Time- and Dose-Dependent Effects of Ionizing Irradiation on the Membrane Expression of Hsp70 on Glioma Cells

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Abstract

The major stress-inducible Hsp70 (HSPA1A) is overexpressed in highly aggressive tumor cells including glioblastoma multiforme and presented on their plasma membrane. Ionizing irradiation increases cytosolic and membrane-bound Hsp70 (mHsp70) on tumor cells. Depending on its intracellular or membrane localization, Hsp70 either promotes tumor growth or serves as a target for the innate immune system. To define the optimal window for a therapy with Hsp70-targeting NK cells the kinetics of the mHsp70 density on the plasma membrane of human glioma cells (U87) has been studied after radiotherapy by multiparameter flow cytometry. Cell cycle dependent alterations in the mHsp70 expression were excluded by seeding different cell numbers on day 0. Low dose irradiation (2 Gy) results in a slow upregulation of the mHsp70 density in U87 cells which peaks on day 4 and starts to decline from day 7 onwards. In contrast, higher radiation doses (4 Gy, 6 Gy) resulted in a faster upregulation of mHsp70 expression on days 2 and 1, respectively, followed by a decline from day 5 onwards. Functionally, elevated mHsp70 densities correlate with an improved lysis by Hsp70-targeting NK cells. In summary, the kinetics of changes in the mHsp70 density upon irradiation on tumor cells is time and dose-dependent.

1. Introduction

The therapeutic application of ionizing radiation plays a pivotal role in the treatment of solid tumors either alone or in combination with surgery and chemotherapy. By causing DNA single- and double-strand breaks, predominantly mediated by reactive oxygen species (ROS), ionizing radiation unfolds its cytotoxic activity against tumor cells [1]. Although the major goal of irradiation is the reduction in viable tumor mass to achieve local tumor control and a decreased dissemination, also immunostimulatory effects have been assigned to ionizing radiation [2]. However, it remains to be elucidated
which dose and radiation regimen is optimal to induce abscopal anti-tumor immune effects. Herein, the role of the major stress-inducible heat shock protein 70 (Hsp70) on the membrane of highly aggressive tumor cells was analyzed as a radiation-inducible target which can be recognized by Hsp70-peptide activated NK cells [3].

Beside its intracellular chaperoning functions including assisting proper protein folding, assembly of nascent polypeptides, preventing protein aggregation and thereby controlling the cellular protein homeostasis, members of the HSP70 family fulfil various cytoprotective tasks upon environmental stress. An up-regulated Hsp70 expression after radiation therapy enhances viability of tumor cells by fostering protein damage repair and impairing apoptotic pathways [4]. By stimulating the overexpression of anti-apoptotic proteins (Bcl-2/Bcl-xL), downregulating pro-apoptotic Bax, Bcl-Xs and Bak [5] or blocking TRAIL-induced apoptosis and formation of the death-inducing signaling complex with death receptors DR4 and DR5, cytosolic Hsp70 promotes tumor cell survival, protects against apoptosis and promotes tumor progression [6].

Apart from ionizing radiation also hyperthermia, hypoxia, heavy metals, glucose deprivation and cytotoxic drugs can induce the Hsp70 expression and stimulate antigen release and danger signal expression [7].

In contrast to intracellular Hsp70, extracellular and membrane-bound Hsp70 plays a key role in stimulating both, adaptive and innate immune responses and thereby initiating protective anti-tumor immunity [8]. By its exclusive expression on the surface of tumor, but not normal cells [9], the membrane-bound form of Hsp70 (mHsp70) provides a tumor-specific target for Hsp70-activated NK cells [10]. Therefore, increasing the membrane density of Hsp70 on tumor cells by irradiation combined with an NK cell-based immunotherapy targeting mHsp70 provides a novel strategy to improve clinical outcome and extend overall survival of patients with advanced tumors.
[11]. Herein, we were interested to identify the optimal radiation dose for up-regulating mHsp70 as a target for NK cells.

2. Results

Membrane Hsp70 expression remains stable during a culture period of 7 days

The mHsp70 expression on tumor cells can be affected by the cell cycle. To maintain cells in the exponential growth phase during a culture period of 7 days different cell counts were seeded on day 0: 0.25 x 10^6 cells were seeded for a culture period of 1-4 days, 0.125 x 10^6 cells for 5 days, 0.06 x 10^6 cells for 6 days and 0.01 x 10^6 cells for 7 days. Representative data of the mHsp70 expression on U87 cells during a period of 7 days after sham irradiation (0 Gy) are summarized in Figure 1. No significant differences in the percentage of positively stained cells (Fig. 1A) and in the mean fluorescence intensity (mfi) (Fig. 1B) were detected over the whole period of time. Similar results were obtained for untreated control cells (data not shown). The slightly lower mHsp70 expression density on day 0 can be attributed to the short incubation period of 5h after trypsin treatment.

Irradiation induces an increase in the mHsp70 density on different tumor cell lines

Although the percentage of mHsp70 positivity remains unaltered on U87 (Fig. 2A), HeLa (Fig. 2B) and HepG2 (Fig. 2C) cells on day 1 after sham (0 Gy), 4 Gy and 6 Gy irradiation, the expression density of Hsp70 differed significantly upon radiation. A radiation dose of 6 Gy nearly doubled the mean fluorescence intensity (mfi) values in all three tumor cell lines. In U87 cells the mfi raised from 38.9±15.8 to 47.1±16.0 and 83.5±14.2 after irradiation with 4 Gy and 6 Gy, respectively (Fig. 2D, p=0.01). In HeLa cells the mfi increased from 9.2±1.9 to 20.6±6.8 (Fig. 2E, p=0.03) on 1 day after a radiation dose of 6 Gy and in HepG2 the mfi values increased from 13.9±2.7 to
22.0±2.6 (p=0.04) after a radiation dose of 4 Gy and to 22.3±4.6 (Fig. 2F, p=0.06) after 6 Gy. Similar results were obtained when untreated control cells were used instead of sham treated cells (data not shown).

**Irradiation-induced upregulation of mHsp70 is transient**

Cell viability of U87 cells was not significantly affected by a radiation with 4 Gy and 6 Gy between days 1 to 7. A comparison of the mHsp70 densities on days 1, 4 and 7 after irradiation with 4 Gy and 6 Gy revealed significantly upregulated mfi values on days 1 and 4 and a drop to initial levels on day 7 after irradiation. On day 1, significant differences were detected in sham (0 Gy) versus 6 Gy irradiated cells and in 4 Gy versus 6 Gy irradiated cells (p≤0.01), whilst no statistically significance was observed in sham versus 4 Gy irradiated cells (Fig. 3A). On day 4 after irradiation, the mfi of sham (0 Gy) treated cells was 41.9±0.8 compared to 79.3±12.8 and 75.1±7.7 after irradiation with 4 Gy and 6 Gy. Significant differences were observed in sham versus 4 Gy and sham versus 6 Gy irradiated cells (p≤0.05), whilst no statistically significance was observed in 4 Gy versus 6 Gy irradiated tumor cells (Fig. 3B). On day 7 after irradiation, the mfi values of sham treated cells remained at 38.73±1.41 and the mfi of 4 Gy and 6 Gy irradiated cells dropped to initial levels (37.60±7.24 and 40.02±5.71, respectively). No significant differences in the mfi values were observed in the differently irradiated cells on day 7 (Fig. 3C).

**Irradiation-induced effects on the mHsp70 density are dose- and time-dependent**

The previous results indicated clear differences in the mHsp70 expression density on U87 cells that depend on the recovery time after irradiation. Figure 4 summarizes the dose- and time-dependent effects on the mHsp70 density. While mfi values of sham (0 Gy) treated cells did not differ significantly over a period of 7 days when different cell counts were seeded (Fig. 1B), an exposure to an irradiation dose of 2 Gy showed a continuous slow increase in mfi values up to a maximum value of 67.4±16.7 on day
that gradually decrease to 39.3±4.3 on day 7 (Fig. 4A). Cells that have been
irradiated with a higher dose of 4 Gy showed a similar kinetic to that after a 2 Gy
irradiation but the mHsp70 density reached a maximum in the mfi of 79.3±12.8 already
on day 4 instead of day 5. The drop in the mfi values thereafter occurred more rapid
than after an irradiation with 2 Gy (Fig. 4B). Following an irradiation with 6 Gy, mHsp70
expression density was significantly increased already on d1 compared to the sham (0
Gy) irradiated cells (23.3±3.7 versus 83.6±14.2; \(p \leq 0.05\)). After an irradiation with 6 Gy
the significantly elevated mHsp70 levels persisted over 3 days and dropped
significantly on day 5 (Fig. 4C). A comparison of all time points and all radiation doses
shown in Figure 4D reflects the different kinetics of the mHsp70 densities in U87 cells.

Increased mHsp70 expression densities are associated with an increased
cytolytic activity mediated by TKD/IL-2 activated NK cells

The density of the mHsp70 expression plays a crucial role for the recognition by
TKD/IL-2 activated NK cells. Therefore, the cytolytic activity of Hsp70 peptide activated
NK cells was tested against sham (0 Gy) and irradiated (4 Gy) tumor cells. The
increase in the mHsp70 density on day 4 after irradiation is shown in Figure 5A. The
lysis of irradiated (4 Gy) tumor cells by TKD/IL-2 activated NK cells is significantly
higher than that of sham (0 Gy) irradiated cells at E:T ratios ranging between 50:1 and
3:1 (Fig. 5B; \(p \leq 0.05\)).

3. Discussion

Compared to normal cells, highly aggressive tumor cells frequently exhibit higher
cytosolic Hsp70 levels which are further upregulated upon environmental stress [12].
Intracellularly, Hsp70 ensures proper protein folding and transport [10] and interferes
with both, intrinsic and extrinsic apoptosis pathway to avoid cell death and thereby
foster tumor cell survival [12-13]. Apart from its cytosolic localization Hsp70 is actively
released in lipid microvesicles such as exosomes [14-15] and is expressed on the plasma membrane of tumor but not normal cells [16]. While cytosolic Hsp70 is mainly responsible for maintaining protein homeostasis [4], mHsp70 mediates dual functions, on the one hand it stabilizes lysosomal and plasma membranes and thereby protects tumor cells from apoptosis [17], on the other hand it provides a tumor-specific target for NK cells that have been pre-activated with an immunogenic Hsp70 peptide TKD plus low dose IL-2 (TKD/IL-2) [8,18]. The specific membrane localization of Hsp70 on tumor but not normal cells can be explained by an interaction of Hsp70 with tumor-specific lipid compounds such as globoyltriaosylceramide Gb3 [16,19-20]. 

In vitro lipid copellation assays revealed that recombinant Hsp70 interacts with the tumor-specific lipid raft component globoyltriaosylceramid Gb3 or with the non-raft lipid compound phosphatidylserine (PS) which translocates to the outer plasma membrane leaflet upon stress [21]. Previously, we could demonstrate that environmental stress not only enhances the synthesis of Hsp70 in the cytosol, but also increases the expression density of mHsp70 on tumor cells [3,20,22]. Since mHsp70 serves as a target for TKD/IL-2 stimulated NK cells [10] in vitro [23] and in vivo [24], time- and dose-dependent effects of irradiation were studied with respect to the mHsp70 density on human tumor cells. A better understanding of the mHsp70 expression kinetics following irradiation might help to identify an optimal therapeutic window for combination therapies consisting of ionizing irradiation and Hsp70 targeting NK cell-based immune therapies. Herein, we could demonstrate that the mHsp70 density correlates the cytolytic activity of Hsp70-targeting NK cells.

Glioblastoma multiforme (GBM) the most common malignant neoplasia of the brain in adults is associated with high mortality rates. Despite multimodal treatment strategies consisting of surgery, radiotherapy and systemic chemotherapy using Temozolomide [25] the median survival remains poor (15-18 months) and recurrence rates are high.
Its intratumoral heterogeneity, concerning genetic alterations and morphology as well as the infiltrative growth and diffuse dissemination into the brain parenchyma often hinder a successful treatment of GBM. Therefore, there is a high medical need for innovative treatment strategies for glioblastoma. The mHsp70 density as a potential NK cell target was measured in the human glioblastoma cell line U87 at different time points after irradiation with doses in the range of 2 Gy to 6 Gy. To rule out cell density dependent effects on the mHsp70 expression and to maintain cells in the exponential growth phase, tumor cells were seeded at different cell densities which were adapted to the different culture periods. In sham as well as non-treated tumor cells the proportion of mHsp70 positive cells remained stable at nearly 100% with a relatively low mfi value over the whole culture period of 7 days. These findings are in line with previous results obtained with other tumor types such as epithelial tumor cells of the head and neck area [26].

Environmental stress such as ionizing irradiation at sublethal doses initiates the synthesis of Hsp70 in tumor and normal cells [10], but only tumor cells show an up-regulated mHsp70 expression [17]. Herein, we have shown for the first time that the kinetics of the mHsp70 expression is dependent on the irradiation dose. While the maximum mHsp70 expression is reached on day 5 after an irradiation with 2 Gy, the peak mHsp70 expression is detected already on day 1 after an irradiation with 6 Gy. Furthermore, after low irradiation doses the decrease to initial mHsp70 levels occurs earlier than after a higher irradiation dose. This means that a low dose irradiation might be associated with a later onset and a shorter peak of the mHsp70 expression compared to a high dose irradiation. These findings are in accordance to results of Diller et al. who showed that the kinetics of the heat shock protein synthesis is transient and proportional to the applied stress [27].
The intracellular Hsp70 levels remain unaltered in sham treated tumor cells, whilst during irradiation with a sublethal dose of 2 Gy and directly after irradiation with lethal doses [28] results in a considerable increase in the Hsp70 synthesis [2,10,16,29]. In line with our assumption that synthesis and expression of Hsp70 is dependent on dose- and time-dependent factors it can be assumed that cytoprotective repair mechanisms that depend on the anti-apoptotic molecular chaperone Hsp70 are also related to the applied stress. Our findings indicate that a high, but yet non-lethal irradiation dose of 6 Gy induces an early onset of the mHsp70 expression after 1 day with a sustained overexpression that persists for 3 days, whereas lower irradiation doses (below 4 Gy) induce a slower up-regulation kinetics with an earlier drop. In an effort to define the optimal timing for the application of ex vivo stimulated mHsp70 targeting NK cells after radiotherapy, it is important to determine the maximum density of mHsp70 on tumor cells. In the case of U87 glioblastoma cells an irradiation dose between 4 Gy and 6 Gy might be optimal to achieve a relative long-lasting upregulation of the mHsp70 expression, in vitro. However, future preclinical studies using fractionated irradiation protocols are necessary to define the optimal irradiation dose which is suitable for an in vivo application of mHsp70-targeting immune cell-based therapies.

There is strong evidence that a lethal dose of 10 Gy which drastically harms tumor cell survival cannot be compensated by cytoprotective repair mechanisms that include the cytosolic overexpression of Hsp70 [30]. After lethal irradiation Hsp70 will be externalized by dying tumor cells as a danger associated molecular pattern (DAMP) with the capacity to initiate anti-tumor immune responses [1,5,31] Hsp70-chaperoned tumor peptides can induce CD8+ T cell mediated immune responses after cross presentation of immunogenic tumor antigens on MHC class I molecules [32], whereas peptide-free Hsp70 in the context of pro-inflammatory cytokines such as IL-2 can further augment the cytolytic and migratory capacity of NK cells that recognize and kill
remaining therapy-resistant mHsp70 positive tumor cells [14,23]. In summary, the applied radiation dose and time kinetics play a critical role in optimizing radiation-induced effects by increasing the density of mHsp70 expression on the surface of surviving residual tumor cells as a target for immune cells. Membrane-bound Hsp70 plays a crucial role in stimulating the innate and adaptive immune system as it is selectively expressed on the cell surface of tumor, but not normal cells [33] and therefore serves as a tumor-specific recognition structure [19]. We could show that Hsp70-targeting NK cells are capable to specifically recognize and kill tumor cells presenting Hsp70 on their plasma membrane via granzyme B mediated apoptosis [9].

4. Material and Methods

Cell line and culture conditions

The epithelial human primary glioblastoma cell line U87 MG (ATCC HTB-14) and the human hepatocellular carcinoma cell line HepG2 (ATCC HB-8065) were grown in complete growth medium, consisting of Dulbecco’s Eagle’s Minimum Essential Medium (DMEM) (Sigma-Aldrich, Steinheim Germany) supplemented with 10% heat inactivated fetal calf serum (FCS) (Sigma-Aldrich), 1% antibiotics (10,000 IU/ml penicillin, 10 mg/ml streptomycin, Sigma-Aldrich), L-glutamine (Sigma-Aldrich), MEM non-essential amino acid solution 100x (Sigma-Aldrich) and sodium pyruvate (Sigma-Aldrich). The epithelial human cervix carcinoma cell line HeLa (ATCC CCL-2) was grown in complete growth medium, consisting of RPMI-1640 (Sigma-Aldrich) Germany supplemented with 10% heat inactivated FCS (Sigma-Aldrich), 1% antibiotics (10,000 IU/ml penicillin, 10 mg/ml streptomycin, Sigma-Aldrich), L-glutamine (Sigma-Aldrich) and sodium pyruvate (Sigma-Aldrich). After reaching confluency, adherent growing tumor cells were trypsinized for 2 minutes at 37°C in trypsin ethylene diamine-tetra-acetic acid (EDTA) (Sigma-Aldrich). Single cell suspensions with different cell counts
were seeded in 15 ml supplemented medium in T-75 ventilated culture flasks. Tumor cells were routinely checked for mycoplasma contamination.

Irradiation

Tumor cells were irradiated with a single dose of 0 Gy (sham), 2 Gy, 4 Gy and 6 Gy using the Gulmay RS225A irradiation machine (Gulmay Medical Ltd., Camberley, UK) at a dose rate of 0.90 Gy/min (15 mA, 200 kEV) or were kept untreated.

Flow cytometry of mHsp70

The mHsp70 expression was determined by flow cytometry on a FACSCalibur flow cytometer (BD Biosciences). Single cell suspensions of sham irradiated and irradiated U87 cells (0.4 x 10⁶ cells per vial) were collected at different time-points after radiation. After a washing step in phosphate-buffered saline (PBS)/10% fetal calf serum (FCS) cells were incubated either with fluorescein-isothiocyanate (FITC)-conjugated mouse monoclonal antibody specific for mHsp70 (cmHsp70.1, IgG1, multimmune GmbH, Munich, Germany) or with an isotype matched FITC-labeled control antibody on ice in the dark for 30 min. Only viable cells (propidium iodide negative cells) were gated and the proportion of positively stained cells and mean fluorescence intensity (mfi) values were analyzed on a FACSCalibur flow cytometer (Becton Dickinson, Heidelberg, Germany.) The mfi is a relative value of the total fluorescence intensity of cmHsp70.1-FITC antibody stained, viable cells subtracted by the intensity of the signal intensity obtained after staining of the cells with an isotype-matched IgG1-FITC control antibody. Fluorescence data were analyzed and plotted by using CellQuest software (Becton Dickinson, Heidelberg, Germany).

Europium assay

The cytolytic activity of human NK cells cultured in RPMI-1640 medium supplemented with 10% heat inactivated fetal calf serum (FCS), 1% antibiotics (10,000 IU/ml penicillin,
10 mg/ml streptomycin), L-glutamine, sodium pyruvate and Hsp70 peptide TKD (2 µg/ml) and IL-2 (100 IU/ml) at a cell density of 5x10^6 peripheral blood lymphocytes (PBL) for 4 days was determined against sham (0 Gy) and 4 Gy irradiated tumor cells at different effector to target (E:T) ratios ranging from 50:1 to 3:1 by using a standard 3.5-hour Europium assay. Mean values of an experiment in triplicates are shown. The specific lysis was calculated according to the equation: (experimental release – spontaneous release) / (maximum release – spontaneous release).

**Statistical analysis**

Each sample was measured in at least 3 independent experiments. The Gaussian distribution of the data was tested by using the Normality test. Comparative analysis was performed by using the Student t-test, evaluating the relevance between variables. Data were presented as mean values with standard deviation. Differences in p values ≤0.05 were considered as statistically significant.

5. **Conclusions**

In contrast to intracellular Hsp70, extracellular and mHsp70 play key roles in stimulating both, adaptive and innate immune responses and thereby might provide protective anti-tumor immunity [8]. By its exclusive expression on the surface of tumor cells, but not normal cells [9] mHsp70 serves as a tumor-specific target for activated NK cells [10]. In this study we present strong evidence that the radiation dose plays a pivotal role on the kinetics of the mHsp70 density of human glioblastoma cells. A low dose irradiation is associated with a later onset and a shorter peak of the mHsp70 expression compared to a higher irradiation dose. Therefore, hypofractionated irradiation schemes with higher doses might be beneficial to generate an extended therapeutic window for mHsp70-targeting immunotherapies.
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Figure Legends

**Figure 1.** Percentage (A) and mean fluorescence intensity (mfi) (B) of mHsp70 positive U87 glioblastoma cells after sham irradiation (0 Gy) on days 0, 1, 2, 4, 5, 6 and 7. Bars represent the mean value and the corresponding standard error of the mean (SEM) of n=3 independent experiments.

**Figure 2.** Percentage of mHsp70 positive tumor cells U87 (A), HeLa (B) and HepG2 (C) cells on day 1 after irradiation with 0 Gy (sham), 4 Gy and 6 Gy. Mean fluorescence intensity (mfi) of mHsp70 on U87 (D), HeLa (E) and HepG2 (F) cells on day 1 after irradiation with 0 Gy (sham), 4 Gy and 6 Gy. Bars represent the mean value and the corresponding standard error of the mean (SEM) of n=3 (U87), n=4 (HeLa), and n=3 (HepG2) independent experiments. Significance: *p≤0.05, **p≤0.01.

**Figure 3.** Mean fluorescence intensity (mfi) of mHsp70 in U87 cells after sham (0 Gy), 4 Gy and 6 Gy irradiation on days 1 (A), 4 (B) and 7 (C). Bars represent the mean values and the corresponding standard error of the mean (SEM) of n=3 independent experiments. Significance: *p≤0.05, **p≤0.01.

**Figure 4.** Dose- and time-dependent radiation effects on the mHsp70 mean fluorescence intensity (mfi) on human glioblastoma cells U87 over 7 days (d0-d7) with different radiation doses 2 Gy (A), 4 Gy (B), 6 Gy (C). Bars represent the mean values and the corresponding standard error of the mean (SEM) of n=3 independent experiments. Significance: *p≤0.05. (D) Summary of the kinetics of the mHsp70 density after sham (0 Gy), 2 Gy, 4 Gy, 6 Gy) over a period of 7 days (d0-d7). For clarity statistical significance was only shown in Figures 4 A, B, C but not in Figure 4D.

**Figure 5.** Comparative analysis of the cytolytic activity of TKD/IL-2 activated NK cells targeting mHsp70 positive tumor cells. Sham (0 Gy) and 4 Gy irradiated tumor cells on
day 4 were used as target cells. (A) mHsp70 mean fluorescence intensity (mfi) on sham (0 Gy) and 4 Gy irradiated tumor target cells. (B) Lytic activity of NK cells against sham (0 Gy) and 4 Gy irradiated tumor target cells. The effector to target (E:T) ratios are ranging from 50:1 to 3:1. The results represent mean values and the corresponding standard error of the mean (SEM) of n=2 independent experiments in triplicates. Significance: *p≤0.05
References


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