Homogenous and Microbeam X-Ray Radiation Induces Proteomic Changes in the Brains of Irradiated Rats and in the Brains of Nonirradiated Cage Mate Rats

Richard Smith1, Jiaxi Wang2, Colin Seymour1, Cristian Fernandez-Palomo1, Jennifer Fazzari1, Elisabeth Schütke3, Elke Bräuer-Krisch4, Jean Laisse5, Christian Schroll6, and Carmel Mothersill1

Abstract
To evaluate microbeam radiation therapy (MRT), for brain tumor treatment, the bystander effect in nonirradiated companion animals was investigated. Adult rats were irradiated with 35 or 350 Gy at the European Synchrotron Research Facility using homogenous irradiation (HR) or MRT to the right brain hemisphere. The irradiated rats were housed with nonirradiated rats. After 48 hours, all rats were euthanized and the frontal lobe proteome was analyzed using 2-dimensional electrophoresis and mass spectrometry. Proteome changes were determined by analysis of variance (\(P < .05\)). Homogenous irradiation increased serum albumin, heat shock protein 71 (HSP-71), triosephosphate isomerase (TPI), fructose bisphosphate aldolase (FBA), and prohibitin and decreased dihydrolipoyl dehydrogenase (DLD) and pyruvate kinase. Microbeam radiation therapy increased HSP-71, FBA, and prohibitin, and decreased aconitase, dihydropyrimidinase, TPI, tubulin DLD, and pyruvate kinase. Cage mates with HR irradiated rats showed increased HSP-71 and FBA and decreased pyruvate kinase, DLD, and aconitase. Cage mates with MRT irradiated rats showed increased HSP-71, prohibitin, and FBA and decreased aconitase and DLD. Homogenous irradiation proteome changes indicated tumorigenesis, while MRT proteome changes indicated an oxidative stress response. The bystander effect of proteome changes appeared antitumorigenic and inducing radioresistance. This investigation also supports the need for research into prohibitin interaction with HSP-70/71 chaperones and cancer therapy.

Keywords
antitumorigenesis, bystander effect, homogenous irradiation, microbeam irradiation, proteomics

Introduction
Microbeam radiation therapy (MRT) is an experimental radiotherapy concept characterized by the delivery of multiple, parallel and very narrow (~25-75 \(\mu m\)) microplanar X-ray beams.1 The technique was initially investigated in the 1950s and there is now international effort to develop its clinical implementation.2 Specifically, these doses are delivered in parallel beams, which focus the high dose along narrow microplanar tracks, thus sparing large volumes of interjacent normal tissue from radiation exposure.3,4 Microbeam radiation therapy therefore allows the application of peak X-ray doses, which are greater than the doses usually used for cancer therapy, by 1 or 2 orders of magnitude (eg, 150-4000 Gy).1 In theory, this offers a non-invasive means for treating brain tumors by delivering targeted radiation doses to precise locations within the brain (ie, the
Materials and Methods

**Rat Husbandry and Homogenous (HR) and Microbeam (MRT) Irradiation**

Rat (adult male, Wistar; 260-280 g), husbandry, radiation procedures, anesthesia and experimental ethical guidelines, and licensing have all been previously described. Briefly, the rats were supplied by Charles River Laboratories (L’Abrele, France) and held in the animal facility at the European Synchrotron Research Facility (ESRF), Grenoble, France, under European Union directive 2010/63/EU. All experimental procedures were carried out on anesthetized rats (as described elsewhere under guidelines issued by the French government and specifically covered by licenses 380325 and B3818510002 and approved by the International Evaluation Committee for Animal Welfare and Rights. A total of 45 rats were used in this investigation; 5 completely untreated rats taken directly from the ESRF animal facility, 4 rats exposed to a sham irradiation + 4 cage mates, 4 rats exposed to 35 Gy HR + 4 cage mates, 4 rats exposed to 350 Gy HR + 4 cage mates, 4 rats exposed to 35 Gy MRT + 4 cage mates, and 4 rats exposed to 350 Gy MRT + 4 cage mates. The procedures for positioning the anesthetized rats on the goniometer and for homogenous field irradiation (HR), using a broad beam, and microbeam radiation (MRT), using a 100 × 14 mm high monochromatic anteroposterior beam, in the ESRF insertion device (ID) hutch beamline, have also been previously described. Microbeam radiation therapy irradiation was of the right hemisphere using an array composed of 50 quasi-parallel rectangular planar microbeams 25-μm thick with a 200 μm center-to-center distance. Homogenous irradiation entrance and MRT peak entrance radiation doses of 35 or 350 Gy were delivered and confirmed using Gafchromic films.

In addition to completely untreated control rats, which never left the ESRF animal facility, a sham irradiation control was included in the study, the purpose being to determine whether proteomic responses occurred as a result of anesthesia and the handling and confinement associated with the HR and MRT irradiation. These sham irradiated rats were anesthetized and positioned in the ID hutch 17 in the same way and for the same period of time as the irradiated rats, but no radiation dose was administered.

**Cage Mate Pairing, Tissue Collection, and Proteomic Analysis**

Following irradiation (or sham irradiation), the rats were transported back to the ESRF animal facility and placed in individual cages with a single marked, nonirradiated rat. After 48 hours, the rats were deeply anesthetized, beheaded, and dissected. Brain tissue samples (5 mm × 5 mm × 3 mm) were taken from the center of the irradiation array in the right hemisphere of the irradiated rats and the matching area from sham irradiated rats, cage mate rats, and completely untreated control
rats. These samples were placed in aluminum foil and immediately frozen in liquid nitrogen and then stored at −80°C.

The brain samples from these rats were then analyzed using 2-dimensional (2-D) gel electrophoresis (2-DE). Thirty microgram samples of brain tissue were homogenized in 300-μL ice-cold 2-D lysis buffer; the buffer composition is detailed elsewhere. The homogenate was clarified, by centrifugation (18 000g for 5 minutes at 4°C) and desalted, using commercially available Pierce desalt columns (Fisher Scientific, Markham, Ontario, Canada). Total protein concentration was then determined using a commercially available protein assay kit (Bio-Rad Laboratories Inc, Mississauga, Ontario, Canada) and 100 μg of total protein was subjected to 2-DE. All 2-DE was carried out using the Protean 2-DE system, first-dimension immobilized pH gradient (IPG) strips (pH 3-10), second-dimension gels (Bio-Rad Laboratories, Inc) and rehydration, and IPG equilibration and second-dimension running buffers (all purchased from Bio-Rad Laboratories, Inc). The full 2-DE protocol, including IPG rehydration, first dimension isoelectric focusing, second dimension Laemmli/12% acrylamide electrophoresis, gel fixing and staining with SYPRO-ruby stain (Bio-Rad Laboratories, Inc), and gel image capture, using a Fluor-S Max gel imager (Bio-Rad Laboratories, Inc), has been described elsewhere.

Each tissue sample was run individually on a single gel. Any gels which were not of an acceptable quality for image analysis were completely rerun using a new 30-mg tissue sample from the rat brain. The gel images were then analyzed using Phoretix 2-D (V 2004) analytical software (Phoretix International, Newcastle, United Kingdom). It is important to stress this analysis was carried out as a blind study. The gels were given a random code number so that, during analysis, it was not known whether the gel was from an irradiated, sham irradiated, cage mate, or control rat. The gels were also calibrated for molecular size (Mr) and isoelectric point (pI) using protein markers (Bio-Rad Laboratories, Inc) and the gel calibration facility of the Phoretix 2-D software.

Two-Dimensional Gel Statistical Analysis

Once gel image analysis had been completed, the gel images were organized by treatment. Protein spots were then quantified by normalized volume, a parameter which combines spot size and intensity to give an index of expression. To fully illustrate any changes in protein expression, individual normalized spot volumes were divided by the mean normalized spot volume of the corresponding protein from untreated control rats to give a “fold-change” index of expression. These fold changes were then compared using analysis of variance followed by least square difference analysis; a P < .05 was considered statistically significant. Consistent protein expression and the results of this statistical analysis were used to select protein spots for excision and identification. In other words, the protein spots selected for protein identification were those which were present on all the gels in the experiment and which showed a statistical difference in normalized volume when compared to the completely untreated control rats. By these criteria, 11 protein spots were selected (Figure 1). Note that no protein spots were found which were consistently absent (ie, downregulated to the point of nondetection) in any experimental treatment.

In-Gel Tryptic Digestion and Matrix-Assisted Laser Desorption Ionization QStar XL Mass Spectrometry Analysis

The full details of In-gel tryptic digestion and matrix-assisted laser desorption ionization (MALDI) mass spectrometry (MS) analysis have been described. In summary, selected protein spots were excised, destained, reduced, treated with iodoacetamide, dehydrated, dried, and digested. The digested proteins were desalted and concentrated in formic acid and analyzed using a high-resolution QSTAR XL MALDI quadrupole time of flight (TOF) mass spectrometer (AB Sciex) coupled to a MALDI 2 ion source with a solid state Nd:YAG laser. Data acquisition was done by Analyst 1.1 Software (Applied Biosystems/MDS Sciex, Foster City, California, USA). Mass spectra in TOF MS and mass spectrometry/mass spectrometry (MS/MS) mode were in a mass range of 50 to 2500 m/z with a resolution of 8000 FWHM and a 60 V focusing potential was used throughout. The MS/MS data analysis was performed using MASCOT and peptide sequences were compared with the MS protein sequence database (MSDB). Tolerances for peptide information were 0.6 Da for MS and MS/MS, and all
information was manually inspected based on y and/or b series ions (N-terminal carboxyl and bond-cleavage fragments from the C-terminal sides of the peptide, respectively).

## Results

### Two-Dimensional Electrophoresis and Protein Identification

Figure 1 illustrates a representative 2-D gel from the right hemisphere of a completely untreated control rat and indicates the protein spots selected for identification. Table 1 lists the protein identities as well as the theoretical (as listed by the protein database) and experimentally derived (determined by 2-D gel calibration) molecular sizes (Mr; kDa) and isoelectric point (pI; pH units). A total of 11 protein spots showed consistent resolution on the gels across the entire experiment and statistically significant changes in expression (as defined by normalized spot volume).

The protein spots which were identified in this investigation as aconitase hydratase, dihydrolipoyl dehydrogenase (DLD), glial fibrillary acidic protein (GFAP), heat shock cognate 71 kDa (HSP-71), prohibitin, serum albumin (S-Alb), triose-phosphate isomerase (TPI), and tubulin alpha-1A were the same identities as those derived from the corresponding protein spots in our previous investigation.14 The spots identified as DLD, fructose bisphosphate aldolase (FBA), pyruvate kinase, and tubulin alpha-1A were unique to this investigation. In our previous study, we were not alerted to any change in the expression of these proteins. However, the slightly different pI values suggest some modification. The experimentally derived Mr of the protein spots identified as S-Alb and HSP-71 were less than the theoretical Mr (Table 1), which suggests it is more likely these were breakdown products rather than intact molecules. It may therefore be more accurate to say that any changes in these proteins were more likely to be changes in protein turnover than changes in protein expression. The experimentally derived Mr of GFAP was greater than the theoretical Mr (Table 1). However, GFAP has been resolved at a similar Mr, 45 to 55 kDa,21 and an even greater Mr, 68 kDa.22 Again this suggests GFAP was most likely an intact molecule.

Table 2 quantifies the changes to specific proteins in the right brain hemisphere of irradiated rats, as a result of HR or MRT, and in the right brain hemisphere of nonirradiated cage mate rats housed with the irradiated animals.

### Proteomic Responses to Broad Beam Homogenous (HR) and Microbeam (MRT) Irradiation

After 48 hours of HR, S-Alb, HSP-71, FBA, prohibitin, and TPI were found to increase in the right hemisphere of the rat brain. Serum albumin, HSP-71, and FBA were increased by 35 and 350 Gy, whereas prohibitin and TPI were unaffected by 35 Gy HR but were increased by 350 Gy. Two proteins were found to be reduced by HR after the same postirradiation interval: pyruvate kinase and DLD. Pyruvate kinase was decreased by 35 and 350 Gy HR, whereas DLD was unaffected by 35 Gy HR but decreased by 350 Gy HR.

Serum albumin was unaffected 48 hours after 35 or 350 Gy MRT but, as was the case with HR, HSP-71, FBA, and prohibitin were increased 48 hours after 35 and 350 Gy MRT. Microbeam radiation therapy also had a similar effect to HR on pyruvate kinase and DLD; 48 hours after irradiation with 35 or 350 Gy, pyruvate kinase was reduced, and 48 hours after irradiation with 350 Gy, MRT DLD was reduced.

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**Table 1. Results of Identification Analysis of the Proteins Indicated in Figure 1.**

<table>
<thead>
<tr>
<th>Gel Spot</th>
<th>Protein Description</th>
<th>Accession Code</th>
<th>MSDB Database</th>
<th>Theor Mr (kDa)</th>
<th>Theor pI (pH units)</th>
<th>Exp Mr (kDa)</th>
<th>Exp pI (pH units)</th>
<th>MOWSE Score</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Aconitase hydratase</td>
<td>ACON_MOUSE</td>
<td></td>
<td>85.5</td>
<td>7.09</td>
<td>77.9</td>
<td>8.61</td>
<td>77 (28)</td>
</tr>
<tr>
<td>2</td>
<td>Dihydrolipoyl dehydrogenase</td>
<td>DLDH_MOUSE</td>
<td></td>
<td>54.3</td>
<td>7.08</td>
<td>52.4</td>
<td>7.96</td>
<td>56 (38)</td>
</tr>
<tr>
<td>3</td>
<td>Dihydropyrimidinase-related protein</td>
<td>DPYLS_RAT</td>
<td></td>
<td>61.5</td>
<td>6.27</td>
<td>59.1</td>
<td>8.22</td>
<td>58 (26)</td>
</tr>
<tr>
<td>4</td>
<td>Fructose-bisphosphate aldolase</td>
<td>ALDOC_HUMAN</td>
<td></td>
<td>39.5</td>
<td>6.13</td>
<td>41.1</td>
<td>8.23</td>
<td>59 (44)</td>
</tr>
<tr>
<td>5</td>
<td>Glial fibrillary acidic protein</td>
<td>GFAP_CARAU</td>
<td></td>
<td>42.6</td>
<td>4.85</td>
<td>53.4</td>
<td>8.66</td>
<td>58 (46)</td>
</tr>
<tr>
<td>6</td>
<td>Heat shock cognate 71 kDa protein</td>
<td>HSP7C_BOVIN</td>
<td></td>
<td>71.2</td>
<td>5.25</td>
<td>38.4</td>
<td>8.88</td>
<td>59 (43)</td>
</tr>
<tr>
<td>7</td>
<td>Prohibitin</td>
<td>PHB_BOVIN</td>
<td></td>
<td>29.8</td>
<td>5.43</td>
<td>30.0</td>
<td>6.59</td>
<td>106 (39)</td>
</tr>
<tr>
<td>8</td>
<td>Pyruvate kinase</td>
<td>KPYM_RAT</td>
<td></td>
<td>57.8</td>
<td>6.30</td>
<td>55.2</td>
<td>8.34</td>
<td>52 (43)</td>
</tr>
<tr>
<td>9</td>
<td>Serum albumin</td>
<td>ALBU_BOVIN</td>
<td></td>
<td>69.3</td>
<td>5.59</td>
<td>28.8</td>
<td>6.31</td>
<td>59 (39)</td>
</tr>
<tr>
<td>10</td>
<td>Triosephosphate isomerase</td>
<td>TPIS_BOVIN</td>
<td></td>
<td>26.7</td>
<td>6.20</td>
<td>26.7</td>
<td>8.60</td>
<td>70 (40)</td>
</tr>
<tr>
<td>11</td>
<td>Tubulin alpha-1A</td>
<td>TBA1A_RAT</td>
<td></td>
<td>50.1</td>
<td>4.84</td>
<td>50.1</td>
<td>6.70</td>
<td>52 (22)</td>
</tr>
</tbody>
</table>

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**Footnotes:**

1. The protein spots indicated in Figure 1.
2. Theor, theoretically determined, that is, as listed on the MSDB database.
3. Exp, experimentally determined, that is, as derived by 2-DE gel calibration.
4. Figures in parenthesis indicate MOWSE score required for significant homology.
Table 2. Fold Change (HR or MRT or Cage Mates to HR or MRT Normalized Spot Volume/Mean Handling Control Normalized Spot Volume) in the Expression of 11 Proteins Indicated in Figure 1.a

<table>
<thead>
<tr>
<th>Protein</th>
<th>Direct Irradiation; Homogenous Radiation</th>
<th>Direct Irradiation; Microbeam Radiation Therapy</th>
<th>Cage Mates; Homogenous Radiation</th>
<th>Cage Mates; Microbeam Radiation Therapy</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>35 Gy</td>
<td>350 Gy</td>
<td>35 Gy</td>
<td>350 Gy</td>
</tr>
<tr>
<td>Heat shock protein 71</td>
<td>2.1 b (1.4/2.9)</td>
<td>1.3 b (1.2/1.7)</td>
<td>2.5 b (1.9/3.4)</td>
<td>3.3 b (1.7/4.3)</td>
</tr>
<tr>
<td>Aconitase</td>
<td>0.8 (0.7/1.0)</td>
<td>1.0 (0.8/1.1)</td>
<td>0.6 b (0.5/0.7)</td>
<td>0.5 b (0.3/0.7)</td>
</tr>
<tr>
<td>Dihydrolipoyl dehydrogenase</td>
<td>1.0 (0.8/1.1)</td>
<td>0.2 b (0.2/0.3)</td>
<td>0.9 (0.5/1.1)</td>
<td>0.3 b (0.2/0.5)</td>
</tr>
<tr>
<td>Dihydropyrimidinase</td>
<td>0.9 (0.6/1.2)</td>
<td>1.0 (0.6/1.1)</td>
<td>0.9 (0.6/1.1)</td>
<td>0.4 b (0.2/0.6)</td>
</tr>
<tr>
<td>Fructose bisphosphate aldolase</td>
<td>1.4 b (1.3/1.4)</td>
<td>1.3 b (1.2/1.6)</td>
<td>2.3 b (2.2/2.5)</td>
<td>1.3 b (1.2/1.8)</td>
</tr>
<tr>
<td>Glial fibrillary acidic protein</td>
<td>1.0 (0.6/1.4)</td>
<td>1.0 (0.8/1.1)</td>
<td>1.1 (0.9/1.3)</td>
<td>1.0 (0.7/1.3)</td>
</tr>
<tr>
<td>Heat shock protein 71</td>
<td>2.1 b (1.4/2.9)</td>
<td>1.3 b (1.2/1.7)</td>
<td>2.5 b (1.9/3.4)</td>
<td>3.3 b (1.7/4.3)</td>
</tr>
<tr>
<td>Prohibitin</td>
<td>1.2 (0.9/1.5)</td>
<td>2.1 b (1.4/3.3)</td>
<td>1.7 b (1.5/1.9)</td>
<td>1.6 b (1.2/2.2)</td>
</tr>
<tr>
<td>Pyruvate kinase</td>
<td>0.5 b (0.3/0.7)</td>
<td>0.3 b (0.3/0.3)</td>
<td>0.4 b (0.2/0.6)</td>
<td>0.2 b (0.2/0.2)</td>
</tr>
<tr>
<td>Serum albumin</td>
<td>2.5 b (2.0/3.1)</td>
<td>1.0 (0.6/1.4)</td>
<td>2.0 b (1.6/2.5)</td>
<td>1.3 (0.8/1.8)</td>
</tr>
<tr>
<td>Tubulin</td>
<td>0.9 (0.9/1.0)</td>
<td>0.8 (0.7/1.1)</td>
<td>0.2 b (0.1/0.4)</td>
<td>0.3 b (0.3/0.3)</td>
</tr>
</tbody>
</table>

Abbreviations: HR, homogenous irradiation; MRT, microbeam radiation therapy.
Data shown as mean and, in parentheses, the min/max fold change (n = 5 handling control rats and n = 4 irradiated or cage mate rats).
A significant change (P < .05) in protein expression compared with the control rats.

which was unaffected by HR, was also reduced by 35 and 350 Gy MRT. However, in direct contrast to 35 Gy HR, there was a reduction in TPI 48 hours after 35 Gy MRT. Aconitase and dihydropyrimidinase, which were also unaffected by HR, were both reduced 48 hours after exposure to 350 Gy MRT.

**Homogenous Radiation: Cage Mate Responses**

Heat shock protein 71 and FBA were increased and pyruvate kinase was decreased in the brain of cage mate rats paired with rats exposed to either 35 Gy or 350 Gy HR. In other words, the cage mate rat changes in these proteins were identical to those seen in the irradiated rats. Dihydrolipoyl dehydrogenase in cage mate rats showed a decrease when paired with rats exposed to 35 Gy HR. Therefore, while not identical (DLD was reduced by both 35 and 350 Gy HR), the HR cage mate response was also similar to directly irradiated HR rats. TPI, serum albumin, and prohibitin, which showed increases in HR rats, were not affected in cage mate rats. Despite being unaffected by HR, aconitase was decreased in cage mate houses with rats exposed to 35 Gy HR.

**Microbeam Radiation Therapy: Cage Mate Responses**

Heat shock protein 71 was increased in cage mate rats paired with rats exposed to 35 and 350 Gy MRT. Thus, HSP-71 was increased in all irradiated and all cage mate rats. Prohibitin was also increased in cage mate rats paired with rats exposed to 35 and 350 Gy MRT (ie, the same response as seen with MRT exposure). This cage mate response was unique to MRT; cage mates to HR rats showed no change in prohibitin. Fructose bisphosphate aldolase was increased in cage mates to 35 and 350 Gy MRT rats. This was the same response as seen in cage mate rats to 35 Gy and 350 Gy HR-exposed rats. However, only 35 Gy MRT resulted in increased FBA, so the increase seen in cage mate rats to 350 Gy MRT-exposed rats occurred despite there being no change in FBA in the irradiated animals.

Aconitase and DLD were downregulated in the brain of cage mate rats, after pairing with rats exposed to 350 Gy MRT. This was similar to the downregulation of aconitase in cage mates paired with rats exposed to 35 and 350 Gy HR and identical to the downregulation of DLD in cage mate pairs with rats exposed to 350 Gy HR.

**Discussion**

**Methodology**

It should be recognized that tumor control involves the use of 2 MRT microbeam arrays which intersect orthogonally at the site of the tumor. However, the investigation described here was intended as an extension of our previous study, which involved MRT-induced bystander effects within the irradiated animal. Here, it was essential to ensure 1 hemisphere only was irradiated so that any proteomic changes in the completely nonirradiated hemisphere were due to a bystander effect. Therefore, a single MRT beam was used. Since the objective of the present study was to compare bystander effect proteomic changes which occur within the irradiated rat with bystander effect proteomic changes induced in completely nonirradiated rats, we deliberately chose to use an identical radiation protocol, as well as an identical protein extraction method and 2-DE format. However, as stated before, we concede that alternative protein extraction methods or narrower pH ranges may
reveal additional proteomic changes and the proteins reported here are not assumed to be an exhaustive list.

Sham irradiation was not found to affect any of the protein spots resolved on the gels of the sham irradiated or the cage mate rats which were paired with the sham irradiated rats. There were also no protein spots which were unique to, or absent from, the gels from sham irradiated rats or their cage mates. This suggests that anesthesia and the restraints necessary for irradiation did not exert a significant effect on the brain proteome.

**Homogenous Radiation: Proteomic Evidence for Radiation Induced Cancer Progression**

The increase in S-Alb and TPI and the decrease in DLD in irradiated rats 48 hours after HR exposure were identical to the responses, which occurred between 4 and 12 hours after irradiation with the same 350 Gy HR doses. Increased S-Alb albumin and TPI and decreased DLD have all been associated with tumor development or cancer progression. The present investigation therefore supports the suggestion that focal high-dose HR (particularly 350 Gy) may induce tumorigenesis in the rat brain.

The increase in prohibitin 48 hours after HR and MRT irradiation also showed considerable consistency to the increases in the same proteins at up to 12 hours after irradiation, with HR or MRT, to 48 hours after irradiation. Therefore, the present investigation provides additional evidence that prohibitin is increased in the mammalian brain following irradiation. However, the association between prohibitin and cancer is less well defined. As a result, the role of prohibitin in terms of causing cancer cell proliferation or suppression is regarded as controversial.

The increase in FBA 48 hours after HR and MRT was not seen 4 to 12 hours after X-ray exposure, which suggests that a 48-hour interval following irradiation was required to induce this change. In addition to X-ray exposure, FBA has been shown to be increased by gamma and ultraviolet radiation. However, like prohibitin, increased FBA is not always associated with cancer. An increase has been seen in colorectal cancer and in a head and neck cancer cell line (FaDuDD), under hypoxia, but a decrease in FBA has been recorded in squamous cell carcinoma and in HT-29 cells expressing the cancer stem cell marker CD133.

**Microbeam Radiation: Proteomic Evidence for Induced Oxidative Stress Response**

Previously, in the brain of MRT irradiated rats, the increase in S-Alb and TPI and the decrease in DLD, following HR, were also accompanied by the HR- and MRT-induced increase in aconitase, another well-documented indicator of tumorigenesis. However, in the present investigation, there was no change in aconitase 48 hours after HR irradiation. Furthermore 48 hours after MRT, there was a reduction in aconitase expression. Collectively, these results suggest that the HR-induced effect on aconitase is limited to somewhere between 12 and 48 hours and that 48 hours after irradiation MRT had the opposite effect on aconitase to that seen between 4 and 12 hours after irradiation.

There is evidence that reducing aconitase activity is a factor in the metabolic rearrangement seen in cancer cells and, as a result, a reduction in expression may be an indicator or poor prognosis in gastric cancers. However, there is little doubt that radiation in general and radiotherapy in particular result in oxidative stress. Aconitase itself is not only highly susceptible to oxidation, but aconitase activity has also been shown to decrease, within 3 days after gamma radiation, specifically because of mitochondrial damage due to the action of ROS. Dihydropyrimidinase and tubulin, proteins which were not found to show any change in expression between 4 and 12 hours after irradiation, but which did show a decline in the present study, 48 hours after irradiation, are also known to be susceptible to oxidation and loss. Moreover, 35 Gy MRT also resulted in a decline in TPI, that is, again the opposite effect to that caused by the same dose HR. Triosephosphate isomerase reduction is known to stimulate antioxidant metabolism and acts to prevent excessive ROS accumulation. A decline in pyruvate kinase, as was consistently the result 48 hours after HR and MRT, irrespective of radiation dose, also results in the resistance to ROS, by inhibiting TPI via accumulation of phosphoenolpyruvate, that is, pyruvate kinase inhibition, via the resulting substrate accumulation, initiates a feedback mechanism which acts on TPI. We therefore propose these results could be indicative of a radiation-induced oxidative stress response. Specifically, following MRT exposure, the decline in oxidation sensitive aconitase, dihydropyrimidinase, and tubulin combined with the counteracting antioxidant response of TPI and pyruvate kinase could be indicative of the “oxidative balance” which occurs between oxidative damage and antioxidant mechanisms.

**Bystander Effect-Induced Proteomics: Antitumorigenesis and Radioresistance**

The increase in HSP-71, in the brain of cage mate rats, was the only proteomic change in the present investigation which was similar to those induced by a bystander effect in the nonirradiated hemisphere of an MRT irradiated rat brain. Perhaps most significantly the brains of cage mate rats did not include the proteomic changes previously associated with oxidative stress-mediated potentially antitumorigenic apoptosis; specifically, TPI, prohibitin, and tubulin did not show a decrease and GFAP did not show an increase. Indeed, the increase in prohibitin, in the nonirradiated cage mate rat brain, was the opposite response to the decrease seen in the nonirradiated hemisphere of rats following cranial MRT irradiation. It is also significant that the cage mate brain reductions in DLD, aconitase, and pyruvate kinase and the increase in FBA, while being identical to the reductions seen in irradiated rat brains, were not part of the bystander effect proteome changes in the nonirradiated hemisphere of rats following cranial MRT.
irradiation.\textsuperscript{14} Therefore, we conclude the present investigation provides proteomic evidence for a fundamental difference between the proteomic response in the brains of nonirradiated cage mate rats housed with irradiated rats and the proteomic response in the nonirradiated hemisphere of a rat brain irradiated by MRT.\textsuperscript{14}

Despite the uncertainty between prohibitin expression and cancer progression (see above, Homogenous Radiation; Proteomic Evidence for Radiation Induced Cancer Progression), increased prohibitin expression can be the basis for tumor suppression.\textsuperscript{43} Among its roles, prohibitin is a regulator of mitochondrial assembly.\textsuperscript{46} One recent review has therefore highlighted the importance of prohibitin in preventing tumorigenesis, by stabilizing mitochondria and improving the efficacy of cancer treatment.\textsuperscript{47} and 2 others have highlighted prohibitin mediated processes as targets for chemotherapy.\textsuperscript{48,49}

Specific examples of chemotherapeutic agents and increased irradiation.\textsuperscript{14} Therefore, we conclude the present investigation suggests a bystander effect in a nonirradiated hemisphere of an MRT irradiated rat brain,\textsuperscript{14} as to whether the advantages of any potentially antitumorigenic proteomic changes could be negated by a detrimental effect on neurological function, also applies to the intermammalian bystander effect.

Heat shock protein 71 is the constitutively expressed member of the HSP-70 family.\textsuperscript{68} Despite its inconsistency or contradictory association with cancer,\textsuperscript{69} HSP-71 is (i) well documented as a chaperone, (ii) critical to ensuring the correct cellular response to neuronal stress tolerance, and (iii) protects against all of the neurological disorders listed above.\textsuperscript{14} Heat shock protein 70 is also a positive predictor of survival following non-small cell lung cancer chemotherapy.\textsuperscript{70} We therefore propose a similar integrative role for HSP-71 in the responses of the brain of a cage mate rat housed with an irradiated rat as we have for the nonirradiated hemisphere of an MRT irradiated rat brain.\textsuperscript{14} We also propose that determining whether FBA and prohibitin act in a tumor suppressive and radioresistant capacity, in companion animals to those exposed to radiation, now warrants specific experimental attention.

**Conclusion**

This investigation has demonstrated proteomic evidence for a RIBE in nonirradiated rats housed with rats exposed to HR or MRT. We propose these responses are potentially protective, specifically antitumorigenic, and possibly confer radioresistance. The increase in prohibitin was similar to the result of a bystander effect in the nonirradiated hemisphere of an MRT irradiated rat brain. This investigation therefore further supports the existing need for research into the role of prohibitin.
in cancer therapy particularly with the interaction of HSP-70/71 chaperones.

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ORCID ID
Richard Smith http://orcid.org/0000-0002-8950-2386
Cristian Fernandez-Palomo http://orcid.org/0000-0002-5095-262X

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