

GM-CSF modulates the migratory phenotype of vaccine-induced T cells by enhancing CXCR3 expression

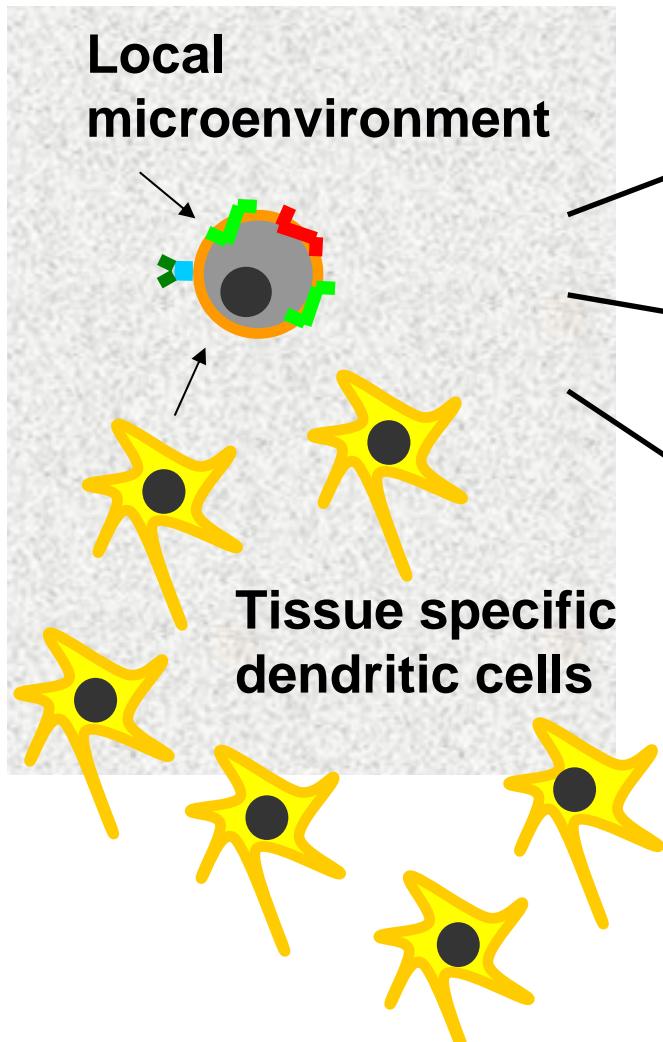
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Background



Selective chemokine receptor



CXCR3

CCR4

CCR9

Inflammatory
tissue

Tumor tissue

Skin

Small intestine

Aim of the study

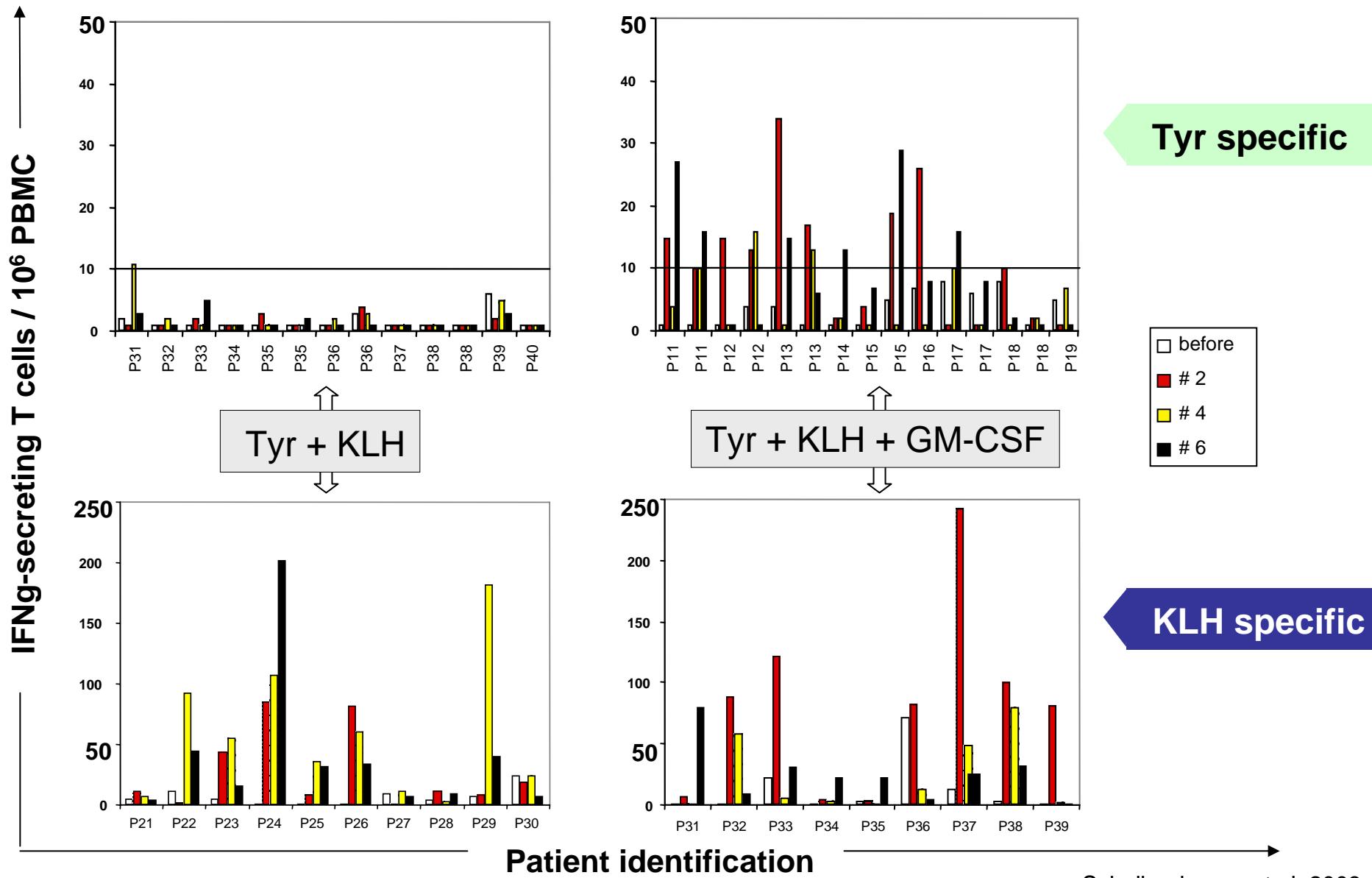
Analysis of

- Expression of chemokine receptors CXCR3 and CCR4 on vaccine-induced T cells.
- Influence of GM-CSF as vaccine adjuvant on chemokine receptor expression.

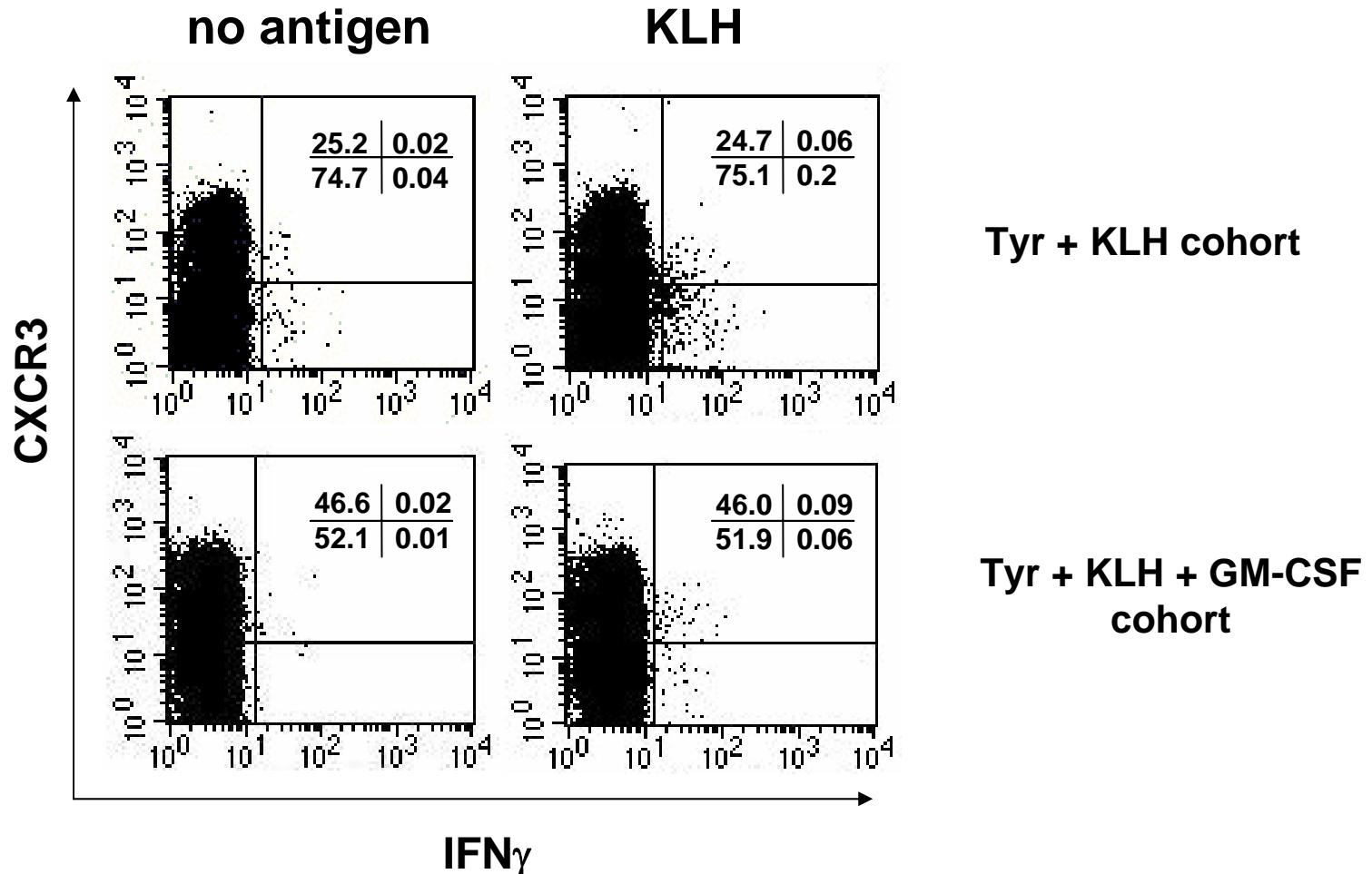
Materials & Methods

- **Patient samples:** stage III/ IV melanoma
 - 2 cohorts
 - 1) Tyrosinase peptide + KLH + GM-CSF
 - 2) Tyrosinase peptide + KLH
- **T cell response assessment**
by IFN γ flow cytometry analysis

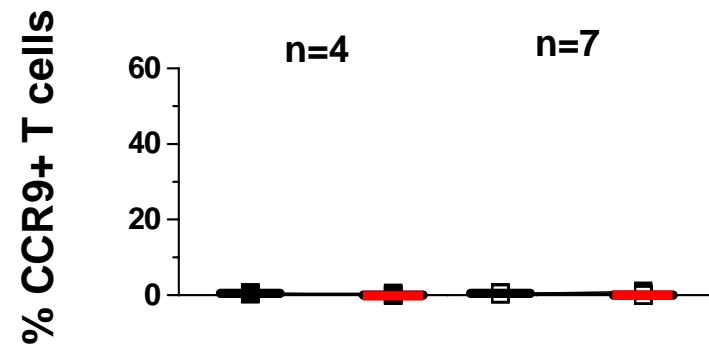
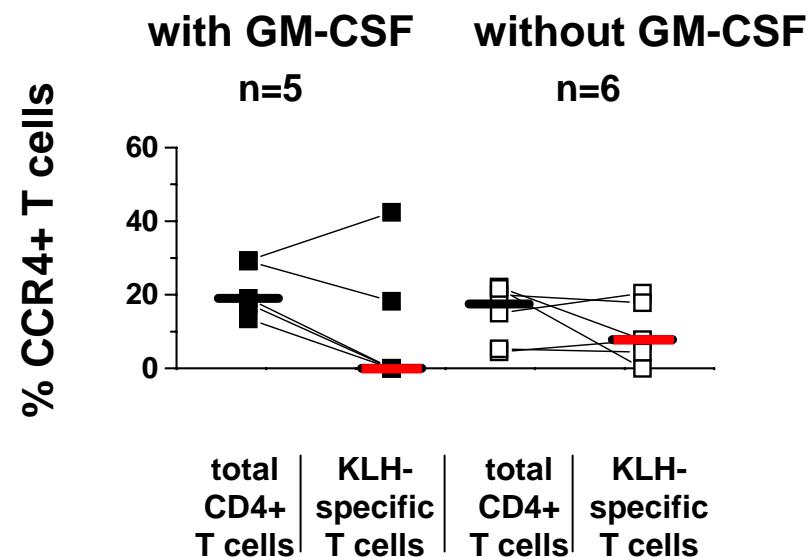
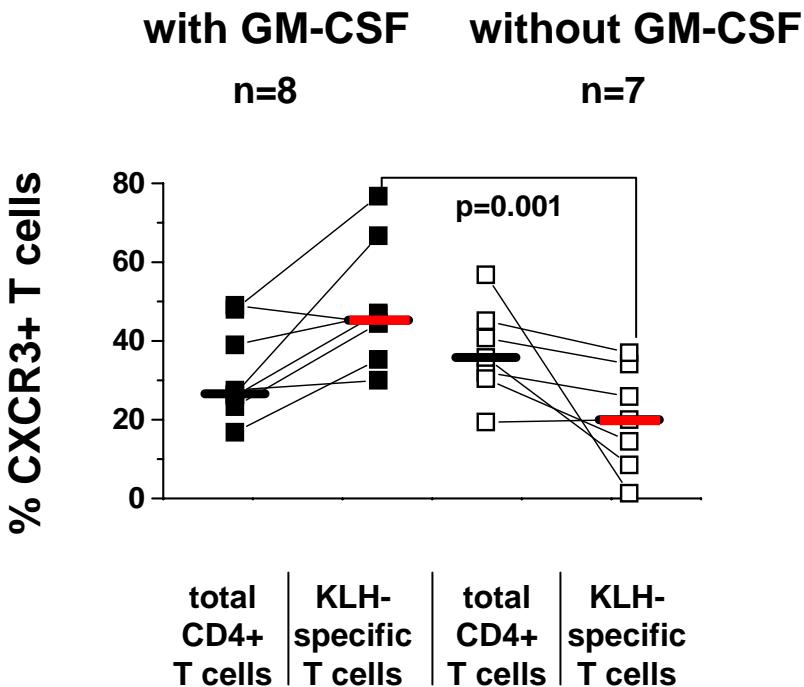
Phase I trial of tyrosinase peptide with adjuvants GM-CSF and KLH



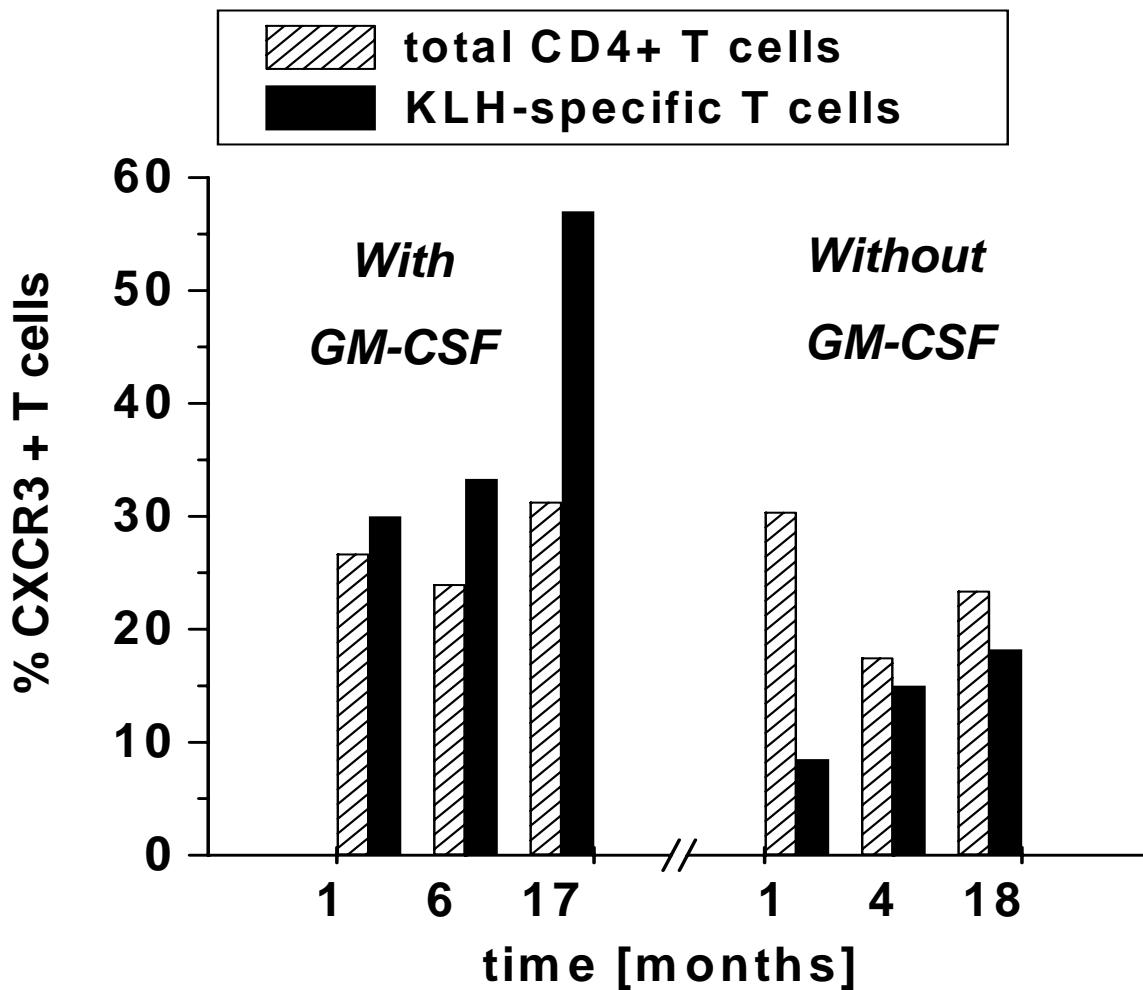
CXCR3 expression on KLH-specific T cells



CXCR3, CCR4 and CCR9 expression on KLH-specific T cells



Follow up of CXCR3 profile of KLH-specific T cells



Summary

- These results show for the first time in humans that it is possible to modulate chemokine receptor expression on T cells generated by vaccination by modifying the local vaccine milieu.
- Immunization protocols that induce T cells expressing CXCR3 may be of considerable value for improvement of immune therapy strategies.



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