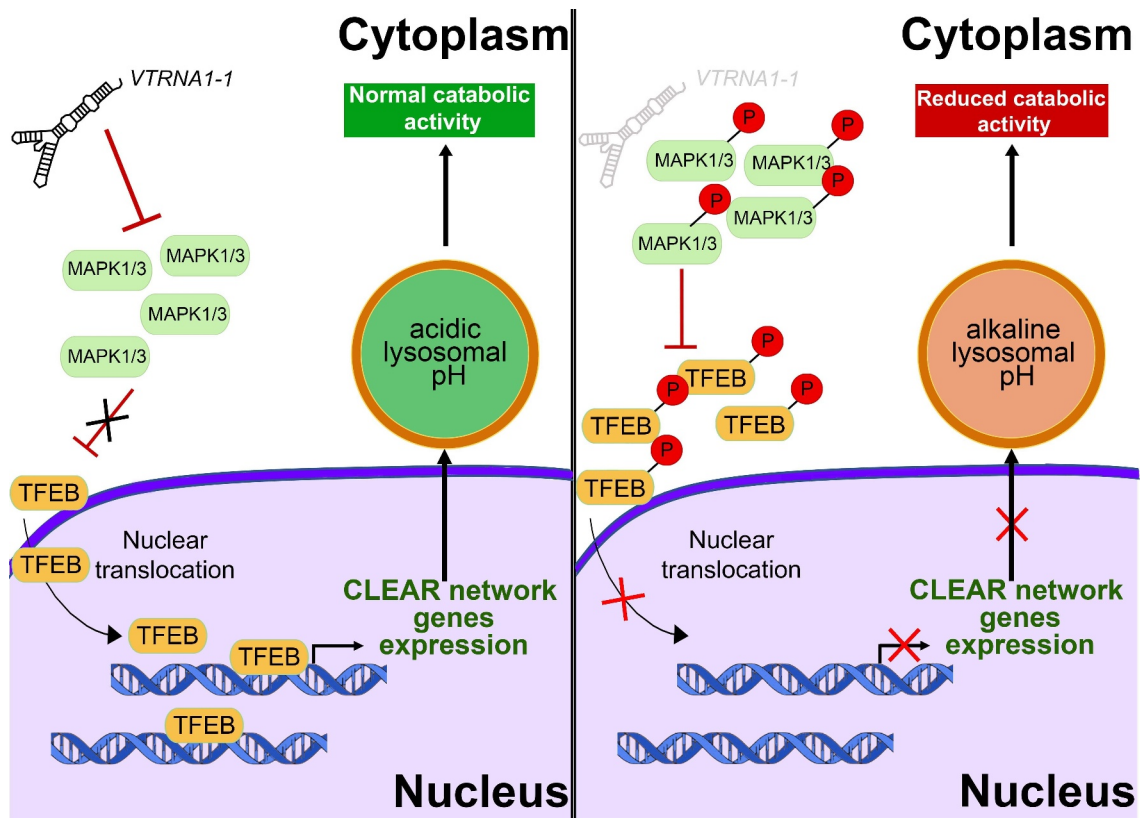


Figure 5. Model for lysosome and autophagy mediated clearance regulation. In *VTRNA1-1/vtrna1-1* WT cells low MAPK1/3 phosphorylation level allows TFEB nuclear translocation leading to CLEAR network genes expression and ensuring intracellular catabolic compartment stability and function (left). Reduced *VTRNA1-1* levels (right) lead to the hyperphosphorylation of MAPK1/3 and TFEB, and consequently to its reduced nuclear translocation, leading to a downregulation of the CLEAR network genes and lysosomal compartment dysfunction.

complex. Accumulating evidence reveals a key role of the small non-coding *VTRNA1-1/vtRNA1-1* in conferring cellular resistance against cell death via chemoresistance [7,12], viral defense and apoptosis [4,8,13,51]. This led us to further investigate the involvement of *VTRNA1-1* in tumorigenesis and cancer progression. In agreement with its previously described function, we observed a modulation of *VTRNA1-1* expression in response to apoptosis induced by physical and mechanical stress in primary human hepatocytes (**Figure S1 A, C**). We further describe that *VTRNA1-1* is involved in tumor cell proliferation, tumorigenesis and chemoresistance, via its newly described role in supporting the function of the intracellular catabolic machinery (**Figures 1, Figures 2D-E and 4A-C**). However, we did not observe that the sensitivity of Huh-7 cells to apoptosis was altered by the lack of *VTRNA1-1* (**Figure 2A, B and Figure S2A, B**). This is in clear contrast to several other immortalized cell lines, such as the Burkitt Lymphoma cell lines (BL2, BL41), the breast cancer model HS578T, the human embryonic kidney cells (HEK293), the lung carcinoma cells A549 and HeLa cells [8,13]. It is safe to conclude that high *VTRNA1-1* levels are correlated with pro-survival characteristics in numerous human cell lines of different tissue origins (**Figure S5**). Thus, it is possible that the well-documented pro-survival biological function of *VTRNA1-1* manifests itself differently and is cell line or tissue dependent and a generalized mechanism of *VTRNA1-1* should not be based on data obtained in one cellular system. A recent study in Huh-7 cells reported that *VTRNA1-1* was able to contribute in the regulation of the autophagic flux, by directly interacting with the autophagy receptor protein SQSTM1/p62 [9]. In our study, we were able to confirm that the loss of *VTRNA1-1* leads to elevated LC3B-II levels suggesting autophagosome accumulation (**Figure 2C and Figure S2C**). Previously these observations were interpreted as evidence for an increase of the autophagic flux in the absence of *VTRNA1-1* [9]. Our *in vitro* and *in vivo* experiments provide further insight into *VTRNA1-1* biology and elaborate the suggested role of SQSTM1/p62 as post-transcriptional regulator of autophagy [9]. Although the decreased steady-state level of SQSTM1/p62 was initially attributed to the lost interaction with *VTRNA1-1* [9], we found that depletion of *VTRNA1-1* resulted in transcriptional downregulation of several catabolism-related genes, including also SQSTM1/p62 (**Figure 3E and F**). Our data demonstrated that reduced expression of SQSTM1/p62 is transcriptionally controlled and it is not necessarily dependent on its interaction with *VTRNA1-1*.

Lysosomes and autophagy are two tightly linked and evolutionarily conserved processes that provide a constant supply of cellular energy mainly through recycling of damaged organelles and proteins [27]. As previously described by Horos et al., *VTRNA1-1* depletion leads to mature autophagosomes accumulation, suggesting an increased autophagic flux [9]. However, autophagosomes accumulation, in the absence of any specific chemical modulator of the flux progression, may come from either autophagy induction or impaired lysosomal autophagy-mediated clearance. Lysosomes are intracellular vesicles responsible for such catabolic and clearance processes

in cell homeostasis, and their function requires the concerted action of hydrolases, acidification pumps for maintaining acidic the intraluminal pH and lysosomal membrane proteins to ensure the integrity of this specific vesicular compartment [28]. Investigating the lysosome-mediated clearance process, we demonstrated that deletion of *VTRNA1-1* caused an alkalization of the intraluminal lysosomal pH, leading to a marked impairment of their proteolytic and degradative activity (**Figure 2D, E**). LAMPs are structural proteins important for maintaining lysosomal membrane integrity and stability by segregating the acidic environment of the lumen from the rest of the cell [52]. Under normal conditions, LAMP1 glycosylation protects the inner lysosomal membrane from cleavage and degradation from the low pH of the intraluminal environment [53]. Knockout of *VTRNA1-1* leads to an impairment of CQ- and BafA-induced LAMP1 accumulation (**Figure 2C**). Additionally, an already stressed and intrinsically unstable status of the catabolic compartment in *VTRNA1-1* KO cells is indicated by the increased glycosylation of LAMP1 in untreated growth conditions (**Figure 2C**). Concurrently, *VTRNA1-1* depletion was indeed able to induce a clear switch of the lysosome intraluminal pH toward a more alkaline environment, affecting the catabolic and proteolytic activity (**Figure 2E**). These findings seem to attribute the autophagosomes accumulation evident in *VTRNA1-1* KO cells to the lysosomal compartment dysfunction.

The expression and activity of the lysosomal components are tightly coordinated to allow it to function under physiological and pathological conditions by the CLEAR genes network, as well as by TFEB nuclear translocation and transcriptional control [54]. Investigating the potential cause for the compromised lysosomal function, we uncovered that TFEB nuclear translocation was decreased in *VTRNA1-1* KO cells, with a significant downregulation of the CLEAR network genes *in vitro* and *in vivo* (**Figure 3C-F**). Genetic complementation of *VTRNA1-1* in KO cells restored not only cellular proliferation (**Figure 1A, C**) but also CLEAR gene expression (**Figure 3E**), further suggesting their interdependence with *VTRNA1-1* levels. Given the well-known tight link between the MAPK pathway and TFEB activation [39], we found that *VTRNA1-1* depletion markedly induced an increased phosphorylation of the downstream effector of the MAPK kinase cascade MAPK1/3 (**Figure 3B**), which finally suggested that *VTRNA1-1* KO triggers a compensatory overactivation of the MAPK pathway (**Figure 5**). Similarly, in HeLa cells we recently demonstrated that deletion of *VTRNA1-1*, but not of the paralog *VTRNA1-3*, modulated several signaling pathways, including the MAPK cascade [13]. The link between vault RNA expression and MAPK signaling was recently substantiated in cultured mouse cortical neurons [55]. Further, we demonstrated that all these phenomena reported here can potentiate the antitumor effect of conventional chemotherapeutic drugs. Specifically, the vast majority of chemotherapeutic agents, due to their hydrophobic and weak base chemical characteristics, often get passively sequestered within the acidic lumen of lysosomes, becoming unable to reach their intracellular molecular

target [56]. Interestingly, the lack of *VTRNA1-1* and the consequently elevated alkaline lysosomal pH lead to a diminished cytotoxicity of the lysosomal disruptor LLOMe, mainly due to its decreased lysosomotropism (Figure S4). On the other hand, the similarly reduced lysosomotropism of sorafenib observed in the KO cells *in vitro* and *in vivo* revealed an increased availability of sorafenib at its target site and therefore a stronger antitumor effect (Figure 4C).

Taken together, our study uncovered the involvement of *VTRNA1-1/vtRNA1-1* in tumor cell proliferation and tumorigenesis, demonstrating its crucial role to maintain the catabolic compartment and chemoresistance (Figure 5). This novel pro-survival function may identify *vtRNA1-1* as a new therapeutic target to overcome chemosensitivity caused by passive lysosomal sequestration of antitumor drugs and augment tumor-patients outcomes.

Materials and methods

Cell culture

Huh-7 (kindly provided by Matthias Hentze; EMBL, Heidelberg, Germany) and HeLa cells (ATCC®-CCL-2™) were cultured in low glucose (5 mM) DMEM (Thermo Fisher, 21,331,046) supplemented with 10% heat inactivated FBS (10,082,147, Thermo Fisher) and 100 U/ml PenStrep glutamine (Thermo Fisher, 10,378,016). The *VTRNA1-1* KO and WT Huh-7 cell lines were kindly provided by M. Hentze (EMBL, Heidelberg, Germany). We derived a Huh-7 *VTRNA1-1* complementation cell line using lentiviral transduction (below) in the Huh-7 KO cells. After successful lentiviral transduction, positively infected cells were selected by treating them with 1 µg/ml puromycin for at least 1 week. Single cell clones were obtained by serial dilution. Normal liver tissue was obtained from surgical resections from consented patients at the University Hospital of Bern. Hepatocytes were then isolated by enzymatic perfusion as previously described [57] and kept in culture at 37°C in a humidified incubator with 5% of CO₂.

Patient-derived tissues

Primary human liver tissue for hepatocyte isolation [57] and HCC tumors from liver were obtained from patients of the University Hospital Bern (Inselspital). Informed consent was obtained prior to surgery in compliance with the local ethics regulations and under approval of local ethics commission (Project-ID Nr. 2019–00157).

Lentiviral transduction

Lentiviral particles were generated in human HEK 293 T cells (kindly provided by M. Ruepp, King's College London), which were transiently transfected with lentiviral plasmids containing cDNAs coding for *VTRNA1-1*, together with the packaging plasmids pSPAX and the envelope plasmid pVSV-G (kindly provided by S. Geley, Medical University Innsbruck). After

48 h and 72 h lentiviral supernatant was collected, sterile filtered (Sigma-Aldrich, Whatman Puradisc FP30, 0.2 mM) and supplemented with polybrene (Sigma, 107,689) to a final concentration of 4 µg/ml and added to the target cells overnight.

Transfection and treatments

Transfections were done using Lipofectamine 3000 (Thermo Fisher, L3000008) for plasmid DNA. Doxorubicin (Sigma-Aldrich, D1515) was diluted in water to 5 mM and used at 1 µM for 48 h. Bafilomycin A₁ (BafA; Sigma-Aldrich, B1793) was diluted in DMSO to 200 µM and used at 100 nM for 4 h. Chloroquine (CQ; Sigma-Aldrich, C6628) was diluted in DMSO to 50 mM and used at 20 µM for 4 h. Rapamycin (Sigma-Aldrich, R0395) was diluted in ethanol to 2 mM and used at 200 nM for 4 h. Leu-Leu methyl ester hydrobromide (LLOMe; Sigma-Aldrich, L7393) was diluted in DMSO to 1 M and used from 0.001 to 1.4 mM for 6 h. Sorafenib tosylate (MedChemExpress, HY-10201A) was used from 5 to 14 µM for 24 h *in vitro* and 100 mg/kg *in vivo* daily. FITC-dextran (Sigma-Aldrich, FD40S) was directly diluted in the medium to 0.1 mg/mL. MTT (Thermo Fisher, M6494) was diluted in PBS (137 mM NaCl, 2.7 mM KCl, 12.5 mM Na₂HPO₄, 1.8 mM KH₂PO₄, pH 7.4) to 5 mg/mL. For high starvation, cells were washed twice with PBS and starved in low glucose DMEM lacking amino acids (USBiological, D9800-13) and serum. To overexpress the TFEB variant with pronounced nuclear localization characteristics [45] the plasmid pCIP-caTfeb (Addgene, 79,013; deposited by Reuben Shaw) was transfected into Huh-7 *VTRNA1-1* KO cells.

Cell proliferation and viability assay

Cell proliferation was detected using 3-(4,5-dimethylthazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT; Thermo Fisher, M6494) or via automated cell counting. For the MTT assay approximately 5×10^3 and 1.5×10^4 cells/well, for proliferation or viability respectively, were seeded into 96-well plates and allowed to attach overnight. After 24, 48, 72 and 96 h, cell viability was assessed using MTT assay. The MTT solution (5 mg/mL) was added to each well for 3 h. The resulting formazan crystals were dissolved in DMSO and the optical density was measured at 570 nm using a Tecan infinite M1000Pro plate reader. Three independent experiments were conducted in triplicates. To assess proliferation rates via automated cell counting, 1×10^5 cells/well were seeded on 6 well plates. After 24 h, 48 h and 72 h cells were washed twice with PBS, detached using trypsin and counted using a TC10™ Automated Cell Counter (Bio-Rad).

Colony formation assay

Approximately 5×10^2 cells were placed in a 6-well plate in DMEM containing 10% FBS for 2 weeks. Colonies were fixed with methanol and stained with 0.1% crystal violet in 20% methanol for 30 min. Each assay was conducted in triplicate.

RNA extraction and quantitative real-time RT-qPCR (qPCR)

RNA was isolated from cells as well as tissue samples by Tri-reagent (Zymo Research, R2050-1-200) following the manufacturer's protocol. Reverse transcription (RT) of total RNA was carried out using SuperScript™ IV One-Step RT-PCR System (Invitrogen, 18,090,010), according to the manufacturer's protocol, using random primer hexamers for cDNA amplification. Subsequent quantitative PCR (qPCR) on the cDNA was carried out using GoTaq® qPCR Master Mix (Promega, A6002), according to the manufacturer's protocol (Table S1 lists the used primers). The QIAgility robot was used to pipette all the reagents into the qPCR reactions and qPCR amplification was carried out using Rotor Gene 6000 (QIAGEN), according to the manufacturer's instructions. qPCR analyses were done in Robotix software (QIAGEN), and differential mRNA transcript abundances were calculated using the $\Delta\Delta C_t$ method as described [58].

Northern blot

2–10 µg total RNA was separated on an 8% denaturing polyacrylamide gel (7 M Urea, in 1x TBE buffer). The gel was subsequently electroblotted onto a nylon membrane (Amersham Hybond N⁺; GE Healthcare, RPN203B) as described [59]. The membranes were hybridized using ³²P-labeled probes listed in Table S1 and exposed to phosphorimaging screens. An autoradiogram was developed using a Typhoon FLA1000 phosphorimager.

Western blot

Total protein cell lysates were obtained using RIPA buffer (10 mM Tris/Cl, pH 8, 1 mM EDTA, 1% Triton X-100 [Sigma-Aldrich, T8787], 0.1% deoxycholate [Sigma-Aldrich, D6750], 0.1% SDS, 140 mM NaCl) containing protease inhibitor cocktail (Sigma-Aldrich, 11,697,498,001), phosphatase

inhibitors (Thermo Scientific, A32957) and 0.5% NP-40 (Sigma-Aldrich, 56,741). For nuclear-cytoplasmic fractionation, cells were treated as described previously [60]. Western blots were performed according to standard protocols using the antibodies listed in Table 1.

GST-assay

Huh-7 and HeLa cells were transiently transfected with the GST-BHMT plasmid as previously shown by Mercer et al. [61]. Briefly, BHMT transfected cells were cultured in starvation DMEM medium lacking the essential amino acids (Arg, Cys, His, Ile, Leu, Lys, Met, Phe, Thr, Trp, Tyr, and Val) (USBiological, D9800-27) and FBS for 6 h in the presence of E64d (6 mM; Sigma-Aldrich, E8640-250UG) and leupeptin (11 mM; Sigma-Aldrich, L-0649). Afterward, proteins from the cells were extracted with RIPA buffer. Total protein of cell extracts was determined by Bradford method (ThermoScientific, 23,227) and GST-BHMT was precipitated with glutathione agarose (Sigma, G4510). Precipitates were then mixed in SDS-PAGE sample buffer, resolved by SDS-PAGE and visualized by western blotting and chemiluminescence.

Lysosomal pH measurements by flow cytometry

Lysosomal pH measurements were based on an adjusted protocol recently published by Gavini et al [37]. In brief, HCC cell lines were seeded and cultured in cell culture medium containing FITC-Dextran (0.1 mg/mL; Sigma-Aldrich, FD40S) for three days. After this period FITC-Dextran was removed by aspirating the medium and fresh cell culture medium was added with the respective treatments as indicated. Lysosomal pH from samples belonging to the standard curve scale (pH ranging from 4 to 6) and experimental-treated ones were analyzed by FACS. Triplicate samples were taken for each condition analyzed.

Subcutaneous HCC cell line xenograft mouse models

Eight- to 12-week-old male *rag2*^{-/-} *il2rg*/*γ_c*^{-/-} mice (in house breeding) were used for the subcutaneous xenograft model. 2×10^6 HCC cells were injected in a 1:1 ratio with Matrigel® (BD Biosciences, 356,234). HCC patient-derived specimens were transplanted subcutaneously. Tumor volume was measured using a digital caliper thanks to a modified ellipsoid formula: volume = (4/3) × π × (length/2) × (width/2) × (height/2). Upon reaching a volume of 250 mm³, animals were randomly divided into different groups as described: administration of the vehicle (50% Cremophor EL and 50% ethyl alcohol mixture at 12–24 mg/mL – control group and sorafenib (100 mg/kg) daily by oral gavage. The tumor volume and the body weight were recorded every second day for fourteen days. At the end of experiment, the entire tumor was carefully removed, and some snap-frozen samples were stored in liquid nitrogen and kept at –80°C until further use. Other parts of the tissue were fixed in 4% formaldehyde for histological analysis. All animal experiments were conducted in accordance to Swiss Guidelines of Care and Use of Laboratory Animals.

Table 1. Antibodies used in this study.

Antibody	Source	Identifier
ACTB/β-actin-peroxidase	Sigma Aldrich	A2228
BCL2	Abcam	ab182858
MAPK1/ERK2- MAPK3/ERK1	Cell Signaling Technology	9911
GAPDH	Cell Signaling Technology	21185
GST (B-14)	Santa Cruz Biotechnology	sc-138
LAMP1	Santa Cruz Biotechnology	sc-17,768
LC3	Novus Biologicals	NB600-1384
MAP2K1/MEK1- MAP2K2/MEK2	Cell Signaling Technology	9911
MTOR	Cell Signaling Technology	29835
MKI67/Ki67	Abcam	ab16667
SQSTM1/p62	Cell Signaling Technology	80255
RPS6KA/p90RSK	Cell Signaling Technology	9911
PARP1	Santa Cruz Biotechnology	sc-8007
p-MTOR	Cell Signaling Technology	55365
TFEB	Cell Signaling Technology	4240
Histone H3	Millipore	06-570
TUBA4A/α-tubulin	Sigma Aldrich	T6074
TFE3	Proteintech	14,480-1-AP
MITF	Proteintech	13,092-1-AP
anti-rabbit IgG, HRP linked	Cell Signaling Technology	70745
anti-mouse IgG, HRP linked	Dako	P0260

Immunohistochemistry

Tissue samples were fixed in 4% formaldehyde, processed, embedded in paraffin and sectioned with Leica Microtome. Sections were stained using MKI67/Ki67 antibody (Abcam, ab16667). For each sample five randomly chosen fields were imaged by Leica camera and analyzed by ImageJ software. Coverslips were mounted with VECTASHIELD Antifade Mounting Medium and images were taken using an automated inverted microscope (Leica DMSI4000 B).

Statistical analysis

Images were quantified with ImageJ. Data are displayed as mean \pm SD or SEM; Student's t test was used; n values are indicated in the respective Figure legends. p-values are indicated in the Figure, $p < 0.05$ was considered statistically. GraphPad Prism v8 was used to create plots.

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Disclosure statement

The authors declare no competing interests.

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