Retraction: Fatty Acid Synthase is a Novel Therapeutic Target in Multiple Myeloma

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Retraction: Fatty acid synthase is a novel therapeutic target in multiple myeloma


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Finbarr Cotter
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Summary

This study investigated the biological significance of the inhibition of fatty acid synthase (FAS) in multiple myeloma (MM) using the small molecule inhibitor Cerulenin. Cerulenin triggered growth inhibition in both MM cell lines and MM patient cells, and overcame the survival and growth advantages conferred by interleukin-6, insulin-like growth factor-1, and bone marrow stromal cells. It induced apoptosis in MM cell lines with only modest activation of caspase -8, -9, -3 and PARP; moreover, the pan-caspase inhibitor Z-VAD-FMK did not inhibit Cerulenin-induced apoptosis and cell death. In addition, treatment of MM cells with Cerulenin primarily up-regulated apoptosis-inducing factor/endonuclease G, mediators of caspase-independent apoptosis. Importantly, Cerulenin induced endoplasmic reticulum stress response via up-regulation of the Grp78/IRE1α/JNK pathway. Although the C-Jun-NH2-terminal kinase (JNK) inhibitor SP600215 blocked Cerulenin-induced cytotoxicity, it did not inhibit apoptosis and caspase cleavage. Furthermore, Cerulenin showed synergistic cytotoxic effects with various agents including Bortezomib, Melphalan and Doxorubicin. Our results therefore indicate that inhibition of FAS by Cerulenin primarily triggered caspase-independent apoptosis and JNK-dependent cytotoxicity in MM cells. This report demonstrated that inhibition of FAS has anti-tumour activity against MM cells, suggesting that it represents a novel therapeutic target in MM.

Keywords: multiple myeloma, fatty acid synthase, apoptosis, JNK.
Cerulenin also induces endonuclease triggers apoptosis, at least in part, via a caspase-independent, released from the mitochondria to cytosol, suggesting that Cerulenin triggers synergistic cytotoxicity and cell death. During Cerulenin-induced apoptosis, both Poly (ADP-ribose) polymerase (PARP), and endonuclease G (Endo G) are released from the mitochondria to cytosol, suggesting that Cerulenin triggers apoptosis, at least in part, via a caspase-independent, AIF/Endo G pathway. Importantly, Cerulenin also induces endonuclease triggers apoptosis, at least in part, via a caspase-independent, released from the mitochondria to cytosol, suggesting that Cerulenin also triggers synergistic cytotoxicity and cell death in MM, confirming that FAS represents a novel therapeutic target in MM.

Materials and methods

Reagents

The FAS inhibitor, Cerulenin, was provided by EMD chemicals (San Diego, CA, USA). Cerulenin, (2R, 3S)-2,3-epoxy-4-oxo-7,10-trans, trans-dodecadienamide (C₁₀H₁₈N₂O₂), a natural product of Cephalosporium caerules, is a specific inhibitor of FAS enzyme across a broad phylogenetic spectrum (Vance et al, 1972; Otsuka, 1976; Moore et al, 1999). It was dissolved in dimethyl sulphoxide (DMSO) at 50 mmol/l stock solution and stored at −20°C. Reconstituent human IL-6, IGF-1 (R&D Systems, Minneapolis, MN, USA) were reconstituted with sterile phosphate-buffered saline (PBS) containing 0.1% bovine serum albumin (BSA). Pan-caspase inhibitor Z-VAD-FMK (Bachem, Bubendorf, Switzerland) was dissolved in methanol, stored at −20°C, and used at 25 μmol/l. JNK inhibitor SP600215 (Calbiochem, San Diego, CA, USA) was stored at 20°C, and used at 5–10 μmol/l. Tunicamycin (Sigma Chemical, St Louis, MO, USA) was dissolved in DMSO at 10 mg/ml stock solution and stored at 4°C, and used at 5 μg/ml. Melphanal and Doxorubicin were purchased from Sigma Chemical. Bortezomib was obtained from Millenium Pharmaceuticals Inc., Cambridge, MA, USA.

Cells

Dex-sensitive (MM.1S) and resistant (MM.1R) human MM cell lines were kindly provided by Dr Steven Rosen (Northwestern University, Chicago, IL, USA). RPMI-8226 and U266 human MM cell lines were obtained from a Millenium Type Culture Collection (Manassas, VA, USA). RPMI 8226-Doxx40 (Doxorubicin-resistant) and RPMI 8226-LRS (Mel-resistant) human MM cell lines were kindly provided by Dr William Dalton (Lee Moffitt Cancer Center, Tampa, FL, USA). OPM1 and OPM2 were provided from Dr P. Leif Bergsagel (Mayo Clinic, Tucson, AZ, USA). Each of these cell lines were cultured in RPMI-1640 containing 10% fetal bovine serum (FBS; Sigma Chemical), 2 μmol/l l-glutamine, 100 U/ml penicillin and 100 μg/ml streptomycin (GIBCO, Grand Island, NY, USA). Fresh peripheral blood mononuclear cells (PBMCN) obtained from three healthy volunteers were separated from heparinized peripheral blood by Ficoll-Hypaque (Pharmacia, Piscataway, NJ, USA) density sedimentation.

Primary MM cells and BMSCs from MM patients, and plasma cells from healthy volunteers

Tumour cells freshly isolated from MM patients and plasma cells from healthy volunteers (>90% CD138⁺) were purified by CD138 positive selection using CD138 (Syndecan-1) Micro Beads and the Auto MACS magnetic cell sorter (Miltenyi Biotec Inc., Auburn, CA, USA). BM mononuclear cells separated by Ficoll-Hypaque were cultured in Dulbecco’s modified Eagles medium (Sigma Chemical) supplemented with 15% heat-inactivated FBS, 2 μmol/l l-glutamine, 100 U/ml penicillin and 100 μg/ml streptomycin for 3–6 weeks to generate BMSCs. Approval for these studies was
obtained from the Dana-Farber Cancer Institute Institutional Review Board. Informed consent was obtained from all patients in accordance with the Declaration of Helsinki protocol.

**Growth inhibition assay**

The growth inhibitory effect of Cerulenin in MM cell lines, primary MM cells, PBMC and normal plasma cells was assessed by measuring 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide (MTT; Sigma) dye absorbance. Cells were plated in 96-well microtitre plates at a density of 2–3×10^4 cells per well, and each plate was incubated for 24 and/or 48 h, with MTT added to each well for the last 4 h. The absorbance of each well was measured at 570/630 nm using a spectrophotometer (Molecular Devices, Sunnyvale, CA, USA).

To measure proliferation of MM cells and BMSC, the rate of DNA synthesis was measured, as described previously (Hideshima et al, 2000). MM cells were incubated in 96-well culture plates (Costar, Cambridge, MA, USA) in the presence of Cerulenin and/or recombinant cytokines IL-6 or IGF-1 for 48 h at 37°C. Cells were pulsed with 18.5 kBq/well of [3H]-thymidine ([3H]-Tdr; Perkin Elmer, Boston, MA, USA) during the last 8 h of culture, harvested onto glass filters with an automatic cell collector (Cambridge Technology, Cambridge, MA, USA), and counted using the LKB Betaplate scintillation counter (Wallac, Gaithersburg, MD, USA). MTT and [3H] uptake analysis were performed in triplicate or quadruplicate cultures.

**Immunoblotting**

The MM cells were harvested, washed twice with PBS, and lysed using lysis buffer: 50 mmol/l Tris–HCl (pH 7.4), 150 mmol/l NaCl, 1% Nonidet P-40, 10 mmol/l sodium pyrophosphate, 5 mmol/l EDTA, 1 mmol/l EGTA, 2 mmol/l NaVO_4, 5 mmol/l NaF, 1 mmol/l phenylmethylsulphonyl fluoride (PMSF), 5 μg/ml leupeptin and 5 μg/ml aprotinin, as described previously (Hayashi et al, 2002). Subcellular proteins were extracted from 2×10^7 viable cells using a mitochondria isolation kit (Pierce, Rockford, IL, USA). The whole cell lysates or fractionated proteins (20 μg per lane) were subjected to sodium dodecyl sulphate–polyacrylamide gel electrophoresis, transferred to nitrocellulose membranes (Bio-Rad Laboratories, Hercules, CA, USA), and immunoblotted with anti-PARP, caspase-8, caspase-3, caspase-9, AIF, voltage-dependent anion channel (VDAC), ASK1, GRP78/Bip, IRE1, JNK, p-ATF-2, p-c-Jun and glyceraldehyde-3-phosphate dehydrogenase (GAPDH) (Cell Signaling, Beverly, MA, USA); as well as with anti-α-tubulin antibodies (Abs) (Santa Cruz Biotechnology, Inc. Santa Cruz, CA, USA), anti-Fatty Acid Synthase Abs (BD Bioscience, San Jose, CA, USA), anti-Endo G (ProSci Inc., Poway, CA, USA) and anti-XBP-1 (Abcam, Inc. Cambridge, MA, USA) Abs.

**Flow cytometric analysis**

For detection of apoptotic cells, mitochondrial protein staining was performed with pycoerythrin-labeled anti-Apo2·7 antibody (Immunotec, Marseille, France). Cells were treated with various concentrations of Cerulenin for 24 h, washed with PBS, and then incubated with Apo2·7 for 30 min. Apo2·7 staining was analysed using the RXP cytomics software on an Epics flow cytometer (Couler Immunology, Hialeah, FL, USA).

**Immunocytochemistry**

Cytospins of cells were prepared on glass slides and fixed with 50% methanol/50% acetone at 4°C. The slides were blocked in PBS with 5% FBS at room temperature (RT), and then incubated with primary antibody overnight at 4°C. Fluorescein isothiocyanate (FITC)-labeled secondary antibody reaction for 30 min was then performed. Coverslips were mounted on the glass slides with VectaShield antifade/4′,6-diamidino-2-phenylindole (DAPI), which were analysed by Nikon E800 fluorescence microscopy (Nikon Inc., Melville, NY, USA).

**Statistical analysis**

Statistical significance of differences observed in drug-treated compared with control cultures was determined using the Mann–Whitney test. The minimal level of significance was P < 0.05. All statistical analyses were determined using GraphPad Prism software (GraphPad Software, Inc. San Diego, CA, USA).

**Isobologram analysis**

The interaction between Cerulenin and Bortezomib, Melphalan, and Doxorubicin was analysed using CalcuSyn software program (Biosoft, Ferguson, MO, USA) to determine whether the combination was additive or synergistic, as described previously (Chou & Talalay, 1984; Raje et al, 2004). This program is based upon the Chou-Talalay method, which calculates a combination index (CI). Analysis is performed based on the following equation: CI = (D1/((D)x1 + (D)x2) + (D)/((D)x1/(D)x2), where (D)1 and (D)2 are the doses of drug 1 and drug 2 that have x effect when used in combination; whereas (D)x1 and (D)x2 are the doses of drug 1 and drug 2 that have the same x effect when used alone. A CI c. 1 represents the conservation isobologram and indicates additive effects, whereas CI c. ≤0.9 indicates synergism.

**Results**

**FAS expression in various cells**

We first examined baseline expression of FAS in various cells. FAS protein was expressed in all MM cell lines (Fig 1A and B; lane 5), as well as in primary tumour cells from MM patients (Fig 1B; lane 4, Fig 1C). Importantly, FAS expression was higher in both MM cell lines and primary tumour cells than in normal plasma cells, as assessed by Western blotting (Fig 1B) and confirmed by immunocytochemical analysis (Fig 1D).

**Cerulenin inhibits growth of MM cells**

We next examined the effect of FAS inhibition by Cerulenin (C_{12}H_{17}NO_{3}; Fig 2A) on growth of MM cells and normal cells,
Cerulenin significantly inhibited the growth of drug-sensitive MM.1S, U266, RPMI8226, OPM1 and OPM2 MM cell lines, with a 50% inhibitory concentration (IC50) at 24 h of 24 ± 16, 22 ± 7, 24 ± 03, 37 ± 03 and 21 ± 53 l mol/l, respectively, and IC50 at 48 h of 12 ± 59, 11 ± 12, 17 ± 08, 11 ± 45 and 9 ± 71 l mol/l respectively (Fig 2B and C). Cerulenin also inhibited growth of Dex-resistant MM.1R, Mel-resistant RPMI-LR5, Dox-resistant RPMI-Dox40 MM cell lines, with IC50 at 24 h of 22 ± 59, 86 ± 21 and 33 ± 29 l mol/l, and IC50 at 48 h of 10 ± 52, 22 ± 73 and 16 ± 52 l mol/l respectively (Fig 2B and C).

Cerulenin overcomes the growth stimulatory effects of cytokines, and inhibits MM cell growth in culture with BMSCs

It has been demonstrated that IL-6 and IGF-1 induce both growth and inhibition of apoptosis in MM cells (Jourdan et al, 2000; Mitsiades et al, 2002, 2004), therefore, we examined whether FAS
inhibition can overcome the effects of exogenous IL-6 and IGF-1. Both IL-6 (10 ng/ml) and IGF-1 (50 ng/ml) triggered increased growth in MM.1S and U266 cells, which was inhibited by Cerulenin \( (P < 0.05) \) (Fig 3A and B). As we previously have shown that the BM microenvironment conferred growth and drug resistance in MM cells (Hideshima et al., 2004), we next examined the effect of FAS inhibition on MM cell growth in the BM milieu. MM.1S and U266 cells were cultured for 48 h with or without BMSC, in the presence or absence of Cerulenin. MM cell adherence to BMSC enhanced \[^{[3]H}\]-thymidine uptake in MM cells, which was inhibited by Cerulenin \( (P < 0.05) \) (Fig 3C). These results indicate that Cerulenin also blocks the growth stimulatory effect of the BM microenvironment on MM cells.

**Cerulenin induces apoptosis via activation of caspase-independent pathway**

To analyse the mechanism of cytotoxicity triggered by Cerulenin in MM cells, we next performed Apo2-L analyses and immunoblotting in MM.1S and U266 cells. Cerulenin induced apoptosis in a dose-dependent manner, evidenced by Apo2-L analysis (Fig 4A, Fig S1C). Cerulenin also triggers caspase -8, -9, -3, and...
PARP cleavage in MM.1S and U266 (Fig 4B, Fig S1A). Although the pan-caspase inhibitor Z-VAD-FMK blocked Cerulenin-induced caspase and PARP cleavage in both MM.1S and U266 cells (Fig 4B, Fig S1A), it did not inhibit Cerulenin-triggered apoptosis (Fig 4D, Fig S1C) and cell death (Fig 4C, Fig S1B). During Cerulenin-induced apoptosis in MM cells, both AIF and Endo G were significantly released from the mitochondria to cytosol in a time-dependent manner (Fig 4E). These results suggest that Cerulenin triggers apoptosis, at least in part, via a caspase-independent, AIF/Endo G pathway.

Cerulenin induces ER stress response via up-regulation of GRP78, IRE1α, JNK pathway

As some reports have demonstrated that FAS inhibitor induced ER stress response in tumour cells (Little et al, 2007), we next examined whether FAS inhibition by Cerulenin triggers activation of GRP78, IRE1α, JNK pathway as an ER stress response in MM cells. The ER stress inducer Tunicamycin significantly up-regulated Grp78, IRE1α, ASK1, JNK and p-JNK in MM.1S cells (Fig 5A). In addition, Cerulenin treatment also induced up-regulation of Grp78, IRE1α and p-JNK, as well as its downstream molecules p-ATF-2 and p-c-Jun (Fig 5A). Importantly, JNK inhibitor SP600215 significantly blocks Cerulenin-induced cytotoxicity in MM.1S and U266 cells, (P < 0.05) (Fig 5B, Fig 5A). We also confirmed that SP600215 markedly inhibited JNK signaling pathway in MM.1S cells by Western blotting (Fig 5E); however, it did not inhibit apoptosis and caspase cleavage triggered by Cerulenin in both MM.1S and U266 cells (Fig 5C and D, Fig S2B and C). In addition, combination pretreatment with JNK inhibitor and ZVAD-FMK did not enhance the inhibitory effect of JNK inhibitor alone, indicating that JNK activation is the major pathway mediating Cerulenin-induced cytotoxicity in MM cells (Fig S3). These results strongly indicate that targeting FAS by Cerulenin induces ER stress response via up-regulation of Grp78/IRE1α/JNK pathway, associated with JNK-dependent, non-apoptotic cell death in MM cells.

Cerulenin combined with Bortezomib, Melphalan, or Doxorubicin induces synergistic cytotoxicity against MM cells

We further examined whether Cerulenin enhances the growth inhibitory effect of other conventional agents in MM cells. MM.1S, U266, and primary MM cells were cultured for 24 h with Bortezomib (0–3.75 nmol/l), Melphalan (0–10 μmol/l) or Doxorubicin (0–400 nmol/l) combined with Cerulenin (0–25 μmol/l). Cerulenin enhanced growth inhibition triggered by Bortezomib, Melphalan and Doxorubicin, as analysed by MTT assay (Fig 6, Fig S4). Next, we analysed the interaction between Cerulenin and these agents by isobologram analysis using CalcuSyn software program (Raje et al, 2004). Based upon the Chou-Talalay method to calculate the CI, we showed synergistic cytotoxic effects between

Fig 3. Cerulenin overcomes the protective effects of IL-6, IGF-1 and BMSCs on MM cell growth. (A, B, C) MM.1S and U266 cells were cultured for 48 h with the indicated concentrations [0 μmol/l ( ), 3.125 μmol/l ( ), 6.25 μmol/l ( ), 12.5 μmol/l ( )] of Cerulenin, in the presence or absence of IL-6 (1 or 10 ng/ml: A), IGF-1 (10 or 50 ng/ml: B), or BMSC (C). Cell growth was assessed by [3H]-thymidine uptake. Cerulenin inhibits MM cell growth and overcomes the stimulating effect of IL-6 (A) or IGF-1 (B) (P < 0.05), and BMSC (P < 0.05) (C). Values represent mean ± SD of quadruplicate cultures.
Cerulenin and Bortezomib, Melphalan, and Doxorubicin, with a maximum CI of 0.567, 0.188 and 0.529, respectively, in MM.1S cells; 0.63, 0.404 and 0.079, respectively, in primary MM cells; and 0.879, 0.591, and 0.301, respectively, in U266 cells. These data demonstrate that these combination therapies enhance Cerulenin-induced cytotoxicity synergistically in MM cells, due to the fact that these agents induce caspase-dependent apoptosis whereas Cerulenin triggers primarily caspase-independent apoptosis.

**Discussion**

This report demonstrated that FAS inhibition by Cerulenin induced cytotoxicity in drug-sensitive and drug-resistant MM cell
lines and in primary patient MM cells, with an IC\textsubscript{50} at 24 h of 21.53–86.21 µmol/l. In contrast, we observed no cytotoxicity against PBMNC and normal plasma cells, indicating selective and potent cytotoxicity against MM tumour cells. Immunoblot and immunocytochemical analysis revealed that FAS protein was more highly expressed in all MM cell lines and primary MM cells than in normal plasma cells, consistent with prior reports of FAS overexpression in cancer cells (Omura, 1976; Milgraum et al.,

Fig 5. Cerulenin triggers ER stress response and JNK activation. (A) MM.1S cells were cultured with Cerulenin (50 µmol/l) for indicated times. Induction of ER stress response by Cerulenin was examined by Western blotting. MM.1S cells treated with 5 µg/ml Tunicamycin (TM) for 24 h were used as a positive control of ER stress induction. Total cell lysates (20 µg protein/lane) were analysed by anti-GRP78, IRE1\textalpha, ASK1, p-JNK, JNK, p-ATF-2, p-c-Jun, XBP-1 and \textalpha-tubulin Abs. (B, C) MM.1S cells were treated with the indicated dose of Cerulenin for 24 h, with or without JNK inhibitor SP600215 (5 µmol/l or 10 µmol/l) pretreatment for 1 h. Cytotoxicity was determined by MTT assay (B). Values represent mean ± SD of quadruplicate cultures. JNK inhibitor significantly blocks Cerulenin-induced cytotoxicity (P < 0.05) (B). The percentage of apoptotic cells was determined by flow-cytometric analysis for APO2\textsuperscript{+} staining (C). (D, E) MM.1S cells were cultured with Cerulenin (50 µmol/l) for the indicated times, with or without SP600215 (10 µmol/l) pretreatment for 1 h. Total cell lysates (20 µg/lane) were subjected to Western blotting using anti-caspase -8, -9, and -3, PARP, p-JNK, JNK, p-ATF-2, p-c-Jun and \textalpha-tubulin Abs. FL, CF indicate the full length and cleaved form respectively.
The BM microenvironment confers drug resistance in MM cells via at least two different mechanisms: (i) cell adhesion mediated drug resistance (CAM-DR) through adhesion of MM cells to fibronectin and BMSC; and (ii) activation of phosphatidylinositol 3-kinase (PI3-K)/Akt and/or Janus kinase 2 (JAK2)/signal transducers and activators of transcription 3 (STAT3) signaling due to high levels of cytokines in the BM milieu (such as IL-6 and IGF-1) (Damiano et al., 1999; Hazlehurst et al., 2000; Hideshima et al., 2001, 2004). Importantly, treatments targeting the BM microenvironment as well as the MM cell can overcome drug resistance in both preclinical and clinical studies (Mitsiades et al., 2004; Hideshima et al., 2007). Our data indicate that Cerulenin overcomes the survival and growth advantages conferred by IL-6, IGF-1 and BMSC, strongly suggesting its utility to overcome conventional drug resistance in MM.

Although caspases are the key executors of apoptosis, recent studies suggest that caspase activation is not the sole pathway for inducing apoptosis or necrosis (Jaattela & Tschopp, 2003; Abraham & Shaham, 2004; Lockshin & Zakeri, 2004). For example, caspase-independent cell death in MM cells is triggered by AS2O3 and inosine monophosphate dehydrogenase inhibitor VX-944 treatment (McCafferty-Grad et al., 2003; Ishitsuka et al., 2005). The AIF/Endo G pathway is a major caspase-independent apoptotic cascade (Daugas et al., 2000; Ahn et al., 2004; Cande et al., 2004; Cregan et al., 2004). In this pathway, enhanced Bcl-2 family expression induces mitochondrial membrane permeability, thereby releasing AIF and Endo G from the mitochondria to the cytosol and nucleus, with subsequent chromatin condensation and cell death. The present study showed that MM cell apoptosis triggered by Cerulenin was primarily mediated via a caspase-independent pathway. Although cleavage of caspase -8,-9,-3 and PARP was induced by Cerulenin treatment, Z-VAD-FMK did not block completely Cerulenin-induced apoptosis and cell death.
Moreover, both AIF and Endo G were released from the mitochondria to cytosol during Cerulenin-induced apoptosis. These results suggest that Cerulenin triggers apoptosis, at least in part, via a caspase-independent AIF/Endo G pathway, with only modest activation of caspase -8,-9,-3 and PARP. As most conventional and novel drugs induce apoptosis in MM cells via activation of caspases, we next examined the effect of combining Cerulenin and other drugs. Cerulenin combined with Bortezomib, Melphalan or Doxorubicin triggered synergistic MM cell cytotoxicity, indicating that it may be useful to sensitize or overcome drug resistance.

The ER stress response is a choreographed series of cellular events activated by specific insults that result in altered ER function (Kauffman, 2004; Xu et al., 2005). The combined effect of this response is the activation of genes associated with an adaptation response. Upon prolonged stress, the adaptation mechanism of the ER stress response is saturated, resulting in cell death. Several studies have demonstrated important associations between lipid synthesis pathways and the ER stress response (Cox et al., 1997; van der Sanden et al., 2004; Sriburi et al., 2004; Tessitore et al., 2004). Inhibition of phospholipid synthesis, especially that of phosphatidylcholine, induces ER stress-related pathways (van der Sanden et al., 2004). Fatty acid synthesis in general, and FAS activity in particular, occurs in the ER (Swinnen et al., 2003). Importantly, because of the direct connection between ER stress activity and phospholipid synthesis, previous reports demonstrated that pharmacologic blockade of FAS activity induced ER stress in tumour cells via activation of PERK/eIF2α and IRE1α/XBP-1 pathways (Swinnen et al., 2003; Little et al., 2007). In addition, Nakamura et al. (2006) reported that activation of the ER stress pathway was associated with survival of MM cells, as they contained a well-developed ER due to M-protein production. Our data and other reports indicated that increased FAS expression in tumour cells is important for ER function; and conversely, suggest a role for ER stress in the antitumour effects of FAS inhibitors. The present study showed that Cerulenin treatment induced upregulation of Gpr78, IRE1α and p-JNK, as well as its downstream molecules p-ATF-2 and p-c-Jun. IRE1α can activate the ASK1/JNK mitogen-activated protein kinase pathway (Urano et al., 2000; Xu et al., 2005; Li et al., 2006), and activation of JNK is a common response to many forms of stress known to influence the cell-death machinery (Davis, 2000). Although studies show that multiple pathways can contribute to ER-stress-induced apoptosis (Li et al., 2006), the mechanisms by which unresolved ER stress causes cell death are not fully characterized (Szegedi et al., 2006; Zhao & Ackerman, 2006).

Bandyopadhyay et al. (2006) proposed a model for apoptotic signaling in which inhibition of FAS leads to accumulation of malonyl-CoA, which in turn inhibits carnitine palmitoyltransferase-1 (CPT-1) and results in up-regulation of ceramide, followed by induction of the proapoptotic genes BNIP3, tumor necrosis factor (TNF)-related apoptosis-inducing ligand (TNFSF10), and death-associated protein kinase 2 (DAPK2). BNIP3 not only induces apoptotic cell death, but also is implicated in necrosis and autophagy (Vande Velde et al., 2000; Daido et al., 2004). TNFSF10 has been shown to induce apoptosis in a wide range of transformed cells (Kelley & Ashkenazi, 2004). In addition, a human single-chain fragment variable (HW1) antibody against TRAIL receptor 2 induced autophagic cell death predominantly via the JNK pathway in a caspase-independent manner (Park et al., 2007). DAPK2 is associated with ceramide-mediated apoptosis (Pelled et al., 2002; Yamamoto et al., 2002), but also induces another cell death pathway mediating autophagic or type II cell death (Inbal et al., 2002; Bialik & Kimchi, 2004) without caspase activation (Inbal et al., 2002). Our data and others suggest that inhibition of FAS by Cerulenin activates multiple death signal pathways in MM cells, including caspase-independent apoptosis and JNK dependent non-apoptotic cell death. JNK and its downstream molecules ATF-2 and c-Jun induce apoptosis in various tumours (Bossy-Wetzel et al., 1997; Podar et al., 2007). Moreover, caspase-independent JNK activation promotes autophagic cell death induced by TNF-α (Jia et al., 2006) and other chemicals (Borsello et al., 2003; Yu et al., 2004; Park et al., 2007). JNK can activate the mitochondrial pathway including AIF, Endo G, cytochrome C and Smac (Weston & Davis, 2007). However, the exact role of c-Jun in mediating cell differentiation, growth, survival and apoptosis is not fully understood (Jochum et al., 2001; Vogt, 2001).

In summary, our data showed that inhibition of FAS by Cerulenin induced primarily caspase-independent apoptosis via upregulation of AIF/Endo G pathway, and JNK-dependent cell death in MM cells. This report demonstrated that inhibition of FAS had anti-tumour activity against MM cells, identifying FAS as a novel therapeutic target. Taken together, our data provide the preclinical rationale for clinical trials targeting FAS to improve patient outcome in MM.

Conflict of interest disclosure
The authors declare no competing financial interests.

Author’s contributions
YO designed, performed, analysed research and wrote the paper; TH and NR designed, analysed research and wrote the paper; HI, SV, TK, HY performed and designed research; SE performed research and analysed data; SP, IB, LS, PR performed research; KCA participated in design, coordination and performance of study, assisted in writing the paper and funded the study.

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**Supplementary material**

The following supplementary material is available for this article online:

**Fig S1.** Cerulenin induces apoptosis in U266 cells. (A) U266 cells were cultured with Cerulenin (50 μmol/l) for the indicated times, with or without Z-VAD-FMK (50 μmol/l) pretreatment for 1 h pretreatment. Total cell lysates (20 μg/lane) were subjected to Western blotting using anti-caspase -8, -9, -3, PARP and α-tubulin Abs. FL, CF indicated as full length and cleaved form, respectively. (B, C) U266 cells were treated with indicated dose of Cerulenin for 24 h, with or without Z-VAD-FMK (25 or 50 μmol/l) 1 h pretreatment. Cytotoxicity was determined by flow-cytometric analysis for Apo2.7 staining (C).

**Fig S2.** JNK inhibitor blocks Cerulenin-induced cytotoxicity in U266 cells. (A, B) U266 cells were cultured with indicated dose of Cerulenin for 24 h, with or without JNK inhibitor SP600215 (5 or 10 μmol/l) pretreatment for 1 h. Cytotoxicity was determined by MTT assay (A). Values represent mean ± SD of triplicate cultures. With or without Z-VAD-FMK (25 or 50 μmol/l) 1 h pretreatment for 1 h. Cytotoxicity was determined by MTT assay (A). Values represent mean ± SD of triplicate cultures. Although ZVAD-FMK does not affect cytotoxicity by Cerulenin, JNK inhibitor significantly blocks Cerulenin-induced cytotoxicity (P < 0.05).

**Fig S3.** Analysis of inhibition by JNK inhibitor SP600215 and/or ZVAD-FMK in MM cells. (A, B) MM.1S and U266 cells were cultured with indicated dose of Cerulenin for 24 h, with or without JNK inhibitor SP600215 (10 μmol/l) and/or ZVAD-FMK (50 μmol/l) pretreatment for 1 h. Cytotoxicity was determined by MTT assay. Values represent mean ± SD of quadruplicate cultures. Although ZVAD-FMK does not affect cytotoxicity by Cerulenin, JNK inhibitor significantly blocks Cerulenin-induced cytotoxicity (P < 0.05).

**Fig S4.** Cerulenin enhances Bortezomib, Melphalan, and Doxorubicin-induced cytotoxicity in U266 cells. U266 cells were cultured for 24 h with indicated concentrations of Bortezomib (A), Melphalan (B), and Doxorubicin (C) [control media (□), 1.25 nmol/l Bortezomib (.), 2.5 nmol/l Bortezomib (▲), 3.75 nmol/l Bortezomib (■); control media (□), 2.5 μmol/l Melphalan (▲), 5 μmol/l Melphalan (■); control media (□), 100 nmol/l Doxorubicin (▲), 200 nmol/l Doxorubicin (■), 400 nmol/l Doxorubicin (▲)] in the presence or absence of Cerulenin (0–25 μmol/l). Cell growth was assessed by MTT assay. Data represent mean ±SD of triplicate cultures. Combination index (CI) of Cerulenin and these agents was analysed using CalcuSyn software.

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