Heat Shock Protein 25-Enriched Plasma Transfusion Preconditions the Heart against Doxorubicin-Induced Dilated Cardiomyopathy in Mice

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ABSTRACT

Extracellular heat shock proteins (eHsps) in the circulation have recently been found to activate both apoptotic and protective signaling in the heart. However, the role of eHsps in doxorubicin (Dox)-induced heart failure has not yet been studied. The objective of the present study was to determine how Dox affects circulating eHsp25 in blood plasma and how eHsp25 affects Dox-induced cardiomyopathy. Wild-type mice [HSF-1(+/+) were pretreated with 100 μl of heterozygous heat shock factor-1 [HSF-1(+/-/)] mouse plasma (which contained 4-fold higher eHsp25 compared with wild-type mice), HSF-1(+/+) plasma, or saline, before treatment with Dox (6 mg/kg). After 4 weeks of this treatment protocol, HSF-1(+/+) plasma-pretreated mice showed increased eHsp25 in plasma and improved cardiac function (percentage of fractional shortening 37.3 ± 2.1 versus 26.4 ± 4.0) and better life span (31 ± 2 versus 22 ± 3 days) compared with the HSF-1(+/+) plasma or saline-pretreated mice. Preincubation of isolated adult cardiomyocytes with HSF-1(+/+) plasma or recombinant human Hsp27 (rhHsp27) significantly reduced Dox-induced activation of nuclear factor-κB and cytokine release and delayed cardiomyocyte death. Moreover, when cardiomyocytes were incubated with fluorescence-tagged rhHsp27, a saturation in binding was observed, suggesting that eHsp25 can bind to surface receptors. Competitive assays with a Toll-like receptor 2 (TLR2) antibody reduced the rhHSP27 binding, indicating that Hsp25 interacts with TLR2. In conclusion, transfusion of Hsp25-enriched blood plasma protected the heart from Dox-induced cardiotoxicity. Hsp25 antagonized Dox binding to the TLR2 receptor on cardiomyocytes.

Introduction

The clinical utility of doxorubicin (Dox) is severely limited by the incidence of life-threatening dilated cardiomyopathy among patients who receive a cumulative dose exceeding 550 to 600 mg/m² during chemotherapeutic regimens (Minotti et al., 2004; Peng et al., 2005). Even though several pathways such as mammalian target of rapamycin inhibition (Zhu et al., 2009), phosphodiesterase-5 inhibition (Fisher et al., 2005), reduction in circulating progenitor cells (Huang et al., 2010), activation of Toll-like receptors (Nozaki et al., 2004; Riad et al., 2008), impaired metabolism (Maslov et al., 2010), and triggering autophagy (Kobayashi et al., 2010) have been identified, the precise mechanism of Dox-induced cardiotoxicity still remains unclear. We recently identified a novel pathway: that heat shock factor (HSF-1) is activated because of Dox-induced oxidative stress, and this activation enhanced the expression of heat shock proteins including Hsp25 in Dox-treated mouse hearts (Vedam et al., 2010). Systemic stresses, such as genotoxic (i.e., Dox treatment) and proteotoxic (accumulation of denatured proteins) stresses, induce HSF-1 activation and enhance stress-inducible protein expression in eukaryotic cells (Baler et al., 1993; Sarge et al., 1993; Xiao et al., 1999). HSF-1 is activated by diverse forms of stress, and the activation of HSF-1 occurs via a multistep process (Sakamoto et al., 2006). Induction of Hsp25 in Dox-treated hearts was shown to regulate p53 transcriptional activity and expression of the proapoptotic protein Bax.

ABBREVIATIONS: Dox, doxorubicin; HSF-1, heat shock factor 1; Hsp, heat shock protein; eHsp, extracellular heat shock protein; TLR, Toll-like receptor; LV, left ventricular; H&E, hematoxylin and eosin; rh, recombinant human; OG-rhHsp27, Oregon Green-labeled recombinant human HSP27; IL-6, interleukin-6; TNF-α, tumor necrosis factor-α; FS, fractional shortening; DAPI, 4,6-diamidino-2-phenylindole; MAPK, mitogen-activated protein kinase; BBI, beat-to-beat interval; NF-κB, nuclear factor-κB.
(Vedam et al., 2010). Many studies have reported that over-expression of small Hsps, such as Hsp27 (the Hsp25 ortholog), protects the heart (Liu et al., 2007; Fan et al., 2008). Paradoxically, however, there is compelling evidence that an elevated level of Hsp25 or Hsp27 is present in failing hearts (Dohke et al., 2006). Thus, it is not clear whether this increase in Hsp27 in the failing human heart (Hsp25 in murine heart) is the result of an unsuccessful protective mechanism or whether the increased expression of Hsp25 indeed potentiates the loss of cardiomyocytes in the heart.

Extracellular Hsps (eHsps), which are present in blood plasma, are being actively studied for their role in innate immunity (Mullhoff, 2006; Schmitt et al., 2007; Dhodapkar et al., 2008). Dox has been reported to activate monocytes and macrophages in the circulation and induce an immunogenic response through activation of interleukin-1 family receptors such as Toll-like receptors (TLRs) (Riad et al., 2008). Thus, we hypothesized that enhancing circulating eHsp25 would antagonize Dox by competitive binding to TLRs, present on immune cells such as macrophages and monocytes, and have been identified as potential targets for Hsps. Upon binding to these receptors, Hsps were found to activate downstream signaling to enhance cytokine secretion as a part of the immune response. Recent studies have shown that cardiomyocytes express TLRs (Petersen et al., 2005; Boyd et al., 2006), and Hsps can bind to these receptors (Roelofs et al., 2006; Kim et al., 2009). In particular, Hsp60 has been found to competitively bind to TLR4 and enhance apoptosis in cardiomyocytes (Lewthwaite et al., 2002; Kim et al., 2009). TLR2 has been identified as a receptor for Dox, and its deletion was found to be protective against Dox-induced cardiotoxicity (Nozaki et al., 2004). Independent studies have shown that circulating eHsp25 could interact with TLRs and act as a putative ligand (Kardys et al., 2008). Thus, we hypothesized that enhancing circulating eHsp25 would antagonize Dox by competitive binding to TLR and prevent Dox-induced cardiotoxicity. Hence, the objective of this study was to establish the relationship between the plasma level of Hsp25 (eHsp25) and Dox-induced cardiomyopathy and to determine whether circulating eHsp25 could protect the heart from Dox.

**Materials and Methods**

**Animals and Treatment.** HSF-1 knockout founders were obtained as a generous gift from Dr. Ivo von Benjamin, University of Utah, crossed with BALB/c (Charles River Laboratories, Inc., Wilmington, MA) mice, and genotyped (Vedam et al., 2010). Eight-week-old HSF-1(+/-), HSF-1(+/-), and HSF-1(-/-) mice in the BALB/c background were used in all studies. All the protocols were approved by the institutional animal care and use committee and were performed according to the National Institutes of Health Guide for the Care and Use of Laboratory Animals (Institute of Laboratory Animal Resources, 1996). Blood was drawn by cardiac puncture, and plasma was collected with heparin as an anticoagulant. Immunodepletion of Hsp25 in the plasma was carried out by immunoprecipitating Hsp25 from whole plasma, using an anti-Hsp25 antibody. The absence of Hsp25 in the immunodepleted plasma was confirmed by Western blotting before injection into mice. Dox dosing was via bolus intraperitoneal injection of 6 mg/kg b.wt. per week for a total of 4 weeks. This dose was reported to cause Dox-induced heart failure in mice within 4 weeks (Yoshida et al., 2009). Plasma infusion was performed through tail vein cannulation. A detailed description of all of the experimental methods is available in the supplemental data.

**Survival and Weight Loss Studies.** Two sets of survival studies were performed. In the first set, HSF-1(+/-), HSF-1(+/-), and HSF-1(-/-) mice were injected with Dox, and survival was followed. In the second set, mice were pretreated with saline or HSF-1(+/-) plasma or HSF-1(-/-) plasma followed by Dox. The Kaplan-Meier curve was plotted and analyzed for 50% survival time in each group.

**Two-Dimensional Echocardiography.** M-mode echocardiography was performed using a GE Vivid 7 ultrasound imaging system (software version 3.4.2; GE Healthcare, GE Healthcare, Chalfont St. Giles, Buckinghamshire, UK) equipped with a 13-MHz linear-array transducer. Two-dimensional short axis echocardiography was performed at the level of the midpapillary muscle. All measurements were averaged from a minimum of three cardiac cycles. LV end-diastolic and end-systolic internal diameters were measured. LV mass and fractional shortening (percentage) were calculated as described before (Vedam et al., 2010).

**Immunoblot Analysis.** Immunoblotting in whole-cell lysates and immunoprecipitates were performed using routine procedures (Vedam et al., 2010).

**Histopathology.** To examine histological changes at the end of 4 weeks, hearts were harvested and fixed in 10% neutral buffered formalin, routinely processed, and embedded in paraffin. Paraffin sections were cut into 4-μm serial sections. Counterstaining with hematoxylin and eosin (H&E) was performed.

**Immunofluorescence Staining in Heart Tissue by Confocal Microscopy.** Tissue sections were incubated with both rabbit anti-α-actin antibody (Abcam, Cambridge, MA) and mouse anti-Hsp25 antibody (Sigma-Aldrich, St. Louis, MO), and the slides were stained with both green Alexa Fluor 488-conjugated goat anti-rabbit secondary antibody and red Alexa Fluor 555-conjugated donkey anti-mouse secondary antibody (Invitrogen, Carlsbad, CA) for fluorescent imaging.

**Recombinant Human Hsp27 Expression, Purification, Tagging with Oregon Green Fluorescent Dye and Binding to Cardiomyocytes.** Recombinant human (rh) Hsp27 (low-endotoxin, human ortholog of murine Hsp25) was used in the present work because it has been very well established in cardiology and cell biology. Human Hsp27 was expressed in E. coli (Panasenko et al., 2002). Expression and purification of recombinant human Hsp27 were performed as described before (Panasenko et al., 2002). Denaturation and renaturation of Hsp27 were also performed as described before (van de Klandert et al., 1998). rhHsp27 was labeled with Oregon Green 488 according to the manufacturer’s protocol (F-6153; Invitrogen). The final labeled protein concentration was quantitated by measuring the absorbance at 280 nm and correcting the absorbance for Oregon Green and the extinction coefficient for Hsp27 (Kim et al., 2009). Cardiomyocytes were incubated with Oregon Green-labeled recombinant human Hsp27 (OG-rhHsp27) with increasing concentrations for 30 min at 4°C and washed, and fluorescence was determined as described by Habich et al. (2002). In competitive assays, the cells were preincubated with anti-TLR2 antibody (Abcam). Dox fluorescence in cardiomyocytes was determined with excitation at 546 nm, measuring emission at 565 nm (Shen et al., 2008).

**Cardiomyocyte Beat-to-Beat Interval Studies.** Isolated cardiomyocytes were treated with Dox alone or HSF-1(-/-) plasma + Dox, and spontaneous contractions of a single cardiomyocyte (supplemental video) were recorded until the cardiomyocyte in focus lost its rod morphology (becomes rounded). Contractility was manually counted as a function of time from the recorded video, and beat-to-beat interval was determined.

**Statistical Analysis.** All values are expressed as mean ± S.E. Significance analyses were performed using one-way analysis of variance followed by a post hoc procedure wherever needed, using SPSS software (version 16; SPSS Inc., Chicago, IL). A log-rank test.
was applied in Kaplan-Meier curves. Statistical significance was accepted at a value of \( P < 0.05 \).

**Results**

**Elevation of Circulating Extracellular Hsp25 in Dox-Treated Mice.** HSF-1\((+/+)\), HSF-1\((-/-)\), and HSF-1\((-/-)\) mice were treated with Dox (6 mg/kg b.wt.) or the vehicle (saline) for 4 weeks, and circulating plasma eHsp25 was determined. Figure 1A shows the Western blots of eHsp25 in plasma of HSF-1\((+/+)\), HSF-1\((-/-)\), and HSF-1\((-/-)\) mice, after 4 weeks of Dox treatment. Quantitative comparison of the basal eHsp25 level in plasma, determined before the Dox regimen was started (Fig. 1B), showed higher eHsp25 in HSF-1\((+/+)\) (approximately 4-fold) and HSF-1\((-/-)\) (approximately 3-fold) mice, compared with HSF-1\((+/+)\) mice, suggesting that the circulating eHsp25 level is higher in HSF-1\((-/-)\) mice. Given the fact that HSF-1 knockout animals are supposed to show low levels of Hsps including Hsp25, and indeed it was reported so in different tissues of these mice previously (Yan et al., 2002), a higher plasma level of eHsp25 in HSF-1\((-/-)\) mice indicates that some other unknown factors are activating Hsps expression in circulating cells. Previous studies have shown that mutant and wild-type hetero trimers of HSF-1 are transcriptionally more active than the HSF-1 homo trimers (HSF-1 trimers are the active form of the HSF-1 transcriptome) (Voelmy, 2005), which may be responsible for the increased basal level eHsp25 in heterozygous mice.

Upon treatment with Dox, the eHsp25 level progressively increased with each week in HSF-1\((+/+)\) mice (Fig. 1B; Supplemental Fig. 1), whereas in the HSF-1\((-/-)\) groups, there was no significant further increase from the already higher basal level (Fig. 1B). The increase in HSF-1\((+/+)\) mice started as early as the 2nd week (Fig. 1B; Supplemental Fig. 1). Of interest, Dox-induced cardiotoxicity, determined in terms of cardiac dysfunction, showed a very minimal change in ejection fraction in the HSF-1\((-/-)\) hearts and only a moderate reduction in HSF-1\((-/-)\) hearts (Supplemental Fig. 2), unlike the severe cardiac dysfunction noted in HSF-1\((+/+)\) mice. Furthermore, the survival analysis of these mice indicated an inverse correlation between survival and basal eHsp25 level (Fig. 1, B and C). The 50% death (log-rank test) was 23 ± 3 days in the HSF-1\((+/+)\) group (lowest basal eHsp25) and 36 ± 4 days in the HSF-1\((-/-)\) group, and, strikingly, no significant death (50% death >100 days) was observed in the HSF-1\((-/-)\) group (highest basal eHsp25).

Because inflammatory cells such as macrophages are known to be activated upon treatment with Dox (Krysko et al., 2011) as an innate immunogenic response, we determined whether there was any increased expression of Hsp25 in macrophages of Dox-treated mice. Intraperitoneal cavity macrophages were isolated from Dox- or saline-treated HSF-1\((+/+)\) and HSF-1\((-/-)\) mice (Fig. 1D) and used for Hsp25 determination. Hsp25, determined by Western blotting, increased (3-fold) in macrophages of Dox-treated HSF-1\((+/+)\) mice (Fig. 1E), compared with saline-treated macrophages. However, in the HSF-1\((-/-)\) group, the macrophages of saline-treated mice showed a very high level of Hsp25 (even higher than that for the Dox-treated HSF-1\((+/+)\) group) (Fig. 1E), and no significant increase was produced by Dox treatment. In macrophages, IL-6 increased upon treatment with Dox in both groups, indicating activation of inflammatory cells (Fig. 1F). We have also found that basal IL-6 was higher in HSF-1\((-/-)\) mice than in HSF-1\((+/+)\) mice (Supplemental Fig. 3). These results suggest that the plasma
level of eHsp25 is probably regulated via an immunogenic responsive mechanism, involving monocyte/macrophage cells and that this mechanism is constitutively active in an HSF-1-deficient setting.

**HSF-1(+/−) Plasma Transfusion Improves Survival and Attenuates Cytokine Release.** Three groups of HSF-1(+/+) wild-type mice were infused with either HSF-1(−/−) plasma (100 μl), HSF-1(+/−) plasma, or an equal volume of saline. Twenty-four hours later, all of these groups were divided into two and infused with either Dox (6 mg/kg) or saline (resulting in six groups: saline alone; saline + Dox; HSF-1(+/+) plasma alone; HSF-1(+/−) plasma + Dox; HSF-1(−/−) plasma alone; and HSF-1(−/−) plasma + Dox). This infusion was repeated for 4 weeks, resulting in four doses of HSF-1(−/−) plasma and Dox (Supplemental Fig. 4). Figure 2A shows survival curves for these six groups. Wild-type mice, treated with HSF-1(+/−) plasma + Dox, showed 50% survival at 31 ± 2 days, which was higher than that for either the HSF-1(+/-) plasma + Dox-treated group (23 ± 4 days) or the saline + Dox-treated group (22 ± 3 days). Moreover, only 60% mortality over 50 days (by the end of the study period) was observed in the HSF-1(+/-) plasma-treated mice, whereas in the HSF-1(+/−) plasma + Dox-treated group or saline + Dox-treated group, more than 80% mortality was observed. Body weight loss induced by Dox as a result of its systemic toxicity was also found to be less for the HSF-1(+/-) plasma-transfused group (Supplemental Fig. 5), illustrating the decreased toxicity in this group. The eHsp25-depleted (by immunoprecipitation using an Hsp25-specific antibody) HSF-1(+/-) plasma transfusion group did not show increased survival, confirming that eHsp25 in plasma is responsible for the enhanced survival of that cohort (Supplemental Fig. 6). Moreover, mice injected with purified rhHsp27 before Dox treatment showed an increase in survival, but the increase was not equal in magnitude to that found for the HSF-1(+/-) plasma treatment group, perhaps because of lipopolysaccharide contamination or denaturing of rhHsp27, causing death among the cohort (Kim et al., 2009).

Even though only 100 μl of HSF-1(+/-) plasma was infused four times, a 4-fold increase in eHsp25 was observed (which was not proportional to the infused amount), suggesting that infusion of HSF-1(+/-) plasma enhances the production of eHsp25, potentially through activation of macrophages in the circulation (Fig. 1E). Moreover, the fact that the plasma level of eHsp25 was further elevated (Fig. 2, B and C) in the HSF-1(+/-) plasma + Dox group, despite less systemic toxicity, indicated that HSF-1(+/-) plasma + Dox transfusions primed the secretory mechanism to release more eHsp25 in response to Dox. To test the fact that immune cells are activated upon plasma transfusion, cytokines, namely IL-6 and TNF-α, were determined. Both IL-6 and TNF-α were increased by treatment with HSF-1(+/-) plasma (Fig. 2, D and E). IL-6 showed a 4-fold increase upon treatment with Dox alone, and although there was an increase in IL-6 in the HSF-1(+/-) plasma + Dox-treated groups, the increase was significantly blunted relative to that for the Dox alone-treated group (Fig. 2D). Likewise, HSF-1(+/-) plasma treatment almost completely removed the Dox-induced increase in TNF-α (Fig. 2E).

![Figure 2](jpet.aspetjournals.org)
Cardioprotection against Dox in HSF-1(+/−) Plasma-Transfused Mice. Figure 3A shows representative echocardiograms of four groups, obtained four weeks after the last dose of Dox. In the HSF-1(+/−) plasma alone- or saline alone-treated groups, cardiac function remained unchanged. The saline + Dox-treated group showed the “dilated cardiomyopathy” phenotype, demonstrated by significant contractile dysfunction observed in the two-dimensional M-mode echocardiograms. The HSF-1(+/+) plasma + Dox-treated group also showed a similar phenotype (data not shown). However, in the HSF-1(+/−) plasma + Dox-treated group, better contractile function relative to that for the other groups was observed (Fig. 3A). The quantitative percentages of fractional shortening (FS), obtained for all the four groups over the study period, are summarized in Fig. 3B. The percentage of FS was 49.4 ± 2 for saline-infused groups. In the saline + Dox-treated group, the percentage of FS decreased to 26.4 ± 4, whereas in the HSF-1(+/−) plasma + Dox-treated group, the percentage of FS was 37.3 ± 2.1, significantly higher than that in the saline + Dox-treated group. Likewise, the LV thickness was significantly higher in the HSF-1(+/−) plasma + Dox group, relative to that in the Dox-treated group (0.66 ± 0.05 and 0.63 ± 0.05 mm, respectively) (Fig. 3C). These results illustrate that the increased survival in the HSF-1(+/−) plasma-pretreated group (Fig. 2A) is probably due to reduced cardiotoxicity. Histopathological evaluation (H&E staining) of LV tissue of all four groups was performed to determine whether there was any reduced tissue damage in the HSF-1(+/−) plasma-pretreated group. The saline or HSF-1(+/−) plasma alone-treated group showed normal myofibrillar structure with striations, branched appearance, and continuity with adjacent myofibrils. The Dox alone-treated group showed extensive myofibrillar degeneration (Fig. 4A). Tissue sections from the HSF-1(+/−) plasma + Dox-treated groups showed normal myofibrillar structure with mild degenerative changes and continuity with adjacent myofibrils (Fig. 4A), and the morphology of cardiac muscle fibers was relatively well preserved. Quantitative assessment of Dox-induced myocardial damage showed a significant decrease in HSF-1(+/−) plasma-pretreated groups (Fig. 4B).
HSF-1(+/−) Plasma Transfusion Reduces HSF-1 Activation and Hsp Expression and Suppresses Apoptosis in Dox-Treated Hearts.

Western blots and quantitative results for the four major Hsps (Hsp90, Hsp70, Hsp60, and Hsp25, determined in heart tissues after 4 weeks of Dox injection) are presented in Fig. 5. Dox treatment induced all major Hsps and significantly increased them in the heart. Such an enhanced expression of Hsps, as a response to stress has been reported to be due to HSF-1 activation in the heart (Turakhia et al., 2007; Vedam et al., 2010). However, in the HSF-1(+/−) plasma + Dox-treated group, the Hsp levels were very much reduced compared with those in the saline + Dox-treated groups, indicating that HSF-1(+/−) plasma pretreatment inhibited Dox-induced HSF-1 activation in the heart. There was no significant induction of Hsps in the saline alone- or plasma alone-treated groups (Fig. 5, A–C).

Although all major Hsps were up-regulated upon treatment with Dox, we focused on Hsp25 because it has been identified as a regulator of p53 activity (Vedam et al., 2010). Immunohistochemical tissue staining was performed in all four groups of hearts to determine whether distribution of Hsp25 expression in the Dox-treated hearts was uniform. Figure 5D summarizes the images of LV tissues stained for Hsp25. As seen, uniform positive staining was confirmed in Dox-treated heart tissues, whereas in the HSF-1(+/−) plasma-pretreated group, Dox treatment did not enhance Hsp25. These results demonstrate that the Dox-induced Hsp expression in the heart is inhibited by HSF-1(+/−) plasma pretreatment, suggesting that the cardioprotection observed in the plasma-pretreated group could be due to inhibition of the HSF-1 activation in these hearts. Further experiments were performed to determine whether these changes in Hsp25 expression occurred in cardiomyocytes. Coregistration of cardiomyocyte-specific α-actin and Hsp25 was performed. Double immunofluorescence staining and confocal microscopy analysis were used in heart tissues. First, heart tissues were co-stained with a cardiomyocyte α-actin rabbit antibody (Abcam) and an Hsp25 mouse antibody (Sigma-Aldrich), followed by anti-rabbit green fluorescent and anti-mouse Texas red secondary antibodies, and stained with DAPI for nuclei staining. Figure 5E shows microscopic images of fluorescence measured at different excitation and emission, corresponding to the three fluorescent tags and presented as merged images. In Dox-treated mouse hearts, the merged images showed largely yellow color (red + green), confirming the overexpression of Hsp25 in cardiomyocytes, whereas Hsp25 was found to be decreased in heart tissues of wild-type mice treated with HSF-1(+/−) plasma alone and with HSF-1(+/−) plasma + Dox (Fig. 5E), as indicated by the greatly decreased yellow intensity. These results confirm that HSF-1(+/−) plasma treatment indeed altered Hsp expression in cardiomyocytes.

Phosphorylated isoforms of myocardial Hsp25, which have been found to play key roles in inducing apoptosis in Dox-treated hearts, were also significantly reduced in the HSF-1(+/−) plasma + Dox-treated group (Fig. 5A). It is known that oxidative stress activates MAPKs in the myocardium, and various MAPKs phosphorylate HSF-1 as well as Hsp25 (Vedam et al., 2010). The p38 MAPK level was, as previously demonstrated, higher in Dox-treated mouse hearts than in saline-treated heart tissue (Fig. 6A). In contrast, the increases in MAPK-activated protein kinase-2 and extracellular signal-regulated kinase 1/2 levels induced in the heart by Dox treatment were significantly attenuated when the mice were treated with HSF-1(+/−) plasma (Fig. 6A). Apoptotic markers were also analyzed in all four groups. First, the p53 level, which has been found to induce Bax in an Hsp25-dependent manner (Stacchiotti et al., 2009), was high in Dox-treated mouse hearts (Fig. 7A), whereas its level was significantly lower in HSF-1(+/−) plasma-pretreated hearts. Likewise, the apoptotic index, defined as the Bax/Bcl2 ratio, was higher in the Dox alone-treated group, relative to the HSF-1(+/−) plasma + Dox-treated group (Fig. 7, A and B). To confirm that attenuation of the Hsp25/p53 apoptotic pathway is indeed responsible for the observed cardioprotection in the HSF-1(+/−) plasma + Dox-treated group, the protein-protein interaction between these proteins was determined. p53 was immunoprecipitated from the lysates of all four
groups, and Hsp25 was immunoblotted. Indeed, Hsp25 and S82 phospho Hsp25 showed positive detection; however, relative to the Dox alone-treated group, the blot intensities were very low in the HSF-1 (H11001/H11002) plasma-pretreated group (Fig. 7C). This finding suggests that Hsp25-dependent apoptotic signaling is very much reduced because of a lack of HSF-1 activation, which is required for p53 transactivation (Vedam et al., 2010).

**eHsp25 Antagonizes Dox and Competitively Binds TLR in Cardiomyocytes.** The following experiments were performed with isolated adult mouse cardiomyocytes to determine whether the HSF-1 (H11001/H11002) plasma with higher eHsp25 can protect cardiomyocytes from Dox in vitro. Cardiomyocytes in culture were treated with Dox, and survival was monitored over time. Figure 8A shows the micrographs and Dox fluorescence image of Dox-treated cardiomyocytes. The fluorescence image illustrates that Dox selectively accumulates in nucleus, because of its DNA intercalation property. Viable and healthy isolated mouse adult cardiomyocytes do not typically contract in culture. However, upon addition of Dox, cardiomyocytes start beating, and this can be used as an indicator of Dox toxicity in culture (Schmidt et al., 2007). A delay in the start of beating or prolonged sustained contractility in the presence of Dox and an extended beat-to-beat interval (BBI) is a sign of decreased Dox-induced myocyte death. Cardiomyocyte contraction, after treatment with Dox, was monitored over time until cessation of contractility and transition to the spherical shape common for dead myocytes (supplemental video), and BBI was calculated. Figure 8B shows the quantitative analysis of BBI, as a function of time. Dox-alone-treated cardiomyocytes start beating fast and lose rod morphology much faster than myocytes pretreated with HSF-1 (H11001/H11002) plasma before addition of Dox, clearly illustrating that HSF-1 (H11001/H11002) plasma protects cardiomyocytes from Dox-induced death.

TLR signaling in Dox-treated myocytes in vitro was assessed from the nuclear fraction of NF-κB. Nuclear translocated NF-κB (p65) was determined by Western blotting and the results obtained are presented in Fig. 8C. Of interest, nuclear translocation of NF-κB was significantly inhibited in HSF-1 (H11001/H11002) plasma-pretreated cardiomyocytes, suggesting that HSF-1 (H11001/H11002) plasma pretreatment inhibits the TLR pathway, because TLR activation has been shown to activate the TLR/MyD88/NF-κB pathway. The cytokines IL-6 and TNF-α were also consistently lower in plasma-pretreated myocytes (Fig. 8, D and E). To confirm that eHsp25 binds to the TLR2 receptor on the cell surface, Oregon Green-labeled recombinant human Hsp27 (OG-rhHsp27) was added to isolated cardiomyocytes in culture, and the fluorescence was measured as described previously (Kim et al., 2009). As shown in Fig. 8F, the Oregon Green fluorescence detected showed saturation with an increasing concentration of OG-rhHsp27. Moreover, preincubation with a TLR2 antibody, before addition of OG-rhHsp27, prevented the detected accumulation of Oregon Green fluorescence (Fig. 8F). Together these results indicate that the eHsp25 in HSF-1 (H11001/H11002) plasma inhibits Dox interaction with the TLR2 receptor and thereby reduces the cardistotoxicity.
Roles for different extracellular Hsps in cardiovascular disease have been the focus of several studies (Kim et al., 2009; Yokota and Fujii, 2010). However, to our knowledge, there are no published data on how Dox affects extracellular heat shock proteins in the circulation and how these circulating eHsps influence Dox-induced heart failure. Our data have demonstrated the novel finding that eHsp25 in the circulation protects against Dox-induced cardiomyopathy by preventing Dox-induced myocyte death. Moreover, our data have shown that preconditioning animals in vivo with plasma containing higher levels of Hsp25, led to an increase in systemic plasma eHsp25, primed the animals for a more robust Dox-induced release of eHsp25 (Fig. 2B), preserved cardiac function after Dox treatment, and increased survival. Hsp25 depletion of the plasma used for pretreatment reversed the observed beneficial effects, and treatment with recombinant Hsp27 largely replicated the increase in survival. Thus, it is clear that when Dox is given in a setting of increased eHsp25, the animal is protected against cardiotoxicity. Of interest, Dox treatment of wild-type animals, which induces cardiac dysfunction and death, also produced an increase in eHsp25, perhaps indicating the activation of some endogenous protective mechanism that failed to protect in some animals.

Fig. 6. HSF-1(+/-) plasma pretreatment inhibits Dox-induced MAPKs in the heart. A, Western blots of various MAP kinases for the four groups as indicated on the images. B, quantitative analyses of the blot intensities (n = 3)(**+, p < 0.001; +, p < 0.01). ERK1/2, extracellular signal-regulated kinase 1/2; GAPDH, glyceraldehyde-3-phosphate dehydrogenase.

Fig. 7. HSF-1(+/-) plasma pretreatment inhibits Hsp25/p53 interaction in the heart. A, Western blots of various markers as indicated on the images (n = 3). B, quantitative analyses of Bax/Bcl2 ratio of the blot intensities (**+, p < 0.001). C, Western blots of Hsp25, Ser-82, and Ser-15 phospho Hsp25 in p53 immunoprecipitates from all the four groups. GAPDH, glyceraldehyde-3-phosphate dehydrogenase; AB, control p53 antibody only.
The mechanism by which eHsp25 is produced, either endogenously or in response to Dox, is unclear. In a previous report, Dox was shown to activate stress translation signaling through HSF-1 activation and increased expression of Hsp25 in the myocardium (Vedam et al., 2010). In the present work, we observed that treating HSF-1(-/-) mice with Dox not only increased Hsp25 in the heart but also increased the plasma level of eHsp25 with time, reaching a maximum after 4 weeks when severe cardiac dysfunction was noted. The source of eHsp25 could be excess intracellular Hsps in the cardiomyocytes released into the circulation by some unknown secretory mechanism or macrophage/monocyte release of Hsp25 or a combination of both. In a recent study, Dox was reported to induce an acute immunogenic response by enhancing TLR2 activity in monocytes and macrophages (Krysko et al., 2011). Such a response also showed increased IL-6 in Dox-treated mice. However, it was not determined whether there was a parallel increase in secretion of Hsp25 during this process. We found that both wild-type animals and wild-type cardiomyocytes produce IL-6, TNF-α, and Hsp25 in response to Dox. Thus, it is clear that secretion of Hsp25 accompanies the secretion of immunogenic cytokines. However, when the animals and cardiomyocytes are treated with plasma containing eHsp25, both still demonstrated a Dox-induced increase in IL-6 and TNF-α levels (although blunted in relation to the untreated animals), but only the whole animals showed an increase in Hsp25. These results imply that the source of the eHsp25 is not the cardiomyocyte and that treatment with eHsp25 can regulate the production of Hsp25 in the heart.

How eHsp25/27 could interact with cardiomyocytes or provide cardioprotection in the heart has not been fully elucidated previously, although the plasma level of eHsp27 was studied as a potential marker of atherosclerosis risk in humans (Vivanco et al., 2005; Kardys et al., 2008). Potential correlations between activation of various surface receptors and Dox-induced cardiac injury have been identified (Nozaki et al., 2004). Although activation of receptors such as the hydrocarbon receptor and the androgen receptor reduced Dox-induced cardiotoxicity, activation of other receptors such as CB1 (Mukhopadhyay et al., 2007), ERB1 (Herbst, 2005), and ANG-1A (Leslie et al., 2004) was found to aggravate apoptosis and myocardial remodeling. Genetic knockout and pharmacological inhibition of these receptors have shown cardioprotective effects against Dox-induced heart failure. TLR2 knockout in mice was shown to protect from Dox-induced toxicity, suggesting that activation of TLR2 by Dox could trigger downstream apoptotic signaling in myocytes.
HSF-1 knockout, which leads to eHsp25 expression in the heart provided cardioprotection via decreased p53 transactivation (Vedam et al., 2010). However, as demonstrated herein, treatment of wild-type mice with eHsp25-containing plasma both increased the eHsp25 level and prevented the induction of Hsp25 expression in the heart. Moreover, the heterozygous HSF-1 (+/−) mice, which showed greater survivability when exposed to Dox and were not susceptible to Dox-induced cardiac dysfunction (Supplemental Fig. 2), had a constitutive increase in eHsp25, which was not further increased by the Dox-induced immunogenic activation. Thus, rather than opposing our previous results, our observations can be interpreted in terms of the “heat shock paradox” (DeMeester et al., 2001): that the induction of heat shock proteins can be either beneficial or detrimental, depending on the timing of Hsp induction in relation to injurious insult. If Hsps are increased before injury, either constitutively or pharmacologically induced (as seen in our present work) or by recombinant overexpression (Liu et al., 2007; Fan et al., 2008), there is protection. However, if Hsp25 expression is induced after injury (via HSF-1 activation in response to Dox-induced oxidative stress), it is cytotoxic and aggravates apoptosis in the heart (Vedam et al., 2010). In support of these findings, recent studies by the Knowlton group have demonstrated that TNF-α injury was worsened by HSF-1 activation induced by heat stress after injury (Kobba et al., 2011). Taken together, the accumulated data indicate that overexpression of eHsp25 by any combined therapy during the treatment of cancer using Dox might result in the suppression of cardiomyopathy.

In conclusion, the present work has provided novel insights on the protection of the heart from the effects of Dox by eHsp25-containing plasma. eHsp25 is found to prevent Dox-initiated apoptotic signaling, potentially by antagonizing TLR2. Our recombinant human Hsp27 tagged with Oregon Green showed TLR2 receptor binding, which could be prevented by a TLR2 antibody, clearly showing that eHsp25 antagonized Dox. Furthermore, NF-κB activation (which is caused by TLR2/ligand binding) was decreased in the HSF-1 (+/−) plasma-treated hearts (Fig. 8F), suggesting that pre-treatment of mice with eHsp25-enriched plasma prevents activation of both the HSF-1 and the NF-κB pathways (Fig. 8G). Heterozygous HSF-1 knockout mice with higher plasma level eHsp25 were highly resistant to Dox-induced heart failure consistently. Therefore, the present study has revealed that a potential therapy based on Hsp25/27 could be developed for clinical applications in the Dox treatment setting and for other applications wherein an antagonist of TLR2 might be beneficial.

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Authorship Contributions

Participated in research design: Krishnamurthy, Kanagasabai, Druhan, and Ilangoovan.

Conducted experiments: Krishnamurthy and Kanagasabai.

Contributed new reagents or analytic tools: Druhan.

Performed data analysis: Krishnamurthy and Ilangoovan.

Wrote or contributed to the writing of the manuscript: Krishnamurthy, Druhan, and Ilangoovan.

References


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Hsp25 Enriched Plasma Transfusion Preconditions the Heart against Doxorubicin-induced Dilated Cardiomyopathy in Mice

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Material and Methods

Animals and treatment

Eight weeks old wild type (HSF-1+/+, Balb/C, Charles River) and knock out HSF-1(both HSF-1+/−, HSF-1+/−) mice, were used. HSF-1 knock out mice were obtained as a generous gift from Dr. Ivor Benjamin from the University of Utah, bred in our animal facility and genotyped (Vedam et al., 2010). All the protocols were approved by the Institutional Animal Care and Use Committee and were performed according to the National Institutes of Health Guide for the Care and Use of Laboratory Animals. Animals were randomized into six groups. Group1: Wt-Con, vehicle (saline) treated HSF-1+/+ wild type mice; Group2: Wt- Dox, 6 mg/kg Dox treated HSF-1+/+ wild type mice; Group3: Homo-Con, vehicle (saline) treated HSF-1−/− knockout mice; Group4: Homo-Dox, 6 mg/kg Dox treated HSF-1−/− knockout mice; Group5: Hetero-Con, vehicle (saline) treated HSF-1−/− knockout mice; Group6: Hetero-Dox, 6 mg/kg body weight Dox treated HSF-1−/− knockout mice. Groups such as, Wt-Dox, Homo-Dox and Hetero-Dox were chronically treated with Dox (Novaplus, TX, USA) by four intraperitoneal injections of 6 mg/kg bw (cumulative dose of 24 mg/kg bw). Injections were performed once in a week, and the mice were observed until eight weeks. In the second set of experiments mice will be randomized into four groups. HSF-1+/+ wild type mice treated with saline; HSF-1+/+ wild type mice treated with plasma taken from HSF-1+/− knockout mice + Saline; HSF-1+/+ wild type mice treated with plasma taken from HSF-1+/− knockout mice + Dox; HSF-1+/+ wild type mice treated with Dox+saline. Once the heart failure was confirmed in the Dox-treated mice by echocardiogram, the animals were deeply anaesthetized using isoflurane. Blood was drawn by using cardiac puncture and plasma collected with heparin as an anticoagulant, centrifuged at 3000 rpm for 10 min and stored in −80°C until determination of eHSP25 level. The hearts from these animals excised, flash frozen in liquid nitrogen and stored in −80°C. Frozen hearts tissues were grounded using mortar and pestle, and
finally suspended in ice cold 1X PBS. The suspension was centrifuged at 1500 rpm for 10 min and the pellet were lysed using 500 μl of 1X RIPA buffer (Santa Cruz, USA) supplemented with 1X protease inhibitor cocktail (Sigma Aldrich, USA), 1 mM PMSF; 1 mM Na$_3$VO$_4$ and incubated on ice for 45 min. After centrifuging at 12,000 rpm at 4°C for 10 min the supernatant was used as a whole myocardial extract.

Death Curve Study
HSF$^{+/\pm}$ mice injected with saline (n=8), HSF-1$^{+/\pm}$ mice injected with HSF-1$^{+/\pm}$ mice plasma+saline (n=8), HSF-1$^{+/\pm}$ mice injected with HSF-1$^{+/\pm}$ mice plasma + Dox (n=10), HSF-1$^{+/\pm}$ mice injected with Dox+saline (n=10) and HSF-1$^{+/\pm}$ mice injected with HSF-1$^{+/\pm}$ plasma+Dox (n=10) were used for the study. The Dox treated group was injected with 6 mg/kg bw of Dox once a week for four weeks and the control mouse were treated with saline. The Kaplan-Meier curve was plotted to study the survival rate in the animals belonging to different groups.

Weight Loss Study
HSF-1$^{+/\pm}$ mice injected with saline (n=8); HSF-1$^{+/\pm}$ mice injected with HSF-1$^{+/\pm}$ mice plasma+saline (n=8); HSF-1$^{+/\pm}$ mice injected with HSF-1$^{+/\pm}$ mice plasma + Dox (n=10); HSF-1$^{+/\pm}$ mice injected with rhHsp27+Dox (n=8); and HSF-1$^{+/\pm}$ mice injected with Dox+saline (n=10) were followed for their weight lose measured every week and the percentage weight loss was plotted.

Two-Dimensional Echocardiography
Mice were habituated to handling to prevent stress that may have occurred during the experiment. To perform echocardiography on conscious animals (Yang et al., 1999), mice were gently held in supine position in the palm of the hand. The left hemi-thorax was shaved and a 1–2 mm-thick layer of pre-warmed hypoallergenic ultrasonic transmission gel (Parker Laboratories, Fairfield, New Jersey) was applied to the thorax. M-mode echo was performed using a GE Vivid 7 ultrasound imaging system (GE Healthcare, Milwaukee, WI, software version 3.4.2) equipped with a 13-MHz linear-array transducer. Gas anesthesia (oxygen with isoflurane) was administered via a facial mask and maintained at a minimum dose of (1–2%, Abbot; Chicago, IL). After induction of anesthesia, the mice will be placed in a shallow left-lateral decubitus position. Spontaneous breathing was maintained throughout the echocardiographic studies via isoflurane blowing. After collecting data from anesthetized mice, the mask was removed while the probe kept in place until the mice become conscious. 2D short axis echo will be performed at the level
of mid-papillary muscle. Frame rate will be set at 300 frames/second. All measurements were averaged from minimum of 3 cardiac cycles. LV end-diastolic and end-systolic internal diameter (LVIDd and LVIDs, respectively) were measured. LV fractional shortening (%) will be calculated as (LVIDd – LVISd)/LVIDd x 100. LV mass was calculated as 0.80 x 1.04 + 0.6. LVRWd: LV lateral wall thickness in diastole, LVPWd: LV posterior wall thickness in diastole (Vedam et al., 2010).

**Immunoblot analysis**

Whole-cell lysates were prepared by lysing the cells in RIPA lysis buffer. Protein concentration was measured by means of the BCA protein assay (Pierce). Proteins were separated in an 8–12% SDS–PAGE and electroblotted on PVDF membranes (Invitrogen). Immunoblot analysis used specific antibodies and enhanced chemiluminescence (ECL)-based detection (Pierce). Antibodies used were: Hsp25 (Heat shock protein 25; catalogue # SPA-801 Stressgen, MI), Hsp25 Ser15 (catalogue # SPA-525, Stressgen), and Hsp25 Ser82 (catalogue # SPA-524), Hsp60 (catalogue # SPA-807), serum albumin (catalogue # ab19195) p53 (catalogue # sc-126 Santa cruz Biotechnology), Bax (catalogue # 2772 cell signaling), Bcl2 (catalogue # 2876 cell signaling), MAPKAP2 (catalogue # KAP-MA105), MAPK (catalogue # 9218 cell signaling), ERK1/2 (catalogue # KAP-MA001 assay design), P-p38MAPK (catalogue # 9211S), AKT (catalogue # 4691 cell signaling), NFκBp65 (catalogue # 3034P cell signaling). Signal was visualized by the enhanced chemiluminescence (ECL) method according to the manufacture’s protocol (Pierce). Western signal was digitized and quantified with UN-SCAN-IT gel 6.1 software.

**Co-Immunoprecipitation of Hsp25 and p53**

To study the association of p53 and Hsp27, immunoprecipitation of p53 and immunoblotting of Hsp25 (and vice versa), were carried. To the normalized protein samples of whole cell lysate (1 g), 40 μl of Pre-clearing matrix (catalogue # sc-45057) was added, mixed gently in a rocker at 4°C for 1 h. After gentle rocking, 2 μgs of p53 (Santa cruz) or Rabbit monoclonal Hsp25 antibody and 500 μls of PBS were added and incubated at 4°C in a rocker for an hour. The beads were centrifuged and pelleted at 1500 rpm for 5 min and washed twice with 500 μl of PBS. After the final wash, the pre-cleared lysates were added to IP antibody-IP matrix complex beads and left overnight at 4°C once again on a rocker. Beads were pelleted at 1500 rpm and washed thrice with RIPA buffer and after the final wash, 15 μls of 4 X sample loading buffer added, samples boiled at 95°C for 5 min, centrifuged at 1500 rpm. The protein samples were electrophoresed and blotted to confirm the coimmunoprecipitation.
Histopathology

To examine histological changes at the end of four weeks, hearts were harvested and fixed in 10% neutral buffered formalin, routinely processed, and embedded in paraffin. Paraffin sections were cut into 4µm serial sections. Counter-staining of haematoxylin and eosin (H&E) was performed.

Immunofluorescence staining in heart tissue by confocal microscopy

Tissue was deparaffinized, rehydrated, washed, and blocked with normal goat serum. After rinsing with TBST, tissue was incubated with both 1:100 dilution of anti-rabbit α actin antibody (catalogue # ab46805 abcam) and 1:200 dilution of anti-mouse HSP25 antibody (catalogue # H-0273 Sigma) in 3% BSA/TBST at 4°C overnight. Slides were washed in TBST three times; both green Alexa Fluor 488 conjugated goat antirabbit secondary antibody and red Alexa Fluor 555 conjugated donkey-antimouse secondary antibody (Molecular Probes, Inc., Eugene, OR) were incubated for 30 min. Slides were washed with TBST three times; nuclei were costained with 0.3 μM 4’,6-diamidino-2-phenylindole hydrochloride, and examined using an Olympus FV1000 spectral confocal microscope.

Expression and purification of human recombinant Hsp27

Expression was performed in Escherichia coli BL21 (DE3) pUBS520. E. coli was cultured with aeration, on Luria–Bertani (LB) media containing ampicillin (100 μg/mL), to an attenuation (600 nm) of 0.4. Isopropyl thio-β-D-thiogalactoside (IPTG) was added to a final concentration of 0.3 mM and culture was continued for a further 2 h at 30°C. The cells were harvested, frozen and used for isolation of recombinant human wild-type Hsp27 (rhHsp27). The initial stage of purification of Hsp27 was performed as described previously (Somara and Bitar, 2004). Briefly, the crude extract of Hsp27 in lysis buffer (50 mM Tris/HCl, pH 8.0, 100 mM NaCl, 1 mM EDTA, 0.5 mM phenylmethanesulfonyl fluoride, 14 mM β-mercaptoethanol) was fractionated with (NH₄)₂SO₄ (0–30% saturation) and passed through the Ni-affinity column, and eluted with elution buffer containing EDTA.

Denaturation and Renaturation of Hsp27.

Denaturation and renaturation of Hsp27 was performed according to (Somara and Bitar, 2004). Recombinant wild-type Hsp27 in buffer B was freeze dried. The samples of freeze-dried Hsp27 was dissolved in 50 mM phosphate (pH 7.5), containing 100 mM Na₂SO₄, 0.02% β-
mercaptoethanol and 6 M urea, up to a final protein concentration of 6 mg/mL, and then stored on ice for 2 h. After incubation, the samples were diluted six fold in the same buffer, minus urea and β-mercaptoethanol, and dialyzed against two changes of the same buffer overnight.

**Binding study**

This study was performed using the approach of Kim et al. (Kim et al., 2009). Recombinant human HSP27 (rhHSP27, Low-Endotoxin) was labeled with Oregon green 488 (Molecular Probes) according to the manufacturer's protocol. The unconjugated dye was separated by dialyzing the protein solution overnight in PBS at 4°C. The final labeled protein concentration was determined by measuring the absorbance at 280 nm corrected by the absorbance for Oregon green and the extinction coefficient for HSP27. For binding assays, cardiac myocytes were incubated with Oregon-green labeled recombinant human HSP27 (OG-rhHSP27) in increasing concentrations for 30 min at 4°C.

**Fluorescence determination in Dox treated cardiomyocytes**

Adult cardiac myocytes were isolated from HSF-1 wild type and knock out mice as follows. Mice were injected intraperitoneally with heparin (2000 units/mouse), followed by a mixture of 100 mg/kg ketamine and 8 mg/kg xylazine. After 20 min, the heart was removed from the chest, cannulated via the ascending aorta, and mounted on a modified Langendorff perfusion apparatus. The heart was perfused for 4 min with tyrode buffer (mM: NaCl 140; KCl 2.5; CaCl₂ 1.8; MgCl₂ 1.0; glucose 10; HEPES 5; pH 7.4 with NaOH) at 37°C, and myocytes were isolated using standard procedure. The cells were plated in lamin coated plate using tyrode buffer (plus 0.5% BSA and 50 μM CaCl₂) for 1 h to allow attachment. Plating was performed on 22-mm² glass coverslips (Corning, NY) coated with 100 μl/coverslip of 50-nM mouse laminin (Invitrogen, Grand Island, NY) for 30–40 min. The plating density was ~60 cells/mm² with total of 20,000 cells/coverslip. Myocytes were treated with 0.5 μM Dox for 1 hr, and fixed using ethanol and used for fluorescence measurements using Nikon microscope when excited at 546 nm and emitted at 565 nm (Shen et al., 2008).

**Cardiomyocytes contractility study experiment**

Briefly, the heart was mounted on a Langendorff apparatus and perfused with modified MEM (Sigma, St. Louis, MO, 37°C, bubbled with 95% O₂–5% CO₂). Blendzyme type IV (0.077 mg/ml) (Roche Applied Science, Indianapolis, IN) was then added to the perfusate. After 7–20 min, the heart was taken down, the ventricles minced, and myocytes were dissociated by
trituration. Subsequently the myocytes were filtered, centrifuged, and resuspended in MEM containing 200 µM Ca^{2+}. Myocytes were used within 6 h after isolation. Isolated cardiomyocytes were treated with 0.5 µM Dox alone or 10% HSF-1^{+/−} mouse plasma + 0.5 µM Dox and spontaneous contractions of single cardiomyocyte were recorded until the myocyte in focus loses its shape (becomes rounded).

**Isolation of peritoneal macrophages**

Thioglycolate-elicited peritoneal macrophages from control and Dox treated mice were obtained as follows. 3% thioglycolate solution was obtained by dissolving 30g of dehydrated Brewer thioglycolate medium powder (Sigma-Aldrich) in 1000 mL deionized water and autoclaved for 20 minutes at 15 pounds of pressure (121°C). The preparation was kept in the dark under sterile conditions at room temperature for at least 3 months before use. Peritoneal exudate cells were elicited by i.p. injection of 2 ml of 3% sterile thioglycolate. After 4 days, cells consisting mostly of macrophages (over 90%) were harvested by peritoneal lavage using 5 ml RPMI medium supplemented with 5% FBS. Cells were plated in culture plates containing RPMI medium supplemented with 10% fetal bovine serum, 100 IU/ml penicillin and 100 μg/ml streptomycin. After 2 h of incubation at 37°C, non adherent cells were removed by vigorous washing, maintained in RPMI medium for 24hrs, and washed again with PBS before using for further experiments.

**Statistics Analysis**

All values were expressed as mean ± SE. To compare echocardiography data, lactate dehydrogenase and creatine phosphokinase one-way analysis of variance (ANOVA) followed by a post hoc procedure was performed using Statistical Package from the Social Sciences (SPSS) software package version 16. Survival curves after Dox injection were created by the Kaplan–Meier method and compared by a Log-Rank test. Statistical significance was accepted at a value of \( P<0.05 \).
References


Supplementary Figure 1

Supplementary Fig. 1: Western blots of eHsp25 in plasma from Dox-treated HSF-1+/+ (WT, Balb/c) for 4 weeks, blotted with authentic sample (AS) provided by the antibody supplier.
Supplementary Fig. 2: (A) Echocardiographic analysis of cardiac function in Dox-treated HSF-1+/+, HSF-1+- and HSF-1-- mice. (B) Quantitative data for echocardiographic measurements of %FS and LV mass. Abbreviations are as defined in materials and methods. **P<0.01 compared with respective control (n=10).
**Supplementary Fig. 3:** IL-6 and TNF-α, measured in wild type (HSF-1+/+) and heterozygous mice (HSF-1+/−) plasma (n=6) (***p<0.001).
Supplementary Fig. 4: Schematic illustration of the experimental protocol followed. HSF-1+/+ mice first injected with plasma or saline (↑), Dox was injected after 24 h (↑). Then after two days, echo (↑) was used to measure the cardiac function. This was repeated for three more weeks, while the echo measurements continued for 8 weeks.
Supplementary Fig. 5: Whole body weight change as function of post drug treatment time in different groups as indicated on the figure. Significant weight loss was observed in Dox treated mice. However, pre-treating the mice with HSF-1 +/- plasma (with higher eHsp25), did not show significant weight loss upon treated with Dox (*p<0.001).
Supplementary Fig. 6: Survival of rhHsp27 (50µg in 100µl of PBS) + Dox and eHsp25 depleted Plasma of HSF-1+/− mice + Dox survival groups. Injection procedure were as in Fig.2A. 50% survival rates were 23±3 days and 28±4 days respectively for eHsp25 depleted HSF−/− plasma infused and rhHsp27 infused mice cohorts. While the Hsp25 depleted plasma did not show significant increase compared to Dox alone infused mice (22±3 days) (Fig.2A), the rhHsp27 infused mice cohort showed very little enhancement.