

Sensitization of chondrosarcoma cells with PARP inhibitor and impact on cancer stem cells

Mathieu Césaire, Antoine Gilbert, Julie Laurence, Nicolas Elie, Florence Giffard, Siamak Haghdoost, Louis-Bastien Weiswald, François Chevalier

▶ To cite this version:

Mathieu Césaire, Antoine Gilbert, Julie Laurence, Nicolas Elie, Florence Giffard, et al.. Sensitization of chondrosarcoma cells with PARP inhibitor and impact on cancer stem cells. ERRS 2021, Nov 2021, CAEN, France. hal-03757108

HAL Id: hal-03757108 https://hal.science/hal-03757108

Submitted on 22 Aug 2022

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Centre de Recherche sur les Ions, les Matériaux et la Photonique



Sensitization of chondrosarcoma cells with PARP inhibitor and impact on cancer stem cells

<u>Mathieu Césaire¹, Antoine Gilbert¹, Julie Laurence¹, Nicolas Elie², Florence Giffard³, Siamak Haghdoost¹,</u> Louis-Bastien Weiswald⁴ and François Chevalier¹

- **1 ARIA Laboratory, UMR6252 CIMAP Caen, France**
- 2 CMAbio³ Centre de Microscopie Appliquée à la biologie, Caen, France
- **3 UMR INSERM U1086 ANTICIPE, Axe BioTICLA, Caen, France**
- 4 Plateforme ORGAPRED, UMR INSERM U1086 ANTICIPE, Axe BioTICLA, Caen, France

Chondrosarcoma is a primary tumour of bone and cartilage. One of the standard treatments is conventional X-ray therapy but chondrosarcomas of high malignancy are resistant. One of the mechanisms of radioresistance concerns the compartment of cancer stem cells (CSC) via their self-renewal and tumorogenesis capabilities. Radiosensitizing molecules, such as PARP inhibitors (PARPi) and carbon ions irradiation (C-ions), could overcome radioresistance and act on the CSC [1–3]. We investigated the cell survival and the impact on CSC of a chondrosarcoma line L835 in 2D and 3D culture models after X-ray (225 KV) compared to **C-ions** (Ganil, 73 keV/ μ m) in combination or not with PARPi. For the study of cell survival, we carried out clonogenic tests. For the study of the CSC, we wanted to determine the ability of cells to form tumorspheres in wells containing a depleted culture medium. We also performed a kinetic analysis on **3D models of spheroids**, using DNA damage tests (γ -H2AX foci) and immunohistochemistry of ALDH, an enzyme more expressed in CSC. Vares, G. et al. Radiother Oncol 2020, 150, 253-261

with the support of the Normandy region within the CHOXTRACC project



RÉGION NORMANDIE



Chevalier, F. et al. Technol Cancer Res Treat 2019, 18, 1533033819871309.

Césaire, M. et al. *J Bone Oncol* 2019, *17*, 100246.

Aim of this study

1- explore the differential cellular responses of chondrosarcomas to conventional radiotherapy and hadrontherapy

2- estimate the capacity of PARP inhibitors in radio-sensitizing chondrosarcoma cells

3- estimate the impact of hadrontherapy and PARP inhibitors on chondrosarcoma cancer stem cells







L835 tumorspheres increased after combination of Xrays and Olaparib from 2 Gy to 6 Gy

The same effect was observed with Cions at 1 Gy and 3 Gy We didn't observe this effect at 2 Gy with only 8 tumorspheres and other experiments are needed





Université de Caen Normandie