

Scientists discover cells critical to childhood leukemia

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Scientists at The Royal Melbourne Hospital and the University of Melbourne in Australia have discovered the cells that cause a common type of childhood leukaemia - T cell Acute Lymphoblastic Leukaemia (T-ALL). Targeting of these cells may lead to improved treatments for this disease and help prevent relapse.

The team, led by Dr Matthew McCormack and Dr David Curtis of the Rotary <u>Bone Marrow</u> Research Laboratories and the University's Department of Medicine at The Royal Melbourne Hospital, made the discovery whilst studying mice prone to developing this <u>leukaemia</u>.

The results have been published online today by the prestigious international journal *Science*.

The team found that with irradiation treatment in animal models, over 99 per cent of cells in the thymus were killed, but these stem cell-like cells persisted and rapidly recovered. This suggests that these cells may survive therapy and be responsible for relapsed disease following treatment.

Currently, children with T-ALL are given extended therapy over two to three years in an attempt to stop a relapse. More targeted therapy on the thymus cells could reduce the length and toxicity of treatment and prevent relapse.

Dr McCormack, a leading international expert on childhood leukaemia,



said: "The cellular origins of this leukaemia are not well understood. Our discovery that these cells are similar to normal <u>stem cells</u> explains why they are capable of surviving for long periods. It also explains why they are remarkably resistant to treatment."

Approximately 50 new cases of T-ALL are diagnosed every year in Australia, two thirds of these in children or adolescents. Adults also contract T-ALL, and the majority succumb to resistant or relapsed disease.

Dr Curtis, a Clinical Haematologist and head of the Leukaemia Research Program at The Royal Melbourne Hospital, said: "The identification of these cells provides an important target for the development and testing of new treatments for patients with T cell Acute Lymphoblastic Leukaemia."

The team will now focus on novel treatments capable of killing these <u>cells</u>, which may lead to clinical trials within the next five years.

Provided by University of Melbourne

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