

## Figure legends

**Figure 1 - Schematic illustration of our protocol combining *in silico* screening and experimental verification for identification of target protein.**

**Figure 2 - Experimental verification by *in vitro* biotinylated incednine pull-down assay.**

Lysates from Ms-1 overexpressing Bcl-xL were incubated with biotin (Biotin) or biotinylated incednine (BI) and avidin beads for 3 h. The beads were washed, and co-precipitated proteins were eluted with 2 mM biotin. The eluted proteins were subjected to western blotting using the indicated antibodies. “Positive candidates” means proteins which were predicted to bind to incednine, and “Negative candidates” means proteins which were predicted not to bind to incednine.

# Tables

**Table 1 List of proteins identified to bind to incednine in our previous binding experiments**

Protein	Uniprot ID	Kegg ID
poly 4- hydroxylase, beta submit	P07237	5034
N-acylaminoacyl peptide hydrolase	P13798	327
Heat shock protein 70	P08107	3303/3304
Protein Phosphatase A2	P67775	5515
Similar to DNA damage-binding protein 1	Q16531	1642
Deoxyhypusin synthase isoform alpha	P49366	1725
Methionine adenosyltransferase alpha/beta	P31153/Q00266/Q9NZL9	4144/4143/27430
4-alpha-glucanotransferase	P35573	178
Actin alpha 4	O43707	81
Eukaryotic Initiation factor 4A3	P38919	9775
Deoxycytidine kinase	P27707	1633
ATP synthase H <sup>+</sup> transporting, mitochondrial F1complex, alpha	P25705	498
prohibitin	P35232	5245
proteasome alpha 7subuit	O14818	5688
proteasome(prosome,macropain) subunit alpha type 8	Q8TAA3	143471
centaurin,beta 2	Q15057	23527
heterogeneous nuclear ribonucleoprotein A/B	Q99729	3182
heterogeneous nuclear ribonucleoprotein K	P61978	3190
heterogeneous nuclear ribonucleoprotein D	Q14103	3184
heterogeneous nuclear ribonucleoprotein A2/B1	P22626	3181
heterogeneous nuclear ribonucleoprotein A1	P09651	3178
heterogeneous nuclear ribonucleoprotein M	P52272	4670
small nuclear ribonucleoprotein polypeptide D2 family	P62316	6633
mitochondrial ribosomal protein L2	Q5T653	51069
mitochondrial ribosomal protein L20	Q9BYC9	55052
mitochondrial ribosomal protein L3	Q6IBT2	11222
mitochondrial ribosomal protein L40	Q9NQ50	64976
mitochondrial ribosomal protein L46	B2RD75	26589
mitochondrial ribosomal protein L49	B2R4G6	740
mitochondrial ribosomal protein L1	A6NG03	65008
mitochondrial ribosomal protein L37	Q9BZE1	51253
small nuclear ribonucleoprotein-associated protein B and B'	P14678	6628
cAMP-dependent protein kinase, regulatory subunit alpha 1	P10644	5573
phosphoribosyl pyrophosphate synthetase-associated protein 1	B2R6M4	5635
peptidylprolyl isomerase-like 2	Q13356	23759
thymoprotein isoform beta, gamma	P42167	7112
fructose-bisphosphate aldolase A	P04075	226
brain creatine kinase	P12277	1152
enolase 1	P06733	2023
Ewing sarcoma breakpoint region 1	Q5THL0	2130
fusion(involved in t(12;16) in malignant liposarcoma)	Q6IBQ5	2521
GDP dissociation inhibitor 2	Q5SX88	2665
nucleosome assembly protein 1-like 1	P55209	4673
nucleosome assembly protein 1-like 4	Q99733	4676
phosphoglycerate dehydrogenase	O43175	26227
triosephosphate isomerase 1	P60174	7167
clathrin heavy chain 1	Q00610	1213
clathrin heavy poly peptide -like 1	P53675	8218
glutamyl-prolyl tRNA synthetase	P07814	2058
retinoblastoma binding protein 7	Q16576	5931
retinoblastoma binding protein 4	Q09028	5928
tripartite motif-containing 28 protein	Q13263	10155
high glucose-regulated protein 8	Q9Y5A9	51441

**Table 2 Representative proteins selected from each cluster and negative candidates for experimental verification**

Cluster No.	Representative Protein
1	ITPR1 (inositol 1,4,5-triphosphate receptor, type 1)
2	DAPK1 (death-associated protein kinase 1)
3	PIK3CG (phosphoinositide-3-kinase, catalytic, gamma polypeptide), PIK3C2B (phosphoinositide-3-kinase, class 2, beta polypeptide)
4	PARP14 (poly (ADP-ribose) polymerase family, member 14)
5	PIP5K3 (phosphatidylinositol-3-phosphate/phosphatidylinositol 5-kinase, type III)
6	PLCB1 (phospholipase C, beta 1)
7	CHD4 (chromodomain helicase DNA binding protein 4)
8	KIF1A (kinesin family member 1A), KIF21B (kinesin family member 21B)
9	ACACA (acetyl-Coenzyme A carboxylase alpha)
10	GTF2IRD2 (GTF2I repeat domain containing 2)
11	RGPD5 (RANBP2-like and GRIP domain-containing protein 5)
Negative	Proteins predicted not to bind to incednine
1	BECN1 (Beclin-1)
2	PARP1 (poly (ADP-ribose) polymerase family, member 1)
3	KIF5B (kinesin family member 5B)

## **Additional files**

**Additional file 1 (Additional file1.pdf) – Validation work for eIF4A3, PDI, PP2A and Hsp70**

**Additional file 2 (Additional file2.pdf) – Proteins computationally predicted to bind to incednine (grouped into 11 clusters)**

**Additional file 3 (Additional file3.pdf) – A structure of biotinylated incednine**

**Additional file 4 (Additional file4.pdf) – Preparation of Flag-tagged Incednine**

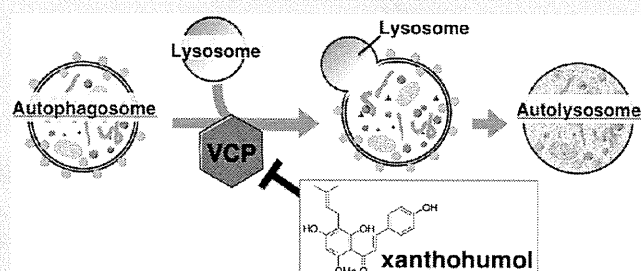
# Xanthohumol Impairs Autophagosome Maturation through Direct Inhibition of Valosin-Containing Protein

Yukiko Sasazawa,<sup>†,#</sup> Shuhei Kanagaki,<sup>†,#</sup> Etsu Tashiro,<sup>†</sup> Toshihiko Nogawa,<sup>‡</sup> Makoto Muroi,<sup>‡</sup> Yasumitsu Kondoh,<sup>‡</sup> Hiroyuki Osada,<sup>‡</sup> and Masaya Imoto<sup>\*,†</sup>

<sup>†</sup>Faculty of Science and Technology, Department of Biosciences and Informatics, Keio University, Yokohama 223-8522, Japan

<sup>‡</sup>Chemical Biology Core Facility, Chemical Biology Department, RIKEN Advanced Science Institute, Saitama 351-0198, Japan

**ABSTRACT:** Autophagy is a bulk, nonspecific protein degradation pathway that is involved in the pathogenesis of cancer and neurodegenerative disease. Here, we observed that xanthohumol (XN), a prenylated chalcone present in hops (*Humulus lupulus* L.) and beer, modulates autophagy. By using XN-immobilized beads, valosin-containing protein (VCP) was identified as a XN-binding protein. VCP has been reported to be an essential protein for autophagosome maturation. Using an *in vitro* pull down assay, we showed that XN bound directly to the N domain, which is known to mediate cofactor and substrate binding to VCP. These data indicated that XN inhibited the function of VCP, thereby allowing the impairment of autophagosome maturation and resulting in the accumulation of microtubule-associated protein 1 light chain 3-II (LC3-II). This is the first report demonstrating XN as a VCP inhibitor that binds directly to the N domain of VCP. Our finding that XN bound to and inactivated VCP not only reveals the molecular mechanism of XN-modulated autophagy but may also explain how XN exhibits various biological activities that have been reported previously.



Macroautophagy (herein referred to as autophagy) is an evolutionarily conserved pathway for degradation of intracellular components including organelles, which is critical for the maintenance of cellular homeostasis. Initially, the cytoplasmic components are sequestered by a unique membrane, referred to as an isolation membrane. Dynamic membrane organization is activated from small membrane particles to autophagosomes by the recruitment of autophagy related genes (ATGs) and microtubule-associated 1 light chain 3 (LC3).<sup>1</sup>

The next stage involves the fusion of autophagosomes with lysosomes and subsequent formation of autolysosomes. The inner membrane of the autophagosomes and the cytoplasm-derived materials contained in the autophagosomes are then degraded by lysosomal hydrolases.<sup>2</sup> The amino acids, which are produced by protein degradation, are then returned to the cytoplasm by lysosomal membrane permeases for reuse. Autophagy occurs in all cells at low basal levels under normal conditions to maintain homeostasis. It has been reported that aberrance of autophagy is involved in the pathogenesis of many diseases including neurodegenerative disease,<sup>3,4</sup> cancer,<sup>5</sup> muscle atrophy, and type 2 diabetes.<sup>6</sup>

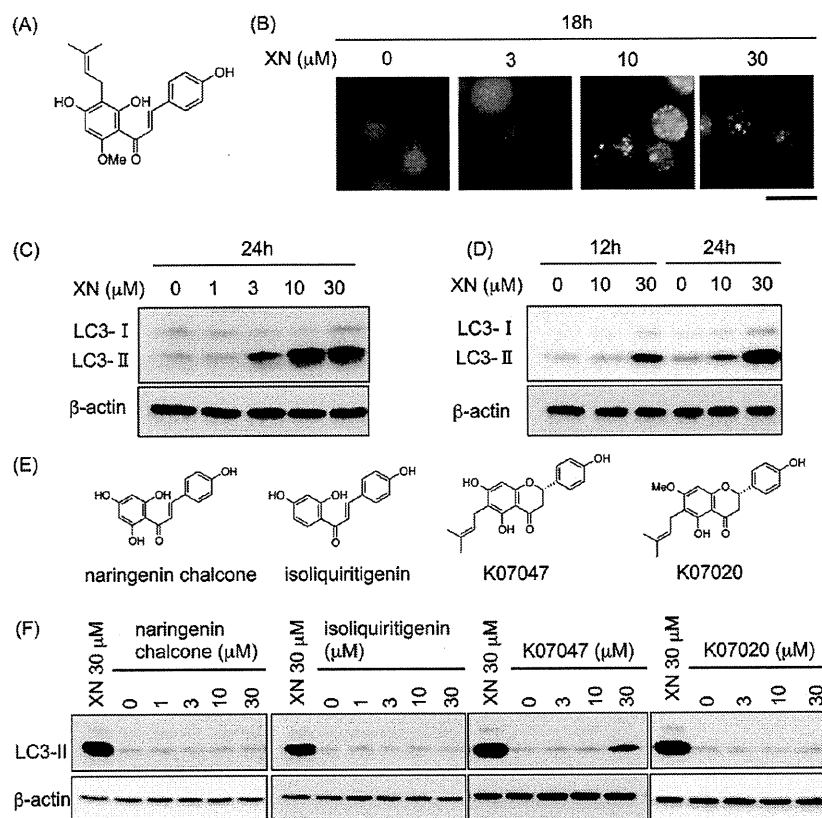
Despite identification of more than 30 ATGs,<sup>7,8</sup> the molecular mechanism of autophagy is still not fully understood. Studying autophagy through chemical genetics could be an ideal approach to gaining a better understanding of autophagy signaling pathways. Most compounds that have been reported to be regulators of autophagy are distributed between two major groups. One group induces autophagy by inhibiting

PI3K/Akt/mTOR signaling,<sup>9</sup> which is the major inhibitory signal that suppresses autophagy. The other group of regulators suppresses autophagy by inhibiting class III PI3K,<sup>10</sup> which is the homologue of yeast VPS34 and is required for the onset of autophagy.

In this study, we explored the mechanism of autophagy and identified additional small compounds that could modulate this process. This was done by screening for a small compound from an in-house natural product library using EGFP-LC3 stably expressing HeLa cells, and we identified xanthohumol (XN) as an autophagy modulator. Xanthohumol (30-[3,3-dimethyl allyl]-20,40,4-trihydroxy-60-methoxychalcone) is the principal prenylated chalcone of the female inflorescences of the hop plant ("hops"), an ingredient of beer.<sup>11</sup> Human exposure to XN is primarily through beer consumption. Several studies have reported on the potential health benefits of XN, including inhibition of diacylglycerol acyltransferase,<sup>12,13</sup> apoptosis induction,<sup>14</sup> NF-kappa B inhibition,<sup>15</sup> and ER stress induction.<sup>16</sup> However, there are no reports that show the relevance of XN to autophagy. Thus, to understand the mechanism by which XN modulates autophagy, we attempted to identify the target protein of XN responsible for the regulation of autophagy.

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**Figure 1.** XN modulated autophagy. (A) Structure of xanthohumol (XN). (B) GFP-microtubule-associated protein 1 light chain 3 (LC3) stably expressing HeLa cells were treated with various concentrations of XN for 18 h. Cells were fixed with 3% (w/v) paraformaldehyde and observed under a fluorescence microscope (scale bar, 20  $\mu\text{m}$ ). (C) HeLa cells were treated with various concentrations of XN for 24 h. Cell lysates were immunoblotted with anti-LC3B antibody.  $\beta$ -Actin was immunoblotted as a loading control. (D) A431 cells were treated with various concentrations of XN for the indicated time. Cell lysates were immunoblotted with anti-LC3B antibody.  $\beta$ -Actin was immunoblotted as a loading control. (E) Structures of naringenin chalcone, isoliquiritigenin, K07047, and K07020. (F) A431 cells were treated with various concentrations of naringenin chalcone, isoliquiritigenin, K07047, or K07020 for 24 h. Cell lysates were immunoblotted with anti-LC3B antibody.  $\beta$ -Actin was immunoblotted as a loading control.

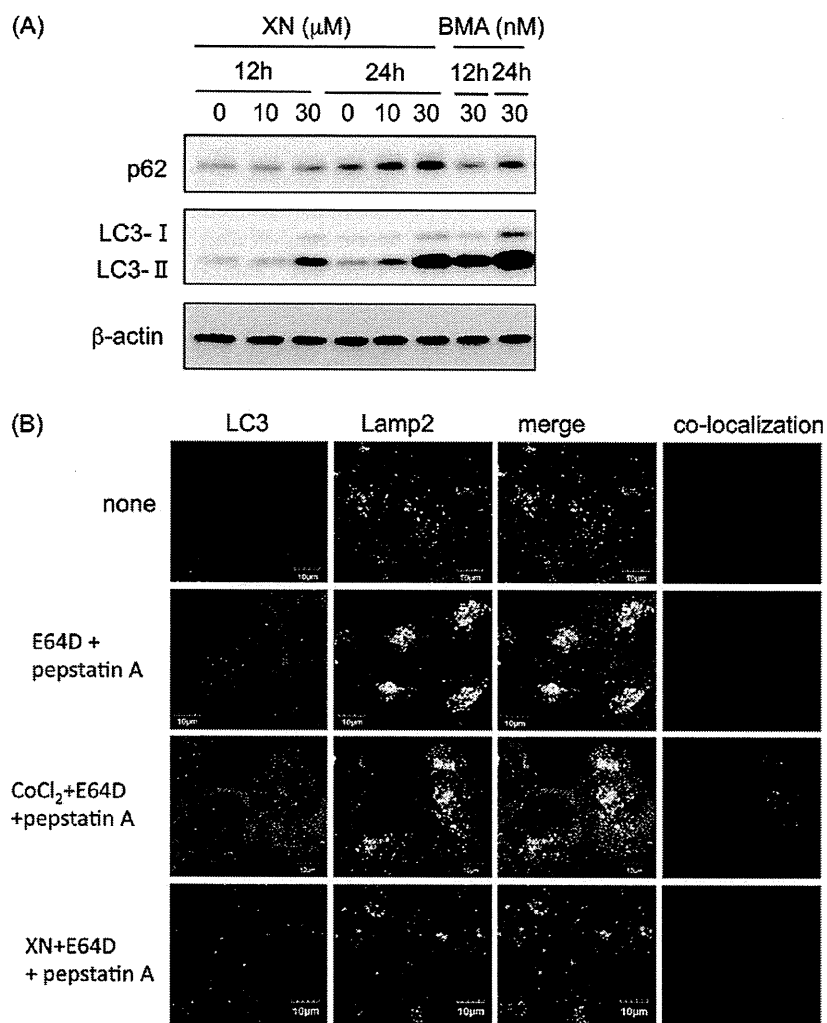
## RESULTS AND DISCUSSION

### Xanthohumol Inhibited Autophagosome Maturation.

In order to identify small compounds that could modulate autophagy and to explore the mechanism of autophagy through chemical genetics, we screened for a small compound from an in-house natural product library. As LC3-II is incorporated into the inner and outer surfaces of autophagosomes, the expression of a green fluorescence protein (GFP)-LC3 fusion protein can be used to identify GFP puncta representing autophagosomes.<sup>17</sup> Using this system to identify compounds that modulate autophagy, we searched for compounds that could increase the number of GFP-LC3 puncta in GFP-LC3 stably expressing human cervical carcinoma HeLa cells and found that xanthohumol (XN) showed this activity (Figure 1A). In untreated cells, GFP-LC3 was observed predominantly as diffuse green fluorescence in the cytoplasm. However, in XN-treated cells, characteristic punctate fluorescent patterns were observed, indicating that XN modulates autophagy in a dose-dependent manner, as shown in Figure 1B. Modulation of autophagy by XN was further confirmed by the detection of LC3-II, which is a phosphatidylethanolamine (PE) conjugated form of LC3, as a faster-migrating band when separated by SDS-PAGE and immunoblotted. As shown in Figure 1C, treatment of HeLa cells with XN for 24 h induced an increase in LC3-II levels in a dose-dependent manner. Similarly, XN increased LC3-II expression levels at 30  $\mu\text{M}$  over 12–24 h in

human epidermoid carcinoma A431 cells (Figure 1D). Next, we examined the effect of two other chalcones (naringenin chalcone and isoliquiritigenin) and two natural flavanones (K07047 and K07020) on LC3-II expression level. As a result, K07047 increased LC3-II levels weakly compared to XN, whereas naringenin chalcone, isoliquiritigenin, and K07020 did not increase LC3-II levels (Figure 1E,F)

The increase in LC3-II expression can be associated with either PE conjugation due to enhanced formation of autophagosomes or a block of LC3-II degradation due to impaired maturation of autophagosomes. To distinguish between these two possibilities, we detected expression levels of p62, a protein that is degraded by autophagy and accumulated when autophagy is impaired. Bafilomycin A1 (BMA) is known to prevent autophagosome maturation by inhibiting autophagosome-lysosome fusion<sup>18</sup> and caused an increase in the expression levels of p62 by inhibiting proteolytic degradation in autolysosomes, as shown in Figure 2A. Treatment with 30  $\mu\text{M}$  XN for 24 h increased the expression levels of p62 as well. These data suggested that the increased LC3-II expression mediated by XN was a consequence of a block of autophagosome maturation. To further confirm that XN inhibited autophagosome maturation, we detected the localization of LC3 and lysosome in the presence of pepstatin A plus E64D, which are the lysosomal protease inhibitors, to inhibit the degradation of LC3 after fusion of autophagosome



**Figure 2.** XN inhibited autophagosome maturation. (A) A431 cells were treated with various concentrations of XN or 30 nM bafilomycin A1 (BMA) for the indicated time. Cell lysates were immunoblotted with anti-p62 antibody.  $\beta$ -Actin was immunoblotted as a loading control. (B) A431 cells were treated with 30  $\mu\text{M}$  XN or 0.3 mM  $\text{CoCl}_2$  in the presence of 30  $\mu\text{M}$  E64D and 30  $\mu\text{M}$  pepstatin A for 24 h. Cells were then fixed with 3% (w/v) paraformaldehyde and immunostained with anti-LC3B and anti-lamp2 antibodies. The cells were observed under confocal microscopy (scale bar, 10  $\mu\text{m}$ ).

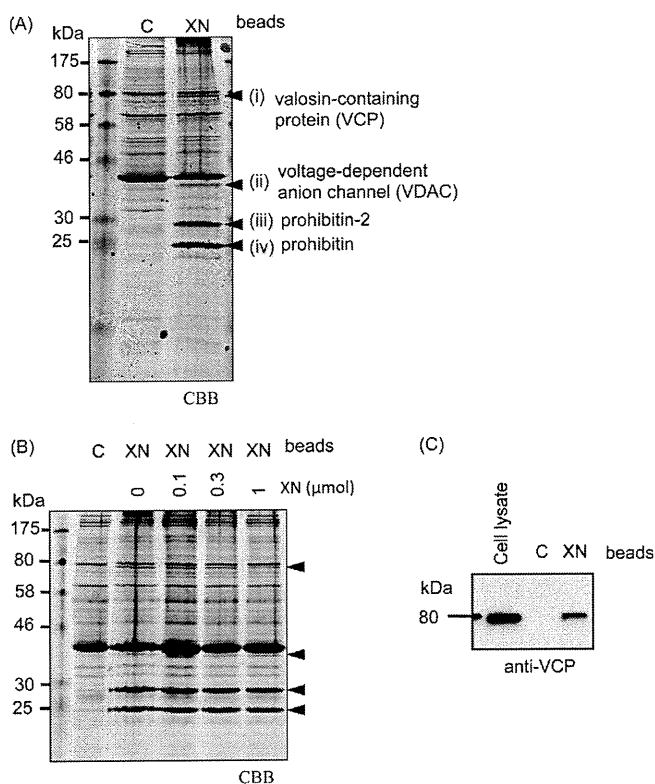
with lysosome. Because  $\text{CoCl}_2$  are known to induce an increase in LC3-II expression levels by accelerating autophagosome formation,<sup>19</sup> we examined the effect of XN on the localization of LC3 and lysosome compared with the effect of  $\text{CoCl}_2$ . As shown in Figure 2B,  $\text{CoCl}_2$  increased the number of LC3-positive puncta co-localizing with lysosome, whereas LC3-positive puncta increased by XN failed to co-localize with lysosome even in the presence of pepstatin A plus E64D. These data strongly indicated that XN impaired autophagosome maturation, resulting in increase in the level of LC3-II.

**Identification of XN-Binding Proteins.** To elucidate the underlying mechanism behind the suppression of autophagosome maturation induced by XN, we attempted to identify the cellular target protein of XN responsible for autophagy modulation. To this end, we used XN-immobilized agarose beads, which were prepared by a photocross-linking method.<sup>20</sup> A431 cell lysates were incubated for 3 h with XN-immobilized beads (XN beads) or control beads as a negative control. The reacted beads were washed, and the co-precipitated proteins were eluted, separated by SDS-PAGE, and stained with Coomassie brilliant blue (CBB). As shown in Figure 3A, four protein bands that specifically co-precipitated with XN beads

were observed. Each protein band was identified by using MALDI-TOF-MS and LC-MS/MS as (i) valosin-containing protein (VCP), (ii) voltage-dependent anion channel (VDAC), (iii) prohibitin-2, and (iv) prohibitin.

Among these proteins, competition was observed only for VCP with 0.1–1  $\mu\text{mol}$  XN as shown in Figure 3B. VCP has been reported to play a role in the maturation of autophagosomes.<sup>21,22</sup> VCP, also known as p97, is one of the best-characterized type II AAA (ATPases associated with diverse cellular activities) ATPases. VCP plays critical roles in a broad range of diverse cellular processes, including ER associated degradation *via* the ubiquitin-proteasome system,<sup>23,24</sup> cell cycle regulation,<sup>25</sup> and DNA repair.<sup>26</sup> Recently, it was reported that VCP is essential for autophagosome-lysosome fusion and formation of autolysosomes in human cell lines.<sup>21,22</sup> Therefore, we speculated that VCP might be the target of XN, and the binding of XN to VCP was confirmed by immunoblotting of co-precipitated protein from XN-beads using anti-VCP antibody (Figure 3C).

**XN Bound Directly to the N Domain of VCP.** Next, to determine whether XN could bind directly to VCP, we performed an *in vitro* binding assay using purified recombinant



**Figure 3.** Identification of XN-binding proteins. (A) A431 cell lysates were incubated with control beads or XN beads for 3 h. The reacted beads were washed, and the eluted proteins were subjected to SDS-PAGE and stained by Coomassie brilliant blue (CBB). The co-precipitated proteins for XN beads were identified by using MALDI-TOF-MS and LC-MS/MS. (B) A431 cell lysates were preincubated with 0.1–1  $\mu\text{mol}$  of XN as a competitor for 1 h and then incubated with control beads or XN beads for 3 h. The reacted beads were washed, and the eluted proteins were subjected to SDS-PAGE and stained by CBB. (C) A431 cell lysates were incubated with control beads or XN beads for 3 h. The reacted beads were washed, and the eluted proteins were immunoblotted with anti-valosin-containing protein (VCP) antibody.

GST-tagged VCP protein. Unlike GST, GST-VCP was co-precipitated only with XN-beads, as shown in Figure 4A. Moreover, competition was observed for VCP in the presence of 0.5  $\mu\text{mol}$  of XN (Figure 4B), indicating that XN binds directly to VCP. On the other hand, competition was not observed for binding of XN-beads and VCP in the presence of XN analogues such as naringenin chalcone, isoliquiritigenin, K07047, and K07020 up to 0.5  $\mu\text{mol}$  (Figure 4C), indicating that these analogues bind to VCP very weakly or fail to bind to VCP at least through the XN binding site. Because these analogues fail to induce LC3-II expression level or induce it very weakly, these observations further confirm the importance of the XN binding to VCP for impairment of autophagosome maturation. In addition, because naringenin chalcone and isoliquiritigenin did not bind to VCP, the prenyl and/or *O*-methyl group of XN is thought to be important for binding to VCP.

VCP is composed of a substrate and cofactor binding N domain followed by two AAA ATPase domains, termed D1 and D2, and forms a hexameric double-ring structure.<sup>27,28</sup> It has been demonstrated that both D1 and D2 domain contain Walker A and Walker B motifs that mediate ATP binding and hydrolysis, respectively. However, these two ATPase domains

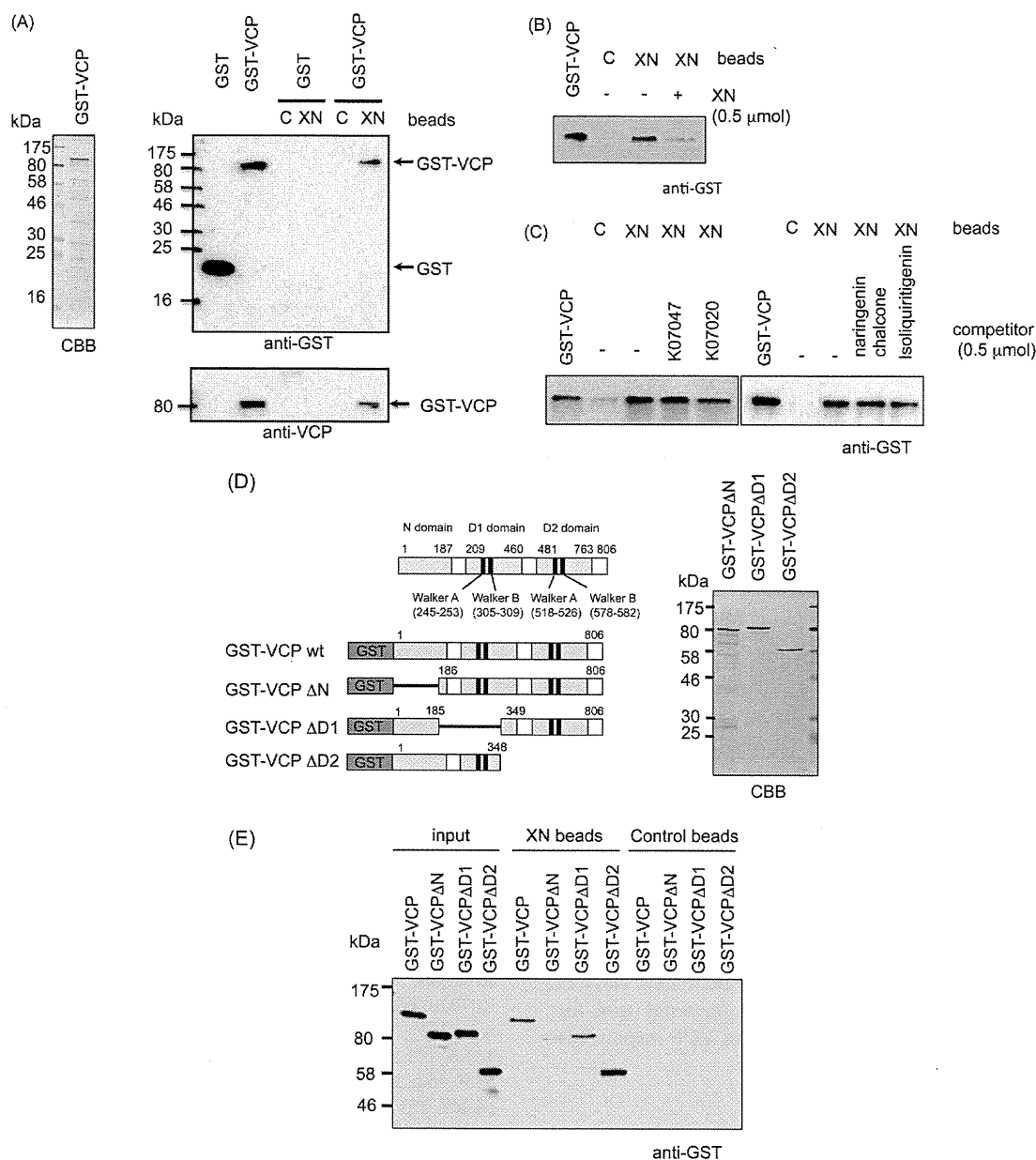
are not catalytically equivalent: D2 domain has the major ATPase activity at physiological temperatures, whereas D1 is involved in the regulation of heat-induced ATPase activity.<sup>29</sup> D1 also plays a major role in hexamerization.<sup>30,31</sup> To determine which domain in VCP is essential for interaction with XN, we prepared three GST-tagged VCP mutants that lacked the N-terminal domain (1–185) (GST-VCP $\Delta$ N), D1 domain containing Walker A and Walker B motifs (186–348) (GST-VCP $\Delta$ D1), and D2 domain containing Walker A and Walker B motifs (349–806) (GST-VCP $\Delta$ D2) (Figure 4D). GST-VCP $\Delta$ D1 and GST-VCP $\Delta$ D2 were co-precipitated with XN beads, whereas GST-VCP $\Delta$ N was not, as shown in Figure 4E. These results indicated that XN bound to the N domain of VCP.

**XN Inhibited VCP Function.** Next, we examined whether this binding of XN to the N domain of VCP could inhibit VCP function. The structural alteration of the N domain of VCP has been reported to induce impaired maturation of autophagosome as well as impaired ER associated degradation (ERAD),<sup>32</sup> and loss of VCP-mediated ERAD activity leads to accumulation of unfolded protein in the ER, resulting in induction of ER stress.<sup>33</sup> Therefore, we examined the effect of XN on the expression of the ER stress markers CHOP and GRP78. As shown in Figure 5A, treatment with 30  $\mu\text{M}$  XN for 12–24 h increased the protein levels of CHOP and GRP78 significantly in A431 cells, suggesting that XN inhibited VCP-mediated ERAD. On the other hand, Hirabayashi *et al.* reported that inhibition of VCP function by using dominant negative VCP induced cytoplasmic vacuolation.<sup>34</sup> These vacuoles are reported to be a result of abnormal budding and enlargement of the ER.<sup>35</sup> We also observed the presence of microscopic vacuoles not only in VCP knockdown A431 cells by using siRNA (Figure 5B) but also in XN-treated A431 cells (Figure 5C). The successful knockdown of VCP using siRNA and resultant up-regulation of LC3-II was confirmed by immunoblotting, as shown in Figure 5B right. Moreover, the XN analogue K07047, which modulated autophagy weakly, also induced vacuolization weakly compared with XN. On the other hand, other analogues including naringenin chalcone, isoliquiritigenin, and K07020, which had no effect on modulation of autophagy, did not induce vacuolization (Figures 1F and 5C). Taken together, these data indicated that XN bound to the N domain of VCP directly, thereby suppressing VCP function.

Apart from autophagy, XN has been reported to inhibit mitogen/antigen-induced T cell proliferation, development of cell-mediated cytotoxicity, and production of Th1 cytokines by inhibiting NF- $\kappa$ B.<sup>36</sup> Moreover, XN has been shown to inhibit the growth of a wide variety of human cancer cell lines by inhibiting proliferation and inducing apoptosis.<sup>37,38</sup> These previous observations regarding XN suggested the following two possibilities: one possible explanation is that various proteins were interfered with by XN and various biological phenomena were affected, and the other is that XN modulated a specific protein, which was involved in the various biological processes. Our finding that XN modulated the function of VCP may explain how XN exhibited the above-mentioned effects, because VCP is reported to play important roles in the degradation of I $\kappa$ B, resulting in enhancement of NF- $\kappa$ B signaling,<sup>39,40</sup> or because the expression level of VCP is correlated with progression, prognosis, and recurrence of certain types of cancer.<sup>41,42</sup>

Two types of VCP inhibitors have been reported in the literature. The first type of inhibitor is classified as a VCP

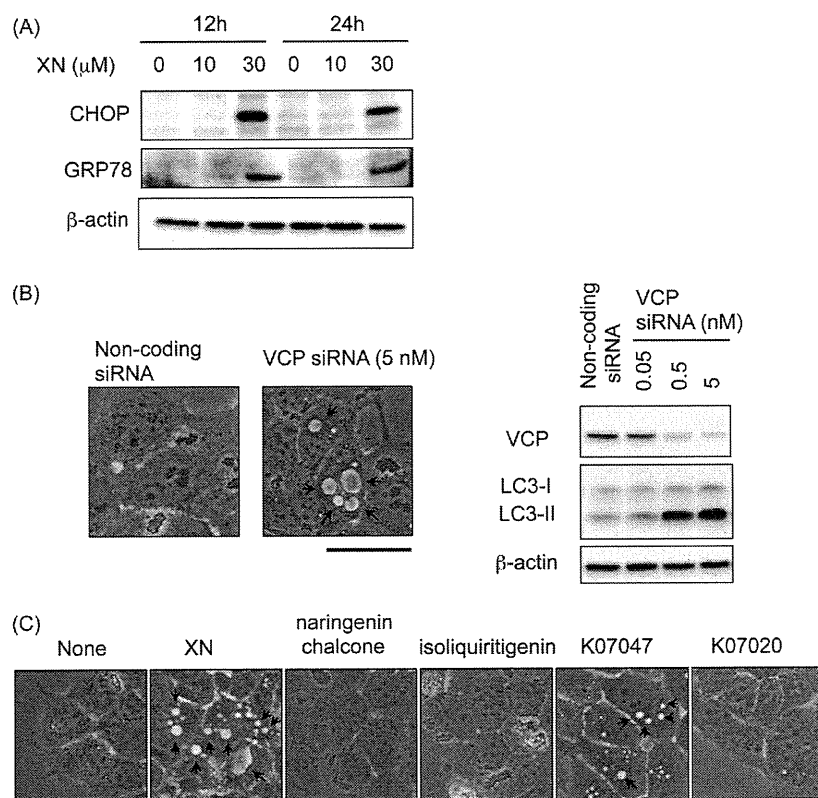




**Figure 4.** XN bound directly to the N domain of VCP. (A) (left) CBB staining of purified GST-VCP protein. (right) Purified GST and GST-tagged VCP were incubated with control beads or XN beads for 3 h. The reacted beads were washed, and the eluted proteins were immunoblotted with anti-GST and anti-VCP antibodies. (B) Purified GST-tagged VCP was preincubated with 0.5  $\mu\text{mol}$  of XN as a competitor for 1 h and then incubated with control beads or XN beads for 3 h. The reacted beads were washed, and the eluted proteins were immunoblotted with anti-GST antibody. (C) Purified GST-tagged VCP was preincubated with 0.5  $\mu\text{mol}$  of naringenin, chalcone, isoliquiritigenin, K07047, or K07020 as a competitor for 1 h and then incubated with control beads or XN beads for 3 h. The reacted beads were washed, and the eluted proteins were immunoblotted with anti-GST antibody. (D) (left) Schematic illustration of GST-VCP, GST-VCP $\Delta\text{N}$ , GST-VCP $\Delta\text{D1}$ , and GST-VCP $\Delta\text{D2}$ . (right) CBB staining of purified GST-VCP $\Delta\text{N}$ , GST-VCP $\Delta\text{D1}$ , and GST-VCP $\Delta\text{D2}$ . (E) Purified GST-VCP $\Delta\text{N}$ , GST-VCP $\Delta\text{D1}$ , and GST-VCP $\Delta\text{D2}$  were incubated with control beads or XN beads for 3 h. The reacted beads were washed, and the eluted proteins were immunoblotted with anti-GST antibody.

ATPase inhibitor, which most likely binds to a site in the D2 ATPase domain. 2-Anilino-4-aryl-1,3-thiazoles were discovered by high-throughput screening (HTS) as inhibitors of VCP ATPase activity, and these were reported to inhibit VCP-associated protein degradation.<sup>43</sup> Syk inhibitor III was reported to be an irreversible inhibitor of VCP ATPase activity by interacting with Cys522 within the D2 ATPase domain of VCP and the ubiquitin-fused reporter protein.<sup>44</sup>  $N^2,N^4$ -Dibenzylquinazoline-2,4-diamine (DBeQ) was identified as a selective, potent, reversible, and ATP-competitive VCP inhibitor by screening a library of chemical compounds.<sup>45</sup> DBeQ blocks

multiple processes that have been shown by siRNA to depend on VCP, including degradation of ubiquitin fusion degradation and ERAD as well as autophagosomal maturation. The second type of VCP inhibitor is Eeyarestatin I (Eer I), which binds to the D1 domain of VCP without affecting ATPase activity.<sup>46</sup> Eer I was found to directly associate with the ER membrane and VCP and inhibited VCP-associated deubiquitinating enzymes, thereby inhibiting VCP-dependent protein degradation. However, so far, VCP inhibitors that bind to the N domain of VCP have not yet been reported. Therefore, XN is the first example of such an inhibitor that binds to the N domain of VCP and



**Figure 5.** XN inhibited VCP function. (A) A431 cells were treated with various concentrations of XN for the indicated time. Cell lysates were immunoblotted with anti-CHOP and anti-GRP78 antibodies.  $\beta$ -Actin was immunoblotted as a loading control. (B) (left) A431 cells were observed under a microscope 72 h after transfection with non-coding siRNA or VCP siRNA (arrows, vacuoles; scale bar, 25  $\mu$ m). (right) A431 cells were transfected with VCP siRNA or noncoding siRNA for 72 h. Cell lysates were immunoblotted with anti-VCP and anti-LC3B antibodies.  $\beta$ -Actin was immunoblotted as a loading control. (C) A431 cells were treated with 30  $\mu$ M XN, naringenin chalcone, isoliquiritigenin, K07047, or K07020 for 24 h and then observed under microscope (arrows, vacuoles; scale bar, 25  $\mu$ m).

inactivates VCP. Thus, XN is proposed to be a new class of VCP inhibitors, which may be used as a powerful tool for identifying the cofactor or substrate protein of VCP responsible for autophagy regulation.

## METHODS

**Reagents.** Naringenin chalcone was obtained as a generous gift from Kikkoman Corporation. Isoliquiritigenin, E64D, and peptatin A were purchased from Sigma-Aldrich Co.

**Cell Line.** Human epidermoid carcinoma A431 cells were grown in Dulbecco's modified Eagle medium supplemented with 5% (v/v) calf serum, 100 U mL<sup>-1</sup> of penicillin G (Sigma-Aldrich Co.), and 0.1 mg mL<sup>-1</sup> of kanamycin (Sigma-Aldrich Co.) at 37 °C in a 5% CO<sub>2</sub>–95% air atmosphere. Human cervical carcinoma HeLa cells were grown in Dulbecco's modified Eagle's medium supplemented with 8% fetal bovine serum, 100 U mL<sup>-1</sup> of penicillin G, and 0.1 mg mL<sup>-1</sup> of kanamycin at 37 °C in a 5% CO<sub>2</sub>–95% air atmosphere. HeLa/GFP-LC3 stable cell lines were established as previously described.<sup>47</sup>

**Isolation of XN from Hop and XN Analogues from Microbial Origin.** XN was isolated from commercially available hop extract obtained from Hopsteiner. The extract (200 mg) was purified by using preparative octadecylsilyl (ODS) HPLC (UG 80, 20 mm, 250 mm; SHISEIDO) with 70% (v/v) aqueous MeOH to obtain pure XN (147 mg). The structures were identified by spectroscopic data (NMR and MS). UV (MeOH)  $\lambda_{\max}$  (log  $\epsilon$ ) 369 (4.56); ESIMS  $m/z$  355 [M + H]<sup>+</sup>; <sup>13</sup>C NMR  $\delta$ 192.8, 165.5, 161.8, 161.2, 157.4, 142.0, 136.0, 130.3 (2C), 128.6, 125.5, 121.7, 116.2 (2C), 106.3, 106.2, 105.0, 56.1, 25.8, 21.6, and 17.9. K07020 and K07047 were isolated from 14 L of culture broths of *Streptomyces* sp. HK-803 and *Streptomyces spiroverticillatus* JC-8444 by UV absorption and mass spectra guided separation to afford 10.6 and 9.3 mg as a pale-yellow powder, respectively. The

structures were identified by spectroscopic data (NMR and MS). K07020: UV (MeOH)  $\lambda_{\max}$  (log  $\epsilon$ ) 226 (4.45), 288 nm (4.33); ESIMS  $m/z$  353 [M – H]<sup>-</sup>; <sup>13</sup>C NMR  $\delta$ 192.9, 164.6, 163.8, 161.8, 158.8, 131.6, 131.5, 128.9 (2C), 123.9, 116.2 (2C), 110.0, 105.7, 93.5, 80.0, 55.9, 46.2, 26.0, 22.7, and 17.9. K07047: UV (MeOH)  $\lambda_{\max}$  (log  $\epsilon$ ) 226 (4.36), 293 nm (4.24); ESIMS  $m/z$  339 [M – H]<sup>-</sup>; <sup>13</sup>C NMR  $\delta$ 197.8, 165.9, 162.6, 162.5, 159.0, 131.5, 131.2, 129.0 (2C), 123.9, 116.3 (2C), 109.6, 103.2, 95.4, 80.4, 44.2, 25.9, 21.8, and 17.8.

**DNA Constructs.** Human cDNA for VCP were amplified from A431 cell cDNA and subcloned into pGEX-2T (GE Healthcare UK Ltd.) to prepare GST fusion proteins in bacteria. Expression vectors encoding GST-fused VCP mutants ( $\Delta$ N, 1–185 aa deletion;  $\Delta$ D<sub>1</sub>, 186–348 aa deletion; and  $\Delta$ D<sub>2</sub>, 349–806 aa deletion) were generated by PCR using pGEX-2T/VCP as a template.

**Fluorescence Microscopy.** For fluorescence microscopy, HeLa cells stably expressing GFP-LC3, which were grown on coverslips, were treated with chemicals for the indicated time at 37 °C. Cells were fixed with 3% (w/v) paraformaldehyde in PBS at RT. The cells were then washed with PBS and observed under a fluorescence microscope (Olympus).

**Western blotting.** Cells were lysed with RIPA buffer [25 mM HEPES, 1.5% (v/v) TX-100, 1% (w/v) sodium deoxycholate, 0.1% (w/v) SDS, 0.5 M NaCl, 5 mM EDTA, 50 mM NaF, 100 mM Na<sub>3</sub>VO<sub>4</sub>, 0.1 mg mL<sup>-1</sup> leupeptin, 1 mM PMSF; pH 7.8]. Proteins were separated by SDS-PAGE, transferred to a PVDF membrane (Millipore), and probed with specific antibodies. This was followed by detection using the ECL Western blotting detection system (Millipore) and LAS-1000 (Fuji Film). The primary antibodies used were as follows: anti-LC3B (L7543, Sigma-Aldrich Co.), anti- $\beta$ -actin (AC-74, Sigma-Aldrich Co.), anti-p62 (5114, Cell Signaling Technology), anti-VCP (ab 11433, Abcam), anti-GST (B-14, Santa

Cruz Biotechnology), anti-GRP78 (H-129, Santa Cruz Biotechnology), and anti-CHOP (MA1-250, Thermo Fisher Scientific Inc.) antibodies. The secondary antibodies were horseradish peroxidase-conjugated anti-mouse IgG and anti-rabbit IgG (GE Healthcare UK Ltd.).

**Immunofluorescent Microscopy.** Immunofluorescent microscopy was carried out as previously described.<sup>48</sup> Fluorescence images were obtained using a confocal laser scanning microscope system FV1000 (Olympus).

**Detection of Binding Proteins for XN Beads.** XN beads were prepared as previously described.<sup>20</sup> A431 cells were harvested, washed with PBS, and then resuspended in binding buffer [50 mM HEPES, 150 mM NaCl, 2.5 mM EGTA, 1 mM EDTA, 1 mM DTT, 0.1 mM PMSF, NP40 1% (v/v) and protease inhibitor cocktail tablets (Roche); pH 7.5]. After cells were lysed by homogenization with sonication, the insoluble material was removed by centrifugation, and the supernatant was collected as cell lysate. The cell lysate (3 mg of protein) was then incubated with XN beads (20  $\mu$ L) for 3 h at 4 °C. The reacted beads were washed with binding buffer, and the binding proteins were eluted with SDS-PAGE sample buffer, separated by SDS-PAGE, and visualized by CBB staining. Identification of the proteins was performed using MALDI-TOF-MS and LC-MS/MS as previously described.<sup>49</sup>

**In Vitro XN Beads Pull-Down Assay.** GST fusion proteins, which were expressed in the *Escherichia coli* BL21 strain and purified using Glutathione Sepharose 4B (GE Healthcare UK Ltd.), were incubated with XN beads in 1 mL of binding buffer for 3 h. The beads were washed with binding buffer and eluted with SDS-PAGE sample buffer. The eluted proteins were then subjected to SDS-PAGE. For the competition assay, each compound was added 1 h before incubation with XN beads.

**RNA Interference.** siRNA double-stranded oligonucleotides designed to interfere with the expression of VCP (sense 5'-UAGAACAGAACUCCCUUGGAAGGUG-3'; Invitrogen) and non-coding siRNA (Invitrogen) as a negative control were used. Reverse transfection was demonstrated by using HiPerFect (QIAGEN) according to the manufacturer's instructions. Briefly, A431 cells were trypsinized, resuspended in antibiotic-free medium, mixed with OPTI-MEM (Gibco) including siRNA and HiPerFect, and then seeded onto a 12-well plate. 72 h after transfection, cells were observed under microscope and lysed for Western blotting.

## ■ ASSOCIATED CONTENT

### Accession Codes

Uni-Prot accession codes are described as following; valosion-containing protein (TERA\_HUMAN), P55072; voltage-dependent anion channel (VDAC1\_HUMAN), P21796; Prohibitin 2 (PHB2\_HUMAN), Q99623; Prohibitin (PHB\_HUMAN), P35232

## ■ AUTHOR INFORMATION

### Corresponding Author

\*E-mail: imoto@bio.keio.ac.jp.

### Author Contributions

<sup>#</sup>These authors contributed equally to this work.

### Notes

The authors declare no competing financial interest.

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## ORIGINAL ARTICLE

# Inostamycin enhanced TRAIL-induced apoptosis through DR5 upregulation on the cell surface

Kohta Yamamoto<sup>1</sup>, Masafumi Makino<sup>1</sup>, Ramida Watanapokasin<sup>2</sup>, Etsu Tashiro<sup>1</sup> and Masaya Imoto<sup>1</sup>

Tumor necrosis factor-related apoptosis-inducing ligand (TRAIL) has been considered as a possible therapeutic agent for cancer treatment. This is because of its selective cytotoxicity against various cancer cells without a detrimental effect on normal cells. However, recent studies have reported that the potential application of TRAIL in cancer therapy is limited, as many cancer cells have been found to be resistant to TRAIL. Therefore, small molecule compounds that potentiate the cytotoxicity of TRAIL would be strategic candidates for therapeutic applications in combination with TRAIL. Here we found that a combined treatment of inostamycin and TRAIL synergistically induced caspase-dependent apoptosis in HCT116 cells. Inostamycin upregulated DR5, and a knockdown of DR5 suppressed the apoptosis that was synergistically induced by co-treatment with inostamycin and TRAIL. Moreover, inostamycin increased the expression of DR5 on the cell surface. Therefore, inostamycin-increased cell surface expression of DR5 may have contributed to the enhancement of TRAIL-induced apoptosis. Our study suggests that combined treatment with inostamycin and TRAIL may offer a strategy to overcome TRAIL resistance in tumor cells.

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**Keywords:** apoptosis; DR5; inostamycin; TRAIL

## INTRODUCTION

Tumor necrosis factor-related apoptosis-inducing ligand (TRAIL), also known as Apo2L, is a type II transmembrane protein. TRAIL was originally identified on the basis of sequence homology to the Fas ligand and tumor necrosis factor.<sup>1,2</sup> TRAIL is hypothesized as being a potentially good therapeutic agent for cancer treatment. This is because TRAIL has been shown to induce apoptosis in a variety of tumor cell lines more efficiently than in normal cells.<sup>3–5</sup> TRAIL exerts its function by binding its receptors, which are expressed on the surface of target cells. To date, four different types of membrane-bound death receptors for TRAIL has been identified: TRAIL-R1/DR4, TRAIL-R2/DR5, TRAIL-R3/DcR1 and TRAIL-R4/DcR2. Both DR4 and DR5 have a conserved cytoplasmic region called the 'death domain'. This is a homolog to Fas and tumor necrosis factor-R1, and is required for TRAIL-induced apoptosis.<sup>6–8</sup> On the other hand, TRAIL also binds to DcR1 and DcR2 that sequester the ligand, but are unable to initiate an apoptosis signal. Thus, DcR1 and DcR2 are considered to be the decoy receptors.<sup>7,9</sup> The apoptosis signal that is induced by TRAIL has been shown to be similar to that which is induced by Fas. The TRAIL homotrimer induces trimerization of DR5 or DR4 on the surface of target cells, which leads to the formation of death-inducing signaling complex. The trimerization of the death domains results in the recruitment of an adaptor molecule Fas-associated protein with death domain (FADD), which in turn

recruits and activates caspase-8. In type I cells, activation of caspase-8 is sufficient for the subsequent activation of effector caspase-3. On the other hand, in type II cells, amplification through the mitochondrial pathway is initiated by the cleavage of Bid by caspase-8. Truncated-Bid causes the loss of mitochondrial membrane potential and caspase-9 cleavage, resulting in the activation of caspase-3 and subsequent cellular apoptosis.

The potential application of TRAIL in cancer therapy is currently limited, as many cancer cells have been found to be resistant to the cytotoxic effects of TRAIL. This resistance may be because of the low expression of pro-apoptotic molecules: death receptors or caspase-8. Alternatively, the resistance may be because of the high expression of anti-apoptotic molecules: decoy receptors, FLICE-like-inhibitory protein, inhibitor of apoptosis protein and Bcl-2. Because of the limitations of TRAIL-induced cytotoxicity, the combination of TRAIL with other small molecule compounds has been postulated as strategy to potentiate the cytotoxicity of TRAIL and its therapeutic applications. Indeed, it was reported that several chemotherapeutic agents and natural products, such as CDDP,<sup>10</sup> etoposide,<sup>10,11</sup> doxorubicin,<sup>12</sup> PS-341 (bortezomib),<sup>13</sup> tunicamycin,<sup>14</sup> rottlerin,<sup>15</sup> brandisianins,<sup>16</sup> silibinin<sup>17</sup> and sodium butyrate,<sup>18</sup> were succeeded to cause the sensitization of TRAIL-resistant tumor cells to TRAIL-induced apoptosis.

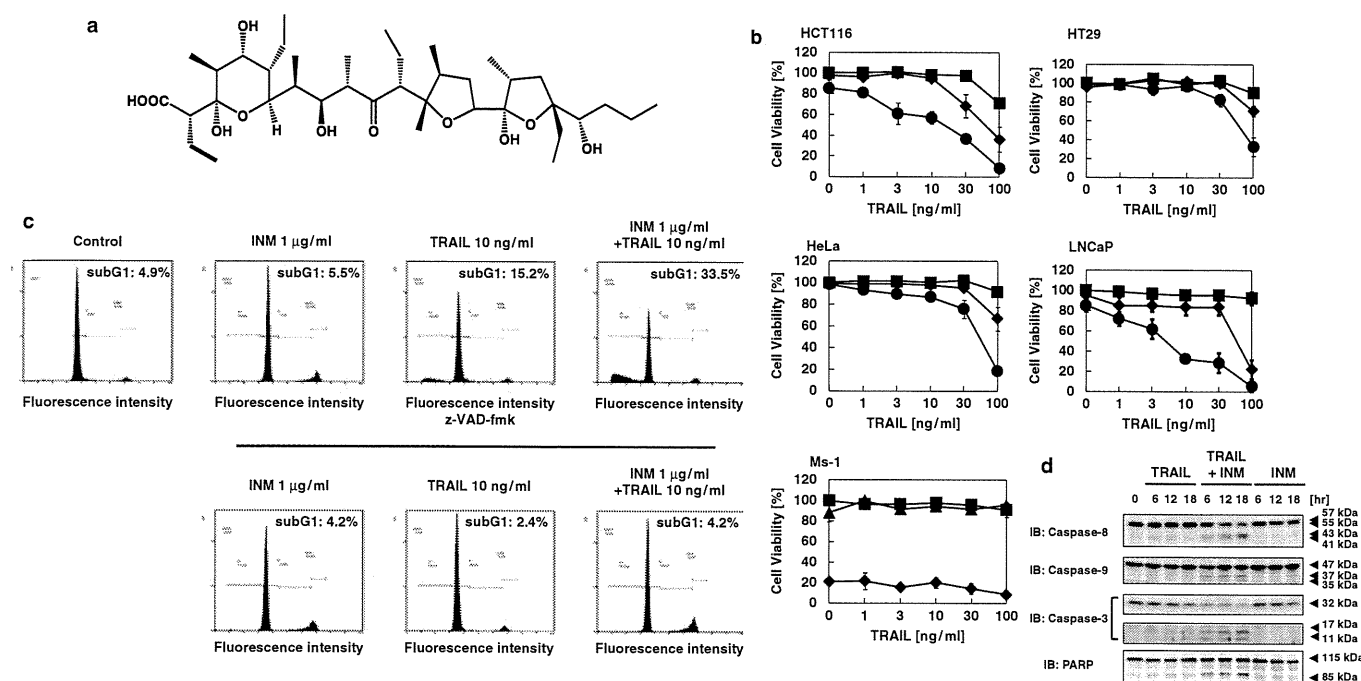
In this study, we have screened candidate small molecule compounds that synergistically induce apoptosis in the presence of

<sup>1</sup>Department of Biosciences and Informatics, Faculty of Science and Technology, Keio University, Yokohama, Japan and <sup>2</sup>Department of Biochemistry, Faculty of Medicine, Srinakharinwirot University, Bangkok, Thailand

Correspondence: Dr E Tashiro, Department of Biosciences and Informatics, Faculty of Science and Technology, Keio University, 3-14-1 Hiyoshi, Kohoku-ku, Yokohama, 223-8522, Japan.

E-mail: tashiro@bio.keio.ac.jp

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**Figure 1** Co-treatment with inostamycin (INM) and tumor necrosis factor-related apoptosis-inducing ligand (TRAIL) synergistically induced apoptosis in HCT116 cells. (a) Structure of inostamycin.<sup>19</sup> (b) Co-treatment with INM and TRAIL induced a synergistic cytotoxicity in HCT116 cells, HeLa cells, HT29 cells and LNCaP cells, but not in Ms-1 cells. HCT116 cells, HeLa cells, HT29 cells, LNCaP cells or Ms-1 cells were treated with the indicated concentration of TRAIL and 0.0 (square), 0.01 (triangle), 0.1 (diamond-shaped) or 1 (circle)  $\mu$ g ml<sup>-1</sup> of INM for 18 h. The cytotoxicity was evaluated using a Trypan Blue Exclusion assay. (c) The cytotoxicity induced by the co-treatment with INM and TRAIL was completely suppressed by z-VAD-fmk. HCT116 cells were pre-treated with or without 100  $\mu$ M of z-VAD-fmk, and then treated further with 10 ng ml<sup>-1</sup> of TRAIL and 1  $\mu$ g ml<sup>-1</sup> of INM. After an 18-h incubation period, the sub-G<sub>1</sub> population was measured by flow cytometer. (d) Co-treatment with INM and TRAIL activated caspase. HCT116 cells were treated with or without 10 ng ml<sup>-1</sup> of TRAIL and 1  $\mu$ g ml<sup>-1</sup> of INM for the indicated periods. Cell extracts were prepared for western blotting to detect the expression of caspase-8, caspase-9, caspase-3 and poly(ADP-ribose) polymerase (PARP).

TRAIL. As a result, we found that co-treatment with inostamycin and TRAIL synergistically induced caspase-dependent apoptosis in colorectal cancer HCT116 cells. Inostamycin was isolated from *Streptomyces* species. MH816-AF15,<sup>19</sup> its structure was shown in Figure 1a. Inostamycin weakly upregulated DR4 and strongly upregulated DR5 expression. Additionally, the synergistic apoptosis that was induced by this co-treatment was completely suppressed in DR5 small interfering RNA (siRNA)-transfected but not in DR4 siRNA-transfected HCT116 cells. Furthermore, inostamycin increased cell surface expression of DR5. Taken together with these results, upregulation of cell surface DR5 by inostamycin possibly contributes to the enhancement of TRAIL-induced apoptosis.

## MATERIALS AND METHODS

### Materials

Inostamycin was prepared as previously described,<sup>19</sup> TRAIL was purchased from Millipore (Billerica, MA, USA). Tunicamycin and z-VAD-fmk were purchased from SIGMA (St. Louis, MO, USA). Rabbit monoclonal anti-poly(ADP-ribose) polymerase, rabbit polyclonal anti-caspase-9, anti-caspase-3, anti-Bcl-xL and mouse monoclonal anti-caspase-8 were purchased from Cell Signaling Technology (Beverly, MA, USA). Rabbit polyclonal anti-survivin was purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). Mouse monoclonal anti-X-linked inhibitor of apoptosis protein (XIAP) was purchased from BD Transduction Laboratories (San Diego, CA, USA). Mouse monoclonal anti-Bcl-2 was purchased from Dako (Carpentaria, CA, USA). Rabbit polyclonal anti-DR5 was purchased from Proscience (Poway, CA, USA). Rabbit polyclonal anti-DR4 was purchased from Millipore. Mouse monoclonal anti-C/EBP homologous protein (CHOP) was purchased from Thermo scientific (Rockford, IL, USA). Mouse monoclonal anti-DR5 phycoerythrin (PE)-conjugated was purchased from eBioscience (San Diego,

CA, USA). Horseradish peroxidase-conjugated anti-mouse IgG and anti-rabbit IgG secondary antibodies were purchased from GE Healthcare (Little Chalfont, UK).

### Cell culture

Human colorectal carcinoma HCT116 cells, HT29 cells and human small cell lung cancer Ms-1 cells were maintained in Roswell Park Memorial Institute medium (Nissui, Tokyo, Japan) supplemented with 5% fetal bovine serum. Human cervical cancer HeLa cells were maintained in Dulbecco's modified Eagle's medium (Nissui, Tokyo, Japan) supplemented with 8% fetal bovine serum. Human prostate adenocarcinoma LNCaP cells were maintained in Roswell Park Memorial Institute medium supplemented with 10% fetal bovine serum—Roswell Park Memorial Institute medium.

### Trypan blue exclusion assay

HCT116 cells, HT29 cells, HeLa cells, LNCaP cells or Ms-1 cells were seeded in 48-well plates and incubated overnight, before experiments were conducted. The cells were treated with TRAIL with or without inostamycin for 18 h. Floating and adherent cells were collected and resuspended in phosphate-buffered saline (PBS<sup>-</sup>). Cells were mixed with trypan blue and counted under the microscope.

### Propidium iodide staining

Propidium iodide staining and flow cytometry were used to determine the degree of cellular apoptosis. HCT116 cells were seeded in 6-well plates and incubated overnight before experiment. The cells were treated with TRAIL with or without inostamycin for 18 h. Floating and adherent cells were collected and resuspended in PBS<sup>-</sup>. Cold ethanol was added in a dropwise manner while vortexing to fix the cells. Fixed cells were collected and resuspended in PBS<sup>-</sup> containing 50  $\mu$ g ml<sup>-1</sup> propidium iodide. Flow cytometry was done using

EPICS (Beckman Coulter, Fullerton CA, USA). The percentage of sub-G<sub>1</sub> cells was used to quantify apoptotic cells.

### Western blotting

Cells were lysed with a lysis buffer containing 25 mM HEPES (pH 7.8), 1.5% Triton-X 100, 1.0% sodium deoxycholate, 0.1% SDS, 0.5 M NaCl, 5 mM EDTA, 50 mM NaF, 100 μM Na<sub>3</sub>VO<sub>4</sub>, 0.1 mg ml<sup>-1</sup> leupeptin and 1 mM phenylmethylsulfonyl fluoride. Proteins were separated by SDS-polyacrylamide gel electrophoresis and transferred to a polyvinylidene fluoride membrane (Millipore). After the membranes had been incubated with primary and secondary antibodies, the immune complexes were detected with an Immobilon Western kit (Millipore), and the luminescence was detected with a LAS-1000 mini (Fujifilm, Tokyo, Japan).

### Real-time reverse transcriptase (RT)-PCR analysis

Total RNA was isolated with TRIzol (Invitrogen, Carlsbad, CA, USA). Reverse transcription was performed with M-MLV reverse transcriptase (PROMEGA, Madison, WI, USA) according to the manufacturer's instructions. Real-time reverse transcription (RT)-PCR was performed using SYBR Premix Ex Taq (Takara, Shiga, Japan). Primer sequence was as follows: DR5, 5'-CACCA GGTGTGATTACAGTG-3' (sense) and 5'-TACGGCTGCAACTGTGACTC-3' (antisense); CHOP, 5'-GCGCATGAAGGAGAAAGAAC-3' (sense) and 5'-TCACCATTCGGTCAATCAGA-3' (antisense); glyceraldehyde-3-phosphate dehydrogenase, 5'-AGGTCGGAGTCAACGGATTT-3' (sense) and 5'-TAGTTG AGGTCAATGAAGGG-3' (antisense).

### RNA interference

siRNA for control (12935-300), CHOP (5'-CCUCACUCUCCAGAUUCCA GUCAGA-3') DR4 (HSS112945) and DR5 (HSS112939) were purchased from Invitrogen. HCT116 cells were transfected with siRNA using HiPerFect (QIAGEN, Hilden, Germany), according to the manufacturer's instructions.

### Cell surface staining of DR5

HCT116 cells were seeded in 6-well plates and incubated overnight before the experiment. Cells were then treated with inostamycin for 12 h. Cells were then collected and resuspended in PBS<sup>-</sup> including PE-conjugated DR5 antibody. After incubation for 1 h at 4 °C, the cells were washed with PBS<sup>-</sup> twice and resuspended in PBS<sup>-</sup>. Flow cytometry analysis was performed with EPICS (Beckman Coulter).

## RESULTS AND DISCUSSION

### Co-treatment with inostamycin and TRAIL synergistically induced apoptosis in HCT116 cells

In this study, we first found that co-treatment with inostamycin and TRAIL synergistically induced cytotoxicity in HCT116 cells. As shown in Figure 1b, although neither 10 ng ml<sup>-1</sup> TRAIL nor 1 μg ml<sup>-1</sup> inostamycin alone showed cytotoxicity in HCT116 cells after 18 h treatment, the cell viability was decreased to about 40% when HCT116 cells were treated with inostamycin and TRAIL in combination. The synergistic cytotoxicity occurred in inostamycin and TRAIL dose-dependent manner. This synergistic effect is cell type-dependent, and is observed in HeLa cells, HT29 cells and LNCaP cells but not in Ms-1 cells (Figure 1b). Flow cytometer analysis demonstrated that the synergistic increase of the Sub-G<sub>1</sub> population was observed in co-treatment with inostamycin and TRAIL in HCT116 cells (Figure 1c). Furthermore, a one hour preincubation with the pan-caspase inhibitor 100 μM z-VAD-fmk suppressed the cell death that was induced by co-treatment with inostamycin and TRAIL (Figure 1c). We also observed the cleavage of caspase-8, caspase-9, caspase-3 and poly(ADP-ribose) polymerase, a well known substrate of caspase-3, at 6 h after inostamycin and TRAIL co-treatment (Figure 1d). In contrast, the cleaved caspase-8, caspase-9, caspase-3 and poly(ADP-ribose) polymerase were only weakly detected when

HCT116 cells were treated with 10 ng ml<sup>-1</sup> of TRAIL alone. Inostamycin alone also could not activate caspase-8, caspase-9 or caspase-3 in HCT116 cells (Figure 1d). These results suggested that co-treatment with inostamycin and TRAIL induced caspase-dependent apoptosis.

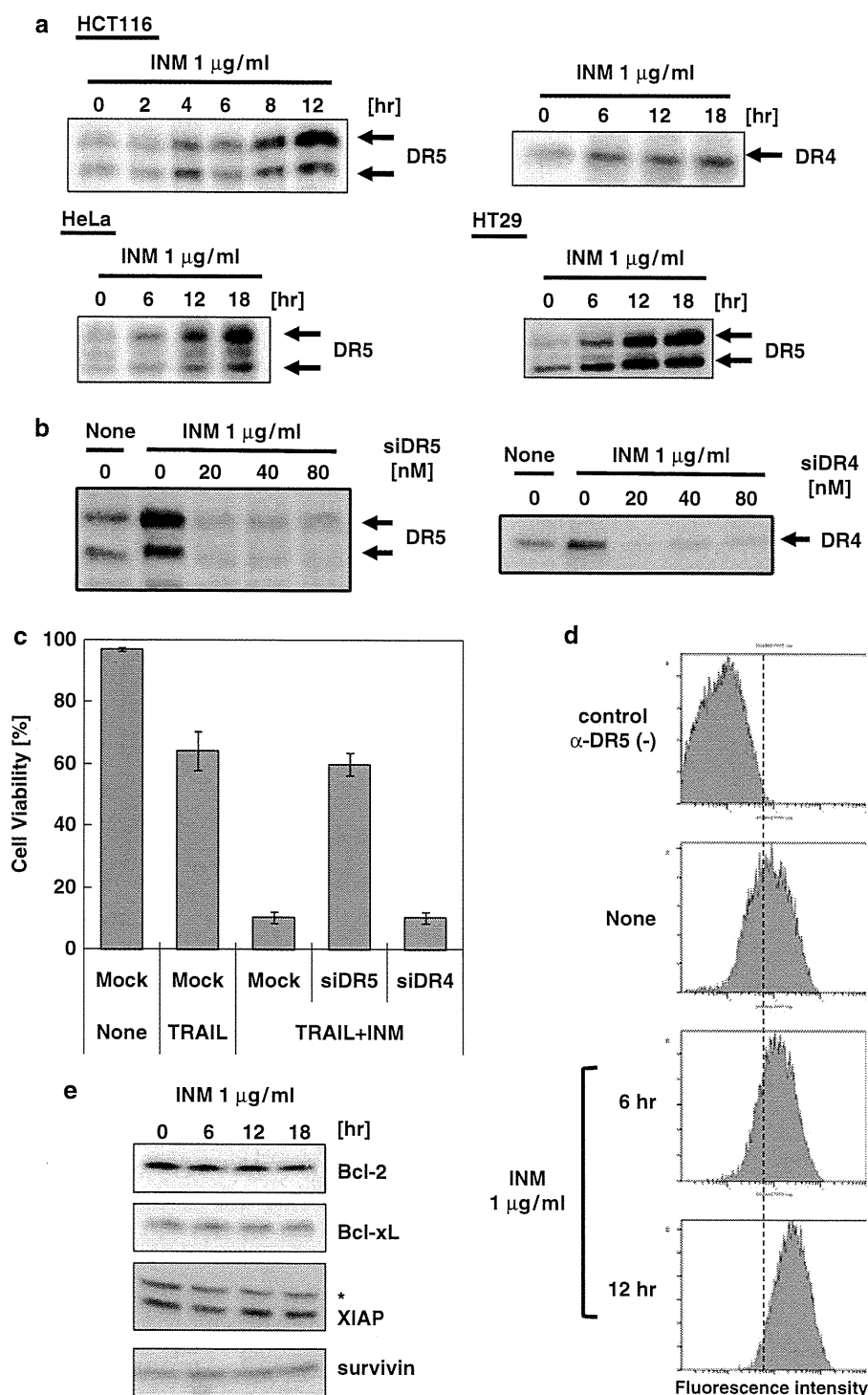
### Inostamycin enhanced TRAIL-induced apoptosis possibly through DR5 upregulation on the cell surface

TRAIL-stimulated death signal is initiated by the binding of TRAIL to DR4 or DR5, which resulted in the subsequent activation of caspase-8. As shown in Figure 2a, western blotting analysis using an anti-DR4 or anti-DR5 antibody showed that inostamycin strongly upregulated the DR5 protein in HCT116 cells, it also slightly upregulated the DR4 protein (Figure 2a). Similarly, inostamycin-induced DR5 upregulation was also observed in HT29 cells and HeLa cells, in which co-treatment with inostamycin and TRAIL induced synergistic cytotoxicity (Figure 2a). Inostamycin-induced DR5 protein was detected as two bands, which were likely products of alternative splice variants of the DR5 gene<sup>20,21</sup> Furthermore, under the condition where inostamycin-upregulated DR4 or -DR5 protein was efficiently reduced by transient transfection of DR4 siRNA or DR5 siRNA (Figure 2b), apoptosis induced by co-treatment with inostamycin and TRAIL was completely suppressed in DR5 siRNA-transfected HCT116 cells, but not in DR4 siRNA-transfected HCT116 cells (Figure 2c). These results indicated that DR5 is indispensable for the apoptosis induced by co-treatment with inostamycin and TRAIL. Moreover, inostamycin induced cell surface expression of DR5 in a time-dependent manner (Figure 2d). On the other hand, the protein expression of four anti-apoptotic molecules, Bcl-2, Bcl-xL, XIAP and survivin, was unaffected by inostamycin in HCT116 cells (Figure 2e). Considering that it increases the cell surface expression of DR5, it is likely that inostamycin-increased cell surface expression of DR5 contributes to TRAIL sensitization.

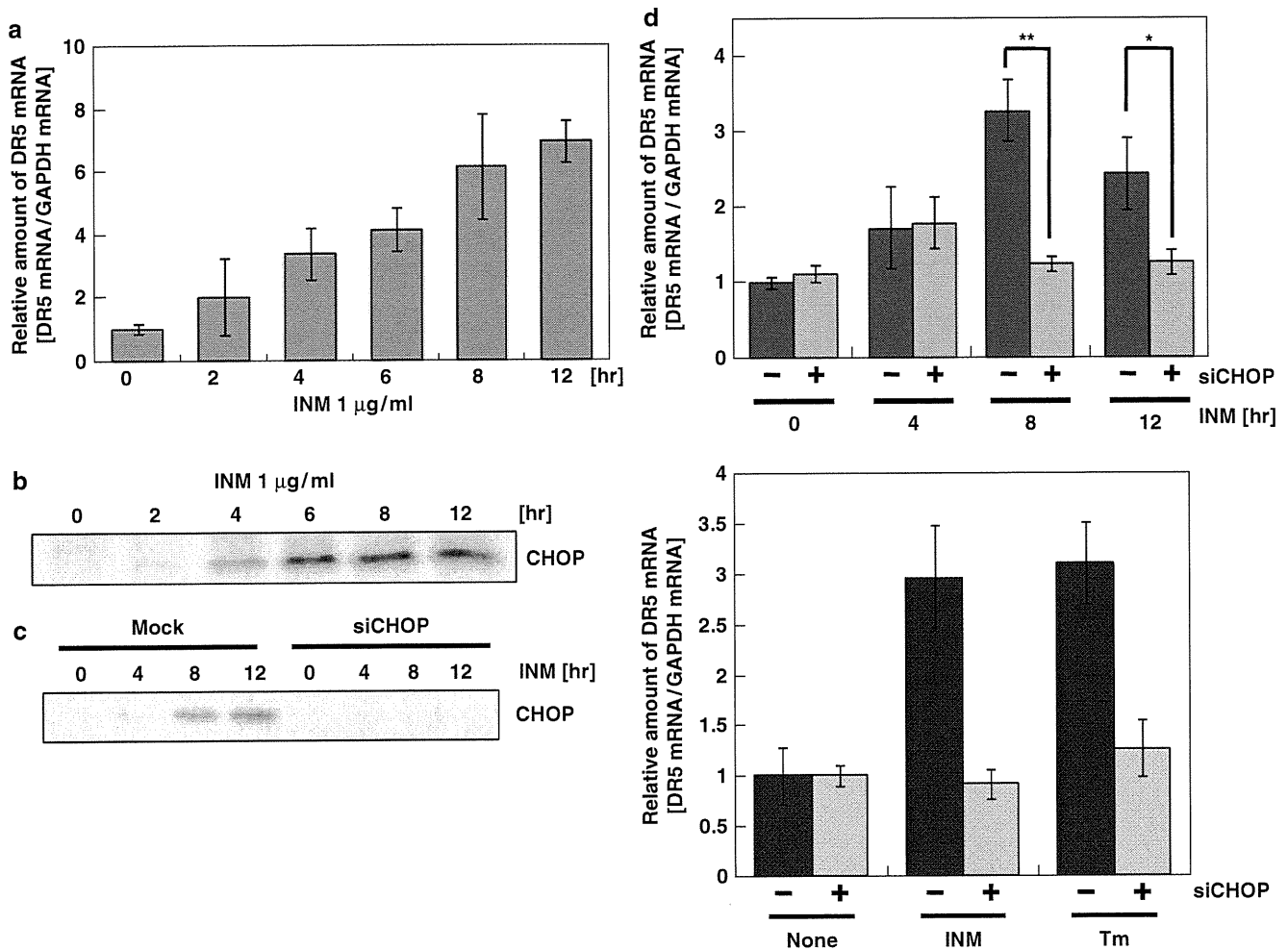
### Inostamycin-induced transcription of DR5 was regulated by CHOP.

As shown in Figure 3a, inostamycin-induced increases in DR5 mRNA levels were evaluated by real-time RT-PCR analysis. This data suggests that inostamycin activates transcription of DR5 in HCT116 cells. Recent studies have reported that tunicamycin-induced<sup>14</sup> and MG132-induced<sup>22</sup> CHOP (C/EBP homologous protein) upregulation activates DR5 transcription through CHOP-binding site in DR5 promoter region. This results in a sensitization of TRAIL-induced apoptosis. As we also found that 1 μg ml<sup>-1</sup> inostamycin increased CHOP protein expression with similar time-course kinetics to DR5 upregulation (Figure 3b), we next examined whether this inostamycin-induced CHOP expression regulated DR5 transcription. When CHOP was successfully silenced by transient transfection of siRNA for CHOP (siCHOP) (Figure 3c), inostamycin-increased DR5 mRNA was markedly suppressed (Figure 3d). Furthermore, as consistent with previous report, tunicamycin-increased expression of DR5 mRNA was also completely suppressed in siCHOP-transfected HCT116 cells (Figure 3d). Next, we examined whether inostamycin-induced DR5 protein upregulation was also suppressed in CHOP-knockdown HCT116 cells. As shown in Figures 4a and b, neither inostamycin-induced DR5 protein expression nor inostamycin-increased cell surface expression of DR5 was suppressed under the conditions where tunicamycin-induced DR5 protein expression was completely suppressed in CHOP-knockdown cells. Furthermore, the transient transfection with siCHOP could not suppress apoptosis induction by the synergistic effect of co-treatment with inostamycin and TRAIL (Figure 4c).





**Figure 2** Inostamycin (INM) increased cell surface expression of DR5 in HCT116 cells. (a) Effect of INM on the expression of DR4 and DR5 in HCT116 cells, and DR5 in HeLa cells and HT29 cells. HCT116 cells, HeLa cells and HT29 cells were treated with  $1 \mu\text{g ml}^{-1}$  of INM for the indicated periods. Cell extracts were prepared for western blotting to detect the expression of DR4 and DR5. (b) Transient transfection of small interfering RNA (siRNA) against DR4 or DR5 suppressed INM-induced DR4 or DR5 expression in HCT116 cells. HCT116 cells were transiently transfected with the indicated concentration of DR4 siRNA (siDR4) or DR5 siRNA (siDR5). After incubation for 24 h, the transfected cells were treated with  $1 \mu\text{g ml}^{-1}$  of INM for 12 h. Cell extracts were prepared for western blotting to detect the expression of DR4 and DR5. (c) Knockdown of DR5 suppressed synergistic apoptosis induction by co-treatment with INM and tumor necrosis factor-related apoptosis-inducing ligand (TRAIL). HCT116 cells were transiently transfected with 20 nM DR4 siRNA (siDR4) or siDR5. After incubation for 24 h, the transfected cells were treated with or without  $1 \mu\text{g ml}^{-1}$  INM together with  $10 \text{ ng ml}^{-1}$  TRAIL for more 24 h. The cell viability was evaluated using a Trypan Blue Exclusion assay. (d) INM increased the cell surface expression of DR5 in HCT116 cells. HCT116 cells were treated with  $1 \mu\text{g ml}^{-1}$  of INM for 6 or 12 h. The cell surface DR5 were stained with or without PE-conjugated DR5 antibody and then further detected with flow cytometer. (e) INM did not affect the expression of anti-apoptotic proteins in HCT116 cells. HCT116 cells were treated with  $1 \mu\text{g ml}^{-1}$  of INM for the indicated periods. Cell extracts were prepared for western blotting to detect the expression of Bcl-2, Bcl-xL, XIAP and survivin. \* indicates non-specific band.

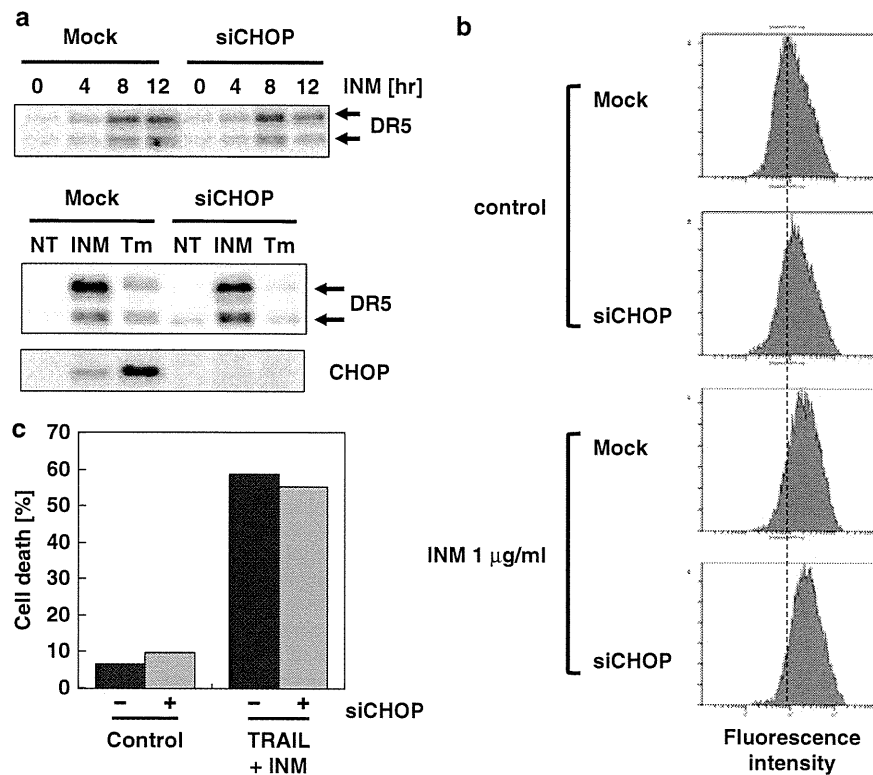


**Figure 3** Inostamycin (INM) activated DR5 transcription through CHOP upregulation. (a) INM induced DR5 upregulation of mRNA levels in HCT116 cells. HCT116 cells were treated with  $1\ \mu\text{g ml}^{-1}$  of INM for the indicated periods. The cells were collected and RNA was extracted. DR5 mRNA levels were evaluated by real-time RT-PCR. DR5 mRNA was normalized with the mRNA levels of glyceraldehyde-3-phosphate dehydrogenase. Data are the means  $\pm$  s.d. among three independent experiments. (b) Effect of INM on the expression of CHOP in HCT116 cells. HCT116 cells were treated with  $1\ \mu\text{g ml}^{-1}$  of INM for the indicated periods. Cell extracts were prepared for western blotting to detect the expression of CHOP. (c) Transient transfection of small interfering RNA (siRNA) against CHOP suppressed INM-induced CHOP expression in HCT116 cells. HCT116 cells were transiently transfected with 20 nM siCHOP. After 24 h incubation, the transfected cells were treated with  $1\ \mu\text{g ml}^{-1}$  of INM for 4, 8 or 12 h. Cell extracts were prepared for western blotting to detect the expression of CHOP. (d) Knockdown of CHOP suppressed INM- and tunicamycin (Tm)-induced DR5 upregulation at mRNA levels. HCT116 cells were transiently transfected with 20 nM siCHOP. After incubation for 24 h, the transfected cells were treated with  $1\ \mu\text{g ml}^{-1}$  of INM for 4, 8 or 12 h or  $10\ \mu\text{g ml}^{-1}$  of Tm for 12 h. The cells were collected and RNA was extracted. DR5 mRNA levels were evaluated by real-time RT-PCR. DR5 mRNA was normalized with the mRNA levels of glyceraldehyde-3-phosphate dehydrogenase (GAPDH). Data are the means  $\pm$  s.d. among three independent experiments. \* $P < 0.05$  and \*\* $P < 0.005$ .

When considered together, our results suggest that inostamycin enhanced TRAIL-induced apoptosis possibly because of a CHOP-independent upregulation of DR5 on the cell surface. At present, the mechanism by which inostamycin upregulates DR5 protein expression remains unclear. However, these results were consistent with our conclusion that the inostamycin-increased cell surface expression of DR5 possibly contributes to the enhancement of TRAIL-induced apoptosis.

In a previous study, various small molecule compounds, such as tunicamycin,<sup>14</sup> were reported to transactivate the DR5 gene by the upregulation of CHOP. The inostamycin-induced increase in DR5 mRNA was suppressed in CHOP-knockdown HCT116 cells; however, DR5 protein expression was not suppressed. Furthermore, knockdown of CHOP by siCHOP suppressed only inostamycin-

increased CHOP-dependent DR5 mRNA expression, but constitutively expressed DR5 mRNA levels were not affected (Figure 3d). Therefore, in siCHOP-transfected HCT116 cells, inostamycin is considered to increase the protein levels of DR5 translated from constitutively expressed DR5 mRNA by posttranslational modulation. Various studies have reported the mechanism of the transcriptional activation of DR5 gene. However, little is known about the mechanism of translation, posttranslational modification, localization or degradation of DR5. Because of the data presented in this study, inostamycin may be a good bioprobe for use in investigating the regulatory mechanism of DR5 protein. Furthermore, elucidation of the mechanism by which inostamycin increases the expression of DR5 on the cell surface may provide a new molecular target for inducing DR5 upregulation to overcome TRAIL resistance.



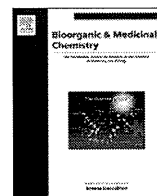
**Figure 4** Inostamycin (INM) enhanced tumor necrosis factor-related apoptosis-inducing ligand (TRAIL)-induced apoptosis possibly through CHOP-independent cell surface DR5 upregulation. (a) Knockdown of CHOP suppressed tunicamycin (Tm)-induced DR5 upregulation but did not suppress INM-induced DR5 upregulation at protein levels. HCT116 cells were transiently transfected with 20 nM siCHOP. After incubation for 24 h, the transfected cells were treated with 1 µg ml<sup>-1</sup> of INM for 4, 8, 12 h or 10 µg ml<sup>-1</sup> of Tm for 12 h. Cell extracts were prepared for western blotting to detect the expression of DR5. (b) Knockdown of CHOP did not suppress the INM-induced upregulation of cell surface DR5. HCT116 cells were transiently transfected with 20 nM siCHOP. After incubation for 24 h, the transfected cells were treated with 1 µg ml<sup>-1</sup> of INM for 12 h. The cell surface DR5 was stained with PE-conjugated DR5 antibody and then detected using a flow cytometer. (c) Knockdown of CHOP did not suppress the induction of apoptosis by co-treatment with INM and TRAIL. HCT116 cells were transiently transfected with 20 nM siCHOP. After incubation for 24 h, the transfected cells were treated with or without 1 µg ml<sup>-1</sup> INM together with 10 ng ml<sup>-1</sup> TRAIL for 18 h. The cells were stained with propidium iodide (PI), and sub-G<sub>1</sub> phase cells were detected using a flow cytometer.

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## Target identification of bioactive compounds

Etsu Tashiro, Masaya Imoto\*

Department of Biosciences and Informatics, Faculty of Science and Technology, Keio University, 3-14-1 Hiyoshi, Kohokuku, Yokohama, Kanagawa 223-8522, Japan

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### ABSTRACT

To fully understand the regulation of cellular events, functional analysis of each protein involved in the regulatory systems is required. Among a variety of methods to uncover protein function, chemical genetics is a remarkable approach in which small molecular compounds are used as probes to elucidate protein functions within signaling pathways. However, identifying the target of small molecular bioactive compounds isolated by cell-based assays represents a crucial hurdle that must be overcome before chemical genetic studies can commence. A variety of methods and technologies for identifying target proteins have been reported. This review therefore aims to describe approaches for identifying these molecular targets.

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### 1. Introduction

To improve our understanding of complex cell systems, functional analysis of proteins has become a significant focus for a growing number of research fields in biology in the post-genome era.<sup>1</sup> Proteins play an intrinsic role in the cellular events of cell growth, survival, migration, and differentiation, through the sequential assembly of protein interactions, which form signal transduction pathways. However, the role of proteins in many cellular events remains unknown. Among a variety of methods to uncover protein function, chemical genetics is a remarkable approach in which small molecular compounds are used as probes to elucidate protein functions within signaling pathways.<sup>2,3</sup> Indeed, several bioactive compounds have led to breakthroughs in understanding the functional roles of proteins. For example, the study of chemical genetics using FK506, which was isolated from a microbial origin as a new immunosuppressant,<sup>4</sup> contributed to the identification of the important role of calcineurin in the immune system, by identifying FKBP12 as a target protein of FK506.<sup>5,6</sup> Another example is the discovery of lactacystin, a new microbial metabolite which induces differentiation in neuroblastoma cells,<sup>7</sup> whose target identification drove the wide application of lactacystin as a proteasomal inhibitor in the field of cell biology.<sup>8–10</sup> There are now a growing number of chemical inhibitors of signal transduction pathways, and these examples demonstrate how chemical genetics can lead us towards understanding cellular events at a molecular level. However, one significant hurdle to developing new chemical probes of biological systems is identifying the target protein of bioactive compounds, discovered using

cell-based small-molecule screening. Identification of the molecular target of bioactive compounds is the major barrier to advancing chemical genetic research. This review therefore aims to describe approaches for identifying these molecular targets.

A variety of methods and technologies for identifying target proteins have been reported. These can be fundamentally categorized into two approaches: direct and indirect.<sup>11</sup> In the direct approach, proteins bound to each compound are purified and directly identified using time-of-flight mass spectrometry (TOF/MS) analysis. Compounds are immobilized using affinity beads or columns to purify the bound proteins (Fig. 1A and Table 1). In this respect, chemical synthesis of the compounds would become a great help for the preparation of the immobilized compounds. However, in the case of natural products, this synthesis is sometimes difficult because of the chiral centers and unique scaffolds in their structures.

On the other hand, the indirect approach offers target candidates by profiling the biological data of the compounds.<sup>11</sup> In the case that the compound was found to perturb some cellular event whose regulatory signaling pathways had been already reported, by clarifying which step within the pathway was affected by the compound, the enzymatic protein engaged in this step could be raised as the target candidate of the compound. In some cases, ‘-omics’ studies (e.g., proteomics, transcriptomics) can help to collect comprehensive data about the biological effects of particular compounds to aid target profiling (Fig. 1B and Table 1).

### 2. Direct approach

Traditional approaches using affinity chromatography, biochemical fractionation, and radioactive ligand binding assays have been successful in identifying the biological targets of multiple

\* Corresponding author.

E-mail address: [imoto@bio.keio.ac.jp](mailto:imoto@bio.keio.ac.jp) (M. Imoto).