

University of Huddersfield Repository

Elkashef, Sara M., Allison, Simon J., Sadiq, Maria, Basheer, Haneen A., Ribeiro Morais, Goreti, Loadman, Paul M., Pors, Klaus and Falconer, Robert A.

Polysialic acid sustains cancer cell survival and migratory capacity in a hypoxic environment

Original Citation

Elkashef, Sara M., Allison, Simon J., Sadiq, Maria, Basheer, Haneen A., Ribeiro Morais, Goreti, Loadman, Paul M., Pors, Klaus and Falconer, Robert A. (2016) Polysialic acid sustains cancer cell survival and migratory capacity in a hypoxic environment. Scientific Reports, 6. p. 33026. ISSN 2045-2322

This version is available at http://eprints.hud.ac.uk/id/eprint/29412/

The University Repository is a digital collection of the research output of the University, available on Open Access. Copyright and Moral Rights for the items on this site are retained by the individual author and/or other copyright owners. Users may access full items free of charge; copies of full text items generally can be reproduced, displayed or performed and given to third parties in any format or medium for personal research or study, educational or not-for-profit purposes without prior permission or charge, provided:

- The authors, title and full bibliographic details is credited in any copy;
- A hyperlink and/or URL is included for the original metadata page; and
- The content is not changed in any way.

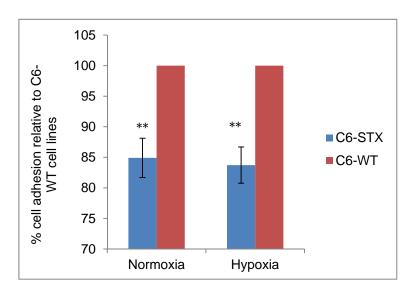
For more information, including our policy and submission procedure, please contact the Repository Team at: E.mailbox@hud.ac.uk.

http://eprints.hud.ac.uk/

Polysialic acid sustains cancer cell survival and migratory capacity in a hypoxic environment

Sara M. Elkashef,^a Simon Allison,^b Maria Sadiq,^a Haneen Basheer,^a Goreti Ribeiro Morais,^a Paul M. Loadman,^a Klaus Pors^a and Robert A. Falconer^a

Supplementary data 1 (S1)



S1. Effect of hypoxia on the polySia-mediated adhesiveness of cancer cells. A polySia-mediated reduction in cell adhesion to Matrigel[©] in C6-STX but not C6-WT cell line was observed under both normoxia and hypoxia. (**P≤0.01)

a Institute of Cancer Therapeutics, Faculty of Life Sciences, University of Bradford, West Yorkshire BD7 1DP, U.K. Tel: +44 (0)1274 235842; Email: r.a.falconer1@bradford.ac.uk

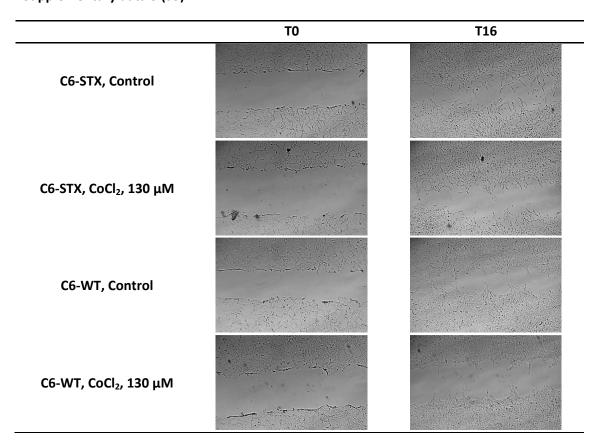
^b Department of Pharmacy, School of Applied Sciences, University of Huddersfield, Huddersfield HD1 3DH, U.K.

Supplementary data 2 (S2)

Cell line and conditions	Apoptotic cells		- Viable cells
	Late apoptotic (necrotic)	Early apoptotic	viable cells
SH-SY5Y (Normoxia)	6.16± 0.15	10.36 ± 2.01	81.94 ± 2.19
SH-SY5Y (Hypoxia)	4.41 ± 0.02	7.29 ± 0.29	85.54 ± 0.43

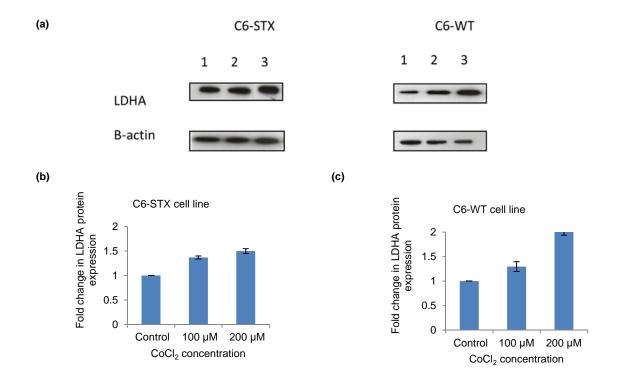
S2. Percentage of annexin V staining of SH-SY5Y cells under hypoxic and normoxic conditions

Supplementary data 3 (S3)



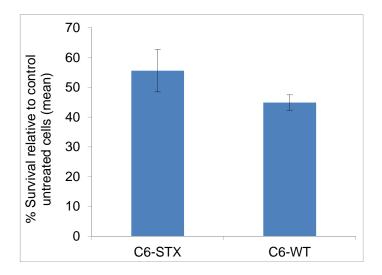
S3. Migration of C6-STX and C6-WT cells following CoCl₂ treatment. Wound size immediately and 16 hours after scratching illustrates wound closure of C6-STX and C6-WT cells with/without 100 μ M CoCl₂.

Supplementary data 4 (S4)



S4. Effect of CoCl₂ treatment on C6 cell lines. (A) Expression of LDHA after treatment of C6-STX and C6-WT cells with different concentrations of CoCl₂. Lane 1 represents control untreated cells, lane 2 after 130 μ M CoCl₂ and lane 3 after 200 μ M CoCl₂. Results are representative of two independent experiments; (B) quantification of fold-change in LDHA expression for C6-STX cells after treatment with 100 μ M CoCl₂ (16 hours) and 200 μ M CoCl₂ (24 hours); (C) quantification of fold-change in LDHA expression of C6-WT after treatment with 100 μ M CoCl₂ (16 hours) and 200 μ M CoCl₂ (24 hours).

Supplementary data 5 (S5)



S5. Effect of HIF-1α induction on the survival of C6-STX and C6-WT cells. (P≥0.05, not significant)