**Brie Definitive Report**

**Tumorigenicity Conferred to Lymphoma Mutant by Major Histocompatibility Complex-encoded Transporter Gene**

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**Summary**

Presentation of antigenic peptides by major histocompatibility complex (MHC) class I molecules requires MHC-encoded molecules of the adenosine triphosphate binding cassette (ABC) family. Defects in these proteins represent a potential risk, since they are essential links in the machinery of T cell–mediated surveillance which continuously scrutinizes peptide samples of cellular proteins. Nevertheless, transfection of the mouse lymphoma mutant RMA-S with the rat ABC gene mtp2 (homologue to mouse HAM2 and human RINGI1), commonly termed TAP-2 genes, led to a marked increase in tumor outgrowth potential in vivo. This occurred despite restored antigen presentation and sensitivity to cytotoxic T lymphocytes, and was found to be due to escape from natural killer (NK) cell–mediated rejection. It has previously been proposed that adequate expression of self-MHC class I is one important mechanism to avoid elimination by NK cells. Our data argue that a defect in the machinery responsible for processing and loading of peptides into MHC class I molecules is sufficient to render cells sensitive to elimination by NK cells. The latter thus appear to function as a surveillance of the peptide surveillance machinery.

MHC class I molecules present short antigenic peptides to CD8+ cytotoxic T cells (1-3). The peptides presented are derived from cytosolic proteins, usually translated endogenously (2). There are at least two genes in the MHC that code for products involved in peptide supply to MHC class I molecules, termed TAP-1 and TAP-2. These products are homologous to the transport proteins in the ATP binding cassette (ABC) family, and are thought to mediate transport from the cytoplasm to the endoplasmic reticulum, either of peptides or a cofactor essential for peptide loading into MHC class I molecules (4-13). The mouse lymphoma mutant RMA-S has a profound (14-16), but not total (17-19), peptide presentation defect in the MHC class I presentation pathway, which is due to a premature stop in the message of the mouse TAP-2 gene HAM2 (19a). The phenotype was restored to normal by transfection of TAP-2 genes from different species, first the mtp2 rat gene (9) and then the HAM2 mouse gene (12) and the RINGI1 human gene (A. Townsend, personal communication); the mtp2 transfectant (RMS-tmt2) was able to process and present influenza virus epitopes to CD8+ T cells (9). This transfectant system has now made it possible to investigate directly the role of TAP genes for in vivo phenomena such as tumorigenicity and transplant rejection. The aim of the present study was to investigate the effects on in vivo immunobiology of RMA-S after restoration of its antigen presentation defect by the rat TAP-2 gene.

**Materials and Methods**

**Animals.** C57BL/6 mice were bred and maintained at the Department of Tumor Biology, Karolinska Institutet. Athymic nu/nu C57BL/6 mice were purchased from Bomholtegaarden, Denmark. All mice were 4-6 wk old at the start of the experiments.

**Tumor Cell Lines.** RMA (derived from RBL-5 after mutagenization, nonselected (20, 21)), RMA-S (derived from RBL-5 after mutagenization and selection with anti-H-2 b antiserum plus complement for loss of MHC class I expression (20, 21)), and RMA-S.mtp2 (RMA-S transfected with mtp2 cDNA; a neomycin-resistant expression vector driven by the β-actin promoter (9)) were grown in RPMI 1640 supplemented with penicillin and streptomycin plus 5% FCS and kept in 50-ml culture flasks at 37°C, 5% CO2. The tissue-cultured medium of RMA-S.mtp2 was supplemented with G418 1 mg/ml to maintain the plasmid.

**In Vivo Rejection Assay.** Untreated and NKI.1 antibody-treated (0.2 ml NKI.1 ascitic fluid intraperitoneally, days -1 and +5) syngeneic C57BL/6 mice were used. Tumor cells were inoculated subcutaneously either in separate flanks or in separate animals, and outgrowth was monitored by palpations twice weekly. Tumors appeared at the site of inoculation, grew progressively, and subsequently killed the mice. In vitro tumor cell lines, phenotypically indistinguishable from the injected tumor cell line, could regularly...
be established from these tumors (data not shown). Macroscopically, metastases were not observed. However, if single cell suspensions were prepared from different organs, e.g., spleen, in vitro tumor cell lines could occasionally be established, which were phenotypically indistinguishable from the injected tumor cell line (data not shown). After initial experiments, the mice were killed when the mean tumor diameter reached 15 mm and no signs of regression were seen. The tumors were examined macroscopically after killing the animals. Mice without tumor growth were observed for at least 8 wk after inoculation.

Short-Term In Vivo Rejection Assay with 125I-IUdR-Labeled Cells. Untreated and NK1.1 antibody-treated (0.2 ml NK1.1 ascitic fluid intraperitoneally 24 h before the test) syngeneic C57BL/6 mice were used. 20 \times 10^6 tumor cells (RMA, RMA-S, and RMA-S.mtp2', respectively) were incubated in 20 ml RPMI plus 5% FCS plus 10 \muCi 125I-IUdR at 37°C for 5 h, washed three times, and diluted to a final concentration of 5 \times 10^6 cells/ml. 200 \mu1 (10^6 cells) of the cell preparations was injected intravenously. The mice were killed after 8 h and the remaining radioactivity was measured in the lungs, the liver, the spleen, and the kidneys, in a gamma counter (Pharmacia-LKB Biotechnology, Stockholm, Sweden). The arithmetic means and SD in each group (five mice/group) from one representative experiment are given.

In Vitro Cytotoxic Assay with Antibody Plus Complement-Depleted Effector Cell Populations. NK cell activity of syngeneic C57BL/6 mice was boosted by administrating 0.2 ml of a 10 mg/ml preparation of tilorone analogue R10.874 DA (Sigma Chemical Co., St. Louis, MO) per os to each animal on day -1. Tilorone augments NK cell activity by induction of IFN in the host (22). Single cell suspensions of spleens from the tilorone-boosted mice were prepared in tissue culture medium. Aliquots of 20 \times 10^6 cells were incubated with 1.5 ml of mAb for 45 min, washed twice in PBS, and finally diluted in tissue culture medium. Reagents used were rabbit complement (Pd-Freez, 5% CO2, washed twice in PBS, and finally diluted in tissue culture medium. Aliquots of 20 \times 10^6 cells were injected intraperitoneally 24 h before the test) syngeneic C57BL/6 mice were used. 20 \times 10^6 tumor cells (RMA, RMA-S, and RMA-S.mtp2, respectively) were incubated in 20 ml RPMI plus 5% FCS plus 10 \muCi 125I-IUdR at 37°C for 5 h, washed three times, and diluted to a final concentration of 5 \times 10^6 cells/ml. 200 \mu1 (10^6 cells) of the cell preparations was injected intravenously. The mice were killed after 8 h and the remaining radioactivity was measured in the lungs, the liver, the spleen, and the kidneys, in a gamma counter (Pharmacia-LKB Biotechnology, Stockholm, Sweden). The arithmetic means and SD in each group (five mice/group) from one representative experiment are given.

Radiolabeled, intravenously inoculated transfecant cells survived well in the lungs (Fig. 2) and liver (data not shown) in normal as well as in NK1.1-treated animals, whereas the mutant cells without mtp2 survived only in the latter. The mtp2 transfection also rendered RMA-S resistant to NK cells in vitro (Fig. 3, A and B).

To our knowledge, this change in tumor cell rejectability is the first direct demonstration that a MHC-encoded transporter gene can have a decisive influence on disease progression in vivo. The result may seem surprising, since one might

![Figure 1](image-url)

**Figure 1.** In vivo outgrowth of RMA (○), RMA-S (△), and RMA-S.mtp2 (△) lymphoma cells in syngeneic C57BL/6 mice. (A) 10^6 tumor cells injected subcutaneously and outgrowth monitored by palpations twice weekly. (B) 10^6 tumor cells injected subcutaneously and outgrowth monitored by palpations twice weekly. Means of three separate experiments, with a total of 12-15 mice in each group are given.

Results and Discussion

RMA-S.mtp2 had a somewhat lower growth rate in vitro as compared with RMA-S (data not shown). Despite this, the transfecant had acquired a malignant phenotype. In contrast with the mutant, the mtp2 transfecant grew out in the majority (70%) of animals after subcutaneous inoculation of 10^6 cells (Fig. 1 A). The mutant failed to grow in more than 20% of the animals even after a 100-fold increase of the inoculum to 10^6 cells, a dose that resulted in 100% tumor outgrowth of transfecant cells (Fig. 1 B). The difference in tumorigenicity between mutant and transfecant cells persisted in T cell–deficient nude mice (data not shown), but disappeared in animals treated with anti-NK.1.1 antibodies (Table 1). The tumorigenicity of the transfecant was due to its capacity to escape from rapid elimination by NK cells.

| Table 1. In Vivo Outgrowth of RMA-S and RMA-S.mtp2+ Lymphoma Cells in Untreated and NK1.1 Pretreated Syngeneic C57BL/6 Mice |
|-------------------|------------------|-------------------|
|                   | Host untreated   | Host NK1.1 pretreated |
| Cells injected    | RMA-S | RMA-S.mtp2 | RMA-S | RMA-S.mtp2 |
| Separate flanks   | 0/8   | 7/8     | 8/8   | 8/8     |
| Separate animals  | 3/13  | 15/15   | 14/15 | 15/15   |

In vivo outgrowth of 10^6 RMA-S and RMA-S.mtp2+ lymphoma cells in untreated and NK.1 antibody-treated syngeneic C57BL/6 mice after subcutaneous inoculations.
Figure 2. Short-term in vivo survival of $^{125}$-UdR-labeled RMA, RMA-S, and RMA-S.mtp2* lymphoma cells in untreated and NKI.1 antibody-treated syngeneic C57BL/6 mice. $10^6$ tumor cells were injected intravenously and the remaining radioactivity in the lungs was determined after 8 h. The arithmetic means and SD in each group from one representative experiment are given.

Figure 3. Sensitivity of RMA, RMA-S, and RMA-S.mtp2* to syngeneic C57BL/6 and allogeneic BALB/c NK effector cells. (A) Syngeneic C57BL/6 (○) and allogeneic BALB/c (○) NK effector cells were tested at effector/target ratios 300, 100, and 33:1. (B) Syngeneic C57BL/6 spleen effector cells were pretreated either with C'-(●), anti-CD4+ C'-(△), anti-CD8+ C'-(▲), anti-asialo-GM1+ C'-(□), or left untreated (○), and tested at effector/target ratios 300, 100, and 33:1.

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References

1. Zinkernagel, R.M., and P.C. Doherty. 1979. MHC restricted cytotoxic T cells: studies on the biological role of polymorphic major transplantation antigens determining T cell restriction-specificity, function and responsiveness. Adv. Immunol. 27:51.

2. Townsend, A.R.M., and H. Bodmer. 1989. Antigen recognition by class I restricted T lymphocytes. Annu. Rev. Immunol. 7:601.

3. Rötzschke, O., and K. Falk. 1991. Naturally-occurring peptide antigens derived from the MHC class I-restricted processing pathway. Immunol. Today. 12:447.

4. Deverson, E.V., I.R. Gow, J. Coadwell, J.J. Monaco, G.W. Butcher, and J.C. Howard. 1990. MHC class II region encoding proteins related to the multidrug resistance family of transmembrane transporters. Nature (Lond.). 348:738.

5. Trowsdale, J., I. Hanson, I. Mockridge, S. Beck, A. Townsend, and A. Kelly. 1990. Sequences encoded in the class II region of the MHC related 'ABC' superfamily of transporters. Nature (Lond.). 348:741.

6. Monaco, J.J., S. Cho, and M. Attaya. 1990. Transport protein genes in the murine MHC: possible implications for antigen processing. Science (Wash. DC). 250:1723.

7. Spies, T., M. Bresnahan, S. Bahram, D. Arnold, G. Blance, E. Mellins, D. Pious, and R. DeMars. 1990. A gene in the human major histocompatibility complex class II region controlling the class I antigen presentation pathway. Nature (Lond.). 348:744.

8. Powis, S.J., A.R.M. Townsend, E.V. Deverson, J. Bastin, G.W. Butcher, and J.C. Howard. 1991. Restoration of antigen presentation to the mutant cell line RMA-S by an MHC-linked transporter. Nature (Lond.). 354:528.

9. Kelly, A., S.H. Powis, L.A. Kerr, I. Mockridge, T. Elliott, J. Bastin, B. Uchanska-Ziegler, A. Ziegler, J. Trowsdale, and A. Townsend. 1992. Assembly and function of the two ABC transporter proteins encoded in the human major histocompatibility complex. Nature (Lond.). 355:641.

10. Spies, T., V. Cerundolo, M. Colonna, P. Cresswell, A. Townsend, and R. DeMars. 1992. Presentation of viral antigen by MHC class I molecules is dependent on a putative peptide transporter. Nature (Lond.). 355:644.

11. Attaya, M., S. Jameson, C.K. Martinez, E. Hermel, C. Aldrich, J. Forman, K. Fischer Lindahl, M.J. Bevan, and J. Monaco. 1992. Ham-2 corrects the class I antigen-processing defect in RMA-S cells. Nature (Lond.). 355:647.

12. Powis, S.J., E.V. Deverson, W.J. Coadwell, A. Ciruela, N.S. Huskisson, H. Smith, G.W. Butcher, and J.C. Howard. 1992. Effect of polymorphism of an MHC-linked transporter on the peptides assembled in a class I molecule. Nature (Lond.). 357:211.

13. Townsend, A.C., Öhlin, H.G. Ljunggren, L. Foster, and K. Karre. 1989. Association of class I major histocompatibility heavy and light chains induced by viral peptides. Nature (Lond.). 340:443.

14. Ljunggren, H.G., N.J. Stam, C. Öhlin, J.J. Neefjes, P. Högblad, M.T. Hemels, J. Bastin, T.N.M. Schumacher, A. Townsend, K. Karre, and H.L. Ploegh. 1990. Empty MHC class I molecules come out in the cold. Nature (Lond.). 346:476.

15. Öhlin, C., J. Bastin, H.G. Ljunggren, L. Foster, E. Wolpert, G. Klein, A.R.M. Townsend, and K. Karre. 1990. Resistance to H-2 restricted but not allo H-2 specific graft and cytotoxic T lymphocyte responses in lymphoma mutant. J. Immunol. 145:52.

16. Esquivel, F., J. Yewdell, and J. Bennick. 1992. RMA/S cells present endogenously synthesized cytotoxic proteins to class I-restricted cytotoxic T lymphocytes. J. Exp. Med. 175:163.

17. Hosken, N.A., and M.J. Bevan. 1992. An endogenous antigenic peptide bypasses the class I antigen presentation defect in RMA-S. J. Exp. Med. 175:719.

18. Rötzschke, O., K. Falk, S. Faath, and H.-G. Rammensee. 1991. On the nature of peptides involved in T cell alloreactivity. J. Exp. Med. 174:1059.

19. Yang, Y., K. Fröba, J. Chambers, J.R. Waters, L. Wu, T. Spies, and P.A. Peterson. 1992. Major histocompatibility complex (MHC)-encoded HAM2 is necessary for antigenic peptide loading onto class I MHC molecules. J. Biol. Chem. 267:11669.

20. Kärre, K., H.G. Ljunggren, G. Piontek, and R. Kiessling. 1986. Selective rejection of H-2-deficient lymphoma variants suggests alternative immune defence strategy. Nature (Lond.). 319:675.

21. Ljunggren, H.-G., and K. Karre. 1985. Host resistance directed selectively against H-2-deficient lymphoma variants. Analysis of the mechanism. J. Exp. Med. 162:1745.

22. Gidlund, M., A. Örn, H. Wigzell, A. Senik, and I. Gresser. 1978. Enhanced NK cell activity in mice injected with interferon inducers. Nature (Lond.). 273:759.

23. Julius, M.H., E. Simpson, and L.A. Herzenberg. 1973. A rapid method for the isolation of functional thymus-derived murine lymphocytes. Eur. J. Immunol. 3:645.

24. Ljunggren, H.G., C. Öhlin, P. Högblad, T. Yamasaki, G. Klein, and K. Karre. 1988. Affenprim and efferent cellular interactions in natural resistance directed against MHC class I deficient tumor grafts. J. Immunol. 140:671.

25. Ljunggren, H.-G., T. Yamasaki, P. Collins, G. Klein, and K. Karre. 1988. Selective acceptance of MHC class I-deficient tumor grafts in the brain. J. Exp. Med. 167:730.

26. Högblad, P., R. Glas, C. Öhlin, H.-G. Ljunggren, and K. Karre. 1991. Alteration of the natural killer repertoire in H-2 transgenic mice: specificity of rapid lymphoma cell clearance determined by the H-2 phenotype of the target. J. Exp. Med. 174:327.

27. Öhlin, C., G. Kling, P. Högblad, M. Hansson, G. Scangos, C. Bieberich, G. Jay, and K. Karre. 1989. Prevention of allogeneic bone marrow graft rejection by H-2 transgene in donor mice. Science (Wash. DC). 246:666.

28. Glas, R., K. Sturmbohl, G.J. Hämmerling, K. Karre, and H.-G. Ljunggren. 1992. Restoration of a Tumorigenic Phenotype by β2-microglobulin transfection to EL-4 mutant Cells. J. Exp. Med. 175:843.

29. Shimizu, Y., and R. DeMars. 1989. Demonstration by class I gene transfer that reduced susceptibility of human cells to natural killer cell mediated lysis is inversely correlated with HLA class I antigen expression. Eur. J. Immunol. 19:447.

30. Quillet, A., F.R. Presse, C. Marchiol-Fourniquelt, A. Harel-Bellan, M. Benbunan, H. Ploegh, and D. Fradelizi. 1988. Increased resistance to non-MHC-restricted cytotoxicity related to HLA-A, B expression: direct demonstration using β2-microglobulin transfected Daudi cells. J. Immunol. 141:17.

31. Storkus, W.J., J. Alexander, J.A. Payne, J.R. Dawson, and P. Cresswell. 1989. Reversal of natural killing susceptibility in target cells expressing transfected class I HLA genes. Proc. Natl. Acad. Sci. USA. 86:2361.

32. Sturmbohl, K., and G.J. Hämmerling. 1990. Reconstitution of H-2 class I expression by gene transfection decreases susceptibility to natural killer cells of an EL-4 class I loss variant.
Eur. J. Immunol. 20:171.

33. Ljunggren, H.G., K. Sturmhöfel, E. Wolpert, G.J. Hämmerling, and K. Kärre. 1990. Transfection of β2-microglobulin restores IFN-mediated protection from NK cell lysis in YAC-1 lymphoma variants. J. Immunol. 145:380.

34. Chadwick, B.S., and R.G. Miller. 1992. Hybrid resistance in vitro. Possible role of both class I and self peptides in determining the level of target cell sensitivity. J. Immunol. 148:2307.

35. Elliott, T. 1991. How do peptides associate with MHC class I molecules? Immunol. Today. 11:386.

36. Townsend, A., T. Elliott, V. Cerundolo, L. Foster, B. Barber, and A. Tse. 1990. Assembly of MHC class I molecules analyzed in vitro. Cell. 62:285.

37. Schumacher, T.N., M.T. Heemels, J.J. Neefjes, W.M. Kast, C.J.M. Melief, and H.L. Ploegh. 1990. Direct binding of peptide to empty MHC class I molecules on intact cells and in vitro. Cell. 62:563.

38. Rock, K.L., S. Gamble, L. Rothstein, C. Gramm, and B. Benacerraf. 1991. Dissociation of β2-microglobulin leads to the accumulation of a substantial pool of inactive class MHC heavy chains on the cell surface. Cell. 65:611.

39. Rock, K.L., C. Gramm, and B. Benacerraf. 1991. Low temperature and peptides favor the formation of class I heterodimers on RMA-S cells at the cell surface. Proc. Natl. Acad. Sci. USA. 88:4200.