

Suppl. Fig. 1 HAM rat disease



Suppl. Fig. 2 Histopathology of affected thoracic spinal lesion. A and C are HE staining and B and D are KB-PAS staining. A high power maginification figure is inserted in D.



Suppl. Fig. 3 Diseases developed in env-pX rats. A and B: Gross (A) and microscopic (B) findings of the affected joints. C: Necrotizing arteritis. D: Myocarditis.
 E: Myositis. F: Dermatitis. G: Sialoadenitis.



Suppl. Fig. 4 Detection of apoptotic death of thymocytes in the thymic cortex of env-pX rats by TUNEL. Atrophied thymus (A) and non-atrophied thymus (B) in env-pX rat. C: Thymus of age-matched non-transgenic control rat.



Suppl. Fig. 5 Immunohistochemistry of infiltrating lymphocytes in the diseased heart of env-pX rat. A: with anti-CD4 antibody, B: with anti-CD8 antibody.



Suppl. Fig. 6 Dermatitis developed in a lethally irradiated WKAH rat reconstituted by BMC from env-pX rats. Macro- (A) and microscopic findings (B).



Suppl. Fig. 7 Necrotizing arteritis in lethally irradiated WKAH rats reconstituted by env-pX SC. A and E: HE staining. B: EVG staining. C and F: PTAH staining. D: immunostaining for CD3.



Suppl. Fig. 8 Necrotizing arteritis in a WKAH rat with an env-pX thymus framework.
A: HE staining. B: EVG staining. Disruption of elastic fibers is evident (arrows). C: PTAH staining. Fibrinoid degeneration of the arterial wall is evident (arrow). D: Detection of the pX gene in DNA extracted from microdissected samples by nested PCR. Lane 1: ladder marker. Lane 2: env-pX lymph node cells as positive control. Lane 3: sample from lymphocytes accumulating at the arteritis lesion in a WKAH rat with an env-pX thymus framework. An expected molecular size of the nested PCR product is 90bp.



Suppl. Fig. 9 Titration of ATF-1DN/BB (A-D) and lacZ/BB (E-H). The undiluted (A and E) and diluted (1:10, B and F; 1:100, C and G; 1:1000, D and H) vector solutions were added to BHK cells (1ml/2x10⁵ cells). Twenty-four h later, cells were stained for anti-ATF-1 antibody (A-D) and for β-galactosidase activity (E-H). The percentage of positive cells was plotted (I). Open and closed circles represent results from A-D and from E-H, respectively.



Suppl. Fig. 10 Susceptibility to the Sindbis virus vector. BHK (A) and rat joint fibroblastic cells (B) were infected with lacZ/BB ($lml/2x10^5$ cells), respectively. Twenty-four h later, cells were stained for β -galactosidase activity.



Suppl. Fig. 11 Development of thymoma in lck-pX rats. A: A lethally expanded tumor in the anterior mediastinum (arrowhead) of a Tg38 male rat. B: Cross section of the tumor. C: Microscopic finding of the tumor (HE staining).
D: A small tumor (arrowhead) locates in the medulla in a 15 week old lck-pX rat before development of any clinical symptoms (HE staining).



Suppl. Fig. 12 Immunohistochemistry and ultrastructure of the thymoma. A: Immunostaining using anti-Tax monoclonal antibody. B: Immunostaining using a monoclonal anti-cytokeratin antibody. C: Electron microscopic findings of tumor cells. D: Desmosomes between tumor cells (arrowhead) and tonofibrils (arrow) in the cytoplasm are clearly evident.



Suppl. Fig. 13 Development of thymomas in normal F344 (non-transgenic recipient) rats after being given total BMC of lck-pX rats. A: A large tumor (arrowhead) in the anterior mediastinum of a recipient rat. B: Microscopically, the tumor resembling the lck-pX thymoma. C and D: Immunostaining using anti-cytekeratin (C) and anti-Tax monoclonal antibodies (D).



Suppl. Fig. 14 Distribution of the UEA-1 positive cells in the lck-pX rat thymus (A) and the lck-pX rat thymoma (B).



Immunocytochemical staining

Suppl. Fig. 15 UEA-1 expression of the BMMC in the lck-pX rat. A: Flow cytometry revealed about 7.5% strongly UEA-1 positive cells in the lck-pX BMC. NC means BMC without UEA-1 staining as a negative control. B: Immunocytochemistry of UEA-1 positive cells (brown) in the lck-pX BMC.



Suppl. Fig. 16 Development of mammary carcinoma in H2-pX rats. Gross (A) and microscopic (B) findings of the mammary carcinoma. B: HE stainig.



Suppl. Fig. 17 Immunohistochemistry in Harderian gland of ETR5 rats, using an anti-Env peptide antibody.

A and B: ETR5 rat, C and D: non-transgenic control rat. Harderian glands (A and C) and ducts of the glands (B and D) are shown. Bars indicate 100μ m.