

Differentiation and Lymphokine Responsiveness of Putative NK Cells in Early Human Development

メタデータ	言語: jpn 出版者: 公開日: 2022-11-07 キーワード (Ja): キーワード (En): 作成者: Taniguchi, Noboru メールアドレス: 所属:
URL	https://doi.org/10.24517/00068041

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1986 Fiscal Year Final Research Report Summary

Differentiation and Lymphokine Responsiveness of Putative NK Cells in Early Human Development

Research Project

Project/Area Number

60480242

Research Category

Grant-in-Aid for General Scientific Research (B)

Allocation Type

Single-year Grants

Research Field

Pediatrics

Research Institution

Kanazawa University

Principal Investigator

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Project Period (FY)

1985 - 1986

Keywords

NK cell activity / NK progenitor cells / Immature NK cells / Fetus / Neonates / Interferon- γ (IFN- γ) / インターロイキン・ γ (IL-2)

Research Abstract

1. Peripheral blood mononuclear cells(MNC) from fetuses of estimated gestational age of 20 wk lacked NK cell activity against K 562 target cells even after 18 hr-treatment with interferon- γ (IFN- γ). Low, but significant levels of basal and IFN- γ -inducible NK cell activity were observed in premature infants of 27-wk-gestation, with a progressive increase of these activities during the last trimester of pregnancy. Contrary to IFN- γ , interleukin-2(IL-2) could induce a marked NK cell activity even in MNC from fetuses of 20-wk-gestational age and in Leu- γ cell population of cord MNC, each of them lacked both basal and IFN- γ -inducible NK cell activity. These ontogenic development and lymphokine responsiveness of human NK cell activity indicated that putative precursors of NK effector cells might be divided into IFN- γ -responsive, more mature inactive forms of NK cells and into putative NK progenitors responding well to IL-2, but not to IFN- γ , which might appear at an earlier stage of fetal development than IFN- γ -responsive ones.
2. In cord blood, Leu- γ cells were comparable in number with adult controls, but Leu- γ cells were very meager. Preliminary studies indicated that low levels of Leu- γ cells in cord MNC might be responsible for the poor ability of cord blood in their anti-CD3-inducible cytotoxicity.
3. Dissociated production of IL-2 and IFN- γ , ample production of IL-2 and poor secretion of IFN- γ , on PHA stimulation is a characteristic of cord MNC. In this report, some experimental data suggesting that IFN- γ production by PHA-stimulated cord MNC to be down-

regulated by preferential activation of suppressor precursors against IFN- γ production, were presented. These PHA-inducible suppressor precursors in cord MNC expressed in large T4 helper phenotype and were radiosensitive in nature.

Research Products (11 results)

	All	Other
	All	Publications (11 results)
[Publications] UENO,Yoshiki: Journal of Immunology. 135. 180-184 (1985)		▼
[Publications] SEKI,Hidetoshi: Journal of Immunology. 135. 2351-2356 (1985)		▼
[Publications] SEKI,Hidetoshi: Journal of Immunology. 137. 3158-3161 (1986)		▼
[Publications] KOIZUMI,Shoichi: Blood. 68. 1065-1073 (1986)		▼
[Publications] SATO,Hiroshi: Journal of Biological Response Modifiers. 5. 191-201 (1986)		▼
[Publications] TANIGUCHI,Noboru: Proceeding of Japan Medical Reproductive Immunology.		▼
[Publications] UENO, Yoshiki: "Differential effects of recombinant human interferon- γ and interleukin-2 on natural killer cell activity of peripheral blood in early human development." Journal of Immunology. 135. 180-184 (1985)		▼
[Publications] SEKI, Hidetoshi: "Mode of in vitro augmentation of natural killer cell activity by recombinant human interleukin 2: a comparative study of Leu- $\text{CD}11\text{a}^+$ and Leu- $\text{CD}11\text{b}^-$ cell populations in cord blood and adult peripheral blood." Journal of Immunology. 135. 2351-2356 (1985)		▼
[Publications] SEKI, Hidetoshi: "Phenotypic and functional characteristics of active suppressor cells against IFN- γ production in PHA-stimulated cord blood lymphocytes." Journal of Immunology. 137. 3158-3161 (1986)		▼
[Publications] KOIZUMI, Shoichi: "Malignant clonal expansion of large granular lymphocytes with a Leu- $\text{CD}11\text{a}^+$, Leu- $\text{CD}7^-$ surface phenotype: in vitro responsiveness of malignant cells to recombinant human interleukin 2." Blood. 68. 1065-1073 (1986)		▼
[Publications] SATO, Hiroshi: "Monoclonal antibody which has the neutralizing activities for human IL-2." Journal of Biological Response Modifiers. 5. 191-201 (1986)		▼

URL: https://kaken.nii.ac.jp/report/KAKENHI-PROJECT-60480242/604802421986kenkyu_seika_hokoku_

Published: 1988-11-09