Heat shock protein 27 is a potential indicator for response to YangZheng XiaoJi and chemotherapy agents in cancer cells

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Abstract. Heat shock protein 27 (HSP27) is a member of the heat shock protein family which has been linked to tumour progression and, most interestingly, to chemotherapy resistance in cancer patients. The present study examined the potential interplay between HSP27 and YangZheng XiaoJi, a traditional Chinese medicine used in cancer treatment. A range of cell lines from different tumour types including pancreatic, lung, gastric, colorectal, breast, prostate and ovarian cancer (both wild-type and resistant) were used. Levels and activation of HSP27 and its potential associated signalling pathways were evaluated by protein array and western blotting. Knockdown of HSP27 in cancer cells was achieved using siRNA. Localisation and co-localisation of HSP27 and other proteins were carried out by immunofluorescence. Cell growth and migration were evaluated in their response to a range of chemotherapeutic agents. The present study first identified, by way of protein array, that YangZheng XiaoJi was able to inhibit the phosphorylation of HSP27 protein in cancer cells. Further demonstration was carried out by way of protein array, and the latter case, HSP27 and HSP70 were two of the gene products that were upregulated in a cisplatin resistant ovarian cancer cell line (19). Thus, this has caused an enormous challenge and increased the sensitivity of cancer cells to the inhibitory effect on cellular migration by YangZheng XiaoJi. YangZheng XiaoJi can act as an agent in first sensitising cancer cells to chemotherapy and secondly to overcome, to some degree, chemoresistance when used in an appropriate fashion in patients who have active HSP27.

Introduction

The heat shock proteins (HSPs) are ubiquitously expressed in almost all cells and across species. The protein family was so named owing to their initial discovery, namely, identified from cells under heat shock and is now known to be involved in a number of other cellular stresses including heat, carcinogens, mechanical and chemical stress (1,2). These proteins serve as molecular chaperones and act as a mechanism to assist protein folding, repair damaged proteins, or assist in degrading the unwanted proteins after stress and injury, thus by doing so the proteins protect the injured cells from potentially lethal damage (3,4). The HSP proteins are thus important regulators in normal physiology, but are also involved in diseases including cardiovascular, wound healing and cancer.

Similarly discovered from cells under heat shock, HSP27 has been found to be widely involved in other types of cell stress, including oxidative response (5-8). The HSP27 is a protein of approximately 27 kDa in size and belongs to the HSP family in which sizes vary between 8-150 kDa. The main cellular function of HSP27 is protecting cells from becoming apoptotic, or anti-apoptotic. Although this role is critically important in homeostasis, it has serious implications in clinical cancer. Firstly, HSP27 has been found to be overexpressed in a variety of human cancers and linked to a poor outcome for those cancer patients (9-13); secondly, high levels of HSP27 in cancer cells have been shown to promote cancer cell growth (14-16); and thirdly, HSP27 was found to be extremely elevated following chemotherapy, most probably as a protective mechanism to chemical stress (17-19). In the latter case, HSP27 and HSP70 were two of the gene products that were upregulated in a cisplatin resistant ovarian cancer cell line (19). Thus, this has caused an enormous challenge

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to cancer treatment and is frequently linked to chemotherapy resistance in cancer.

The key mode of activation of HSP27 is by phosphorylation. Phosphorylation of HSP27 allows itself to form oligomers which may facilitate the chaperone process under cellular stress (20,21). Phospho-HSP27 interacts with other pro-apoptotic proteins such as DAXX and ASK1 and therefore by blocking pro-apoptotic protein interaction blocks the apoptotic pathway (22,23). Other known pathways of HSP27 include interaction with AKT1 and the oestrogen receptor. By interacting with these signalling proteins, phospho-HSP27 blocks some of their actions in cancer cells. The most commonly reported sites of phosphorylation with HSP27 are S15, S78, S82 and S86. Consequently, phosphorylated HSP27 protein levels have been found to be frequently raised in clinical cancer tissues and chemoresistant cancer cells (24,25).

Together, it is suggested that HSP27 is a valid target for cancer treatment. Methods to either reduce the protein levels of HSP27 or the phosphorylation of HSP27, would be desirable in devising cancer treatment (26). A new anti-HSP27 agent, known as OGX427 (Apatorsen) has entered into a late phase clinical trial in patients with various malignant conditions including pancreatic, lung, bladder or prostate cancer, alone or in combination with other chemotherapies (27-30). Early studies have shown a favourable outcome from the new treatment.

There have been clinical trials which have shown that YangZheng XiaoJi (YZXJ), a combination of traditional Chinese medicinal herbs benefit cancer patients; however, how this mechanism of action is achieved remains unknown (31-34). The present study reports an accidental and surprising discovery that a traditional Chinese herbal medicine, known as YangZheng XiaoJi (YZXJ) used in treating patients with cancer, can suppress the phosphorylation of HSP27 and cell functions related to this protein.

Materials and methods

Materials. Antibodies to human HSP27 (sc-13132), caspase-3 (sc-7148), caspase-8 (sc-70501), caspase-9 (sc-17784), phospho-FAK (sc-81493), and GAPDH (sc-32233) were purchased from Santa Cruz Biotechnologies Inc. (Santa Cruz, CA, USA). Therapeutic agents including cisplatin, topotecan, paclitaxel, curcumin, 5-FU were purchased from Sigma-Aldrich (Poole, Dorset, UK). Antibodies to FAK (ab131435) and phospho-HSP-27 (S86) (ab17938) were purchased from Abcam (Cambridge, UK). FITC/TRITC-conjugated Phalloidin and FITC- and TRITC conjugated secondary antibodies were from Sigma-Aldrich. Secondary antibodies (fluorescence- and HRT-conjugated) were also from Sigma-Aldrich. Anti-HSP27 siRNA, control siRNA and transfection reagents were also obtained from Santa Cruz Biotechnologies Inc.

Cells. Human gastric cancer (AGS and HGC27), pancreatic cancer (PANC1), ovarian cancer (SKOV3 and COV504), lung cancer (A549 and SKMES1), breast cancer (MDA MB-231), prostate cancer (PC-3) cells were purchased from LGC Standard/ATCC (Southampton, UK). Ovarian cancer cells A2780 and its cisplatin resistant strain A2780/CP70 were gifts from Imperial College London (Dr Euan Stronach).

YangZheng XiaoJi extracts. An extract from YZXJ, named DME25 was prepared using a DMSO based method that has been described in full (35,36). The herbal medicinal formula, YangZheng XiaoJi was obtained from Yiling Pharmaceuticals (Shijiazhuang, HeBei, China). The formula contained the following 16 ingredients: Panax ginseng C.A. Mey, Astragalus membranaceus (Fisch.) Bge.var. mongolicus (Bge.) Hsiao, Ligustrum lucidum Ait, Curcuma phaeocaulis Val, Ganodema lucidum, Gynostemma pentaphylla (Thunb) Mak, Atractylodes macrocephala Koidz., Scutellaria barbata D.Don, Oldenlandia diffusa (willd.) Roxb, Poria cocos, Duchesnea indica Focke, Solanum lyratum Thunb, Artemisia scoparia (Bge.) Ki, Cynanchum paniculatum Kitag, Eupolyphaga sinensis Walker, and Gallus domesticus Brissin.

For experimental use, the extract was diluted in the respective cell culture media and the dilution range was between 1/100 to 1/2,000.

Immuno-fluorescent staining. Cells, plated with test agents in 8-well chamber slides, were first fixed using 4% formalin, lightly permeabilised with 0.1% Triton X100 for 5 min and blocked with a Tris buffer (25 mM, pH 8.4) that contained 7.5% pre-immunized goat serum for 1 h. Primary antibodies (including anti-HSP27, anti-phospho-HSP27 (S86), anti-caspases), diluted in the blocking buffer was added to the respective slides which were kept in the dark at full humidity on a slow moving platform for 1 h, and then washed thoroughly. FITC-tagged secondary antibodies were then added. After 1 h, the slides were washed thoroughly and mounted using FluorSave™ (Calbiochem, Nottingham, UK). TRITC-conjugated phalloidin was diluted at final concentration of 10 µg/ml and added to the cells together with the secondary antibodies. The slides were examined on an Olympus microscope and photographed using a Hamamatsu digital camera. The staining intensity was determined using ImageJ software.

Knockdown of HSP27. Cells at approximate 70% confluence were rinsed with BSS buffer. Anti-HSP27 siRNA, diluted in RNA free water was first mixed with a transfection reagent and left at room temperature for 30 min before being added to the cells. After 7 h medium was replaced and cells were left to incubate for another 24 h. Expression of HSP27 was tested using RT-PCR.

Cell growth assay. Cancer cells were first seeded in 96-well tissue culture plates and allowed to adhere. The test materials and their combinations were then added to the cells, which were subsequently incubated for a period of 72 h. After removing the culture media, cells were fixed with 4% formalin and stained with 0.5% Crystal violet. After extensive washing, the staining was extracted with 10% acetic acid and plates were read on a multiple channel plate reader at 540 nm (Elx800; Bio-Tek, Swindon, UK). The test regimens included chemotherapeutic agent alone, DME25 alone, DME25 and chemotherapeutic agent combination and DME25 pre-treatment followed by chemotherapeutic agent.

Protein arrays for detecting phosphorylation changes. Cancer cells at 90% confluence in two T75 tissue culture flasks were washed with BSS buffer and then placed into a fresh batch
of DMEM supplemented with 5% FCS. After 5 h, treatment was added, again in 5% FCS. After a 5-h period, cells were removed from the flasks with a cell scraper. The cells were pelleted using a centrifuge at 2,500 rpm for 5 min. Lysis buffer was added to the cell pellets and placed on a spinning wheel for 1 h at 4°C. The lysates were then spun at 12,000 x g for 10 min at 4°C. The supernatants were carefully collected and the insolubles discarded. Based on absorbance the protein concentration in the cell lysates were quantified using the scatter line chart of Microsoft Excel and then adjusted to 2 mg/ml. The samples were stored at -20°C until use. Antibody based protein arrays, namely KAm850, which has 854 capture antibodies spotted on to each array slides (Kinexus Bioinformatics Ltd., Vancouver, Canada) were used in the present study. The following are the key parameters collected and used for the data analysis: Globally Normalized Signal Intensity, Background corrected intensity values are globally normalized. The Globally Normalized Signal Intensity was calculated by summing the intensities of all the net signal median values for a sample. Z Scores, Z score transformation corrects data internally within a single sample. Z Score Difference, The difference between the observed protein Z scores in samples in comparison. Z Ratios, Divide the Z Score Differences by the SD of all the differences for the comparison. 

Electric cell-substrate impedance sensing (ECIS) based analyses on cell adhesion and cell migration. Briefly, 96-well W96E1 microarrays were used on the ECIS Ztheta instrument (Applied Biophysics Ltd., Troy, NJ, USA). Lung cancer cells were added to the wells of the array, followed by immediate tracking of cell adhesion over a range of frequencies (1,000 to 64,000 Hz) using automated modules. The adhesion was analysed using the mathematical modelling methods as previously described. For cellular migration, confluent lung cancer monolayers in the arrays were electrically wounded (2,000 mA for 20 sec each), after which the migration of the cells was immediately tracked, again over a range of frequencies. All the experiments were conducted in triplicate.

Western blotting analysis of HSP27 proteins. Proteins, prepared in a similar fashion to that for protein analysis, were separated on a SDS-PAGE gels, following transfer of
proteins to nitrocellulose membranes, the membranes were probed with the respective antibodies in a set protocol that including extensive washings in between the antibody probing and finally detecting the antibody signals using chemiluminescence reagents. The membranes were visualised using an imager (G-Box; Syngene, Cambridge, UK).

Results

DME25 has a marked effect on the phosphorylation of HSP27. Using a protein array, we found that the YZXJ extract DME25 resulted in a marked reduction of phosphorylation of HSP27, particularly on Serine86 (S86) phosphorylation, in SKMES1 lung cancer and PANC-1 pancreatic cancer cells (Fig. 1A and B, respectively).

Localisation of HSP27 and phospho-HSP27 (S86) in cancer cells, assessed by immunofluorescence. HSP27 was seen broadly in the nucleus and in cytoplasmic region of lung cancer cells (Fig. 2A, top panel). It is very interesting to note that both total HSP27 and phospho-HSP27 was localised in focal adhesion and pseudopodia regions of the cells. Treatment of the lung cancer cells, SKMES1 with DME25 resulted in loss of phospho-HSP27 from the focal adhesion and pseudopodia

Table I. Effects of cisplatin on the growth of cancer cells in combination with DME25.

| Cisplatin | 16 ng/ml | YZXJ | 400 ng/ml | YZXJ | 10 µg/ml | YZXJ |
|-----------|----------|------|-----------|------|----------|------|
| AGS       | 7.9±2.5  | 9.1±0.7 | 20.8±2.0 | 14.5±2.6 | 93.1±0.5 | 91.5±0.5 |
| MIAPACA2  | 50.1±4.5 | 92.1±1.5 | 77.5±1.5 | 93.1±4.4 | 77.8±2.5 | 93.2±2.6 |
| A549      | 20.1±1.7 | 12.2±2.7 | 50.4±1.3 | 44.1±1.7 | 90.4±0.4 | 89.9±0.2 |
| SKMES1    | 1.7±0.3  | 34.6±1.7 | 23.0±1.4 | 46.3±2.0 | 80.5±3.5 | 86.9±2.6 |
| COV504    | NC       | 3.8±6.2  | 27.2±4.7 | 43.4±2.7 | 74.6±2.8 | 85.1±1.3 |
| SKOV3     | NC       | 17.2±1.9 | 0.5±1.5  | 9.8±2.6  | 46.7±1.5 | 66.7±2.7 |

NC, negative control.
regions of the cells, although the changes of total HSP27 did not appear to differ (Fig. 2). The same changes of phospho-HSP27 were seen in pancreatic cancer cells, PANC1 (Fig. 2A, bottom panel) as well as in ovarian cancer cell SKOV3 and gastric cancer AGS cells (data not shown). The inhibition on levels and activation of HSP27 was also demonstrated by western blotting analysis in both the lung and pancreatic cells (Fig. 2B and C, respectively).

Activated HSP27 is co-localised with caspase-9 in cancer cells, which is prevented when cells are treated with DME25. When co-stained for phospho-HSP27 (S86) and caspase-9, it was found that both molecules co-localised in regions of focal adhesion (FAC) and pseudopodia (Fig. 3A and B). It was very interesting to note that when cells were treated with DME25, this pattern of co-localisation appears to break (Fig. 3B).

Phospho-HSP27 and its co-localisation with filamentous actin and caspase-9 in cancer cells and in chemoresistant cancer cells. Using dual fluorescence staining, we have shown that phospho-HSP27 and F-actin did co-localise in FAC regions in both wild-type ovarian cancer A2780 (Fig. 4A, top panels) and in cisplatin resistant A2780-CP70 cells (Fig. 4 bottom panels). The same was seen with lung cancer cells (Fig. 5). In all the cells tested, treatment with DME25 reduced the location of phospho-HSP27 to the FAC region. The co-localisation...
of phospho-HSP27 and caspase-9 in wild-type A2780 and cisplatin resistant A2780/CP70 were also reduced in the presence of DME25 (Fig. 4B).

Effect of DME25 on the growth of ovarian cancer cells. A2780/CP70 showed approximately 8 times less sensitivity to cisplatin compared with the wild-type parental cells (Fig. 6). Co-culture of both cisplatin and DME25 resulted in the cells becoming more sensitive to cisplatin, as seen for lung, gastric, pancreatic and ovarian cancer cells (Tables I-V). However, knocking down HSP27 by siRNA did not significantly influence the sensitivity of the cells to the chemotherapeutic agents (data not shown).

HSP27 shows marked influence on cell migration and acts synergistically with YangZheng XiaoJi on cancer cell migration. In light of the co-localisation of HSP27 and phalloidin at the focal adhesion regions of cancer cells and our previous reports on the impact of DME25 on the migration of various cells including cancer cells, we tested how HSP27 might affect cell migration. It was readily demonstrable that knocking down HSP27 resulted in a sustained reduction of cell migration (Fig. 7A and B). It was further confirmed that knocking down HSP27 increased sensitivity of cancer cells to low dosage chemotherapeutic agents and in particular when DME25 was present (Fig. 7C and D).

Discussion
The current study stemmed from a surprising finding in our search for the potential pathways that are influenced by
HSP27, YangZheng XiaoJi and sensitivity to chemotherapeutic agents. The role of HSP27 in the sensitivity to chemotherapy agents is not without controversy. For example, it has been reported recently that overexpression of HSP27 in pancreatic cancer cells would increase the sensitivity of the cancer cells to gemcitabine (37). Although reasons for this are not clear, the type of drugs used in the study may be an explanation, as these drugs are transported and act in cancer cells via different mechanisms.

HSP27 acts as a cyto-protective molecule during androgen independence in prostate cancer patients and in prostate cancer cell models (38). Likewise, levels of HSP27 correlated with resistance to irinotecan in colorectal cancer cells (39). HSP27, together with HSP70, is linked to the sensitivity of colorectal cancer to the naturally-occurring anticancer agent curcumin (40).

Knocking down HSP27 sensitises glioblastoma multiforme tumour cells to the HSP90 inhibitor, 17-N-allylamino-17-demethoxygeldanamycin (17-AAG) and staurosporine by changes in apoptosis (41). The same was seen in glioma cells, in that silencing HSP27 resulted in an increase in apoptosis in response to temozolomide and quercetin (42).

HSP27, YangZheng XiaoJi and cell migration. The finding that HSP27 is co-localised with phalloidin to the focal adhesion

| 5-FU  | 1.6 µg/ml   | 40 µg/ml   | 1000 µg/ml |
|-------|-------------|------------|------------|
|       | Control     | YZXJ       | Control    | YZXJ       | Control    | YZXJ       |
| AGS   | 13.1±1.6    | 3.6±1.8    | 59.6±0.8   | 56.8±1.6   | 79.2±0.6   | 75.3±2.1   |
| MIA PACA2 | NC        | 57.8±1.5   | 33.4±5.9   | 70.8±3.2   | 64.4±3.5   | 88.6±7.8   |
| A549  | 15.0±3.1    | 12.7±4.1   | 59.2±2.0   | 57.6±3.5   | 70.3±1.8   | 69.9±2.1   |
| SKMES1| 10.5±2.3    | 29.0±1.4   | 51.9±3.4   | 65.0±1.3   | 57.5±2.6   | 68.2±2.1   |
| COV504| 15.6±1.4    | 48.2±1.6   | 40.5±2.4   | 59.1±2.6   | 50.4±0.9   | 63.7±1.6   |
| SKOV3 | 10.4±0.8    | 20.9±2.8   | 32.7±43.5  | 43.9±1.9   | 46.0±1.8   | 60.9±1.4   |

NC, negative control.
regions of cancer cells is interesting. From the immunofluorescence staining, HSP27 particularly the phosphorylated form of the protein, is frequently co-localised with Phalloidin at the focal adhesion regions of cancer cells, where caspase-9 is also seen. It is of interest to note that inclusion of YZXJ extract DmE25 in the study markedly reduce the presence of phospho-HSP27 (S86) at the focal adhesions. Given the key role of focal adhesion in cell-matrix adhesion and cellular migration, it is plausible to suggest that activation of HSP27 (by way of phosphorylation) coordinates with other proteins in the focal adhesion area, such as integrins and the focal adhesion kinase (FAK) to regulate the adhesion and migration of cancer cells. We have recently reported that YZXJ was able to inhibit the phosphorylation of FAK in cancer cells and in endothelial cells (35,43).

Together with the finding that knocking down HSP27 resulted in an increase in cellular migration in cancer cells, it is suggested that HSP27, in particular when combined with YZXJ, is an important regulator of EMT reversal or MET. It has been reported recently that HSP27 is key to IL-6 dependent and IL-6 independent EMT in prostate cancer cells, by acting on IL-6 induced activation of STAT3/Twist and Snail (44,45). It is also involved in EGF induced EMT in prostate cancer cells by influencing the migration of these cells (46,47). Other than the reported effects on cancer cells, HSP27 has also been shown to be a mediator for angiogenesis (48), suggesting that HSP27 is involved in a wider range of processes in cancer development and progression. Taken together, it is clear that HSP27 is a viable target in cancer treatment and that a combined approach, with either YZXJ or another method, would be a valid consideration when devising a therapy in either targeting HSP27 or in certain chemoresistance cases.

Table IV. Effects of topotecan on the growth of cancer cells in combination with DME25.

| Topotecan       | Control   | YZXJ       | Control   | YZXJ       | Control   | YZXJ       |
|----------------|-----------|------------|-----------|------------|-----------|------------|
| AGS            | 7.6±0.6   | 18.3±3.7   | 46.0±5.8  | 65.9±2.6   | 88.1±2.0  | 86.5±2.8   |
| MIAPACA2       | 0.9±0.9   | 65.6±4.1   | 39.5±3.3  | 80.2±2.6   | 66.1±0.9  | 91.6±4.5   |
| A549           | NC        | NC         | 48.9±4.3  | 60.0±2.0   | 68.8±1.3  | 66.3±1.6   |
| SKOV3          | 11±0.7    | 30.2±2.1   | 38.9±1.1  | 50.1±2.7   | 79.9±2.8  | 62.1±2.8   |
| COV504         | NC        | 23.1±0.8   | 24.9±5.1  | 50.12±12.5 | 39.4±1.8  | 79.9±2.8   |
| SKMES1         | 7.8±0.6   | 40.4±2.8   | 62.3±1.4  | 88.4±3.4   | 50.6±1.1  | 72.6±2.7   |

Table V. Effects of cisplatin on the growth of wild-type and cisplatin-resistant ovarian cancer cells in combination with DME25.

| Treatment      | A2780     | A2780/CP70 |
|----------------|-----------|------------|
|                | 160 ng/ml | 4 µg/ml    | 160 ng/ml | 4 µg/ml    |
| Cisplatin only | 12.3±8.2  | 72.5±2.5   | 4.7±4.0   | 40.8±8.6   |
| Cisplatin + YZXJ | 72.4±3.6  | 80.5±3.3   | 26.8±0.4  | 46.3±1.3   |

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