

UNDER STRICT EMBARGO UNTIL: 1400 GMT MONDAY 07 DECEMBER 2015

Cancer study identifies genes that stop onset of leukaemia

Genes that act as brakes to stop the development of an aggressive form of leukaemia have been identified by researchers.

Their findings offer fresh insights into how to tackle the disease and could lead to new therapies that prevent relapses.

Scientists have found that two molecules – Hif-1alpha and Hif-2alpha – work together to stop the formation of leukemic stem cells in an aggressive type of blood cancer called Acute Myeloid Leukaemia (AML).

The cancer occurs when production of new blood cells by the bone marrow goes awry. This leads to the formation of leukemic stem cells, which fuel the disease and provide a constant flow of abnormal leukaemia cells.

The University of Edinburgh study shows that blocking Hif-2alpha – or both Hif-1alpha and Hif-2alpha – accelerates the development of leukaemia.

The findings are surprising because previous research had suggested that blocking Hif-1alpha or Hif-2alpha may stop leukaemia progression.

The work was carried out at the University's MRC Centre for Regenerative Medicine and the Edinburgh Cancer Research Centre. Researchers say that their new results suggest that therapies designed to block these molecules may have no impact or could even worsen disease.

Conversely, designing new therapies that promote the activity of Hif-1alpha and Hif-2alpha could help to treat AML or stop the disease from recurring after chemotherapy.

Around 2,500 people are diagnosed with AML in the UK each year. Chemotherapy drugs can help to eliminate leukaemia cells but have no effect on leukemic stem cells that cause the disease. This means the disease sometimes relapses.

The study is published in the *Journal of Experimental Medicine*. It was funded by Cancer Research UK, Bloodwise, the Kay Kendall Leukaemia Fund and the Medical Research Council.

Ranked among the top universities in the world

Professor Kamil R Kranc, Cancer Research UK Senior Fellow at the MRC Centre for Regenerative Medicine, who led the study, said: "Our discovery that Hif-1alpha and Hif-2alpha molecules act together to stop leukaemia development is a major milestone in our efforts to combat leukaemia. We now intend to harness this knowledge to develop curative therapies that eliminate leukaemic stem cells, which are the underlying cause of AML."

Dr Milica Vukovic, first author of the study, said: "Leukaemia is an umbrella term for a vast number of very complicated and different diseases. Given our findings implicating Hif-1alpha and Hif-2alpha as tumour suppressors in AML, it would be very interesting to investigate their roles in other leukaemias."

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