



(11) **EP 1 717 244 B1**

(12) **EUROPEAN PATENT SPECIFICATION**

(45) Date of publication and mention of the grant of the patent:  
**04.06.2014 Bulletin 2014/23**

(51) Int Cl.:  
**C07K 7/08 (2006.01) C07K 16/18 (2006.01)**  
**C07K 16/40 (2006.01) A61K 38/10 (2006.01)**  
**A61K 38/17 (2006.01) A61P 35/02 (2006.01)**

(21) Application number: **04807007.2**

(86) International application number:  
**PCT/JP2004/018646**

(22) Date of filing: **14.12.2004**

(87) International publication number:  
**WO 2005/056594 (23.06.2005 Gazette 2005/25)**

(54) **AKT ACTIVITY SPECIFICALLY INHIBITING POLYPEPTIDE**

POLYPEPTID, DAS DIE AKT-AKTIVITÄT SPEZIFISCH HEMMT

POLYPEPTIDE INHIBANT SPECIFIQUEMENT L'ACTIVITE D'AKT

(84) Designated Contracting States:  
**DE FR GB IT**

(30) Priority: **15.12.2003 JP 2003416556**  
**28.04.2004 JP 2004134583**

(43) Date of publication of application:  
**02.11.2006 Bulletin 2006/44**

(73) Proprietor: **Japan Science and Technology Agency**  
**Kawaguchi-shi,**  
**Saitama 332-0012 (JP)**

(72) Inventors:  
• **NOGUCHI, Masayuki**  
**Sapporo-shi,**  
**Hokkaido 064-0811 (JP)**  
• **OKADA, Futoshi**  
**Yonagoshi,**  
**Tottori 683-0834 (JP)**  
• **HIROMURA, Makoto**  
**Fukuoka-shi,**  
**Fukuoka 815-0047 (JP)**

(74) Representative: **Williams, Aylsa**  
**D Young & Co LLP**  
**120 Holborn**  
**London EC1N 2DY (GB)**

(56) References cited:  
**WO-A-00/55169 WO-A-96/13514**  
**WO-A-98/53050 WO-A1-96/13514**  
**JP-A- 2002 500 649**

- **FRENCH SAMUEL W ET AL: "A modeled hydrophobic domain on the TCL1 oncoprotein mediates association with AKT at the cytoplasmic membrane." BIOCHEMISTRY 21 MAY 2002, vol. 41, no. 20, 21 May 2002 (2002-05-21), pages 6376-6382, XP002453345 ISSN: 0006-2960**
- **KÜNSTLE GERALD ET AL: "Identification of Akt association and oligomerization domains of the Akt kinase coactivator TCL1." MOLECULAR AND CELLULAR BIOLOGY MAR 2002, vol. 22, no. 5, March 2002 (2002-03), pages 1513-1525, XP002453346 ISSN: 0270-7306**
- **MING XIU-FEN ET AL: "Rho GTPase/Rho kinase negatively regulates endothelial nitric oxide synthase phosphorylation through the inhibition of protein kinase B/Akt in human endothelial cells." MOLECULAR AND CELLULAR BIOLOGY DEC 2002, vol. 22, no. 24, December 2002 (2002-12), pages 8467-8477, XP002453347 ISSN: 0270-7306**
- **CUI TAI-XING ET AL: "Angiotensin II subtype 2 receptor activation inhibits insulin-induced phosphoinositide 3-kinase and Akt and induces apoptosis in PC12W cells." MOLECULAR ENDOCRINOLOGY (BALTIMORE, MD.) SEP 2002, vol. 16, no. 9, September 2002 (2002-09), pages 2113-2123, XP002453348 ISSN: 0888-8809**
- **NARDUCCI M G ET AL: "THE MURINE TCLL ONCOGENE: EMBRYONIC AND LYMPHOID CELL EXPRESSION" ONCOGENE, BASINGSTOKE, HANTS, GB, vol. 15, no. 8, 1997, pages 919-926, XP008070759 ISSN: 0950-9232**

Note: Within nine months of the publication of the mention of the grant of the European patent in the European Patent Bulletin, any person may give notice to the European Patent Office of opposition to that patent, in accordance with the Implementing Regulations. Notice of opposition shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).

**EP 1 717 244 B1**

- LAINE JARMO ET AL: "Differential regulation of Akt kinase isoforms by the members of the TCL1 oncogene family." THE JOURNAL OF BIOLOGICAL CHEMISTRY 1 FEB 2002, vol. 277, no. 5, 1 February 2002 (2002-02-01), pages 3743-3751, XP002453349 ISSN: 0021-9258
- MARTELLI A M ET AL: "A new selective AKT pharmacological inhibitor resuces resistance to chemotherapeutic drugs, TRAIL, all-trans-retinoic acid, and ionizing radiation of human leukemia cells" LEUKEMIA, MACMILLAN PRESS LTD, US, vol. 17, no. 9, September 2003 (2003-09), pages 1794-1805, XP002990091 ISSN: 0887-6924
- NOGUCHI MASAYUKI ET AL: "Proto-oncogene TCL1: more than just a coactivator for Akt." THE FASEB JOURNAL : OFFICIAL PUBLICATION OF THE FEDERATION OF AMERICAN SOCIETIES FOR EXPERIMENTAL BIOLOGY AUG 2007, vol. 21, no. 10, August 2007 (2007-08), pages 2273-2284, XP002453350 ISSN: 1530-6860
- HIROMURA MAKOTO ET AL: "Inhibition of Akt kinase activity by a peptide spanning the betaA strand of the proto-oncogene TCL1." THE JOURNAL OF BIOLOGICAL CHEMISTRY 17 DEC 2004, vol. 279, no. 51, 17 December 2004 (2004-12-17), pages 53407-53418, XP002453351 ISSN: 0021-9258
- VIRGILIO L. ET AL.: 'identification of the TCL1 gene involved in T-cell malignancies' PROC. NATL. ACAD. SCI. USA vol. 91, no. 26, 1994, pages 12530 - 12540, XP002953421
- NARDUCCI M.G. ET AL.: 'The murine TcII oncogene: embryonic and lymphoid cell expression' ONCOGENE vol. 15, no. 8, 1997, pages 919 - 926, XP008070759

**Description****Technical Field**

5 **[0001]** The present invention relates to a polypeptide specifically inhibiting the activity of serine threonine kinase Akt (Protein Kinase B) ; a DNA encoding the peptide; a specific inhibitor of Akt activity or an antitumor agent, which contains the polypeptide as an active ingredient; and the like.

**Background Art**

10 **[0002]** Akt kinase (Protein Kinase B hereinafter referred to as Akt) is a serine threonine phosphorylation enzyme that was found in the beginning of the 1990s one after the other by following the homology between virus v-Akt. To date, it has been identified that there are three subtypes of the Akt. These molecules have around 80% homology, and have attracted attention from the beginning in relation to oncogenic transformation. In particular, it has been found and attracted  
15 attention that they play a central role to inhibit apoptosis in intracellular cytokine signaling (Genes & Dev., 13:2905-2927, 1999; Annu. Rev. Biochem, 67:481-507, 1998; Biochem.J, 335: 1-13, 1998).

**[0003]** This Akt has an approximate molecular weight of 57 kD, is selectively bound to inositol phosphate in the pleckstrin homology domain (PH domain) ; and has a function at the N-terminal, which mainly plays a role to define the localization to cell membranes. It also has a phosphorylation kinase domain at the c-terminal. It has been speculated  
20 that the binding of PIP3 and the like to PH domain by a signal from Phosphatidylinositol 3-kinase (PI3K) to translocate AKT molecules onto the membrane, and to alter the tertiary structure of Akt are involved in the activation.

**[0004]** Phosphorylations of the both two amino acids, threonine 308 (Thr308) and serine473 (Ser473) have been considered to be essential for Akt activation. While it has been known that Thr308 is phosphorylated by phosphoinositide dependent kinase (PDK1), the phosphorylation process of Ser473 has not been fully elucidated; it has been only speculated  
25 the possibility that some of uncertain molecules such as integrin linked kinase (ILK) or PDK2 are involved in the phosphorylation process. Further, recently the possibility of autophosphorylation for Ser473 phosphorylation has been reported.

**[0005]** It has known that the activated Akt promotes the phosphorylation of the molecules involved in the apoptosis inhibition. An amino acid sequence in the vicinity of serine/threonine, which is phosphorylated by this Akt, has been known as RXRXXS/T (J. Biol. Chem., 275: 36108-36115). The molecule such as BAD.Caspase 9.FKHRI (forkhead  
30 transcription factor) has the above amino acid sequence, and has been known as a substrate of Akt under physiological conditions. Inactive BAD is phosphorylated by Akt; binds to 14-3-3 protein in a phosphorylation-dependent manner; and liberates proteins such as active, Bcl-2 or Bcl-XL, which have apoptosis inhibitory action. It is thought that Akt plays a central role for apoptosis inhibitory control through these known functions and unexplained various targets (Cell,  
35 96:857-868, 1999).

**[0006]** As described above, serine/threonine kinase Akt has a function to phosphorylate serine or threonine residue of intracellular protein specifically, and plays a role to mediate signal transduction to multi-organelle by the phosphorylation function. Further, the phosphorylation function of the Akt regulates a variety of intracellular mechanisms, and involved  
40 in the regulation of the various intracellular mechanisms such as, mitosis, cell growth, cell differentiation, control of lipid metabolism, immune response, inflammatory response, control of glycogen metabolism. At the same time, this means that the phosphorylation function of the Akt is involved in a wide range of various diseases and disorders such as cancer, obesity, autoimmune injury, inflammation and diabetes (type 2).

**[0007]** Recently, it has been reported that Akt activation is involved in breast cancer, lung cancer, prostate cancer, ovarian cancer, or hematological malignancies such as leukemia and lymphoid tumor (Annu. Rev. Biochem. 68, 965,  
45 1999). As Akt activity is raised in these malignancies, Akt activation is considered to be a cause of these malignancies. In recent days, these serine/threonine kinase activities are modulated by using short peptides that are derivatives of the HJ loop of a serine/threonine kinase, with which treatments for diseases and disorders as described above have been attempted (published Japanese translation of PCT international publication No: 2002-500649).

**[0008]** On the one hand, TCL1 has been known as a protooncogene. TCL1 was noted that the activity is raised in human T cell prolymphocytic leukemia (T-PLL), and has been known to date that there are three similar subtypes (TCL1,  
50 MTCP1, TCL1b) (Oncogene, 8:2475-2483, 1993; Proc. Natl. Acad. Sci. USA, 91:12530-12534, 1994). It has been known that the expression was activated by translocation of these loci: 14q, 32,  $\chi$ 28 to T cell receptor loci to develop human leukemia (T-PLL). However, it is a small protein of 13-16kD, and does not have the specific function structure that has been known to date, whose function has so far not been elucidated.

**[0009]** The expression of these molecules is comparatively limited under physiological conditions. TLC1 expression is limited to the lymphoid cells of T cells in an early differentiation (CD3-/CD4-/CD8-), and various B cells before the plasma cell differentiation. Further, although the details in the expression of MTCP1 under physiological conditions are unknown, it was identified from the recent analysis results of gene expression that the expression are induced in activated

T cells TCL1b is a recently cloned molecule and exists in the immediate vicinity of TCL locus. It is thought that there are five subtypes in mouse and is only one subtype in human. It has been reported that this gene expression has distinctly high expression in germinal cells in an early differentiation.

**[0010]** Genes of TCL1 are cloned, and a base sequence of 342 and an amino acid sequence of 113 are elucidated (US Patent No: 5,985,598).

**[0011]** However, functions of TCL1 have been unknown at all to date. The present inventors found that protooncogene TCL1 binds to Akt, from the search result of the protein molecules binding to Akt with the use of a library derived from human B cell by two-hybrid screen using a yeast to elucidate the process of Akt activation. That is, the present inventors indicated that TCL1 is bound to Akt, multimerized, and Akt of the multimer is activated; and found that TCL1 is the Akt co-activator promoting the Akt activation (Mol.Cell, 6:395-407, 2000). Further, the present inventors elucidated that TCL1 facilitated the Akt-mediated cell division, inhibition of apoptosis and the like, and is a predisposing factor for leukemia, tumor of human lymphatic system and the like. In subsequent studies, by coimmunoprecipitation assay using recombinant proteins in and out of cells, the present inventors indicated that TCL1 facilitated the polymerization among heterogeneous Akt molecules and phosphorylation of serine472/473 residues of Akt among heterogeneous Akt molecules; and elucidated the molecular mechanism that TCL1 activates the Akt (J. Biological Chemistry, 277[5], 3743-3751, 2002)

**[0012]** Further, the present inventors prepared an amino acid random library of TCL1 oncogene by applying PCR method and identified the amino acid sites required for the binding of Akt to TCL1 and the polymerization of TCL1, and moreover identified the mutated form of TCL1 which lacks dimerization or binding potency to Akt of TCL1. In addition, it was identified that the mutated form of TCL1 lacks Akt activation potency (both in vitro and in vivo), and loses various types of functions in TCL1 such as stabilization of mitochondrial outer membrane, inhibition of apoptosis, and nuclear translocation of Akt (Molecular and Cellular Biology, 22[5], 1513-1525, 2002). D16 and I74 have been identified as amino acid residues mediating the association of TCL1 with Akt. The BC-sheet of TCL1 appears to be essential for TCL1 homodimerisation. The present inventors found that protooncogene TCL1, whose function was unknown so far, is a co-activator of Akt, and is essential for the binding to Akt and the polymerization of TCL1-to-TCL1 when Akt activates.

**[0013]** it is known that the TCL1 protein comprises two discontinuous amino acid sections, amino acids 15-21 and 72-80, and that this protein mediates association with AKT at the cytoplasmic membrane. Site-directed mutations of this region do not affect TCL1 secondary structure but do disrupt AKT interactions. Therefore this amino acid region, which is conserved in other members of the TCL1 oncoprotein family such as TCL1B and MTCP1, may contain a novel AKT binding domain (Biochemistry, 41:6376-6382, 2002).

Patent document 1: Published Japanese translation of PCT international publication No. 2002-500649'

Patent document 2: Specification of U.S. Patent No. 5,985,598

Nonpatent document 1: Genes & Dev., 13: 2905-2927, 1999

Nonpatent document 1: Annu. Rev.Biochem, 67: 481-507, 1998

Nonpatent document 1: Biochem.J, 335: 1-13, 1998

Nonpatent document 1: J. Biol. Chem., 275: 36108-36115

Nonpatent document 1: Cell, 96: 857-868, 1999

Nonpatent document 1: Oncogene, 8:2475-2483,1993

Nonpatent document 1: Proc. Natl. Acad. Sci. USA, 91: 12530-12534, 1994

Nonpatent document 1: Mol.Cell, 6:395-407, 2000

Nonpatent document 1: J. Biological Chemistry, 277[5], 3743-3751, 2002

Nonpatent document 1: Molecular and Cellular Biology, 22[5], 1513-1525, 2002

Nonpatent document 1: Biochemistry, 41: 6376-6382, 2002

#### Disclosure of the Invention

Object to be solved by the present invention

**[0014]** An object of the present invention is to provide a polypeptide specifically inhibiting the activity of serine threonine kinase Akt (Protein Kinase B); a DNA encoding the polypeptide; an antibody specifically binding to the polypeptide; a specific inhibitor of Akt activity or an antitumor agent, which contains the polypeptide as an active ingredient; and the like.

Means to solve the object

**[0015]** The present inventors elucidated that protooncogene TCL1, whose function had been unknown at all, directly binds to Akt that is involved in human malignancy and the like, promotes the activation of Akt, that is, is the co-activator of Akt, moreover, it is responsible for leukemia, tumor of human lymphatic system and the like. It was indicated that the mutated form of TCL1, which does not bind to Akt, lacks the Akt activation potency and loses various types of functions

in TCL1 such as stabilization of mitochondrial outer membrane, inhibition of apoptosis, and nuclear translocation of Akt. From these serial studies; it was found that the position of an amino acid residue10-24 in the amino acid sequence of TCL1 (human) is the binding site to Akt, and cell growth associated with Akt activation and the like are specifically inhibited by using a polypeptide sequence of the amino acid residue; and the present invention was led to complete.

5 [0016] Further, it is also herein described that similar functions exist in the TCL1B and MTCP1 (human) having the similar functions to those of TCL1, and found that the cell growth associated with Akt activation inhibits at the positions of an amino acid residue8-22 in amino acid sequence of TCL1B (human) and of an amino acid residue 5-19 in amino acid sequence of MTCP1 (human). Furthermore, in the present invention, it was also identified that the cell growth associated with Akt activation was inhibited at the positions of an amino acid residue9-24 in amino acid sequence of mouse TCL1 and of an amino acid residue9-24 in amino acid sequence of rat TCL1. Polypeptide in the present invention competitively inhibits the binding of phsphoinositide (phosphatidylinositol) to Akt.

10 [0017] In other words, the present invention comprises a polypeptide consisting of an amino acid sequence (SEQ ID NO: 1 of the sequence listing) corresponding to the position of an amino acid residue10-24 in amino acid sequence of TCL1 (human). Also herein described is an amino acid sequence (SEQ ID NO: 3 of the sequence listing) corresponding to the position of an amino acid residue8-22 in amino acid sequence of TCL1B (human), an amino acid sequence (SEQ ID NO: 5 of the sequence listing) corresponding to the position of an amino acid residue5-19 in amino acid sequence of MTCP1 (human), an amino acid sequence (SEQ ID NO: 7 of the sequence listing) of the position of an amino acid residue9-24 in amino acid sequence of TCL1 (mouse), and an amino acid sequence (SEQ ID NO: 9 of the sequence listing) of the position of an amino acid residue9-24 in amino acid sequence of TCL1 (rat),and specifically inhibiting Akt activity. DNA (SEQ ID NO: 2, of the sequence listing) encod the polypeptide, of the invention. Also herein described are DNA sequences SEQ ID No: 4, 6, 8 and 10.

15 [0018] Further, the present invention comprises a polypeptide derivative consisting of an amino acid sequence wherein one or several amino acids are deleted, substituted or added in an amino acid sequence of the polypeptides, and specifically inhibiting Akt activity; and a DNA encoding the sequences. Also herein described is DNA that hybridizes with DNA in the sequences under stringent conditions and specifically inhibits Akt activity.. Furthermore, the present invention includes a method for producing a polypeptide of the present invention by incorporating the DNA into an expression vector, constructing a recombinant expression vector, and introducing the recombinant vector into a host cell to be expressed.

20 [0019] Further, the present invention includes an antibody that specifically binds to the polypeptide specifically inhibiting the Akt activity of the present invention; moreover, includes utilization of specific inhibitor of Akt activity that contains a polypeptide of the present invention as an active ingredient, and of antitumor agent that contains the polypeptides as an active ingredient for prevention and treatment of malignancy and the like. Furthermore, the present invention includes a method for specifically inhibiting Akt activity by introducing the DNA encoding a polypeptide of the present invention into living cells to express the polypeptide.

25 [0020] That is, specifically, the present invention relates to (1) a polypeptide specifically inhibiting Akt activity, which consists of an amino acid sequence indicated in SEQ ID NO: 1, of the sequence listing; and (2) a polypeptide consisting of an amino acid sequence wherein one or several amino acids are deleted, substituted or added in the amino acid sequence indicated in SEQ ID NO: 1, of the sequence listing, and specifically inhibiting Akt activity.

30 [0021] The present invention also relates to (3) a gene DNA encoding a following protein (a) or (b): (a) a polypeptide consisting of an amino acid sequence indicated in SEQ ID NO: 1; (b) a polypeptide consisting of an amino acid sequence wherein one or several amino acids are deleted, substituted or added in the amino acid sequence indicated in SEQ ID NO: 1, and specifically inhibiting Akt activity; (4) a DNA consisting of a base sequence indicated in SEQ ID NO: 2; or part or whole of these sequences, and encoding a polypeptide that specifically inhibits Akt activity. Also herein described is (5) a DNA hybridizing with the DNA according to "4" under stringent conditions, and encoding a polypeptide that specifically inhibits Akt activity.

35 [0022] The present invention further relates to (6) a recombinant expression vector, which is constructed by integrating a DNA encoding the polypeptide that specifically inhibits Akt activity according to any one of "3"- "5" into a gene expression vector; (7) a method for producing a polypeptide that specifically inhibits Akt activity wherein the recombinant expression vector according to "6" is introduced into a host cell and expressed; (8) an antibody which is induced by using a polypeptide indicated in SEQ ID NO: 1 of the sequence listing and specifically binds to the polypeptide; (9) the antibody according to "8" wherein the antibody is a monoclonal antibody; and (10) the antibody according to "8" wherein the antibody is a polyclonal antibody.

40 [0023] Also herein are described the polypeptides of SEQ ID NO: 3, 5, 7, and 9. The present invention still further relates to (11) a specific inhibitor of Akt activity, wherein the polypeptide according to "1" or "2" is an active ingredient; (12) the specific inhibitor of Akt activity according to "11", wherein the polypeptid is a sequence of an amino acid residue 10-24 of an amino acid sequence for human TCL1 protein wherein specific inhibition of Akt activity is the inhibition of binding of phosphoinositide to Akt. Also described herein is (13) the specific inhibitor of Akt activity according to "11", wherein the polypeptid is a sequence of an amino acid residue8-22 of an amino acid sequence for human TCL1B protein;

(14) the specific inhibitor of Akt activity according to "11", wherein the polypeptid is a sequence of an amino acid residue5-19 of an amino acid sequence for human MTP1 protein; (15) the specific inhibitor of Akt activity according to "11", wherein the polypeptid is a sequence of an amino acid residue9-24 of an amino acid sequence for mouse TCL1 protein; (16) the specific inhibitor of Akt activity according to "11", wherein the polypeptid is a sequence of an amino acid residue9-24 of an amino acid sequence for rat MTP1 protein; and (17) the specific inhibitor of Akt activity according to any one of "11"-16", wherein specific inhibition of Akt activity is the inhibition of binding of phosphoinositide to Akt.

**[0024]** Furthermore, the present invention relates to (18) an antitumor agent wherein the polypeptide according to "1" or "2" is an active ingredient; (19) the antitumor agent according to "18", wherein the antitumor agent is an agent for prevention or treatment of malignancy; (20) the antitumor agent according to "19", wherein treatment of malignancy is prevention or treatment of breast cancer, lung cancer, leukemia or lymphoid tumor. Also described herein is (21) a method for specifically inhibiting Akt activity by introducing a DNA encoding the polypeptide that specifically inhibits Akt activity according to any one of "3"-5" into living cells to express the polypeptide.

#### Effect of the present invention

**[0025]** Oncogene TCL1 is a co-activator of Akt (serine threonine phosphorylation enzyme: Protein Kinase B), and the TCL1 directly binds to Akt to promote the activation of Akt. In the present invention, polypeptides in the present invention enable its utilization as a specific inhibitor of Akt activity, by finding that the position binding to Akt in the amino acid sequence of the TCL1 is specified, and the polypeptides consisting of an amino acid sequence of the positions of the TCL1, TCL1B and MTCP1 that bind to Akt specifically inhibits the Akt activity. To date, a specific peptide inhibitor of Akt has been unknown, therefore, a polypeptide of the present invention can be expected the utilization thereof as an entirely new Akt activity inhibitor.

**[0026]** Further, Akt is an intracellular signaling molecule that plays a central role for inhibiting apoptosis. Akt is activated in many cancer cells, and apoptosis is impaired. As a result, apoptosis is decreased; cells are abnormally proliferated, and cancer is developed. Akt has been known to be a central molecule regulating apoptosis, and is an important subject of study; however, agents specifically inhibiting a central molecule Akt in apoptosis inhibitory mechanism have not been developed yet. Specific inhibitor of Akt activity in the present invention is involved in mechanism that contributes to cancer development such as apoptosis regulation, and leads to development of agents for prevention or treatment of malignancy in a background of AKT activation by overexpression of TCL1 gene or abnormality of cancer suppressor gene PTEN. As the Akt activity involved in malignancies such as breast cancer, lung cancer, prostate cancer, ovarian cancer, and hematological malignancy such as leukemia or lymphoid tumor; specific inhibitors of Akt activity in the present invention can be used as an antitumor agent (anticancer agent) for prevention or treatment of various human malignancies caused by activation of the Akt kinases.

#### Brief Description of Drawings

##### [0027]

[Figure 1]

Figure 1 is a set of pictures showing 10 clones of amino acid substitution indicating low interaction (8h positive [+]) with Akt that was constructed and observed in experiments of the examples of the present invention, displaying amino acid sequence of TCL1 side-by-side.

[Figure 2]

Figure 2 is a set of pictures showing the results of  $\beta$ -Gal lifting assay with the use of TCL1 mutants (D16G, K30M, Q46R, I74V, or M106V) induced by site-directed mutagenesis and wild-type TCL1, in experiments of the examples of the present invention.

[Figure 3]

Figure 3 is a set of pictures showing the results of quantitative liquid  $\beta$ -Gal assay with the use of TCL1 mutants (D16G, K30M, Q46R, I74V, or M106V) induced by site-directed mutagenesis and wild-type TCL1, in experiments of the examples of the present invention.

[Figure 4]

Figure 4 is a set of pictures showing the results of in vitro kinase assay for wild-type TCL1 to examine the association of Akt with TCL1 homodimer, which is required for TCL1-induced Akt activation, in experiments of the examples of the present invention.

[Figure 5]

Figure 5 is a set of pictures showing the results of in vitro kinase assay for TCL1 mutants (36-38A TCL1) to examine the association of Akt with TCL1 homodimer, which is required for TCL1-induced Akt activation, in experiments of the examples of the present invention.

[Figure 6]

Figure 6 is a picture showing the results of MTT assay with the use of the peptide consisting of the sequence of 10-24 amino acid residues of TLC1 oncogene to identify that the cell growth associated with AKT activation is specifically inhibited, in experiments of the examples of the present invention.

[Figure 7]

Figure 7 is a set of pictures showing the results of western blotting after the binding test for 10/24 peptide with Akt subtypes, Akt1, Akt2 and Akt3 molecules, by using co-immunoprecipitation assays, in experiments of the examples of the present invention.

[Figure 8]

Figure 8 is a set of pictures showing the results of western blotting after the inhibition tests for Akt activation with 10/24 peptide, by Akt kinase assays with the use of GSK (Glycogen Synthesis Kinase 3) as substrates, in experiments of the examples of the present invention.

[Figure 9]

Figure 9 is a set of pictures showing the results of western blotting with the use of various antibodies for the inhibition effects on the AKT activation of 10/24 peptide in mouse QrSP-11 fibrosarcoma cells and control peptide for phosphorylation (serine 473 residue, threonine 308 residue), in experiments of the examples of the present invention.

[Figure 10]

Figure 10 is a set of pictures observed under the microscope showing the results of examination for the inhibition effects on the membrane translocation and AKT activation of 10/24 peptide with the use of 293 cells (ATCC), in experiments of the example of the present invention.

[Figure 11]

Figure 11 is a picture showing the results of examination for the effects on apoptosis of 10/24 peptide with the use of human T cell leukemia cells (T4), in experiments of the examples of the present invention.

[Figure 12]

Figure 12 is a picture showing the results of examination for in vivo anti-tumor effect by 10/24 peptide with the use of transplanted tumor cells, in experiments of the examples of the present invention.

[Figure 13]

Figure 13 is a set of pictures showing the results of observations in macroscopic appearance, H&E staining, TUNEL staining, and Akt activation; of the experimental mouse tumors examined in vivo anti-tumor effect by 10/24 peptide that was collected on the day 9, with the use of transplanted tumor cells, in experiments of the examples of the present invention.

### Beat Mode of Carrying Out the Invention

**[0028]** The present invention comprises polypeptides: an amino acid sequence of amino acid residue 10-24 in human TCL1 oncogene which specifically inhibits Akt activity. Also herein described is an amino acid sequence of amino acid residue8-22 in human TCL1B, an amino acid sequence of amino acid residue 5-19 in human MTCP1, an amino acid sequence of amino acid residue9-24 in mouse TCL1, and an amino acid sequence of amino acid residue9-24 in rat TCL1, which are specifically inhibits Akt activity. The amino acid sequences is indicated in SEQ ID NO: 1. Also herein is described SEQ ID NO: 3, SEQ ID NO: 5, SEQ ID NO: 7 and SEQ ID NO: 9 of the sequence listing. Further, the present invention comprises polypeptide derivatives consisting of an amino acid sequences wherein one or several amino acids are deleted, substituted or added in the amino acid sequences of the polypeptides, and specifically inhibiting Akt activity. DNA sequences encoding a polypeptide that consists of an amino acid sequence of amino acid residue 10-24 in Human TCL1 oncogene. Also herein described are DNA sequences encoding an amino acid sequence of amino acid residue8-22 in human TCL1B, or an amino acid sequence of amino acid residue5-19 in human MTCP1; an amino acid sequence of amino acid residue9-24 in mouse TCL1; or an amino acid sequence of amino acid residue 9-24 in rat TCL1; The sequence of the invention is indicated in SEQ ID NO: 2. Also herein described are SEQ ID NO: 4, SEQ ID NO: 6, SEQ ID NO: 8, and SEQ ID NO: 10 of the sequence listing. The present description includes a DNA encoding a polypeptide that hybridizes with DNA of the sequences under stringent conditions and specifically inhibits Akt activity.

**[0029]** The polypeptide in the present invention can be synthesized by known method for polypeptide synthesis based on the structure of the polypeptide that consists of amino acid sequences indicated in SEQ ID NO: 1 of the sequence listing; and further, can be produced by genetic manipulation with the use of a DNA sequence encoding the polypeptide. A DNA sequence of the entire gene in Human TCL1 oncogene and an amino acid sequence of a protein encoded by the gene are disclosed in US Patent No: 5,985,598, and the sequences of the gene and protein can be accessed from a database of GenBank by accession number: X82240 and CAA57708. In addition, a vector incorporated a full-length gene (cDNA and genome DNA) of TCL1 is respectively deposited at acceptance number: 75923 and 75924 as a deposit of microorganism according to Budapest Treaty on the International Recognition of the Deposit of Microorganisms for the Purposes of Patent Procedure at a depository institution in USA, American Type Culture Collection (ATCC). Further,

a DNA sequence of mouse TCL1 gene and an amino acid sequence of a protein encoded by the gene, a DNA sequence of rat TCL1 gene and an amino acid sequence of a protein encoded by the genes, can be accessed from a database of NCIB by accession numbers NP\_033363 and XP\_345720 respectively.

5 [0030] A DNA sequence of the entire gene in TCL1B and an amino acid sequence of a protein encoded by the gene are disclosed in a reference (Proc. Natl. Acad. Sci. U.S.A., 96(6), 2949-2951, 1999), and can be accessed from a database of NCBI by accession number: AF\_110465. Further, a DNA sequence of the entire gene in MTCP1 and an amino acid sequence of a protein encoded by the gene are disclosed in a reference (Oncogene 8(9), 2475-2483, 1993), can be accessed from a database of NCBI by accession number: BC\_002600.

10 [0031] A polypeptide in the present invention can be obtained by genetic manipulation performing the following step: producing DNA from an information as the above DNA sequences using synthesis method, or slicing out the DNA of the present invention from a source of TCL1 genes such as the above by using restriction enzyme; incorporating the gene into an appropriate expression vector; and introducing the recombinant vector into host cells to express the polypeptide. A variety of polypeptide derivatives in the present invention can be produced by the following steps: producing DNA of a base sequence encoding the polypeptide; constructing an expression vector by using the DNA; and introducing the expression vector into appropriate known host cells to express the polypeptide. Mutation of DNA sequence encoding a variety of polypeptide derivatives can be conducted by known genetic engineering gene mutation procedure.

15 [0032] To produce a polypeptide in the present invention by genetic manipulation, incorporating a DNA encoding the polypeptide into a known expression vector, constructing a recombinant expression vector, and introducing the vector into host cells to express the polypeptide can be performed. Introduction of the recombinant expression vectors into host cells can be performed with the use of appropriate known methods. For example, as for a host cell of prokaryote, E coli, Bacillus subtilis and Pseudomonas strains can be exemplified; as for a vector when using the prokaryote as host cells, a vector of E coli strains or the like such as pUC19, pBR322 and pBR327 can be used; as for a promoter, tryptophan promoter, PL promoter, lac promoter or tac promoter can be exemplified; as for a marker gene, ampicillin-resistant gene, tetracycline-resistant gene or the like can be used.

20 [0033] As for host cells of eukaryotic microorganism, yeast is widely used in general; as for a vector, for example, YRp7 and the like may be used. When using cultured cells of higher animals as host cells, COS cell, CHO cell (Chinese hamster ovary cell) and the like can be used. As for a promoter, for example, adenovirus-2 major late promoter, SV40 early promoter, SV40 late promoter, and a promoter from cytomegalovirus or Rous sarcoma virus; as for a marker gene, for example, neomycin-resistant gene, methotrexate-resistant dihydrofolate reductase (DHR) gene and the like; can be used. Furthermore, as for host cells, an insect cell such as BmN4 cell, Sf9 cell and Sf21 cell can be used.

25 [0034] In the present invention, by using a polypeptide of the present invention such as a polypeptide consisting of an amino acid sequence of amino acid residue10-24 in human TCL1, the binding of phosphoinositide (Phosphatidylinositol) are inhibited, and as a consequence, Akt (protein Kinase B) activity, cell growth, and anti-tumor effects can be obtained. For this peptide, administration as a recombinant protein, and a method of administration using a virus vector or a mammalian expression vector, can be considered. Induction method for peptide by fusion method with TAT peptide (part of HIV virus protein) can be also used. Moreover, electroporation, and intracellular induction methods that are pharmacologically possible, can be used. Peptide modification in the sense of peptide stabilization, PEG (polyethylene Glycol), FCR (FC Receptor), and production of fusion peptide with other peptide, can be used.

30 [0035] The present description includes DNA encoding a polypeptide that hybridizes with a DNA sequence encoding a polypeptide consisting of an amino acid sequence of amino acid residue10-24 in human TCL1 (a base sequence indicated in SEQ ID NO: 2 of the sequence listing). Also described herein is a DNA sequence encoding a polypeptide consisting of an amino acid sequence of amino acid residue8-22 in human TCL1B (a base sequence indicated in SEQ ID NO: 4 of the sequence listing), a DNA sequence encoding a polypeptide consisting of an amino acid sequence of amino acid residue5-19 in human MTCP1 (a base sequence indicated in SEQ ID NO: 6 of the sequence listing), a DNA sequence encoding a polypeptide consisting of an amino acid sequence of amino acid residue9-24 in mouse TCL1 (a base sequence indicated in SEQ ID NO: 8 of the sequence listing), a DNA sequence encoding a polypeptide consisting of an amino acid sequence of amino acid residue9-24 in rat TCL1 (a base sequence indicated in SEQ ID NO: 10 of the sequence listing), under stringent conditions and specifically inhibits Akt activity.

35 [0036] As for the conditions in the following phrase, "hybridizes ... .. under stringent conditions", for example, hybridization at 42°C and washing treatment at 42°C with a buffer containing 1×SSC, 0.1% SDS, and preferably hybridization at 65°C and washing treatment at 65°C with a buffer containing 0.1×SSC, 0.1% SDS can be exemplified. Further, as for factors affecting the stringency of hybridization, there are a variety of factors other than the above temperature conditions; therefore, those in the art can conduct a stringency equivalent to the above exemplified stringency of hybridization using a variety of factors in combination.

40 [0037] The present invention includes an antibody specifically bound to a polypeptide that specifically inhibits the Akt activity of the present invention. As for the antibody, monoclonal antibody and polyclonal antibody can be exemplified. The antibody can be produced by ordinary method with the use of a polypeptide of the present invention as an antigen. It can be considered that the antibody of the present invention specifically inhibits the binding of TCL1, to Akt by specifically

binding to TCL1 at the Akt-binding site. Also described herein is an antibody that specifically inhibits the binding of TCL1B or MTCPI. Further, the antibody of the present invention can be used to detect the diseases involved in TCL1, gene in tissue cells, serum and the like, by using antigen-antibody reaction with TCL1 polypeptide. In immunological measurement by using the antibody of the present invention, for example, known immunological measurement such as RIA assay, ELISA assay and fluorescent antibody method can be used. Further described herein are antibodies that can be used to detect the diseases involved in TCL1B or MTCPI.

**[0038]** In the present invention, a specific inhibitor of Akt activity uses a polypeptide of the present invention as an active ingredient, and an antitumor agent uses the polypeptide as an active ingredient for prevention and treatment of malignancy and the like. To use a polypeptide of the present invention as an active ingredient for specific inhibitor of Akt activity, and for an antitumor agent of prevention and treatment for malignancy and the like, the polypeptide can be used alone or by adding various mixture ingredients: pharmaceutically acceptable ordinary carriers, binding agent, stabilizing agent, excipient, diluent, pH buffer agent, disintegrating agent, solubilizer, solubilizing agent, isotonic agent and the like for formulation. These specific inhibitors of Akt activity and the prevention or treatment agent for malignancy and the like can be administered orally or parenterally. That is, ordinary administration formulations to use, for example, powder, granule, capsule, syrup, suspension and the like can be administered orally, or, for example, solution, emulsion, suspension and the like can be administered parenterally by injection, and moreover, spray formulation can be administered into nostril.

**[0039]** When a polypeptide of the present invention specifically inhibiting Akt activity is used for prevention/treatment of cancer and the like, a polypeptide specifically inhibiting Akt activity can be directly inoculated into cancer cells by using a reagent without cytotoxicities such as Chariot (Active Motif), which forms noncovalent complexes with macromolecules such as proteins, peptides or antibodies, alters structures of polypeptide, and can deliver polypeptide molecules into cells. In addition, the doses can be appropriately selected by depending on disease type, body weight of patient, administration method, and the like. As for a subject of the administration of a specific inhibitor for Akt activity or of an antitumor agent in the present invention, prevention or treatment for various diseases caused by Akt activation can be exemplified, especially prevention or treatment for malignancies such as breast cancer, lung cancer, prostate cancer, ovarian cancer, and hematological malignancy such as leukemia or lymphoid tumor can be exemplified.

**[0040]** In the present invention, by introducing the DNA encoding a polypeptide that specifically inhibits Akt activity of the present invention, into living cells to express the polypeptide, the Akt activity can be specifically inhibited. As for an expression vector in animal cells for introducing the DNA encoding a polypeptide that specifically inhibits Akt activity, into living cells, any vector can be used as long as the DNA encoding the above polypeptide of the present invention is integrated into a vector for animal cells, as for the vector for animal cells, the expression system is not especially limited as long as the DNA encoding the above polypeptide of the present invention can be expressed in host cells. For example, expression systems derived from chromosome, episome and virus; for example, vectors derived from bacterial plasmid, yeast plasmid, papovavirus such as sv40, vaccinia virus, adenovirus, fowlpox virus, pseudorabies virus, lentivirus and retrovirus; vectors derived from bacteriophage, transposon and the combination of those; for example, vectors derived from genetic components of plasmid and bacteriophage such as cosmid and phagemid, can be exemplified.

**[0041]** These expression systems not only cause the expression but also may contain regulatory sequences regulating the expressions. Further, in an expression vector for animal cells of the present invention, liposomes are also included. In these vectors for animal cells, an adenovirus vector is especially preferable in terms of safety and utility. In prevention and treatment for cancer and the like, administering directly at lesion sites (in situ) is preferable, for example, when using an adenovirus expression vector, the vector suspension can be directly inoculated into the lesion site such as cancer tissues. Further, when using a liposome holding the DNA encoding a polypeptide that specifically inhibits the Akt activity of the present invention, the liposome suspension can be directly inoculated into the lesion site such as cancer tissues.

**[0042]** The present invention will be explained in detail in the following by referring to the examples, but the technical scope of the present invention will not be limited to these.

#### Example 1

[Identification of a binding sequence of Akt and TCL1]

**[0043]** The interaction of an amino acid partial mutation TCL1 clone with Akt was examined by using yeast two-hybrid screening and semiquantitative  $\beta$ -Gal assay (MOLECULAR AND CELLULAR BIOLOGY, Mar. 2002, P.1513-1525)

## EP 1 717 244 B1

[TCL1 (human) amino acid random mutation library screening for identification of the Akt-TCL1 binding site]

(Materials and Methods)

### 5 1.TCL1 library

**[0044]** A full length of human TCL1 in pGAD424 (Clontech) was amplified by PCR with the use of a primer comprising 5'-CCACCAAACCCAAAAAAGAGATCGAATTCATG and 5'-ATTCATAGATCTCTGCAGGTCGACGGATCCTCA, and a random TCL1 amino acid library was produced.

10

### 2. Production of amino acid mutant forms of TCL1

**[0045]** An amino acid-substituted mutant form of TCL1 (D16G, K30M, Q46R, 174V, M106V, 36-38A, or 36A/38Δ) was produced by PCR with the use of the following primer, and native and mutated forms were subcloned into the vector expressing pGAD424 (Clontech), pME18SHA (Mol. Cell 6:395-407), or pCMV Flag (Kodak).

15

**[0046]** Primers used are as follows (mutated codons are shown in lower-case letters) : 5'-ATG GCC GAG TGC CCG ACA CTC GGG GAG GCA GTC ACC GAC CAC CCG ggc CGC CTG TGG GCC for D16G; 5'-GTG TAT TTG GAC GAG atg CAG CAC GCC TGG CTG for K30M; 5'-G ATA AAG GAT AGG TTA cgg TTA CGG GTG CTC TTG for Q46R; 5'-CCA AGC CTG CTG CCT gtc ATG TGG CAG CTC TAC for 174V; 5'-ATC ATC GGA TCC TCA GTC ATC TGG CAG CAG CTC GAG AAG cac GTC CTC C for M106V; 5'-CAG CAC GCC TGG CTG gcc gcg gcc ATC GAG ATA AAG GAT and a reverse complementary sequence for 36-38A; and 5'-GCC TGG CTG gcc TTA ATC GAG ATA and a reverse complementary sequence for 36A/38Δ. Mutated positions of an amino acid-substituted mutant form of TCL1 (D16G, K30M, Q46R, 174V, M106V, 36-38A, or 36A/38Δ) are shown in figure 1.

20

### 25 3. Yeast two-hybrid screening

**[0047]** Screening was performed by yeast two-hybrid system for detection of protein interaction to detect interaction of TCL1 with Akt protein.

**[0048]** Y190 cells (Clontech), by using lithium acetate method; and human Akt2 (Akt2/PAS2-1) and TCL1 random library, in accordance with the previous report (Mol. Cell 6:395-407.Proc. Natl. Acad. Sci. USA 94:11534-11539); were expressed in yeast. In the presence of 3-amino-1,2,4-triazole (SIGMA), around 104 clones from cDNA library were screened. β-galactosidase (β-Gal) activity in His<sup>+</sup> colony was measured by using filter-lift assay. Yeast clones were classified into clone categories: 3h-positive [++], 8h-positive [+], and 24h-negative [-] by the intensity of β-Gal. Ten clones were selected from the each category for sequencing of the nucleotide.

30

### 35 4. Quantitative β-Gal assay

**[0049]** Y190 cells (Clontech) were expressed in yeast together with TCL1 of wild-type, D16G, K30M, Q46R, 174V, or M106V by using Akt2/PAS2-1, with the use of pGAD424 (Clontech) vector. TCL1 mutant was produced by using PCR-based site-directed mutagenesis and/or Quikchange kit (Stratagene). Liquid β-Gal assay was performed for quantitative determination of binding intensity by using ONPG (O-nitrophenyl-β-D-galactopyranoside; Sigma) (Mol. Cell 6:395-407). The indicated values were determined by western blotting analysis (GAL4 activation domain antibody [Ab]: Clontech), and normalized by expressing transformant in each yeast.

40

### 45 [Experiment and results]

**[0050]** Random TCL1 library was produced by PCR-mediated random mutation to determine the amino acid residues required for Akt-TCL1 binding. Each occurrence of the substituted DNA nucleotide was 1.4% for dATP, 3.8% for dTTP, 4.0% for dGTP, and 1.4% for dCTP. The total frequency of nucleotide substitution in this library was 2.7%, and the insertion-deletion frequency was 0.09%. The library size was around  $2.5 \times 10^4$ . The substituted positions were dispersed over 90% or more of the total TCL1 molecules in the sequenced 25 sample clones.

50

**[0051]** Next, a yeast two-hybrid screening was performed to examine the interaction of each clone with Akt2. Yeast clones were classified into three categories based on the blue-color intensity expressed in β-Gal lifting assay (β-Gal positive [++] at 3h, β-Gal positive [+] at 8h, and negative [-]). Nucleotide sequences of the 10 clones in each category were determined.

55

**[0052]** A ++clone included wild-type TCL1, or the mutants at P5, P15, D43, L45, P61, M75 and D88 positions; did not affect β-Gal activity, and its residue did not react to Akt interaction. The observed 10 clones of amino acid substitution indicating low interaction (8h positive [+]) with Akt, and amino acid sequence of TCL1 were displayed side-by-side (Fig. 1).

[0053] Deposition of substitution was evidently found in specific residues of these clones. The substitution was found in at least one of amino acids in D16, K30, Q46, 174 and M106, in 9 clones out of 10 clones. Negative clone [-] was not involved in nucleotide insertion, insertion associated with large deletions, structural shift, and/or a large amount of mutants. Therefore, further analysis was eliminated. The present inventors hypothesized that clones showed Akt low interaction must contain an amino acid residue required for Akt-TCL1 interaction. Therefore, each mutant (D16G, K30M, Q46R, I74V or M106V) was produced in TCL1 by using a site-directed mutagenesis method. In D16G and I74V mutants, the bindings to the Akt mutant were resulted in dramatic decrease in D16G and 174V, as shown in  $\beta$ -Gal lifting assay (Fig. 2) and quantitative liquid  $\beta$ -Gal assay (Fig. 3).

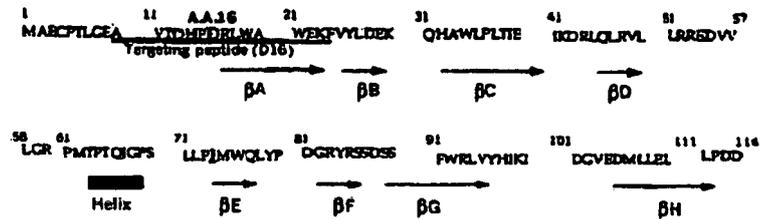
(Association with Akt and TCL1 homodimer, required for TCL1-induced Akt activation)

[0054] Wild-type TCL1 showed an increase in Akt kinase activity in an in vitro kinase assay. This was predominantly correlated with Akt Ser-473 phosphorylation levels. However, D16G TCL1 did not affect the Akt kinase activity even though in a dose escalation experiment (Fig. 4). D16G TCL1 did not affect both GSK-3 $\alpha$  phosphorylation and Akt Ser-473 phosphorylation. D16G TCL1 was evaluated by Akt kinase activity. Similarly, 36-38A TCL1 that binds to Akt but not form homodimer (in an in vitro kinase assay, determined by phosphorylated GSK-3 $\alpha$  and Ser-473 Akt) was not able to enhance the Akt kinase activity (Fig. 5). These indicated that the mutated form of TCL1, which does not bind to Akt, lacks Akt activation potency (both in vitro and in vivo).

(Production of a binding sequence between Akt and TCL1)

[0055] From the result of previous analysis in TCL1 crystal structure (Molecular and Cellular Biology, Mar.2002, p.1513-1525), it was thought that D16 exists at initial position of the first  $\beta$  sheet, and Akt kinase binds on a surface made by the first  $\beta$  sheet and the forth  $\beta$  sheet. Based on these series of studies, it was thought that amino acid binding to Akt in TCL1 protein molecules, amino acid residue sequence 10-24 (Table 1) of TCL1 oncogene in the vicinity of the 16 residue (Asparadic Acid), binds to Akt and can be an inhibitor of Akt activation.

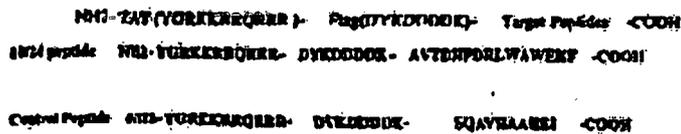
[Table 1]



[0056] Based on the above hypothesis, two peptides, which are a peptide in the vicinity of binding site of the TCL1 and AKT, i.e., a peptide in the vicinity of amino acid residue 10-24 (displays as "10/24") of TCL1, and a control peptide; were produced (Table 2). Peptide were produced by an ordinary peptide synthesizer, and purified by gel filtration or HPLC. Peptides produced by Hokkai BioSystem, and an American company were used.

[Table 2]

Targeting peptides design



## Example 2

[Assay for 10/24 peptide of TCL1]

5 1. Cell growth test by using MTT assay

[0057] Cell growth test was performed by using MTT assay. That is, in experiments of cell growth by using WST-8 reagent [2-(2-methoxy-4-nitrophenyl)-3-(4-nitrophenyl)-5-(2,4-disul fophenyl)-2H-tetrazolium, monosodium salt] (347-07621, Dojin, Kumamoto, Japan) assay, it was identified that amino acid residue sequence 10-24 peptide (NH<sub>2</sub>-AVTDHPDRLWAWKEF-COOH) of this TCL1 oncogene specifically inhibits cell growth associated with AKT activation (Fig. 6).

[0058] In this method, 10/24 peptide was pre-treated at 0-50  $\mu$ M concentration in cell growth test by using T4 cell lines without stimulation, 48 hours later the growth potency was measured by using WST-8 reagent assay, the absorbance at 450 nm was measured by ELISA using a microplate reader (Model 550; BioRad, Tokyo, Japan).

15

2. Binding test by co-immunoprecipitation assay

[0059] To examine the cause of inhibition of the specific cell growth of 10/24 peptide, by using co-immunoprecipitation assay, it was identified that 10/24 peptide specifically binds to Akt kinase (Fig. 7). In this method, AKT kinase was overexpressed in human 293 cells, and the harvested cell lysates was incubated with 10/24 peptide (NH<sub>2</sub>-AVTDHPDRLWAWKEF-COOH) for around 2 hours. Further, the treated cell lysates were added agarose beads bound to the specific antibody against an epitope that was fused with Akt, and co-incubated for 2-3 hours. Then, molecules adhered in the cell lysates were immunoprecipitated with agarose beads bound to this antibody, and examined the binding to Akt kinase by using western blotting with specific antibody.

25

2. Lipid-protein pull down assay

[0060] Lipid-protein pull down assay was performed for 10/24 peptide of TCL1.

30 Method:

[0061] Lipid-protein pull down assay was performed by using PIP Beads (PI (3,4,5) P3 Echelon Bioscience Incorporated). 10/24 NH<sub>2</sub>-AVTDHPDRLWAWKEF-COOH, and  $\beta$ C NH<sub>2</sub>-EKQHAWLPLTIE-COOH as a control, were used. After treatment for 2 hours at 4°C by using 50ng of AKT kinase (unactivated, Upstate Biotechnology, #14-279), 25  $\mu$ l of PIP Beads (PI (3, 4, 5) P3 Echelon Bioscience Incorporated) was added, then washed with fluid containing (10 mM HEPES, pH7.4, 0.25% NP-40.140mM NaCl), and western blotting was performed by using Akt antibody (Cell Signaling) (Fig. 8). In three lanes from the left in the figure, the bindings to AKT were inhibited in a dose-dependent manner at 1-400  $\mu$ M, while control peptides in right lanes in the figure did not inhibit at all. From these results, it was identified that the peptide (NH<sub>2</sub>-AVTDHPDRLWAWKEF-COOH) competitively inhibits the bindings of Phosphoinositide (PI (3,4,5) P3) to Akt kinase. Thus, this is considered to be the inhibition mechanism for Akt activation.

40

Example 3

[Binding test between 10/24 peptide and Akt subtype molecule by using co-immunoprecipitation assay]

45

[0062] Binding test between 10/24 peptide and Akt subtype molecule: Akt1, Akt2, or Akt3, was performed by using co-immunoprecipitation assay.

Method:

50

[0063] The cell lysate overexpressed Akt kinase in 293 cells (ATCC) i.e., Akt1, Akt2 or Akt3 in pCMV6 was expressed in 293 cells (ATCC) by using calcium phosphate method, and the overexpressed cells were harvested, lysed, and pre-treated with Protein G/A agarose mixture (50% v/v, ProG/A, Pharmacia). Akt or a control peptide ( $\beta$ C) was added to the cell lysate at 400  $\mu$ M, inoculated with ProG/A for 3 hours at 4°C, and was added anti Flag M2 antibody (Sigma). The resultant immune precipitant was washed, and then were identified the binding to Akt kinase by western blotting (anti-HA antibody, 3F10, Boehringer Mannheim). The results are shown in figure 7. As shown in the figure, the 10/24 peptide bound to any of the three subtypes of molecules, Akt1, Akt2, and Akt3.

55

## EP 1 717 244 B1

### Example 4

[Inhibition effect test for Akt activation by Akt kinase assay with the use of GSK (Glycogen Synthesis Kinase 3) as substrate]

- 5 **[0064]** It has been known that Akt promotes phosphorylation of GSK (Glycogen Synthesis Kinase 3).  
**[0065]** With the use of the GSK as substrate Akt kinase assay was performed, and inhibition test for Akt activation with 10/24 peptide was performed.

Method:

- 10 **[0066]** In vitro Akt kinase assay was performed by using kit (Cell Signaling, #9840). Recombinant Akt protein extracted from mammalian cells was mixed with 200  $\mu$ M concentrations of peptide, and reacted for 2 hours. Phosphorylation was performed for 4 minutes at 30°C. After analyzing the reaction mixture on SDS gels, GSK phosphorylation was determined by western blotting. The results are shown in figure 8. As shown in the figure 8, the 10/24 peptide was effectively inhibited the phosphorylation potency for GSK peptide of Akt (three lanes from the left in the figure show the inhibition of GSK phosphorylation diluting the black bands). The similar inhibition effect on Akt kinase activity was also identified by using peptide (NH<sub>2</sub>-VTDHPDRLWAWEK -RRR-VTDHPDRLWAWEK -COOH) having repetitive sequence of 11-23 from 10-24 (AVTDHPDRLWAWEKF).

### 20 Example 5

[inhibition effect test for Akt phosphorylation activation in mouse fibrosarcoma cells (QrSP-11)]

- 25 **[0067]** Inhibition effect on Akt activation of AKT phosphorylation (serine 473 residue, threonine 308 residue) of 10/24 peptide and control peptide in mouse QrSP-11 fibrosarcoma cells were examined.

Method:

- 30 **[0068]** QrSP-11 cells were mixed with 50 mM concentrations of peptide for 16 hours. Then the mixture was stimulated with PDGF (PDGF-AB, Sigma, 3226). The cells were lysed in the presence of phosphatase inhibitor, analyzed on SDS gels, and western blotting was performed by ECL (Amersham) with the use of various antibodies (Cell Signaling; anti-Akt #9272, anti-pThr308 #9275L, and anti-pSer473 #9271L). The results are shown in figure 9. As shown in the figure, in the 10/24 peptide treated cells, phosphorylations of both serine473 and threonine308 were inhibited as shown at the right in the figure. That is, phosphorylation inhibition caused by 10/24 peptide was indicated by diluting the black bands at the right compared to the second and forth rows from the left in the figure.

### Example 6

[Inhibition effect test for the activation and membrane translocation of AKT of 10/24 peptide]

- 40 **[0069]** Inhibition effect for the activation and membrane translocation of AKT of 10/24 peptide were tested with the use of 293 cells (ATCC).

Method:

- 45 **[0070]** One mg of AKT was overexpressed in 293 cells (ATCC) with the use of FuGENE6 (Roche Diagnostics). Sixteen hours later the serum was eliminated, and the cells were stimulated with PDGF-AB (Sigma, #3226) for 10 minutes. The cells were fixed with 4% paraformaldehyde, stained with FITC-conjugated anti-HA antibody (12CA5, MBL) or phospho-Ser 473 antibody (587-F11, Cell Signaling), and observed under the confocal microscope (Nikon).  
50 **[0071]** The results are shown in figure 10. As shown in the figure 10, in the 10/24 treated cells (g-i), the inhibition of the membrane translocation and activation of AKT was identified compared with the control peptides (a-f) (around the cells glowed green or yellow as shown in d or f, while this effect was inhibited in the 10/24 peptide as shown in j or i. See the figure). That is, from the results of the experiment, it was identified that the 10/24 peptide inhibits the activation at the same time as the translocating of AKT to the membrane in cells. AKT essentially translocates to the cell surface where it is activated, but in the 10/24 peptide treated cells, it was identified that the translocation to membrane and activation of AKT was inhibited.

## EP 1 717 244 B1

### Example 7

[Induction of apoptosis and anti-tumor effect by 10/24 peptide]

5 **[0072]** The effect on apoptosis of 10/24 peptide was examined with the use of human T cell leukemia cells (T4).

Method:

10 **[0073]** 10/24 peptide (NH<sub>2</sub>- - AVTDHPDRLWAWKEF -COOH) was pre-treated at 0-30  $\mu$ M concentrations with the use of T4 cell lines without stimulation, 48 hours later stained with propidium iodide, and the apoptosis was determined by FACS(Beckton Dickinson). Further, to identify the AKT dependency of the anti-tumor effect, myr-AKT (constitutively activated AKT) was overexpressed. The results are shown in figure 11. As shown in the figure, 10/24 peptide was identified to enhance apoptosis, compared with control peptide. Similar tendency to enhance the apoptosis was also identified during the induction of apoptosis by dexamethasone. As a result of myr-AKT overexpression, apoptosis was inhibited ( $\Delta$  in Fig. 11), and it was identified that 10/24 peptide achieves the effect in AKT dependent manner.

### Example 8

[In vivo anti-tumor effect by 10/24 peptide]

20 **[0074]** In vivo anti-tumor effect by 10/24 peptide with the use of the transplanted tumor cells was examined.

Method:

25 **[0075]** Mouse QrSP-11 fibrosarcoma cells was transplanted to abdominal wall of C57BL/6 mouse, and examined the inhibition effect on tumor growth by peptide. 10/24 peptide or control peptide was injected directly into tumor cells (2  $\mu$ M was administered 3 times a week for each mouse, shown by arrows in the figure), the tumor diameters were measured and the volumes were calculated. The results are shown in figure 12. As shown in the figure, it was identified that tumor growth was effectively inhibited in 10/24 peptide.

### Example 9

[Histological examination of 10/24 peptide treated mouse tumor]

35 **[0076]** Experimental mouse tumors of example 8 were harvested on the day 9, and histologically examined by macroscopic appearance, hematoxylin-eosin (H&E) staining (an observing method for conditions of cell nucleus and the like), TUNEL (Tdt-mediated dUTP nick end labeling, #MK500, Takara) immunostaining (a histologically identifying method for the way of the atypical death of carcinoma i.e., apoptosis), and phospho Akt (Ser473) monoclonal antibody (a method for determining 587F11, Cell Signaling, phosphorylation of AKT kinase, and observing the activation). The results are shown in figure 13.

40 **[0077]** As shown in pictures in the figure, by macroscopic appearance, reduction in tumor size was clearly observed in the 10/24 peptide treated compared to the control; by H&E staining, increased apoptosis was observed in the 10/24 peptide treated; by TUNEL staining, increased cell apoptosis was observed in the 10/24 peptide treated; and by Akt activation, inhibition of Akt activation was observed in the 10/24 peptide treated. That is, by histological examination of the 10/24 peptide treated mouse tumor, it was identified that the 10/24 peptide suppressed tumor growth, increased apoptosis (H&E, TUNEL), and inhibited AKT activity (p473 staining).

### SEQUENCE LISTING

50 **[0078]**

<110> Japan Science and Technology Agency

<120> Akt activity specifically inhibiting polypeptide.

55

<130> P026371EP

<140> 04807007.2

EP 1 717 244 B1

<141> 2004-12-14

<150> JP 2003-416556

<151> 2003-12-15

5

<150> JP 2004-134583

<151> 2004-04-28

10

<160> 20

<170> PatentIn version 3.3

<210> 1

<211> 15

15

<212> PRT

<213> Homo sapiens

<400> 1

20

Ala Val Thr Asp His Pro Asp Arg Leu Trp Ala Trp Glu Lys Phe  
1 5 10 15

<210> 2

25

<211> 45

<212> DNA

<213> Homo sapiens

<400> 2

30

gcagtcaccg accaccggga ccgctgtgg gctgggaga agttc 45

<210> 3

<211> 30

<212> PRT

35

<213> Homo sapiens

<400> 3

40

Met Ala Ser Glu Ala Ser Val Arg Leu Gly Val Pro Pro Gly Arg Leu  
1 5 10 15

45

Trp Ile Gln Arg Pro Gly Ile Thr Glu Asp Glu Glu Glu Arg  
20 25 30

<210> 4

<211> 90

<212> DNA

50

<213> Homo sapiens

<400> 4

55

atggcctccg aagcttctgt gcgtctaggg gtgccccctg gccgtctgtg gatccagagg 60

cctggcatct acgaagatga ggaggggaga 90

EP 1 717 244 B1

<210> 5  
 <211> 25  
 <212> PRT  
 <213> Homo sapiens

5

<400> 5

Met Ala Gly Glu Asp Val Gly Ala Pro Pro Asp His Leu Thr Val His  
 1 5 10 15

10

Gln Glu Gly Ile Tyr Arg Asp Glu Tyr  
 20 25

15

<210> 6  
 <211> 75  
 <212> DNA  
 <213> Homo sapiens

20

<400> 6

atggcaggag aggatgtggg ggctccaccc gatcacctct gggttcacca agagggtatc 60

taccgcgacg aatac 75

25

<210> 7  
 <211> 16  
 <212> PRT  
 <213> Mus musculus

30

<400> 7

Ala Glu Thr Pro Ala His Pro Asn Arg Leu Trp Ile Trp Glu Lys His  
 1 5 10 15

35

<210> 8  
 <211> 48  
 <212> DNA  
 <213> Mus musculus

40

<400> 8  
 gcagagacac ctgcacaccc caaccgcctg tggatctggg agaagcac 48

45

<210> 9  
 <211> 16  
 <212> PRT  
 <213> Rattus norvegicus

50

<400> 9

Pro Glu Thr Pro Pro His Pro Asp Arg Leu Trp Leu Trp Glu Lys His  
 1 5 10 15

55

<210> 10  
 <211> 48  
 <212> DNA

EP 1 717 244 B1

<213> Mus musculus

<400> 10  
ccagagacac cccacacccc cgaccgcctg tggctctggg agaagcac 48

5

<210> 11  
<211> 33  
<212> DNA  
<213> Artificial

10

<220>  
<223> Oligonucleotide primer

<400> 11  
ccaccaaac caaaaaaga gatcgaattc atg 33

15

<210> 12  
<211> 33  
<212> DNA  
<213> Artificial

20

<220>  
<223> Oligonucleotide primer

<400> 12  
attcatagat ctctgcaggt cgacggatcc tca 33

25

<210> 13  
<211> 60  
<212> DNA  
<213> Artificial

30

<220>  
<223> Oligonucleotide primer

35

<400> 13  
atggccgagt gcccgacact cggggaggca gtcaccgacc acccgggccg cctgtgggcc 60

40

<210> 14  
<211> 33  
<212> DNA  
<213> Artificial

<220>  
<223> Oligonucleotide primer

45

<400> 14  
gtgtattgg acgagatgca gcacgcctgg ctg 33

50

<210> 15  
<211> 34  
<212> DNA  
<213> Artificial

55

<220>  
<223> Oligonucleotide primer

<400> 15

EP 1 717 244 B1

gataaaggat aggttacggt tacgggtgct cttg 34

5 <210> 16  
<211> 33  
<212> DNA  
<213> Artificial

10 <220>  
<223> Oligonucleotide primer

<400> 16  
ccaagcctgc tgcctgtcat gtggcagctc tac 33

15 <210> 17  
<211> 49  
<212> DNA  
<213> Artificial

20 <220>  
<223> Oligonucleotide primer

<400> 17  
atcatcggat cctcagtcac ctggcagcag ctcgagaagc acgtcctcc 49

25 <210> 18  
<211> 39  
<212> DNA  
<213> Artificial

30 <220>  
<223> Oligonucleotide primer

<400> 18  
cagcagcct ggctggccgc ggccatcgag ataaaggat 39

35 <210> 19  
<211> 24  
<212> DNA  
<213> Artificial

40 <220>  
<223> Oligonucleotide primer

45 <400> 19  
gcctggctgg ccttaatcga gata 24

50 <210> 20  
<211> 29  
<212> PRT  
<213> Homo sapiens

<400> 20

55



**Patentansprüche**

- 5
1. Polypeptid, das die Akt-Kinaseaktivität spezifisch hemmt, und das aus der in SEQ ID NO: 1 des Sequenzprotokolls angegebenen Aminosäuresequenz besteht.
2. Polypeptid, bestehend aus einer Aminosäuresequenz, bei der eine oder mehrere Aminosäuren in der in SEQ ID NO: 1 des Sequenzprotokolls angegebenen Aminosäuresequenz deletiert ist bzw. sind, und das die Akt-Kinaseaktivität spezifisch hemmt.
- 10
3. Gen-DNA, die ein nachfolgendes Protein (a) oder (b) codiert:
- (a) Polypeptid, bestehend aus der in SEQ ID NO: 1 angegebenen Aminosäuresequenz;
- (b) Polypeptid, bestehend aus einer Aminosäuresequenz, bei der eine oder mehrere Aminosäuren in der in SEQ ID NO: 1 angegebenen Aminosäuresequenz deletiert ist bzw. sind, und das die Akt-Kinaseaktivität spezifisch hemmt.
- 15
4. DNA, bestehend aus der in SEQ ID NO: 2 angegebenen Basensequenz.
5. Rekombinanter Expressionsvektor, der durch Integrieren der DNA nach Anspruch 3 in einen Gen-Expressionsvektor konstruiert wird.
- 20
6. Verfahren zur Herstellung eines Polypeptids, das die Akt-Kinaseaktivität spezifisch hemmt, wobei der rekombinante Expressionsvektor nach Anspruch 5 in eine Wirtszelle eingebracht und exprimiert wird.
- 25
7. Antikörper, der durch die Verwendung des in SEQ ID NO: 1 des Sequenzprotokolls angegebenen Polypeptids induziert wird, und der spezifisch an das Polypeptid bindet.
8. Antikörper nach Anspruch 7, wobei der Antikörper ein monoklonaler Antikörper ist.
- 30
9. Antikörper nach Anspruch 7, wobei der Antikörper ein polyklonaler Antikörper ist.
10. Spezifischer Inhibitor der Akt-Aktivität, wobei das Polypeptid nach Anspruch 1 oder 2 ein Wirkstoff ist.
- 35
11. Spezifischer Inhibitor der Akt-Aktivität nach Anspruch 10, wobei das Polypeptid die Sequenz eines Aminosäurerests 10-24 einer Aminosäuresequenz für humanes TCL1-Protein ist.
12. Spezifischer Inhibitor der Akt-Aktivität nach Anspruch 10, wobei die spezifische Hemmung der Akt-Aktivität die Hemmung der Bindung von Phosphoinositid an Akt ist.
- 40
13. Antitumormittel, wobei das Polypeptid nach Anspruch 1 oder 2 ein Wirkstoff ist.
14. Antitumormittel nach Anspruch 13, wobei das Antitumormittel ein Mittel zur Vorbeugung oder Behandlung von Malignität ist.
- 45
15. Antitumormittel nach Anspruch 14, wobei die Behandlung der Malignität die Vorbeugung oder Behandlung von Brustkrebs, Lungenkrebs, Leukämie oder Lymphoid-Tumor ist.
16. Polypeptid nach Anspruch 1 oder 2 zur Verwendung bei der Behandlung von Krebs.

50

**Revendications**

- 55
1. Polypeptide inhibant spécifiquement l'activité Akt kinase, qui est constitué de la séquence d'acides aminés indiquée dans SEQ ID NO: 1 de la liste de séquences.
2. Polypeptide constitué d'une séquence d'acides aminés dans laquelle un ou plusieurs acides aminés sont délétés dans la séquence d'acides aminés indiquée dans SEQ ID NO: 1 de la liste de séquences, et inhibant spécifiquement l'activité Akt kinase.

## EP 1 717 244 B1

3. ADN génique codant pour une protéine (a) ou (b) suivante :

(a) un polypeptide constitué de la séquence d'acides aminés indiquée dans SEQ ID NO: 1 ;

(b) un polypeptide constitué d'une séquence d'acides aminés dans laquelle un ou plusieurs acides aminés sont  
5 déléétés dans la séquence d'acides aminés indiquée dans SEQ ID NO: 1, et inhibant spécifiquement l'activité Akt kinase.

4. ADN constitué de la séquence de bases indiquée dans SEQ ID NO: 2,

10 5. Vecteur d'expression recombinant, qui est construit par intégration de l'ADN selon la revendication 3 dans un vecteur d'expression génique.

6. Procédé pour produire un polypeptide qui inhibe spécifiquement l'activité Akt kinase, dans lequel le vecteur d'ex-  
pression recombinant selon la revendication 5 est introduit dans une cellule hôte et exprimé.

15 7. Anticorps qui est induit en utilisant le polypeptide indiqué dans SEQ ID NO: 1 de la liste de séquences et se lie spécifiquement au polypeptide.

8. Anticorps selon la revendication 7, l'anticorps étant un anticorps monoclonal.

9. Anticorps selon la revendication 7, l'anticorps étant un anticorps polyclonal.

10. Inhibiteur spécifique de l'activité Akt, le polypeptide selon la revendication 1 ou 2 étant une substance active.

25 11. Inhibiteur spécifique de l'activité Akt selon la revendication 10, le polypeptide étant la séquence des résidus d'acide aminé 10 à 24 d'une séquence d'acides aminés de la protéine TCL1 humaine.

12. Inhibiteur spécifique de l'activité Akt selon la revendication 10, l'inhibition spécifique de l'activité Akt étant l'inhibition  
de la liaison de phosphoinositide à Akt.

30 13. Agent antitumoral, le polypeptide selon la revendication 1 ou 2 étant une substance active.

14. Agent antitumoral selon la revendication 13, l'agent antitumoral étant un agent pour la prévention ou le traitement  
d'une malignité.

35 15. Agent antitumoral selon la revendication 14, le traitement d'une malignité étant la prévention ou le traitement du cancer du sein, du cancer du poumon, d'une leucémie ou d'une tumeur lymphoïde.

40 16. Polypeptide selon la revendication 1 ou 2 pour utilisation dans le traitement de cancer.

Fig.1

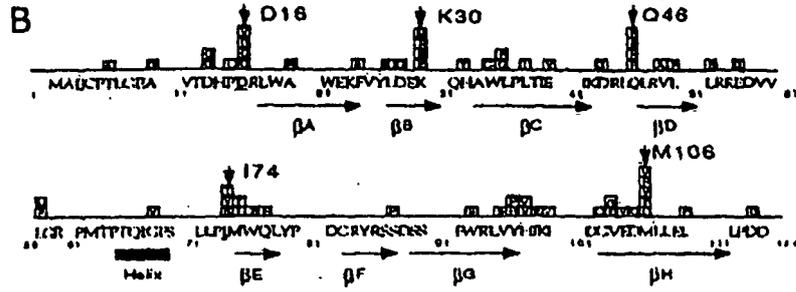


Fig.2

	Clone 1	Clone 2	Clone 3
TCL1 D16G	•	•	•
TCL1 K30M	•	•	•
TCL1 Q46R	•	•	•
TCL1 I74V	•	•	•
TCL1 M106V	•	•	•
TCL1 Wild Type	•	•	•

Fig.3

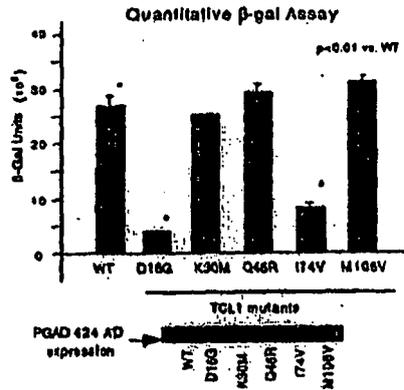


Fig.4

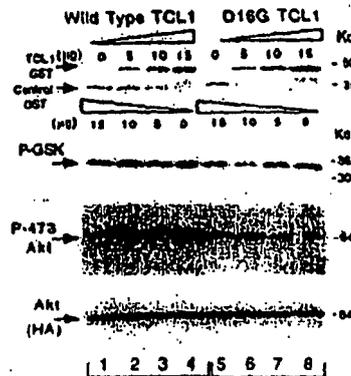


Fig.5

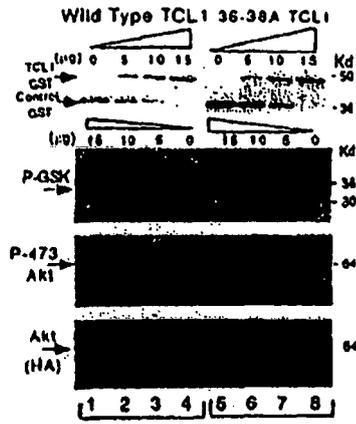


Fig.6

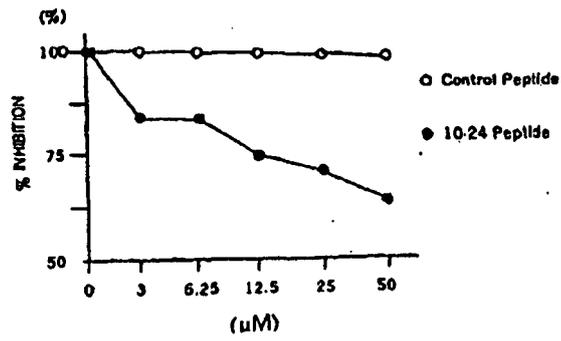


Fig.7

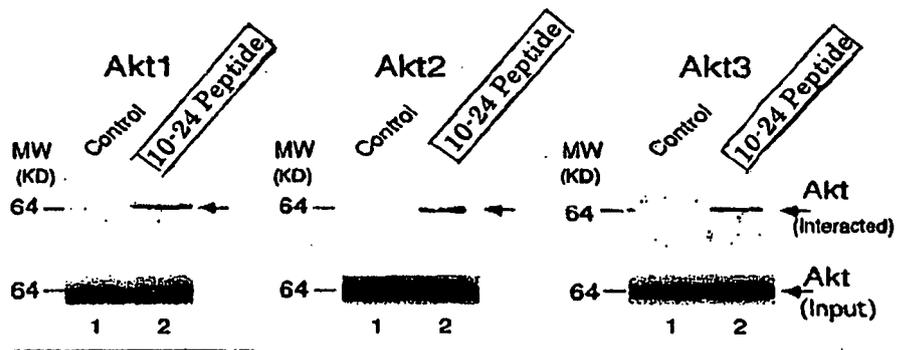


Fig.8

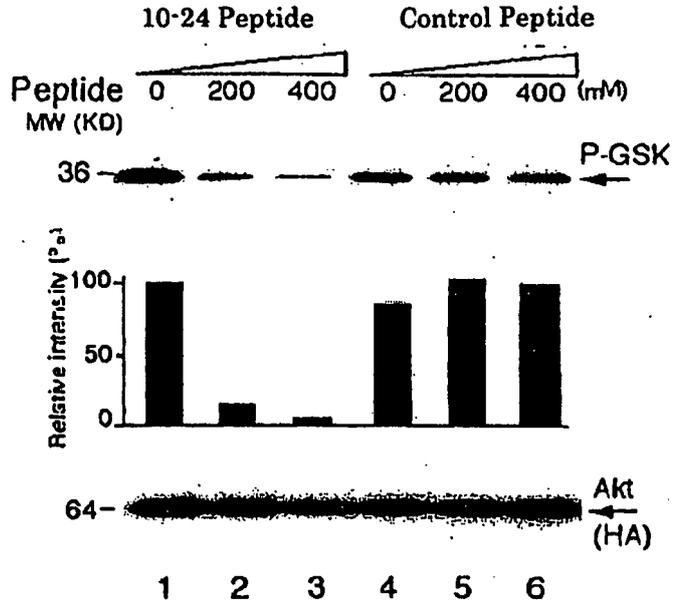


Fig.9

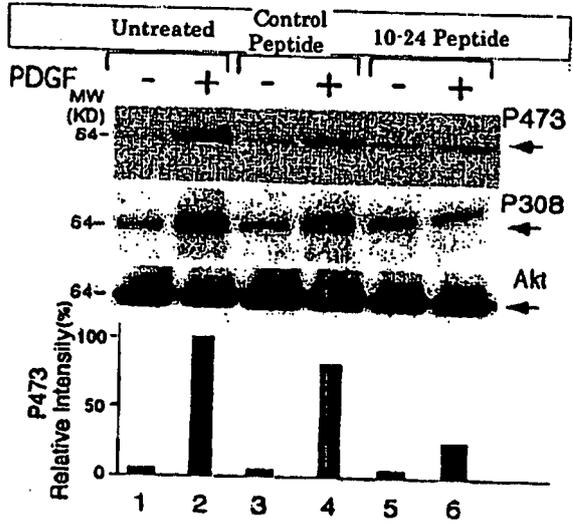


Fig10

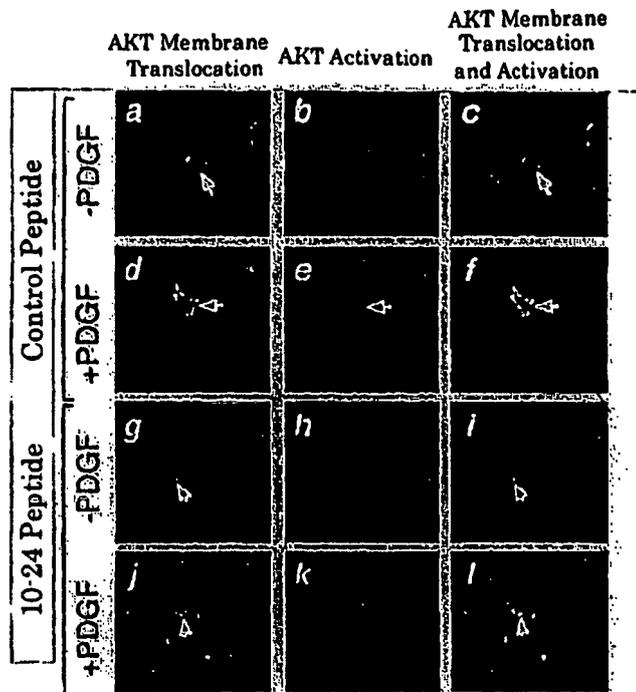


Fig.11

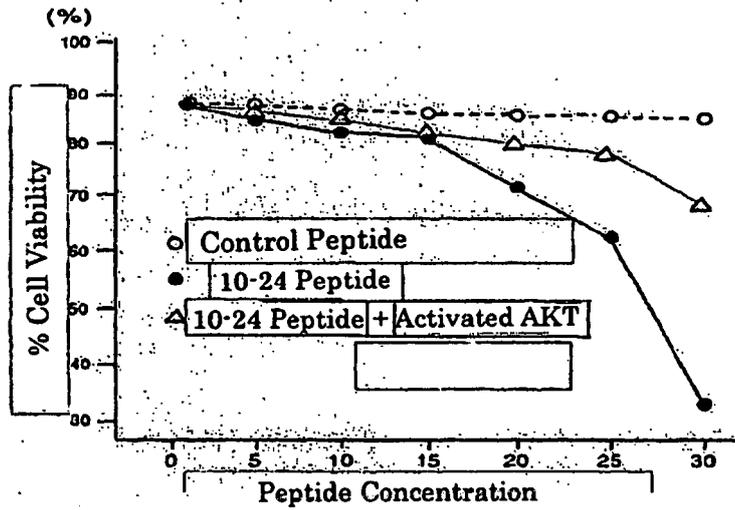


Fig.12

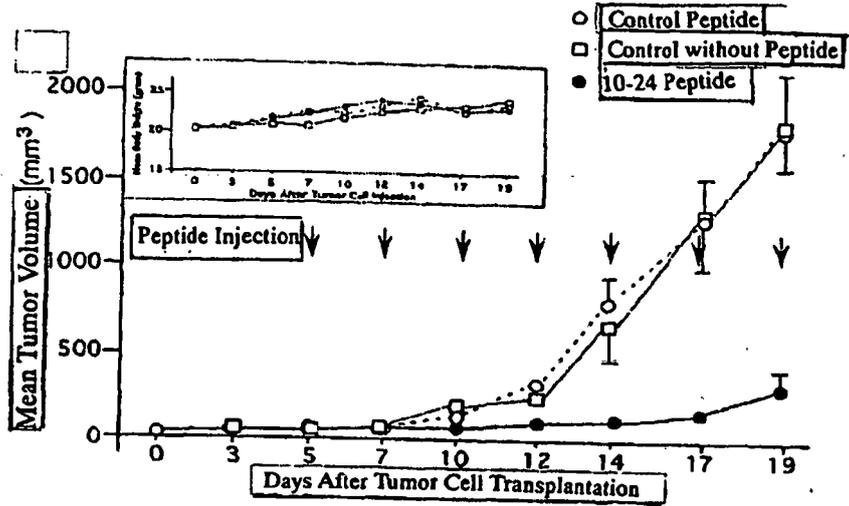
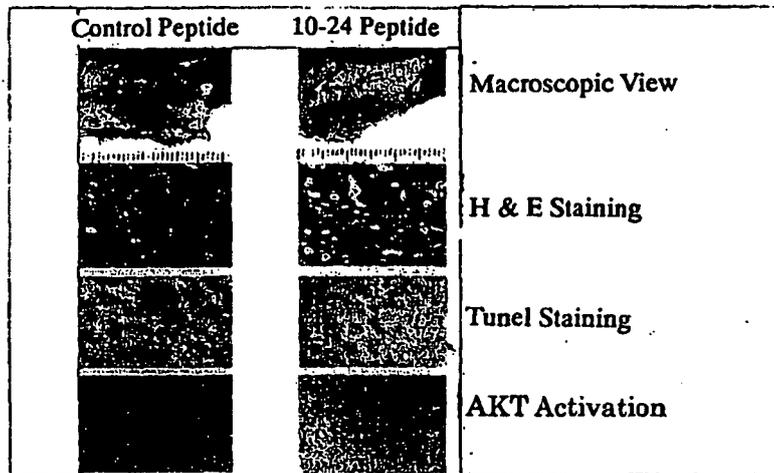


Fig.13



**REFERENCES CITED IN THE DESCRIPTION**

*This list of references cited by the applicant is for the reader's convenience only. It does not form part of the European patent document. Even though great care has been taken in compiling the references, errors or omissions cannot be excluded and the EPO disclaims all liability in this regard.*

**Patent documents cited in the description**

- US 5985598 A [0010] [0013] [0029]
- JP 04807007 B [0078]
- JP 2003416556 A [0078]
- JP 2004134583 A [0078]

**Non-patent literature cited in the description**

- *Genes & Dev.*, 1999, vol. 13, 2905-2927 [0002] [0013]
- *Annu. Rev. Biochem*, 1998, vol. 67, 481-507 [0002]
- *Biochem.J.*, 1998, vol. 335, 1-13 [0002] [0013]
- *J. Biol. Chem.*, vol. 275, 36108-36115 [0005] [0013]
- *Cell*, 1999, vol. 96, 857-868 [0005] [0013]
- *Annu. Rev. Biochem.*, 1999, vol. 68, 965 [0007]
- *Oncogene*, 1993, vol. 8, 2475-2483 [0008] [0013]
- *Proc. Natl. Acad. Sci. USA*, 1994, vol. 91, 12530-12534 [0008] [0013]
- *Mol. Cell*, 2000, vol. 6, 395-407 [0011] [0013]
- *J. Biological Chemistry*, 2002, vol. 277 (5), 3743-3751 [0011] [0013]
- *Molecular and Cellular Biology*, 2002, vol. 22 (5), 1513-1525 [0012] [0013]
- *Biochemistry*, 2002, vol. 41, 6376-6382 [0013]
- *Annu. Rev. Biochem*, 1998, vol. 67, 481-507 [0013]
- *Proc. Natl. Acad. Sci. U.S.A.*, 1999, vol. 96 (6), 2949-2951 [0030]
- *Oncogene*, 1993, vol. 8 (9), 2475-2483 [0030]
- *MOLECULAR AND CELLULAR BIOLOGY*, March 2002, 1513-1525 [0043]
- *Mol. Cell*, vol. 6, 395-407 [0045] [0048] [0049]
- *Proc. Natl. Acad. Sci. USA*, vol. 94, 11534-11539 [0048]
- *Molecular and Cellular Biology*, March 2002, 1513-1525 [0055]