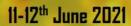
ISBN: 978-81-950236-5-3

2nd Virtual Annual

International Conference on Naturopathy, Nanotechnology, Nutraceuticals and Immunotherapy in Cancer Research



Organized by

School of Life Sciences

&

Association of Cancer Education and Research (ACER)

B.S. Abdur Rahman Crescent Institute of Science & Technology, Chennai, India

In association with

Purdue University, USA

Sponsored by

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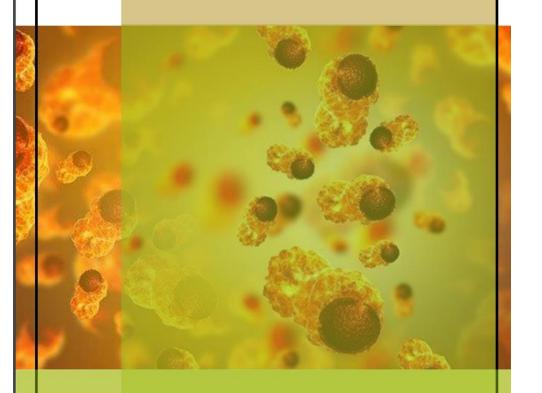












2nd Virtual International Conference on Naturopathy, Nanotechnology, Neutraceuticals and Immunotherapy Cancer Research- 2021



Proceedings of ICN3IC-21, Virtual, 11 June – 12 June 2021 ISBN: 978-81-950236-5-3

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ANALYSIS OF INVASIVE PROPERTIES OF RETINOBLASTOMA AND UVEAL MELANOMA XENOGRAFT USING CHICK CHORIOALLANTOIC MEMBRANE ASSAY

Perumal Jayaraj¹, Nimita Kant², Seema Sen³

¹Department of Zoology, Sri Venkateswara College, University of Delhi

²Department of Zoology, Shivaji College, University of Delhi.

3Department of Ocular Pathology, Dr. R. P. Centre AIIMS, New Delhi.

Corresponding Author's email: nimita@shivaji.du.ac.in

Introduction: Retinoblastoma (RB) and uveal melanoma (UM) are the two most invasive of ocular malignancies. The current research on these often relies on nude mice model and matrigel invasion assay to demonstrate their metastatic behaviour using cell lines. Recent studies have underlined the implication of utilizing chorioallantoic membrane assay (CAM) to understand the metastatic potential of patient derived xenografts (PDXs). In this study the invasive property of RB and UM PDXs were analysed on chick CAM. Methodology: RB and Uveal Melanoma tumors were isolated after compliance from donor patients. The tumour was dissected and 2-4mm of the tumor tissue were placed onto the CAM, On the 17th day of chick embryonic development, the CAM tissue bearing the tumor was excised and fixed in 4% formaldehyde for H/E staining and immunohistochemistry (IHC) analysis. Synatophysin anti human antibody was used for immunohistochemical staining of RB cell invasion in CAM. Result: Examination of CAM by histology revealed that the CM xenograft melanoma invaded into the mesoderm of the chick CAM membrane. Further both the RB and UM tissues induced angiogenesis in the area of in the implantation on the chick CAM. The immunohistochemical analysis revealed anti-human synaptophysin positivity in RB cells which were attached to the ectoderm of the CAM and in cell invading into the CAM mesoderm. Discussion: An increased vascular growth and invasion of both RB and UM PDXs shows that the chick CAM assay model can be a good alternative to study invasive properties of Ocular malignancies.

Keywords: Retinoblastoma, Uveal Melanoma, Chorio allantoic membrane assay, Patient derived Xenograft.

Proceedings of ICN3IC-21, Virtual, 11 June – 12 June 2021 ISBN: 978-81-950236-5-3

CCA187

INVESTIGATION OF THE ROLE OF CYCLOOXYGENASE-2 IN SQUAMOUS CELL CARCINOMA OF THE EYELID USING IMMUNOHISTOCHEMISTRY

Chitra¹, Perumal Jayaraj², Seema Kashyap³, Seema Sen³

²Department of Zoology, Sri Venkateswara College, University of Delhi

¹Department of Biosciences, Jamia Millia Islamia, New Delhi

Corresponding Author's email: jay.aiims@gmail.com

Squamous cell carcinoma (SCC) of the eyelid is a malignant tumor which arises as a result of abnormal growth of squamous cell layer of the skin epithelium. Cyclooxygenases are enzymes which are crucial for lipid metabolism. Cyclooxygenase-2(COX-2) is a mitogeninducible isoform of cyclooxygenase (prostaglandin-endoperoxide synthase). The expression of COX-2 is undetectable during normal, non-pathological conditions but is rapidly induced in various cancers and its inhibition by Non Steroidal Anti-inflammatory Drugs (NSAIDs) is known to reduce the risk of many cancers by suppressing and preventing tumorigenesis. Aim of our study is to investigate the role of COX-2 signalling pathway in human eyelid SCC and its association with clinicopathological parameters. For this, 20 histopathologically confirmed cases of SCC were selected for the study. Cytoplasmic expression of COX-2 was detected in 80% of the SCC cases. COX-2 expression was found to be correlated with size of the tumor; 63% of tumor cases with size >3cm showed positive expression of COX-2. Significant expression of COX-2 was observed in the invasive front as well as infiltrating lymphocytes. COX-2 expression was found to be associated with differentiation as high expression of COX-2 was observed in 60% well differentiated (WD) SCC cases, highlighting significance of COX-2 in differentiation in SCC. For the first time, COX-2 expression has been demonstrated in eyelid SCC, however, study in large cohort is required for further statistical analysis and significant correlation with these parameters.

Keywords: Squamous cell carcinoma, Cyclooxygenase 2, Non-steroidal anti inflammatory drugs.

³Department of Ocular Pathology, All India Institute of Medical Sciences, New Delhi