

# Combinatorial Use of Chitosan-Nanoparticles, Reversine and Ionising Radiation on Breast Cancer Cells Associated with Mitosis Deregulation

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**Abstract:** Breast cancer is the most commonly occurring cancer in women worldwide and the second most common cancer overall. The development of new therapies to treat this devastating malignancy is needed urgently. Nanoparticles are one class of nanomaterial with multiple applications in medicine, ranging from their use as drug delivery systems and the promotion of changes in cell morphology to the control of gene transcription. Nanoparticles made of the natural polymer chitosan are easy to produce, have a very low immunogenic profile and diffuse easily into cells. One hallmark feature of breast cancer is genome instability caused by defects in the Spindle Assembly Checkpoint (SAC), the molecular signalling mechanism that ensures the timely and high-fidelity transmission of the genetic material to an offspring. In recent years the use of nanoparticles to treat cancer cells has gained momentum. This is in part due to the fact that nanoparticles made of different materials can sensitise cancer cells to chemo- and radiotherapy. These advances prompted us to study the potential sensitising effect of chitosan-based nanoparticles on breast cancer cells treated with reversine, a small size inhibitor of Mps1 and Aurora B that induces premature exit from mitosis, aneuploidy and cell death, before and after exposure of the cancer cells to X-rays irradiation. Our measurements of metabolic activity as indicator of cell viability, DNA damage by alkaline comet assay, and immunofluorescence using anti-P-H3 as mitotic biomarker, indicate that chitosan-nanoparticles by themselves elicit cellular responses that affect mitosis and cell viability and can sensitise breast cancer cells to X-ray radiation (2Gy). We also show that such sensitisation effect is not caused by direct damage to the DNA by the nanoparticles. Taken together, our data indicates chitosan-nanoparticles have potential application for the treatment of breast cancer as adjunct to radiotherapy.

**Keywords:** Nanoparticles; SAC inhibition; breast cancer; mitosis deregulation; tumour sensitisation

## 1. Introduction

The high prevalence of breast cancer, the most common type of cancer in women and the second most commonly occurring cancer in the world, represents a global health challenge. Over 2 million new cases of breast cancer worldwide were reported in 2018 alone [1]. Undoubtedly, the development of new therapeutic strategies to treat breast tumours is of utmost importance. Genome instability and chromosome segregation defects are two defining features of solid tumours, the most aggressive types of cancer, and often associated to defects in the Spindle Assembly Checkpoint (SAC), the molecular signalling mechanism of higher organisms that monitors the proper segregation of the genetic material every time a cell divides [2].

Nanoparticles of different composition including mesoporous silica and metals such as gold, titanium and molybdenum have been shown to sensitise cancer cells to radiation [3-7], an effect that may be due to the accumulation of nanoparticles in tumour neo-vessels thus increasing radiation absorption in the tumour tissue without affecting the healthy tissue that surrounds the cancer cells [3].

However, the synthesis of these nanoparticles is laborious and expensive, which limits their broader use in the clinic. In contrast, the synthesis of nanoparticles made of the natural polymer chitosan is easy and cheap, with the aggregated value that chitosan-nanoparticles are non-immunogenic. Chitosan nanoparticles can exert multiple influences on cells, such as inducing changes in cell morphology, impacting cellular junctions, impeding motility, and inducing cell gaps [8,9].

Moreover, studies on rats have shown that nanoparticles based on chitosan isolated from squid can have an anti-cancer protective effect in liver cells and that this can be due to the nanoparticles' antioxidant and anti-lipidemic properties [10-12]. Chitosan nanoparticles have gained much attention due to their good biocompatibility and biodegradability and have been approved by FDA for wound dressing applications [13,14]. The overall shape, size distribution, surface morphology, and surface charge of chitosan nanoparticles dictate the dynamics of their entry and diffusion into the cell [15,16]. Importantly, such physicochemical properties can be easily manipulated to enhance the nanoparticles' permeation and retention in tumour sites as well as to use them as drug carriers.

The fact that inhibition of Mps1 and Aurora B kinase activity results in premature mitosis exit with misattached/unaligned chromosomes, which in turn results in severe chromosome missegregation, aneuploidy and cell death [2] and that nanoparticles made of mesoporous silica and metals can sensitise cancer cells to chemo and radiotherapy, prompted us to investigate the potential sensitisation effect of chitosan-nanoparticles to X-ray irradiation of MCF-7 breast cancer cells treated with the Mps1 and Aurora B inhibitor reversine. Our data show that chitosan-nanoparticles by themselves elicit cellular responses that affect mitosis and cell viability and sensitised MCF7 cells to X-ray irradiation and that this effect is not due to genotoxicity of the chitosan-nanoparticles. These results suggest that this type of nanomaterial can be used to improve the treatment of breast cancer and possibly other types of aggressive tumours as adjunct to radiotherapy.

## 2. Materials and Methods

### 2.1. Materials

Chitosan of low molecular weight and sodium tripolyphosphate were purchased from Sigma-Aldrich, Dorset, England. Bis-Tris 12% precast gels were from Thermo Fisher. Anti-N-terminal Mps1 antibody (ab135819) was from Abcam. Anti- $\beta$ -actin antibody was from Wuhan Fine Biotech. Anti-rabbit IgG AP-linked and Anti-Phosphohistone-Ser 10-H3 were from Cell Signalling Technology (CST)/NEB (#7054).

### 2.2. Nanoparticle Synthesis

CS-NP were manufactured by ionic gelation method using sodium tripolyphosphate [17-19]. For this, 0.3 % (w/v) of low molecular weight chitosan was prepared in acetic acid 1.0 % v/v. An equal volume of sodium tripolyphosphate 0.1% (w/v) was added afterwards. The solution was mixed for 1 h using a magnetic stirrer. The nanoparticles thus formed were filtered through a 0.45 µm membrane (Sarstedt, Nümbrecht, Germany) and stored at 4 °C until use. Fluorescent nanoparticles were prepared as described before [20, 21] with the following modifications: as soon as the nanoparticles were formed, a 0.0125 mg/mL rhodamine 123 (Sigma-Aldrich, Dorset, England) solution was added to the nanoparticle suspension and leave under magnetic stirring for 6 h protected from the light. After this, the nanoparticles were centrifugated at 54000 RCF (Optima XPN-80 ultracentrifuge, Beckman Coulter, Fichtenhain, Germany) to separate dye-labelled and unlabelled nanoparticles. This method relies on the generally accepted assumption that the fluorochrome is absorbed to the nanoparticles surface [20].

### *2.3. Nanoparticle Characterisation*

Nanoparticles tracking analyses was carried out by dynamic light scattering (DLS) measurements using the Z-view instrument (Particle Metrix, Ammersse, Germany) and by transmission electron microscopy (TEM) using a tungsten filament HV at 100 kV (Hitachi, Tokyo, Japan). For the latter study, a fresh nanoparticle suspension was placed on a copper grid and left undisturbed for 5 min. The excess was removed using filter paper and the nanoparticle suspension dried and placed into the microscope for image analysis.

### *2.4. Nanoparticles Intracellular Localisation*

To investigate the cellular distribution of the chitosan-nanoparticles (CS-NP from now onwards), they were labelled with rhodamine 123 (Sigma-Aldrich, Dorset, England) and the MCF-7 cells left to grow up on a glass cover slide. Once the cells were attached, CS-NP loaded with rhodamine 123 were added to the cultured cells and incubated for 3 h. As a control, the equivalent amount of rhodamine 123 alone was added to a separate MCF-7 cells culture. After this, both cell cultures were washed twice with phosphate buffered saline (PBS) 1x (potassium phosphate monobasic 1.05 mM; sodium phosphate dibasic 2.96 mM; sodium chloride 155.17 mM, pH 7.4) for 5 min, fixed with 0.4% paraformaldehyde for 30 min and washed twice with PBS. The nuclei of fixed cells were stained with 4',6-Diamidino-2-Phenylindole, Dihydrochloride 0.5 µM (DAPI) (Fisher Scientific, Loughborough, UK) and mounted on a slide with 10% polyvinyl alcohol (PVA), 2.5% 1, 4-Diazabicyclo-octane (DABCO), 5% glycerol, and 25 mM Tris buffer (PVA-DABCO) mounting medium (Sigma-Aldrich, Dorset, England). The preparations were kept refrigerated until analysis by laser scanning confocal microscopy (LSCM 880 Zeiss, Berlin, Germany) at 63x magnification.

### *2.5. Cell Culture*

One validated cryo-vial sample of the estrogen-dependent human breast cancer cell line (MCF-7) was kindly donated by Dr. Frances Willenbrock, Department of Oncology, University of Oxford, Oxford, UK; cells were kept in Dulbecco's Modified Eagles Medium Nutrient Mixture F-12 HAM (Sigma-Aldrich, Dorset, England), supplemented with 13% Heat Inactivated Fetal Bovine Serum (Sigma-Aldrich, Dorset, England), 1% Penicillin-Streptomycin (Corning, Flintshire, UK) and 1% L-Glutamine (Gibco, Loughborough, UK) at 37 °C in humidified atmosphere containing 5% CO<sub>2</sub>. Mycoplasma contamination tests were performed every four weeks to ascertain the cell were mycoplasma-free.

### *2.6. Cell Irradiation*

1.5x10<sup>6</sup> MCF-7 cells were seeded on T75 flasks (Sarstedt, Nümbrecht, Germany) and 3x10<sup>4</sup> cells/well seeded on 96 well plates (Grenier Bio-one, Stonehouse, UK). Cells between 70-80 % of confluence were treated with CS-NP during 3 h, then the media was removed and replaced with new media containing reversine 5 µM and incubated during 1.5 h. Previous reports by others have shown

that reversine at this relatively high concentration is effective to inhibit the SAC kinases Mps1 and Aurora B in HeLa cells after 24 h exposure [22,23], Following sequential treatment with chitosan-nanoparticles and reversine, the breast cancer cells were X-ray irradiated (2 Gy dose) at the Oxford Institute for Radiation Oncology, University of Oxford, using 250 kV X-rays (0.25 mm copper filtration, resulting in a half-value layer of 1.0 mm Cu) at a dose rate of 0.55 Gy/min. The radiation time for all samples was under 4 min. Dosimetry was performed using Gafchromic EBT3 film (Ashland Advanced Materials, Bridgewater NJ) in the position of the cells and analysed as previously described [24].

### *2.7. Cell Viability Assay*

Cell viability after irradiation procedures was tested by the metabolic principle of the active enzymes from the viable cells to reduce the dye resazurin (Fisher Scientific, Loughborough, UK) to resorufin, thereby generating a quantitative measure of viability and cytotoxicity. The dye was added 30 min after irradiation of cells and incubated at 37 °C for 3 h. The fluorescence was then measured using a microplate reader (Spectra Max I3, Molecular Devices, Wokingham, UK) with an excitation-emission wavelength of 560/590 nm, respectively.

### *2.8. Comet Assay*

The alkaline comet assay [25] was carried out by adopting the protocol originally reported by Singh and collaborators [26]. After 30 min of X-rays irradiation the cells were harvested and counted in a Neubauer Chamber using Trypan Blue (Sigma-Aldrich, Dorset, England). Viable cells were kept on ice and used to make a  $2 \times 10^4$  cell suspension. The cells were mixed with 100  $\mu$ L of 1% (w/v) low melting point agarose and spread on a frosted microscope slide precoated with 1% (w/v) normal melting point agarose. A coverslip was used on the smear and the slides left to solidify for 10 min on ice. The coverslip was removed, and the slides immersed on fresh cold lysis buffer (2.5M NaCl, 100 mM EDTA, 10 mM Tris, 1% dimethyl sulfoxide (DMSO), and 1% Triton X-100 pH10) overnight at 4 °C. Next day, comet slides were placed on a horizontal tank containing electrophoresis buffer (300 mM NaOH, 1 mM EDTA pH 13) and left undisturbed for 20 mins to allow DNA unwinding. After this, electrophoresis was carried out at 19 V, for 30 min. The slides were then neutralised with cold neutralisation buffer (0.4 M Tris-HCl pH 7.5), rinsed with distilled water and stained with Diamond Nuclei Acid Stain (Promega, Wisconsin, USA). The slides were dried at room temperature protected from light and scored using the Comet Assay IV software (Perceptive Instruments, Staffordshire, UK) coupled to an epifluorescent microscope (Axio Scope A 1, Zeiss, Berlin, Germany) with 40x objective magnification.

### *2.9. Mitotic Analysis and Western Blot*

Cells after 30 min of X-ray irradiation were washed two times with cold PBS and fixed with paraformaldehyde 4 %-Triton X-100 0.1% during 30 min at room temperature. The cells were then washed three times with cold PBS and incubated overnight at 4 °C after addition of 50  $\mu$ L/well of a 1:500 dilution of anti-phospho-histone H3 (Cell Signalling Technology, Hertfordshire, UK). After this, the antibody was removed, and the plates washed with PBS. 4  $\mu$ M of Hoechst 33342 was then added and the plates scored using an imaging cytometer (Celigo, Nexcelom Bioscience, Manchester, UK). After SDS-PAGE in Bis-Tris 12% precast gels using MOPS-SDS as the running buffer, western blot analysis was carried out using the primary anti-Mps1 antibody at 1:500 dilution and the secondary anti-rabbit IgG AP-linked antibody at 1:1000 dilution following a standard protocol. The protein  $\beta$ -actin was used as the loading control and detected using an anti- $\beta$ -actin antibody.

### *2.10. Statistical Analyses*

All the experiments were conducted by triplicate and the data analysed using the Graph Pad Prism 8 software. Cell viability and mitotic index data were assessed by analyses of variance (one-way ANOVA) and multiple means comparisons ( $p < 0.0001$ ) according to the Tukey's (also known as

the Honestly Significance Difference, HSD) test  $p < (0.05)$ . This type of statistical analysis enables the calculation of minimal differences between means from pairwise comparisons of data groups, where means differing by more than the HSD value indicate a significant difference. For comet assays the Kruskal-Wallis test was used followed by Dunns ( $p < 0.05$ ) pairs of means comparison.

### 3. Results

#### 3.1. Physicochemical Characterisation of Chitosan Nanoparticles

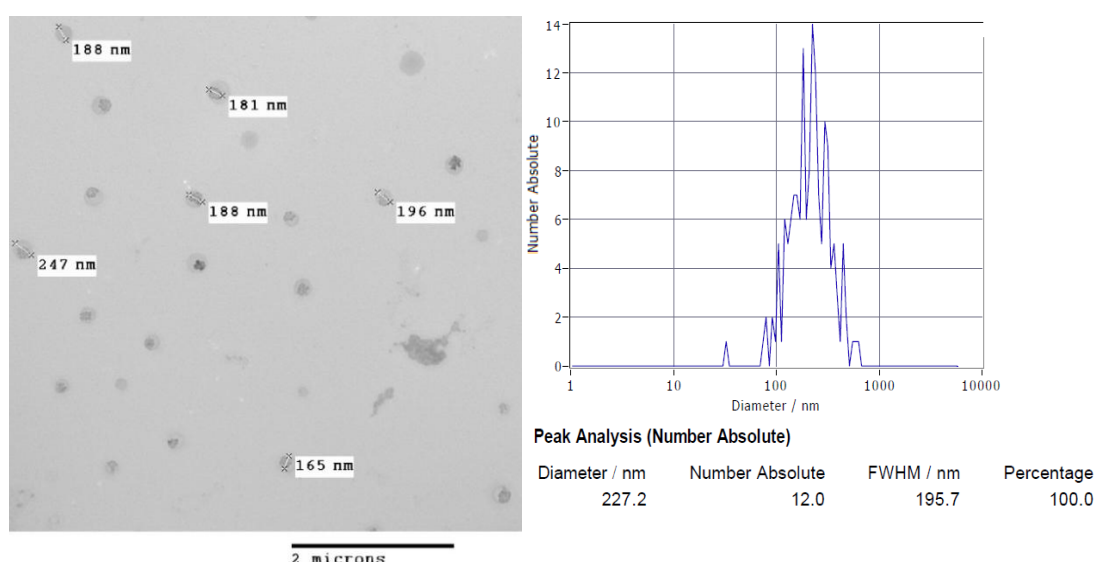
Initially, key physicochemical properties of CS-NP including the nanoparticles concentration, size distribution, and zeta potential were determined. As **Table 1** shows, the CS-NP exhibited a positively charged surface, as reported by the Z potential value, with an average size (hydrodynamic diameter) of around 200 nm.

**Table 1.** Physicochemical properties of the CS-NP. The data corresponds to the average of five independent batches.

Nanoparticles	Physicochemical properties		
	Z-Potential (z-average) mV $\pm$ SD	Size (Hydrodynamic ratio) nm $\pm$ SD	Concentration particles/mL $\pm$ SD
CS-NP	29.6 $\pm$ 9.5	224 $\pm$ 31	5.9E+10 $\pm$ 2E+10
CS-NP[R123]	38 $\pm$ 0.16	227 $\pm$ 97	4.3E+7 $\pm$ 2E+10

#### 3.2. Nanoparticles Morphology

In an attempt to gain direct insight into the morphology of the chitosan-based nanomaterial, the chitosan-nanoparticles were analysed by Transmission Electron Microscopy (TEM). TEM images of CS-NP revealed that the ionotropic gelation method [17-19] used for the synthesis of CS-NP was adequate to produce spherical nanoparticles. This class of nanoparticles had spherical morphology and an average size distribution around 200 nm as estimated by nanoparticle tracking analysis from dynamic light scattering (DLS) measurements (see **Figure 1** for details).

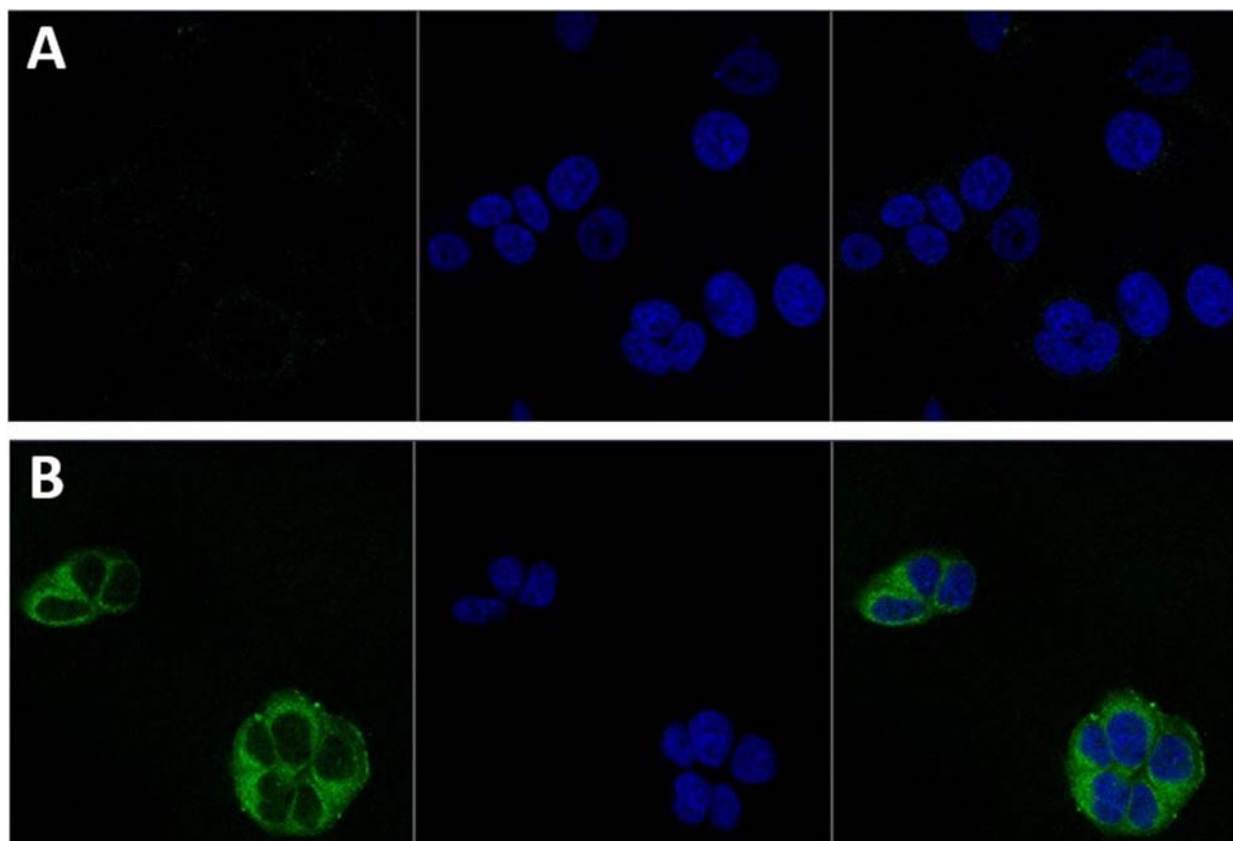


**Figure 1.** Left, TEM image of CS-NP and Right, size distribution plot based on nanoparticle tracking analysis.

#### 3.3. Nanoparticles Cellular Localisation

Because the subcellular localisation of nanoparticles can vary depending on their chemical composition and surface morphology, the subcellular localisation of the CS-NP in MCF-7 cells was

investigated by confocal microscopy. To this aim, chitosan-nanoparticles were labelled with the fluorescent marker rhodamine 123. The physicochemical parameters of CS-NP did not changed with loading the dye (see Table 1 for details). As shown in **Figure 2**, the CS-NP were easily incorporated into the breast cancer cells and readily distributed in the cytoplasm after three hours of exposure. A wide distribution in the cytoplasm of chitosan nanoparticles loaded with rhodamine has also been observed in olfactory ensheathing cells [20]. After three hours of exposure CS-NP were not detected in the nuclei. **Figure 2** also shows that CS-NP did not influence the overall morphology of MCF-7 cells.

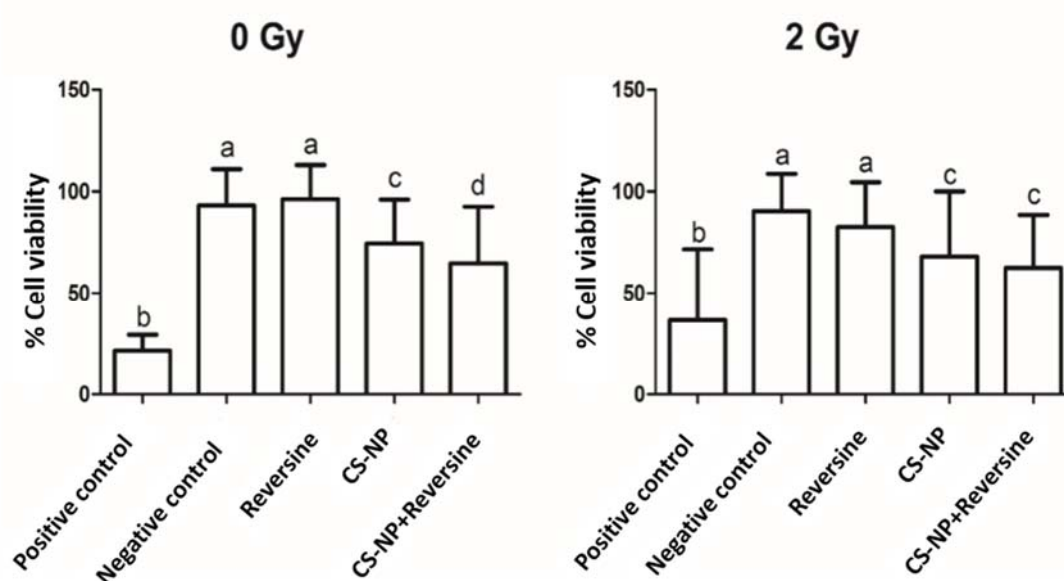


**Figure 2.** Intracellular localisation of CS-NP in MCF-7 cells as determined by light scattering confocal microscopy. A) Cells with rhodamine 123 alone. B) cells after 3 hours exposure to CS-NP. In both rows, the left panel shows the cytoplasmic distribution of the CS-NP (green); the middle panel the DAPI-stained nuclei and the right panel the merge image. All the images were taken at 63x magnification.

#### 3.4. Cell Viability Assay

After determination of the intracellular localisation of CS-NP, we investigated the potential toxicity of CS-NP on the breast cancer cells. To this aim, the viability of MCF-7 cells treated with CS-NP was determined after X-rays irradiation (2 Gy). Non-irradiated (0 Gy) were used as a control. These experiments showed that MCF-7 cells viability was lower in the irradiated cells compared to the control group. In cells treated with CS-NP + reversine a similar decrease of cell viability was observed for both non-radiated and X-rays irradiated cells (**Figure 3**). Reversine inhibits the spindle assembly checkpoint (SAC) kinases Mps1 and Aurora B in a dose-dependent manner [23]. Because in breast cancer cells that were treated with reversine for only 1.5 hours there was not significant effect on cell viability in non-radiated (0 Gy) cells compared to the negative control and only a slight effect on cell viability on X-rays irradiated (2 Gy) cells, it is assumed that the observed effect on cell viability is largely due to the CS-NP themselves. Taken together, the data indicates that CS-NP sensitised MCF-7 cells to biological radiation. Because metal-containing nanoparticles have been reported to abrogate DNA repair and to cause extensive damage of telomeres in MCF-7 cells [27] we

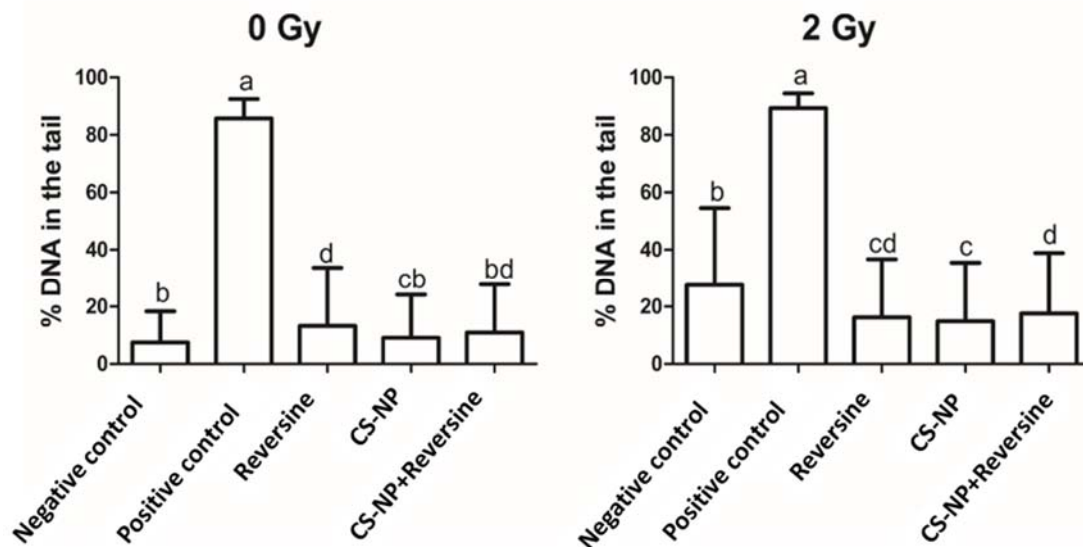
investigated if the sensitising effect of the CS-NP was due to alteration of DNA stability. To address this question, Comet assays were carried out in non-irradiated and X-rays irradiated cells. The results of these studies are described below.



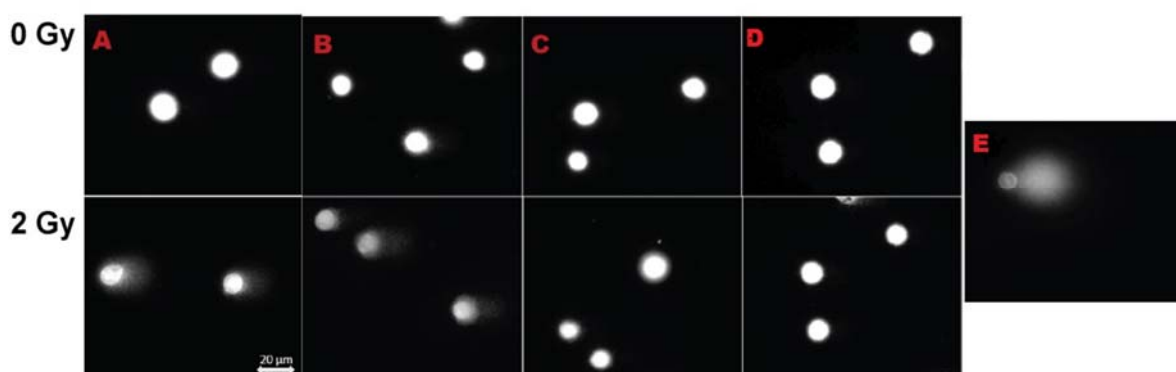
**Figure 3.** Effect of ionising-radiation (2 Gy) on MCF-7 cell viability. Non-irradiated (0 Gy) MCF-7 cells were used as negative control whereas MCF-7 cells treated with  $\text{H}_2\text{O}_2$  100  $\mu\text{M}$  to induce DNA damage following X- rays irradiation, were used as positive control. The bars represent the mean value of three independent experiments and the lines the standard deviation. Bars with same letter are not significantly different (Tukey > 0.05). Non-irradiated (0Gy) samples: negative control, 96.2±9.9; positive control, 36.6±7.8; reversine, 93.1±17.5; CS-NP, 74.5± 21.6; CS-NP+reversine, 64.7±, 26.8. X-rays irradiated (2Gy) samples: negative control, 90.3± 18.8; positive control, 21.3±14.9; reversine, 82.5± 20.5; CS-NP, 68.0±, 29.9; CS-NP+reversine, 62.4± 27.0.

### 3.5. DNA Damage by Comet Assay

The percentage of DNA damage in non-irradiated (0 Gy) and X-ray irradiated (2 Gy) MCF-7 cells, as determined by the alkaline comet assay, is shown in **Figure 4**. As anticipated, in both non-irradiated and irradiated cells, the extent of DNA damage was much lower than in the positive control  $\text{H}_2\text{O}_2$  at 100  $\mu\text{M}$ : 85.7±6.7 (0 Gy) and 89.3± 0.14 for the irradiated (2 Gy) cells. In non-irradiated (0 Gy) MCF-7 cells the extent of DNA damage was as follows: CS-NP, 9.2±0.43; reversine, 13.2± 0.59; CS-NP + reversine, 11.0 ± 0.48. A slight increase of DNA damage was observed when the cells were irradiated (2 Gy): CS-NP, 14.9±0.59; reversine, 16.2± 0.59; CS-NP + reversine, 17.6± 0.61. Such effect was clearer for the negative control, where the percentage of DNA in the tail was higher in the irradiated cells (0 Gy, 7.4±0.30; 2 Gy, 27.6± 0.77). We then used the comet assay to quantify DNA breaks associated with the direct damage to the cells induced by radiation (**Figure 5**). We selected the comet assay because it is generally recognised as one of the most sensitive methods for analysing and quantifying DNA damage in individual eukaryotic cells. Taken together, the data presented in Figures 4 and 5 indicate that the observed sensitisation of MCF-7 cells by the CS-NP was not directly related to genotoxicity. This is in stark contrast to the cellular effects reported in other types of nanoparticles, where genotoxicity resulting from DNA oxidative lesions has been observed [28].



**Figure 4.** Percentage of DNA damage in non-irradiated (0 Gy) and X-rays irradiated (2 Gy) MCF-7 cells as measured by the alkaline comet assay.  $\text{H}_2\text{O}_2$  at 100  $\mu\text{M}$  was used as the positive control. The bars represent the mean value and the lines the standard deviation. Bars with same letter are not significantly different (Dunns > 0.05). Average values for non-irradiated (0 Gy) samples: negative control,  $7.4 \pm 0.30$ ; positive control,  $85.7 \pm 6.7$ ; reversine,  $13.2 \pm 0.59$ ; CS-NP,  $9.2 \pm 0.43$ ; CS-NP+ reversine,  $11.0 \pm 0.48$ . Average values for X-rays irradiated (2 Gy) samples: negative control,  $27.6 \pm 0.77$ ; positive control,  $89.3 \pm 0.14$ ; reversine,  $16.2 \pm 0.59$ ; CS-NP,  $14.9 \pm 0.59$ ; CS-NP+ reversine,  $17.6 \pm 0.61$ .



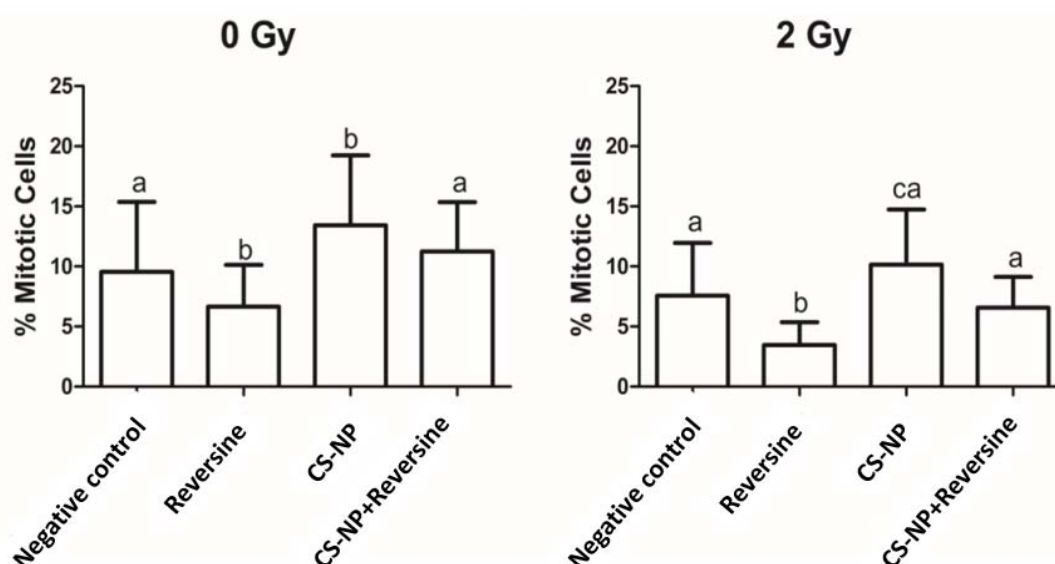
**Figure 5.** Representative images of the cells assayed by the comet assay. The nuclei was stained with diamond nuclei acid stain. Above, non-irradiated cells. Below, irradiated cells where A= negative control; B= cells treated with reversine 5  $\mu\text{M}$ ; C= cells treated with CS-NP; D= cells treated with CS-NP + reversine 5  $\mu\text{M}$ ; E= positive control, consisting of cells treated with 100  $\mu\text{M}$   $\text{H}_2\text{O}_2$ . All the images were taken at 40x magnification.

### 3.6. Mitotic Index

The ATP-binding competitor reversine is known to cause premature exit from mitosis in HeLa, U2OS, and retinal pigment epithelial cells during an unperturbed mitosis and to inhibit colony formation of human acute myeloid leukemia cells [22, 23]. Reversine was first reported as an inhibitor of Aurora B kinase that abolished the Aurora B-dependent phosphorylation of histone H3 [22]. In an attempt to establish whether CS-NP can synergise chemotherapy based on SAC kinases (Mps1 and Aurora kinase B) inhibition by reversine, the number of mitotic cells was estimated by cytometry analysis. Our study shows that in MCF-7 cells treated with reversine, the percentage of cells in mitosis was much lower than in the negative control in both non-irradiated and X-rays irradiated cells (**Figure 6**). Moreover, the use of an anti-phospho-histone H3 antibody revealed that in non-irradiated and X-rays irradiated cells, the number of MCF-7 cells in mitosis following treatment with CS-NP



alone was consistently higher (0 Gy,  $13.4 \pm 5.8$ ; 2 Gy,  $10.1 \pm 4.5$ ) than in cells exposed to reversine alone (0 Gy,  $6.7 \pm 3.4$ ; 2 Gy,  $3.5 \pm 1.8$ ) and slightly higher to that of cells treated with CS-NP + reversine (0 Gy,  $11.24 \pm 4.1$ ; 2 Gy,  $6.7 \pm 2.5$ ). Whether this behaviour indicates mitosis rescue by the CS-NP is uncertain and an interesting aspect that warrant further investigations (**Figure 6**). What is clear from our study is that the combined use of CS-NP + reversine followed by X-rays irradiation (2 Gy) does not lead to a synergistic response in the treated MCF-7 cells and that the sensitisation effect to X-rays irradiation is due only to the exposure to CS-NP.



**Figure 6.** Mitotic index of non-irradiated (0 Gy) and irradiated (2 Gy X-rays) MCF-7 cells. The bars represent the mean values whereas the lines indicate the standard deviation. Bars with same letter are not significantly different (Tukey > 0.05). Mean values of non-irradiated (0 Gy) samples: negative control,  $9.6 \pm 5.8$ ; reversine,  $6.7 \pm 3.4$ ; CS-NP,  $13.4 \pm 5.8$ ; CS-NP+ reversine,  $11.24 \pm 4.1$ . Mean values of X-rays irradiated (2 Gy) samples: negative control,  $7.79 \pm 4.4$ ; reversine,  $3.5 \pm 1.8$ ; CS-NP,  $10.1 \pm 4.5$ ; CS-NP+ reversine,  $6.7 \pm 2.5$ .

### 3.7. Mps1 Expression

As at least in cultured cells, amorphous silica and certain metal-containing nanoparticles can induce genotoxic, cytotoxic and changes in transcriptomic responses including the expression of genes related to oxidative stress responses, protein synthesis and degradative enzymes [29-31]. Because Mps1 is a SAC kinase amplified in MCF-7 breast cancer cells [32] we investigated if exposure to the CS-NP before and after ionising radiation had any effect on Mps1 kinase expression levels. Western blot analysis of SDS-PAGE (Supplementary Figure 1) showed that treatment of MCF-7 cells with CS-NP, reversine, and CS-NP + reversine followed by X-rays irradiation (2 Gy) did not affect the expression of Mps1 kinase in MCF-7 cells. A similar situation was observed in non-irradiated cells. In these studies  $\beta$ -actin was used as the loading control (Supplementary Figure 1).

## 4. Discussion

One important problem in clinical practice is that X-rays irradiation of tumour cells can also induced damage to the normal tissue. This situation has boosted the development of novel approaches including the use of nanotechnology to induce sensitisation to radiation on cancer cells [3]. In this work we evaluated the use of chitosan nanoparticles to sensitise MCF-7 cells to X-rays irradiation. Characterisation of the CS-NP (show in **Table 1** and **Figure 1**) included determination of the CS-NP size, shape, surface morphology, and surface charge. These are fundamental properties

that influence the nanoparticles interaction with biomolecules and the uptake mechanisms used by the cells, which include the opening of tight junctions [6] and clathrin and caveolae pathways [9]. Although the study of the precise uptake mechanism of chitosan-based nanoparticles was not addressed in this work, it is important to note that the CS-NP were detected inside the cells after three hours of exposure and that they were evenly distributed through the cytoplasm (**Figure 2**), arguing in favour of their use to target cancer cells. It is worth noting that independent studies by other researchers have shown that titania NP doped with rare earth elements is an effective strategy to cause radiosensitisation of tumour cells [7, 33]. Whether the radiosensitisation efficiency and/or stability within the cells of CS-NP is higher than that of metallic nanoparticles including rare earth elements-doped titania NP, remains to be established.

Earlier studies on CS-NP showed that high molecular weight chitosan are cytotoxic [34]. However, independent reports by others found no evidence of cytotoxicity of chitosan nanoparticles to MCF-7 cells following exposure for 6 and 24 h even at high concentrations of chitosan [31]. Our data is in agreement with the former reports, as it shows that in MCF-7 cells treated with CS-NP, the nanomaterial has a mild negative effect on the viability of MCF-7 breast cancer cells and that such effect is not due to the induction of DNA damage (**Figures 3-5**). Although producing nanoparticles of different sizes is not relevant to this study, is worth mentioning that CS-NP of a narrower size distribution can be achieved through a combination of changes in the method of nanoparticles synthesis reported here together with separation of CS-NP of various dimensions using size exclusion chromatography. It would be interesting to explore if CS-NP of narrower size distribution can enhance their permeation across the cell membrane and/or their retention in tumour sites.

It is worth noting that in mitotic index determination based on quantification of P-Ser(10)-H3, a well established mitosis marker and Aurora kinase B substrate, the number of mitotic cells treated with CS-NP + reversine was higher than that observed in cells exposed to reversine alone (**Figure 6**), suggesting a possible rescue of the SAC. Such scenario would argue against the development of a therapeutic strategy aiming to induce premature mitosis exit. However, in our hands the potential rescue of the SAC in the treated MCF-7 cells was not paralleled by an improvement in cellular viability. Further combinatorial studies using different dose levels of CS-NP and reversine at extended exposure times are required to clarify the apparent SAC rescue induced by CS-NP. Furthermore, in the light of our current understanding of the mechanism of action of SAC inhibitors, targeting mitosis exit inhibition rather than induction of premature mitotic exit may be advantageous to reduced cytotoxic side effects [35]. This may be particularly effective for mitosis exit inhibitors used in combination with vinca alkaloids (vinblastine, vincristine, vindesine, vinflunine, vinorelbine) and taxanes (docetaxel, taxol), two chemical classes of drugs that through different mechanisms of action, induce mitotic arrest by interfering with microtubule assembly, thus promoting genome instability above the threshold required to kill cancer cells [36]. It will be important to assess to what extent CS-NP can sensitise triple negative breast cancer and other types of tumours of poor prognosis to combinations of small size inhibitors of mitosis together with microtubule poisons when used alone and in conjunction with subsequent x-rays irradiation.

## 5. Conclusions

Chitosan nanoparticles constitute a nanomaterial that has potential to be used as adjunct to radiotherapy given their property to radiosensitise MCF-7 breast tumour cells, possibly as a result of their accumulation (passive or targeted) in the cancer cells. Future work will seek to identify the cell signalling pathway(s) implicated in the sensitisation to radiation of chitosan-nanoparticles, which should provide new molecular insight of cancer cells sensitisation to X-rays radiation. This new knowledge may in turn open up new opportunities for the adequate stratification of breast cancer patients in clinical trials, leading to better therapeutics strategies for the treatment of this disease.

**Author Contributions:** SPO, PRN and VMB-G designed the study. RDT designed the nanoparticles. SPO carried out the synthesis of nanoparticles and most of the experiments. LH and JM assisted on TEM data collection and on confocal microscopy, respectively. EE, MH, and MK assisted on radiation data collection. SPO, PRN and VMB-G worked on data interpretation and prepared the manuscript with input from all the co-authors.

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