

Poster presentation

Open Access

The use of specific anti-growth factor antibodies to abrogate the oncological consequences of transfusion: an in-vitro study

Tahwinder Upile*, Waseem Jerjes, Jaspal Mahil and Colin Hopper

Address: Head & Neck Centre, University College London Hospitals, London, UK

* Corresponding author

from 1st Scientific Meeting of the Head and Neck Optical Diagnostics Society
London, UK. 14 March 2009

Published: 28 July 2009

Head & Neck Oncology 2009, 1(Suppl 1):P16 doi:10.1186/1758-3284-1-S1-P16

This abstract is available from: <http://www.headandneckoncology.org/content/1/S1/P16>

© 2009 Upile et al; licensee BioMed Central Ltd.

Introduction

Peri-operative blood transfusion is associated with reduced prognosis in a number of solid malignancies. We investigate its role in a head & neck squamous cell cancer cell line. Growth of these cell lines was analogous to endothelial growth. Direct exposure to transfusion products exaggerated this effect. It was logical therefore to assess the effects of anti-endothelial antibodies on this interaction.

Materials and methods

Control (HUVEC) and tumour cell lines were exposed to transfusion products. The pre-incubation of the transfusion product with anti-endothelial growth factors was assessed by a growth assay.

Results

The antibody did not directly reduce growth in the tumour cell line, however there was a significant reduction ($p < 0.001$) in tumour cell line growth caused by transfusion products pre-incubation with anti-endothelial growth factor antibody. This was found in several other tumours.

Conclusion

We have shown some of the prognostically deleterious effects of peri-operative transfusion in head & neck cancer patients is caused by the transfusion products release of endothelial growth factors. This is found to be the case in several of the tumour groups (Colonic and Prostate) for which this phenomena has been previously reported. It can now be hypothesized that this is due to the specific

expression of receptors to these growth factors in these tumour types which are not universally found. It would also explain why this phenomenon does not occur for all tumour types.