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## Organic Cation Transporter Preferentially Expressed in Hematopoietic Cells and Leukemias and Uses Thereof

Jeffrey A. Moscow  
*University of Kentucky*, [jmoscow@uky.edu](mailto:jmoscow@uky.edu)

Xin Lu  
*University of Kentucky*, [xin.lu@uky.edu](mailto:xin.lu@uky.edu)

Craig Jordan  
*University of Kentucky*, [jordan.craig@uky.edu](mailto:jordan.craig@uky.edu)

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**Moscow et al.**

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(54) **ORGANIC CATION TRANSPORTER  
PREFERENTIALLY EXPRESSED IN  
HEMATOPOIETIC CELLS AND LEUKEMIAS  
AND USES THEREOF**

(75) Inventors: **Jeffrey Moscow**, Lexington, KY (US);  
**Xin Lu**, Shanghai (CN); **Craig Jordan**,  
Rochester, NY (US)

(73) Assignee: **University of Kentucky Research  
Foundation**, Lexington, KY (US)

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20, 2004, now abandoned.

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20, 2003.

(51) **Int. Cl.**

**C12Q 1/00** (2006.01)  
**G01N 33/53** (2006.01)  
**C07K 14/435** (2006.01)

(52) **U.S. Cl.** ..... **435/4; 530/350**

(58) **Field of Classification Search** ..... None  
See application file for complete search history.

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*Primary Examiner*—Bridget E Bunner

(74) *Attorney, Agent, or Firm*—McDermott Will & Emery  
LLP

(57) **ABSTRACT**

A novel organic cation transporter (OCT) gene, OCT 6, and  
use thereof is described. The OCT6 gene is preferentially  
expressed in human hematopoietic tissues, including CD34+  
cells and leukemia cells. Its narrow tissue distribution, sub-  
strate specificity, and close homology to other cell membrane  
transporters make OCT6 an attractive target for the treatment  
of myeloid diseases.

**7 Claims, 10 Drawing Sheets**

FIG. 1A

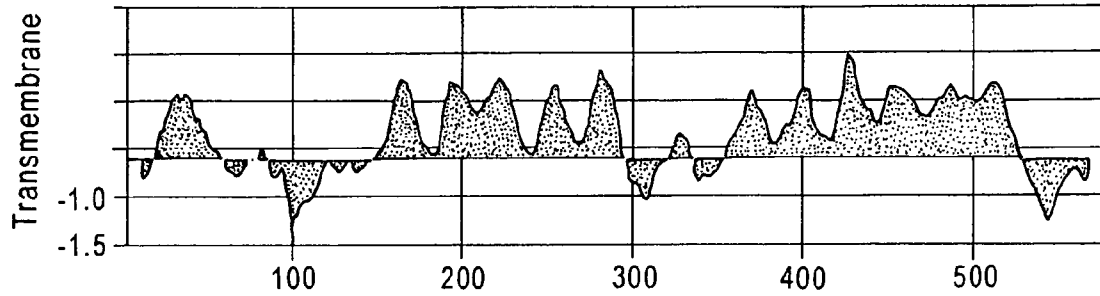
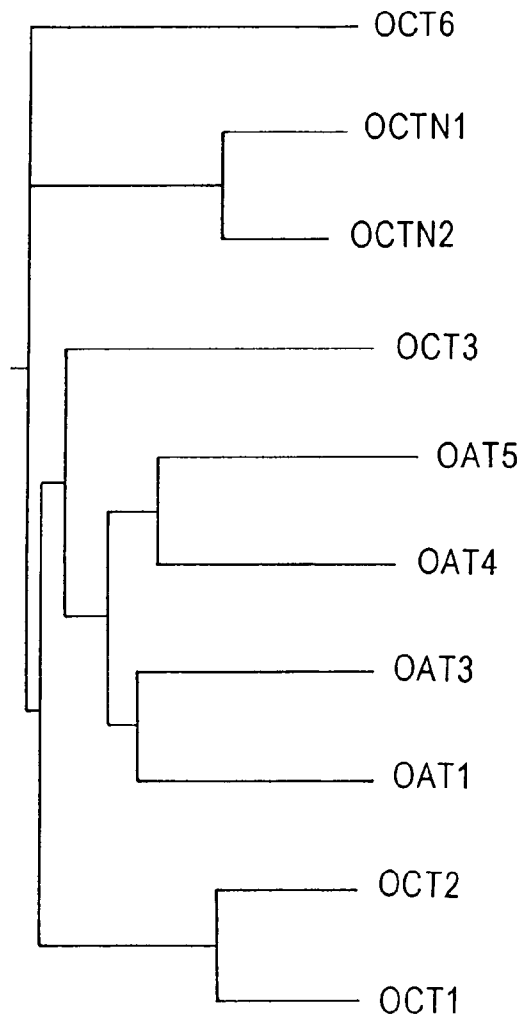


FIG. 1B



SEQ ID NO. 2  
 OCT6 M G S R H F E G I Y D H V G H F G R F Q R V L Y F F L L C Y A F Q N I S C G G F F Y H I A S V F F L A G T P H  
 SEQ ID NO. 4  
 OCTN1 M R D Y D E V I A F L G E W F G P F Q R L I F F L L S A S I I P N G F Y T G L A S V F F L A G T P E  
 SEQ ID NO. 5  
 OCT3 M A Q F V Q V L A E I G D F G R F Q R L I F F L L C Y A S I I L P N G F Y T G L A S V F F L A G T P E  
 SEQ ID NO. 6  
 OCTN2 M R D Y D E V T A F L G E W F G R F Q R L I F F L L S A S I I L P N G F Y T G L A S V F F L A G T P E  
 SEQ ID NO. 7  
 OCT2 M P T T V D D D V L E H G G E S G W F Q K Q A H L V F L L P S L M L L S A T F A P I C V V G I G L A S V F F L A G T P E  
 SEQ ID NO. 8  
 OCT1 M A F E E L L S Q V G G L G E S G R F Q K Q M L L Q V L T F I L P S L M L L S A T F A P I C V V G I G L A S V F F L A G T P E  
 SEQ ID NO. 9  
 OAT5 M A F S K L L E Q V G Q A G G V M G H F Q T L L Q V L T F I L P S L M L L S A T F A P I C V V G I G L A S V F F L A G T P E  
 SEQ ID NO. 10  
 OAT4 M A F S B I L L D R V G G S M G H F Q T L L Q V V A F L G L L P S L M L L S A T F A P I C V V G I G L A S V F F L A G T P E  
 SEQ ID NO. 11  
 OAT3 M A F N D L L Q Q Q G R F Q I H V A F L G L L P S L M L L S A T F A P I C V V G I G L A S V F F L A G T P E  
 SEQ ID NO. 12  
 OAT1 M A F N D L L Q Q Q G R F Q I H V V T L V V L P S L M L L S A T F A P I C V V G I G L A S V F F L A G T P E

50  
 H V C R R P P G N V S Q V V F H N H S N W S L E D T G A L L S S G Q K D Y V T P L L R L R D D G G E I W E L S R  
 H R C R V P D - - - A A N L S S - - - A W R - - - L S A - - - - N N S Q - - - A E Q L - - - N Y T T V P P G L G I S I P P M G  
 H R C R A V A W - - - V K N H T F N - - - A W R - - - L S A - - - - A H T T V - - - N Y T T V P P G L G I S I P P M G  
 H R C R V P D - - - A A N L S S - - - A W R - - - L S A - - - - N N S Q - - - A E Q L - - - N Y T T V P P G L G I S I P P M G  
 H R C R S P G - - - V A E L S L R C G W S P A E E L - - - - A W R - - - L S A - - - - N N S Q - - - A E Q L - - - N Y T T V P P G L G I S I P P M G  
 H R C Q S P G - - - V A E L S L R C G W S P A E E L - - - - A W R - - - L S A - - - - N N S Q - - - A E Q L - - - N Y T T V P P G L G I S I P P M G  
 H R C W V H M - - - V A E L S L R C G W S P A E E L - - - - A W R - - - L S A - - - - N N S Q - - - A E Q L - - - N Y T T V P P G L G I S I P P M G  
 H R C W T H M - - - L D N N G S - - - - A V S T N M T P - - - - - - - N G G L - - - - - - - P W V L P E  
 H H C R P P H - - - N A S T G -  
 H H C R P P A - - - D A N L S K -  
 H H C R P P A - - - A N A N L S K -  
 H . C R  
 100

FIG. 2A

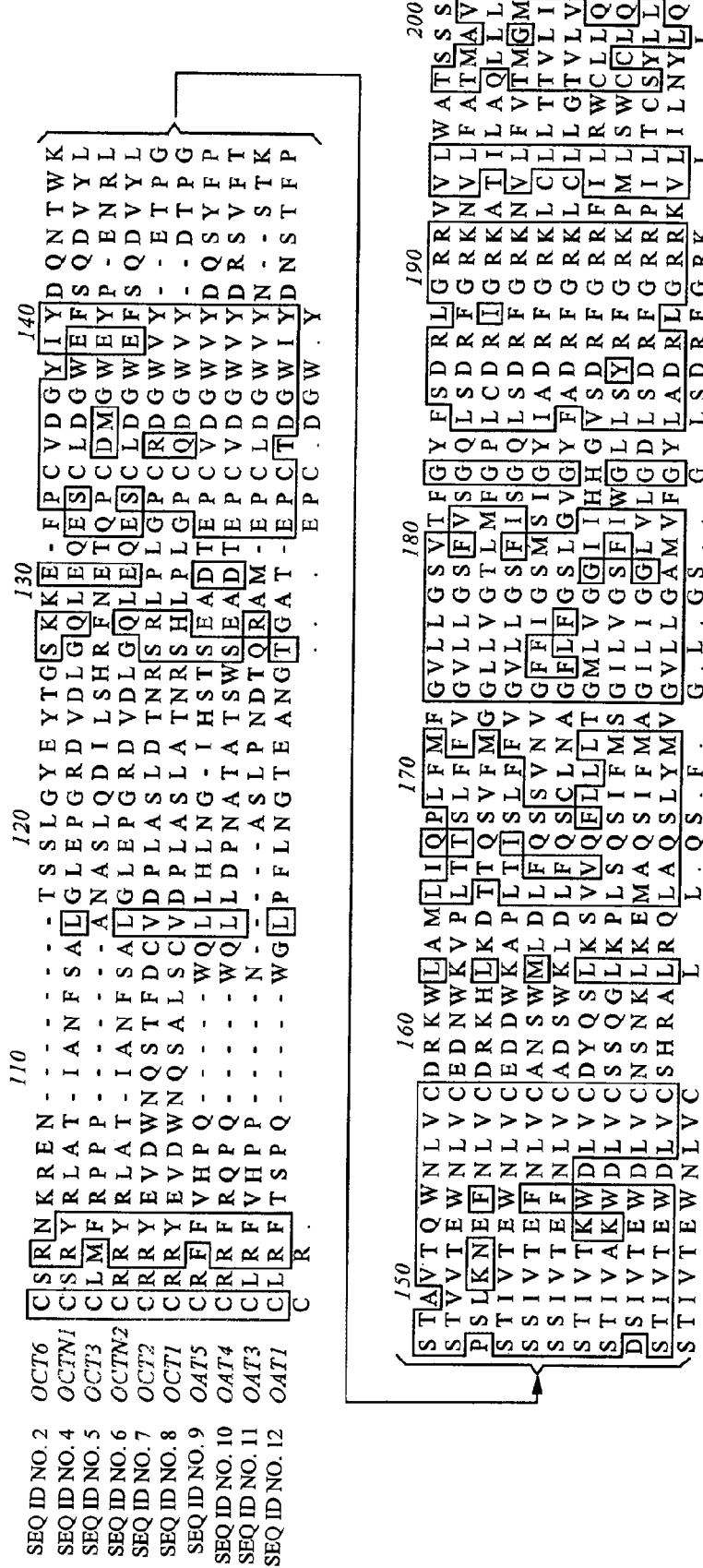


FIG. 2B

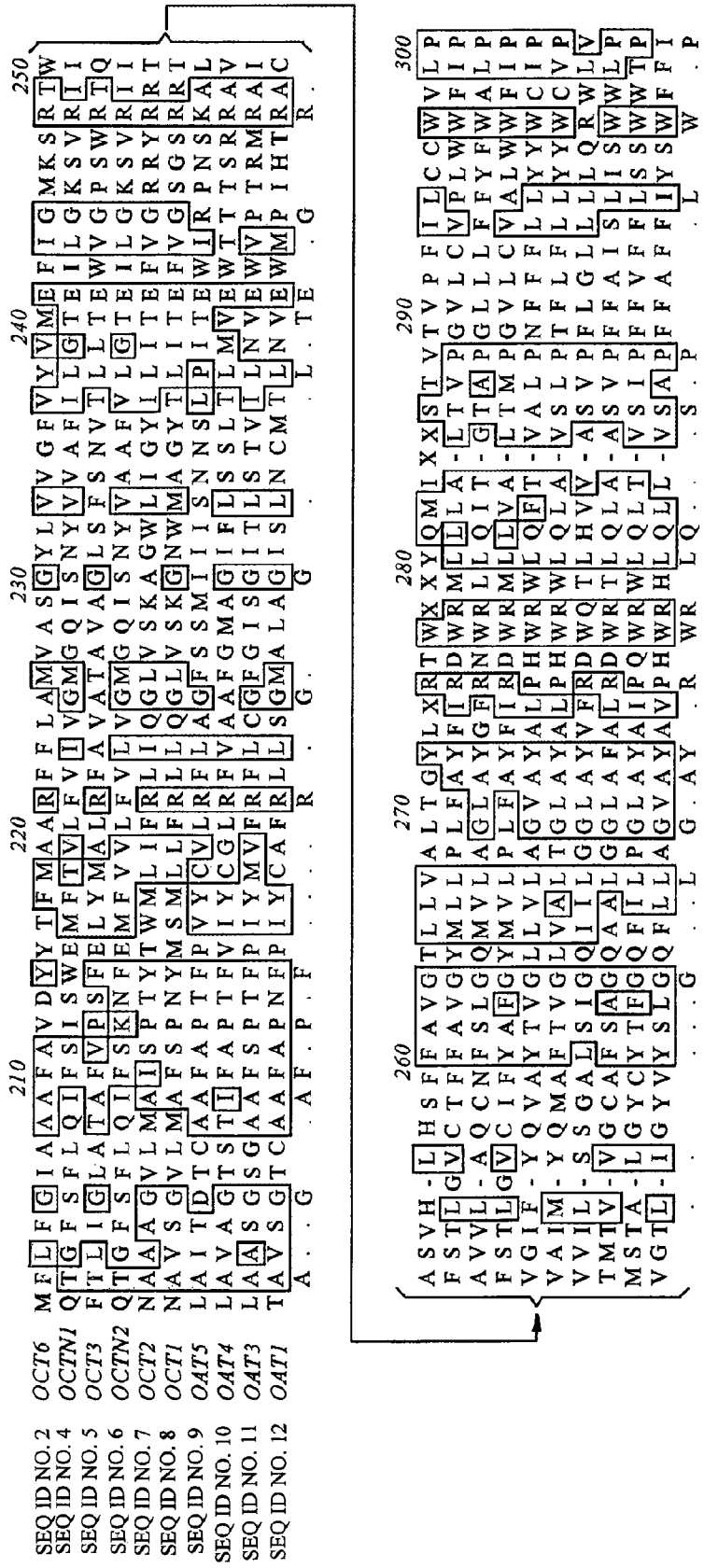


FIG. 2C

SEQ ID NO. 2	OCT6	ES	FWL	SI	GR	Y	E	E	Q	K	I	V	D	I	M	A	K	W	N	R	A	S	S	C	K	L	S	E	L	L	S	L	D	L	Q	G	P	V	S	N	S	P	T		
SEQ ID NO. 4	OCTN1	ES	FWL	SI	GR	Y	E	E	Q	K	I	V	D	I	M	A	K	W	N	R	A	S	S	C	K	L	S	E	L	L	S	L	D	L	Q	G	P	V	S	N	S	P	T		
SEQ ID NO. 5	OCT3	ES	AR	WL	IS	QR	R	E	E	I	Q	K	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A
SEQ ID NO. 6	OCTN2	ES	AR	WL	IS	QR	R	E	E	I	Q	K	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A
SEQ ID NO. 7	OCT2	ES	AR	WL	IS	QR	R	E	E	I	Q	K	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A
SEQ ID NO. 8	OCT1	ES	AR	WL	IS	QR	R	E	E	I	Q	K	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A
SEQ ID NO. 9	OAT5	ES	AR	WL	IS	QR	R	E	E	I	Q	K	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A
SEQ ID NO. 10	OAT4	ES	AR	WL	IS	QR	R	E	E	I	Q	K	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A
SEQ ID NO. 11	OAT3	ES	AR	WL	IS	QR	R	E	E	I	Q	K	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A
SEQ ID NO. 12	OAT1	ES	AR	WL	IS	QR	R	E	E	I	Q	K	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A	A

FIG. 2D





SEQ ID NO. 2	OCT6	A S I L A P	510	F S V D L S S I W I F I P Q L F V G T M A L L S	530	- G V L T L T L K L P P E T	540	L P P E T	G K R
SEQ ID NO. 4	OCTN1	G S I I A P		Y F V I L L G E Y N A A L P M L I Y G S L T V L I G I L T L		L I G I L T L		L P P E S	L G M T G
SEQ ID NO. 5	OCT3	G G I L T P		Y F V I L L G E Y H A A L P M L I Y G S L T V L I G I L T L		L I G I L T L		L P P E T	L G M T G
SEQ ID NO. 6	OCTN2	G S I L S P		Y F V I L L G E Y D R F L P Y I L M V F G V L G L L V A L T A I L T L		L I G I L T L		L P P E S	L G M T G
SEQ ID NO. 7	OCT2	G G I I T P		Y F V I L L G E Y R L R E V W Q A L P L I I Y G I F P I I G - G L I V F L F F L L P P E T		L I G I L T L		L P P E T	L G M T G
SEQ ID NO. 8	OCT1	G G I I T P		Y F V I L L G E Y R L R E V W Q A L P L I I Y G I F P I I G - G L I V F L F F L L P P E T		L I G I L T L		L P P E T	L G M T G
SEQ ID NO. 9	OAT5	G A A L		Y F V I L L G E Y T V F F T L L P P W I I Y G I F P I I G - G L I V F L F F L L P P E T		L I G I L T L		L P P E T	L G M T G
SEQ ID NO. 10	OAT4	G A M G		Y F V I L L G E Y Q A L P P L L P P W I I Y G I F P I I G - G L I V F L F F L L P P E T		L I G I L T L		L P P E T	L G M T G
SEQ ID NO. 11	OAT3	G S M V S P		Y F V I L L G E Y V Q P P L L P P W I I Y G I F P I I G - G L I V F L F F L L P P E T		L I G I L T L		L P P E T	L G M T G
SEQ ID NO. 12	OAT1	G S I V S P		Y F V I L L G E Y L Y P P L F I Y G I F P I I G - G L I V F L F F L L P P E T		L I G I L T L		L P P E T	L G M T G

550	L A T W E E A A K L E S E N E S K S	560	L L T T N N S G L E K T E A I T P R D S G L G E
	L P E T L E Q M Q L L G P H P R S P K S V P S H T R M L K D G Q E R P T I L K S T A F		L L T T N N S G L E K T E A I T P R D S G L G E
	L K D T L Q D L E L R V K G M K H R K I E - - - K N L - - - K E K A - - - R Q E A V T V E S T S L		L L T T N N S G L E K T E A I T P R D S G L G E
	L P D T I E E A E N M Q R P R K K E - - - K N L - - - K E K A - - - R Q E A V T V E S T S L		L L T T N N S G L E K T E A I T P R D S G L G E
	L P D T I K D V E N Q K - - - T A A Q N - - - R Q E A V T V E S T S L		L L T T N N S G L E K T E A I T P R D S G L G E
	L P D T I Q D L E N W S L R A K K P K - - - Q E P E V E K A S Q R I P L Q P H G P G L G S S		L L T T N N S G L E K T E A I T P R D S G L G E
	L P D T I E D L E - - - S R K K G K Q T R - - - E H Q K Y M V P L Q A S A Q E K N G L		L L T T N N S G L E K T E A I T P R D S G L G E
	L P D T I V Q D L E - - - D . . . D . . . E		L L T T N N S G L E K T E A I T P R D S G L G E

FIG. 2F

FIG. 3A

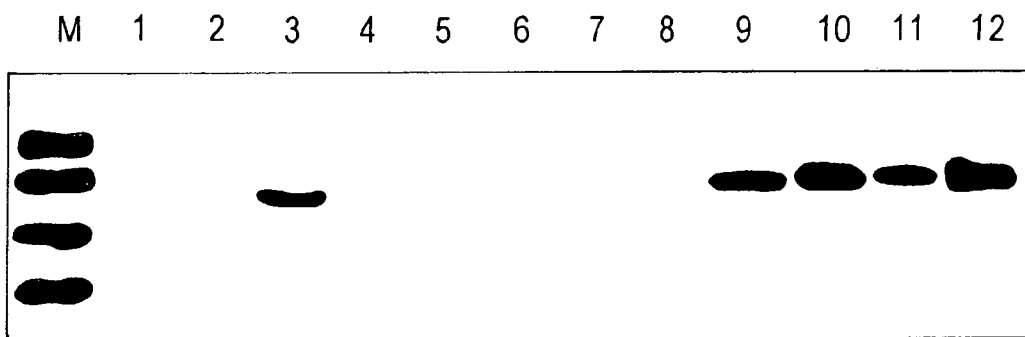


FIG. 3B

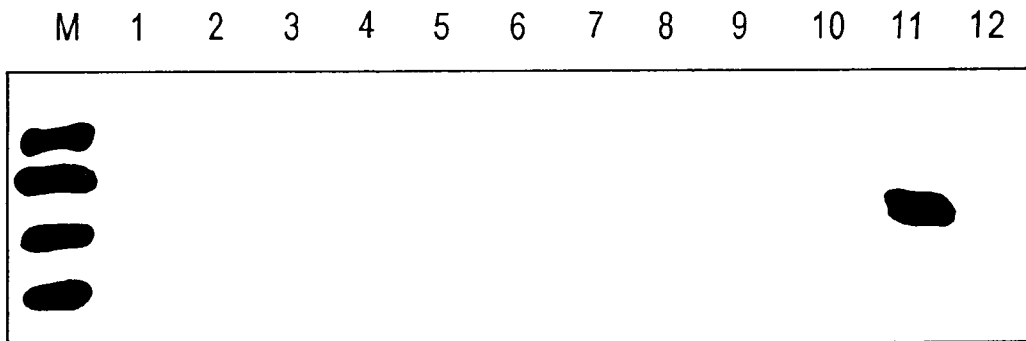


FIG. 4

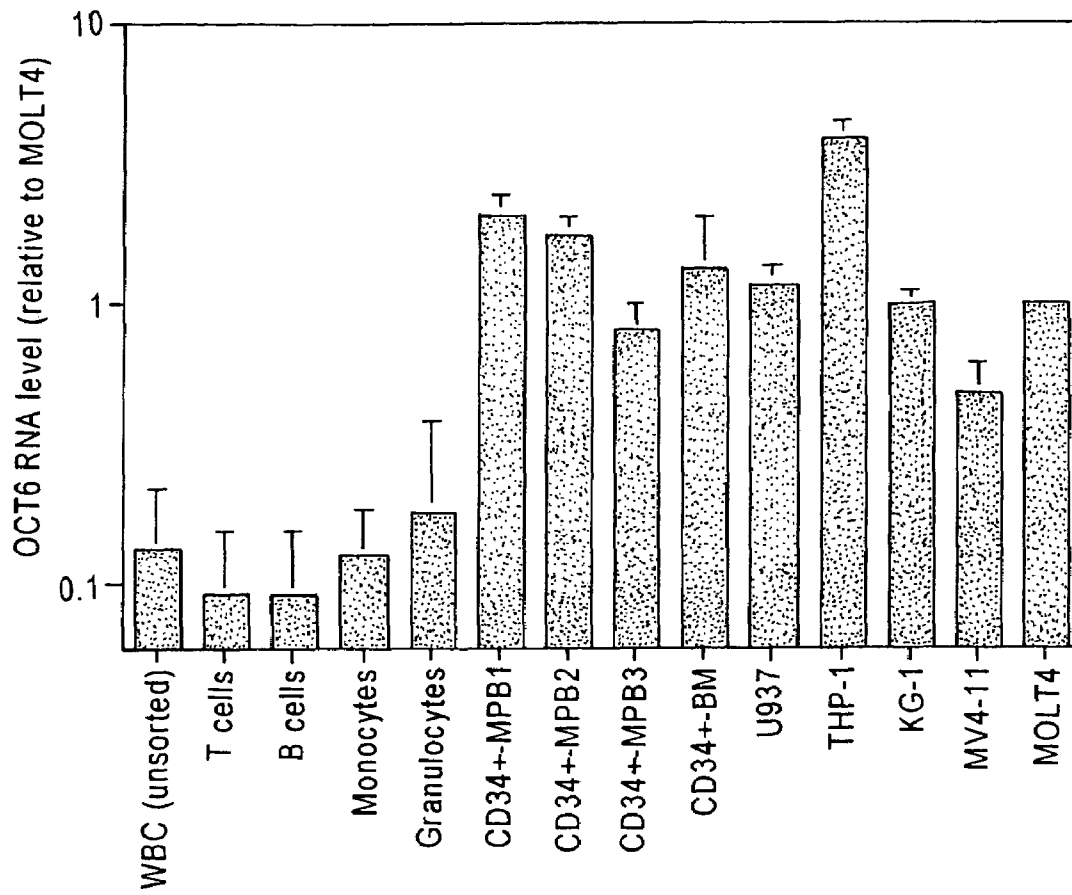
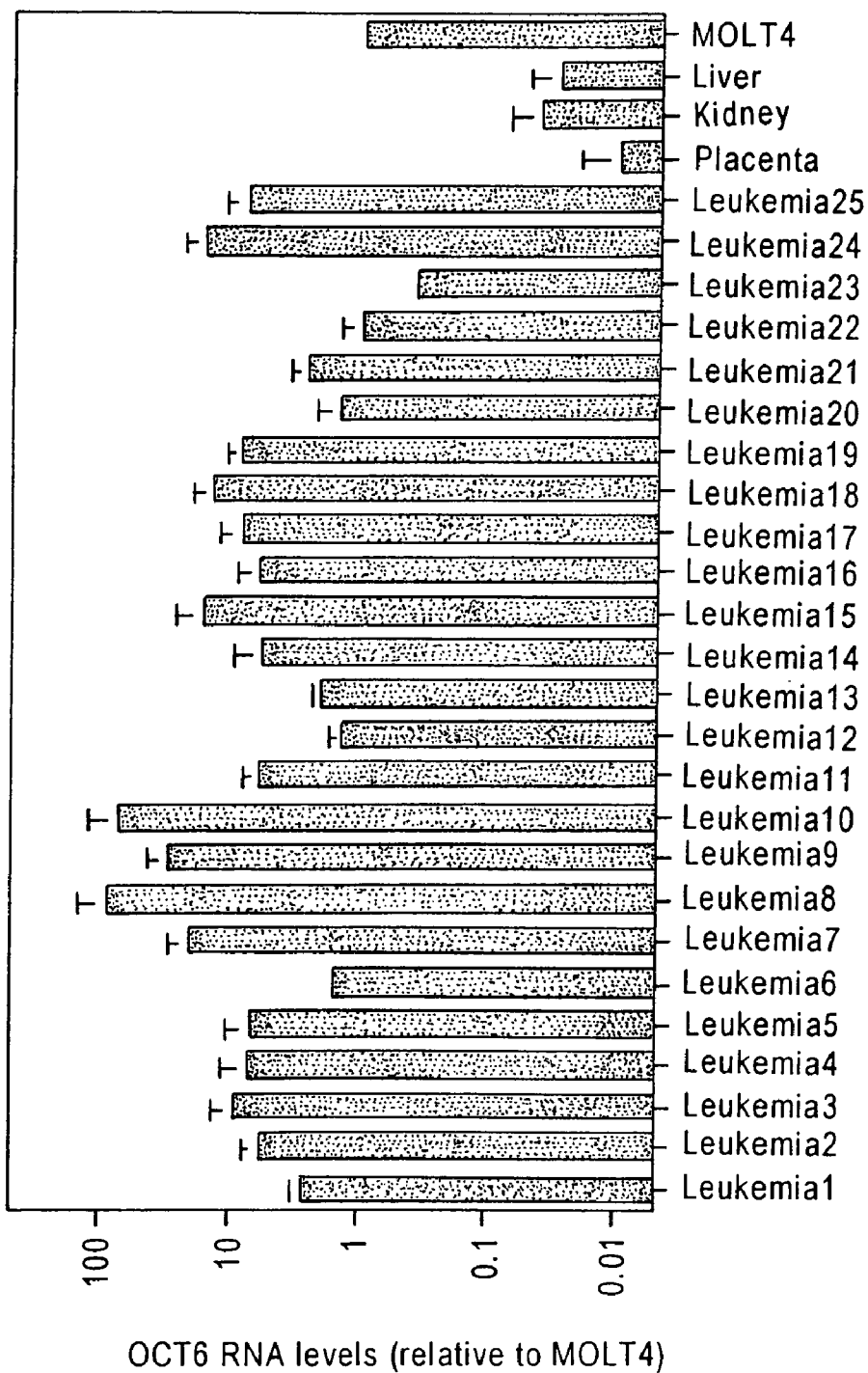


FIG. 5



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**ORGANIC CATION TRANSPORTER  
PREFERENTIALLY EXPRESSED IN  
HEMATOPOIETIC CELLS AND LEUKEMIAS  
AND USES THEREOF**

CROSS REFERENCE TO RELATED  
APPLICATIONS

This application is a divisional application of and claims the benefit of application Ser. No. 10/849,551, filed May 20, 2004 now abandoned, which claims the benefit of U.S. Provisional Application No. 60/471,709, filed May 20, 2003.

FIELD OF THE INVENTION

The invention relates to a gene encoding an organic cation transporter, OCT6, and its use as a target for the treatment of hematological malignancies, and in particular, leukemia. The invention further relates to screening methods for identifying agonists and antagonists/binding partners of OCT6 transport activity.

BACKGROUND OF THE INVENTION

The lipid bilayer of the cellular membrane insulates the intracellular milieu from exposure to hydrophilic compounds. Unlike lipophilic compounds that can diffuse through cellular membranes, water-soluble compounds usually require specific transport mechanisms to gain access to the intracellular space. The regulation of the traffic of polar compounds in both directions across the cellular membrane is a complex process involving several large families of transport proteins.

Most often in cancer research, drug transport is thought of as a mechanism of cellular drug resistance, as drug efflux pumps such as the products of the MDR1 and MRP genes have been shown to be mechanisms of resistance to lipid-soluble anticancer drugs. However, drug transport is a two-way street, and mechanisms also exist for pumping drugs into cells. For polar, water-soluble anticancer agents, drug uptake, and not drug efflux, is the critical determinant of cellular drug accumulation.

Most cancer chemotherapy employs drugs that are lipid-soluble that can easily penetrate the cell membrane of cancer cells. One advantage of using lipid-soluble drugs is that they easily gain intracellular access to different types of cancer cells, so many cancer cells appear to be initially sensitive to these drugs. The disadvantage is that cancer cells learn to increase the activity of drug efflux pumps in the cell membrane to pump lipid-soluble drugs out of the cell, resulting in drug resistance.

In contrast, potential water-soluble anticancer drugs may not survive the preclinical screening process since there is a great deal of variability in the expression of drug transport genes in different types of cancer cells. Variability in transport gene expression may result in variability in accumulation of polar, water-soluble drugs. One approach to more effectively utilize water-soluble anticancer drugs is to identify which of the dozens of transport genes are actually expressed in tumors.

The importance of carrier-mediated anticancer drug uptake is exemplified in reduced folate carrier (RFC) mediated uptake of methotrexate (MTX). Methotrexate (MTX), a reduced folate analogue, is scavenged and retained in cells by mechanisms designed to secure folates from the environment. The major mechanism of MTX uptake at pharmacologic concentrations is the reduced folate carrier (RFC), an OAT trans-

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porter with a  $K_m$  for MTX between approximately 0.8-26  $\mu\text{M}$ . Decreased RFC activity has been observed in several in vitro models of transport-mediated MTX resistance (Biochem. Pharmacol. 11: 1233-1234, 1960). Once rodent and human genes encoding proteins with RFC activity were isolated, the molecular explanations for decreased RFC activity emerged. RFC1 transfection into the transport-deficient MTX<sup>R</sup> ZR75 cell line resulted in a 20-fold increase in 6-hour MTX uptake and a concomitant 250-fold increase in sensitivity to MTX relative to control cell clones, showing that the RFC1 gene reconstitutes RFC activity and has a significant impact on MTX cytotoxicity (Moscow, et al., Cancer Res. 55: 3790-3794, 1995).

In different cell lines, MTX transport deficiency has been ascribed either to mutations in the RFC gene or in decreased expression of the RFC gene product. Several studies have demonstrated that RFC1 gene expression is an important determinant of sensitivity to MTX. In in vitro studies, we have found that RFC1 RNA levels correlate with MTX sensitivity in a panel of non-selected cell lines, including breast cancer cell lines (Moscow et al., Int J Cancer. 72: 184-190, 1997).

A plethora of genes with the ability to transport MTX out of the cell have been reported, including MRP1, MRP2, MRP3, MRP4, the organic anion transporters hOAT2 and hOAT3, and the mitoxantrone-resistance protein (BCRP/MXR). However, despite the multitude of MTX export genes, clinical studies have shown a relationship between the expression of RFC1, the mechanism of MTX uptake, and prognosis in Acute Lymphoid Leukemia (ALL) and osteosarcoma. As a result, RFC1 expression and MTX uptake are now implicated as determinants of clinical sensitivity in several types of tumors. Thus, the role of RFC1 in mediating sensitivity of its cytotoxic drug substrates has become a prototype that illustrates the potential role of transporters, like OAT and OCT genes, in determination of anticancer drug selectivity and toxicity.

However, there is a need to identify additional channels, or transporters, that are found in specific cancers, to enable the targeting of different cancers with anticancer agents that are substrates for those transporters.

SUMMARY OF THE INVENTION

The present invention is directed towards a membrane protein that functions to transport hydrophilic substances across cellular membranes. The protein, OCT6, is a new member of the organic cation transporter (OCT) family (SLC22 gene family). Tissue distribution of this protein is distinct from other OCT protein family members; being detected in leukemia, leukemia blast cells and CD34+ cells.

In one aspect, the present invention provides a novel target for hematological malignancies such as leukemia, an OCT6 transporter.

In another aspect of the present invention there is a method for screening potential substrates that selectively bind the OCT6 transporter. The method involves contacting a cell which overexpresses an OCT6 transporter gene with a test compound and determining whether the test compound is a substrate for the OCT6 transporter.

In another aspect, there is a method for screening potential anti-cancer agents in a cell overexpressing an OCT6 transporter gene. The method comprises determining viability of a cell which expresses OCT6 transporter gene incubated in the presence and absence of a test compound and identifying the test compound as a potential anti-cancer agent if there is cellular influx of the test compound and cell death.

In another aspect of the invention, a test kit is provided for screening candidate drugs for hematologic malignancies comprising a mammalian cell line or cells which overexpress OCT6, a control substrate and a detectable substance.

In still another aspect of the invention, there are immunogenic compositions for treating hematological malignancies. In a preferred embodiment, immunogenic compositions for treating leukemia comprise a substrate that binds selectively to a leukemia cell expressing the OCT6 transporter gene. In another preferred embodiment of the invention, the substrate comprises an antibody that selectively binds to the OCT6 transporter protein. Preferably, the OCT6 transporter protein allows cellular uptake of the substrate which then causes cell death. In one embodiment the substrate is cytotoxic and in another preferred embodiment the substrate is coupled with a cytotoxic agent.

In still another aspect, the present invention provides a method for impairing a leukemia cell comprising contacting the cell with a cytotoxic OCT6 transporter protein. In one embodiment the substrate is a cytotoxin and in another embodiment the substrate is coupled to a cytotoxic agent.

In yet another aspect, the present invention provides a method for treating hematological malignancies comprising administering to a subject in need thereof an immunogenic composition comprising a substrate that binds selectively to a cell expressing the OCT6 transporter gene. In a preferred embodiment the OCT6 transporter protein allows cellular uptake of the substrate which then causes cell death. In another preferred embodiment the substrate is cytotoxic. In another preferred embodiment, the substrate is coupled with a cytotoxic agent.

#### BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1. A. shows the predicted hydrophathy profile of OCT6.

FIG. 1. B. is a dendrogram showing phylogenic relationship between OCT6 (SEQ ID NO:2) and other OCT and OAT proteins, including, OCTN1 (SEQ ID NO:4), OCT3 (SEQ ID NO:5), OCTN2 (SEQ ID NO:6), OCT2 (SEQ ID NO:7), OCT1 (SEQ ID NO:8), OAT5 (SEQ ID NO:9), OAT4 (SEQ ID NO:10), OAT3 (SEQ ID NO:11), and OAT1 (SEQ ID NO:12).

FIG. 2A-F. is the CLUSTLAW alignment of OCT6 and other OCT and OAT proteins. The bottom row represents areas of consensus.

FIG. 3. shows the normal tissue distribution of OCT6 RNA determined by RT-PCR using a cDNA panel. Only 1000x (highest) cDNA concentration is shown. Panel A. 1, salivary gland; 2, thyroid; 3, adrenal; 4, pancreas; 5, ovary; 6, uterus; 7, prostate; 8, skins; 9, peripheral blood leukocytes; 10, bone marrow; 11, fetal brain; 12, fetal liver. Panel B. 1, brain; 2, heart; 3, kidney; 4, spleen; 5, liver; 6, colon; 7, lung; 8, small intestine; 9, muscle; 10, stomach, 11, testis; 12, placenta.

FIG. 4. shows quantitative RT-PCR for the transporter gene OCT6 performed with RNA extracted from peripheral blood leukocytes, CD34+ cells and additional hematopoietic cell lines. Fresh discarded buffy coats that were twice sorted by FACS using CD14 (monocytes), CD15 (granulocytes), CD3 (T-cells) and CD20 (B-cells). Purities of 99% or better were obtained. For peripheral WBC and sorted subsets, the average±SD represent pooled results from samples from 2 individuals performed in triplicate or quadruplicate. For CD34-selected mobilized peripheral blood (MPB), the results from each of 3 individuals are shown. For CD34-selected bone marrow (CD34+BM), the results are from one individual. OCT6 levels were normalized to the expression of

actin RNA, as a control for equivalence of mRNA template. The units, in log scale, are arbitrary and based on a standard curve of OCT6 RT-PCR in serially diluted HL60 RNA. Unity is defined as the level of OCT6 RNA found in MOLT4 cells.

FIG. 5. shows quantitative RT-PCR for the gene OCT6 using RNA extracted from leukemic blasts obtained from patients at the time of initial diagnosis. OCT6 levels were normalized to the expression of actin RNA, as a control for equivalence of mRNA template. The OCT6 RNA levels in placenta, liver, kidney and MOLT-4 cell line were determined concurrently and shown for comparison. The units, in log scale, are arbitrary and based on a standard curve of OCT6 RT-PCR in serially diluted HL60 RNA. Unity is defined as the level of OCT6 RNA found in MOLT4 cells.

#### DETAILED DESCRIPTION OF THE INVENTION

The present invention is based on the discovery and isolation of a new member of the SLC22 gene family (the OCT family of proteins) that is unusual for its distinct pattern of tissue distribution. Rather than the typical high levels of expression in liver, kidney or placenta, high levels of RNA for this transporter were found in some leukemia cell lines, in CD34+ cells, and in circulating leukemia blast cells.

All patents, patent applications and literature cited in this description are incorporated herein by reference in their entirety. In the case of inconsistencies, the present disclosure, including definitions, will prevail.

#### OCT Family

Two families of proteins involved in maintaining homeostasis of charged organic compounds are the organic anion transporters (OATs) which carry the SLC21 designation and the organic cation transporters (OCTs), which carry the SLC22 designation (See Table 1). OATs and OCTs each have characteristic patterns of tissue expression, with predominant expression in a tissue involved in the transport of xerobiotics, i.e., liver, kidney or placenta.

TABLE 1

Organic anion and cation transported genes			
Gene Family	Gene Name	Locus Link	Alternative Names
SLC21	SLC21A1	6577	
	SLC21A2	6578	PGT
	SLC21A3	6579	OATP, OATP1, OATP1b, OATP-A
	SLC21A4	28237	OAT-K1, OAT-K2
	SLC21A5	28236	OATP2, OATP-2
	SLC21A6	10599	LST-1, OATP-C
	SLC21A7	28235	OATP3, OATP-3
	SLC21A8	28234	LST2, OATP8, SLC21A8, OATP-8
	SLC21A9	11309	OATP-B
	SLC21A10	28233	OATP4
	SLC21A11	28232	OATP-D
	SLC21A12	28231	LOC51737, OATP-E, POAT
	SLC21A13	28230	OATP5, OATP-5
	SLC21A14	53919	OATP-F
SLC22	SLC22A1	6580	OCT1
	SLC22A2	6582	OCT2
	SLC22A3	6581	OCT3
	SLC22A4	6583	OCTN1
	SLC22A5	6584	OCTN2, CDSP, SCD
	SLC22A6	9356	NKT, OAT1, OAT-1
	SLC22A7	10864	NLT, OAT2, OAT-2
	SLC22A8	9376	OAT3, OAT-3
	SLC22A9		OAT4, OAT-4

The OAT and OCT carriers result in increased cellular accumulation of their respective substrates, despite the fact

that they are carriers that mediate facilitative diffusion. For carriers, the degree of intracellular accumulation may not exceed the extracellular concentration. However, the presence of the carrier allows uptake in comparison to no uptake in the absence of the carrier, and drugs that bind an intracellular target or which are chemically modified in the cells, e.g., by phosphorylation or polyglutamylation, may be eliminated from the substrate pool and not available for transport back across the cellular membrane.

The first five members of the SLC22 family of transporters, OCT1, OCT2, OCT3, OCTN1, and OCTN2, have been characterized as organic cation transporters. The uptake of many cations, such as tetraethylammonium (TEA), N-1-methylnicotinamide (NMN), choline, procainamide, amantadine and morphine are mediated by these polyspecific transporters. In general, these transporters are potential-dependent, but independent of sodium and proton gradients. These genes are all characterized by the presence of 11 or 12 transmembrane domains, as predicted by hydrophobicity analysis, and all have a large hydrophilic loop between transmembrane domain (TMD) 1 and TMD2.

OCT substrates are shown below in Table 2. Tetraethyl ammonium (TEA) is the classic substrate for OCT transporters. In addition, OCT1, OCT2 and OCT3 transport 1-methyl-4-phenylpyridinium (MPP). Compared to OCT2, OCT1 has a higher affinity for some cations (for example mepiperphenidol and procainamide), a similar affinity for others (for example, decynium 22 and quinidine), and a lower affinity for corticosterone (See Koepsell et al., *Ann. Rev. Physiol.* 60: 243-266, 1998.). OCT3 is an electrogenic transporter for TEA and guanidine. Other physiologic substrates for OCT transporters include dopamine, histamine, epinephrine and norepinephrine, acetylcholine and 5-hydroxytryptamine (Burckhardt, et al., *Am J Physiol Renal Physiol.* 278: F853-66., 2000.), suggesting an important role for these transporters in the central nervous system, in addition to their role in hepatic and renal clearance. Interestingly, despite its cationic nature, recent studies have identified cimetidine as a selective inhibitor, but not a substrate for several organic cation transporters, including rOCT1, rOCT2, rOCT3, hOCTN1, and hOCTN2.

TABLE 2

Common Name	OCT Substrates			
	Gene Name	Cell Type	Substrate	KT (uM)
OCT1	SLC22A1	HeLa	TEA	229
OCT1	SLC22A1	Xenopus	MPP	14.6
OCT2	SLC22A2	Xenopus	Norepinephrine	1900
OCT2	SLC22A2	Xenopus	Histamine	1300
OCT2	SLC22A2	Xenopus	Dopamine	390
OCT2	SLC22A2	Xenopus	Serotonin	80
OCT2	SLC22A2	HEK293	MPP	16
OCT2	SLC22A2	HEK293	Dopamine	330
OCT2	SLC22A2	Xenopus	Amantadine	27
OCT2	SLC22A2	Xenopus	Memantine	34
OCT3	SLC22A3	HeLa	TEA	2500
OCT3	SLC22A3	HRPE	MPP	47
OCTN1	SLC22A4	Fibroblasts	L-Carnitine	6.6
OCTN2	SLC22A5	HEK293	L-Carnitine	4.34
OCTN2	SLC22A5	HEK293	L-Carnitine	4.3
OCTN2	SLC22A5	HEK293	D-Carnitine	10.9
OCTN2	SLC22A5	HEK293	Acetyl-L-carnitine	8.5
OCTN2	SLC22A5	Xenopus	L-Carnitine	4.8
OCTN2	SLC22A5	Xenopus	D-Carnitine	98
OCTN2	SLC22A5	JAR	L-Carnitine	3.5

OCT1 and OCT2 are predominantly expressed in the kidney and liver. These transporters are located on the basolateral surface of renal tubules and, therefore, play a role in the removal of organic cations from the blood. OCT3 is most abundantly expressed in placenta. In addition, other tissue-specific roles have been implicated for these transporters. As noted above, OCTs may play a role in transport of endogenous neuroleptic substrates, and OCT3 has been implicated in the disposition of cationic neurotoxins and neurotransmitters in the brain (Wu, et al., *J Biol Chem.* 273: 32776-86, 1998). Dhillon et al. (*Clin Pharmacol Ther.* 65: 205, 19996) used RT-PCR followed by functional transport studies (TEA) to identify OCT1 expression in a human mammary epithelial cell line (MCF12A). Further, the OCT1 gene has been shown to be up regulated in lactating mammary epithelial cells.

The OCTN1 gene, cloned from a cDNA, shows sequence similarity to organic cation transporter genes, which is highly expressed in kidney as well as trachea, bone marrow and fetal liver. Recombinant OCTN1 expressed in mammalian cells exhibited saturable uptake of TEA that was pH sensitive. Several others suggest that OCTN1 is a renal proton/organic cation antiporter functioning at the epithelial apical membrane. The uptake of pyrilamine, quinidine, verapamil and L-carnitine were increased by expression of OCTN1 in *Xenopus* oocytes.

Another OCT protein family member, OCTN2, cloned from a human placental trophoblast cell line, is expressed widely in human tissues including kidney, placenta and heart. OCTN2 is more closely related to OCTN1 than to OCT1, OCT2 and OCT3 (*Biochem Biophys Res Commun.* 246: 589-95, 1998). Transfection of OCTN2 has demonstrated its role in the transport of TEA and carnitine. OCTN2-mediated transport of TEA is sodium independent, whereas transport of carnitine is sodium-dependent. The role of sodium in OCTN2-mediated carnitine transport not only involves the electrogenic gradient, but the presence of sodium also alters the affinity of OCTN2 for carnitine. Germline mutations of OCTN2 result in primary carnitine deficiency, a syndrome of progressive cardiomyopathy and skeletal myopathy. The symptoms associated with this syndrome are thought to result not only from generalized carnitine deficiency from decreased renal carnitine reabsorption, but also from inability of cardiac and skeletal myocytes, which ordinarily express OCTN2, to accumulate carnitine. This syndrome demonstrates that tissue-specific OCT-mediated transport is essential for accumulation of required cations in specific tissues.

The present invention identifies a new transport protein in the OCT family, OCT6, preferentially expressed in leukemia cell lines, leukemia blast cells and CD34+ cells. The cell surface localization and the transporter function of the OCT6 gene product suggest its usefulness as a target in the diagnosis and treatment of hematologic malignancies.

As used herein, the term "antibody" refers to an immunoglobulin molecule with a specific amino acid sequence evoked in by an antigen, and characterized by reacting specifically with the antigen in some demonstrable way.

As used herein, the term "carrier" refers to a diluent, adjuvant, excipient, or vehicle with which the compositions of the present invention are administered.

As used herein, "compound" refers to any agent, chemical, substance, or substrate, whether organic or inorganic, or any protein including antibodies, peptides, polypeptides, peptoids, and the like.

As used herein, the term "cytotoxin" or cytotoxic agent includes any specific substance, which may or may not be antibody, that inhibits or prevents the functions of cells, causes destruction of cells, or both.

As used herein, the term “derivative” refers to something produced by modification of something pre-existing; for example, a substance or chemical compound that may be produced from another substance or compound of similar structure in one or more steps.

As used herein, the term “fragment” refers to a part of a larger entity, said larger entity comprising by non-limiting example, an antibody, compound or substance.

As used herein, the term “leukemia blast” or “leukemic blast” refers to lymphoblasts, the abnormal immature white blood cells associated with leukemia.

As used herein, the term “monoclonal antibody” is not limited to antibodies produced through hybridoma technology. The term “monoclonal antibody” refers to an antibody that is derived from a single clone, including any eukaryotic, prokaryotic, or phage clone, and not the method by which it is produced.

As used herein, the term “pharmaceutically acceptable carrier” refers to a carrier that may be administered to a subject, together with one or more liver protecting agents and one or more mushroom powder or extract of the present invention, and which does not destroy the pharmacological activity thereof and is nontoxic when administered in doses sufficient to deliver a therapeutic amount of the compound.

As used herein, the term “substrate” refers to a substance, compound, agent, antibody or derivatives and/or fragment thereof, acted upon by the OCT6 transporter protein (e.g., a substance that is taken across the cellular membrane by action of the OCT6 transporter protein).

OCT6 (SEQ ID NO:1) was first identified as a potential OCT gene by assembling and sequencing ESTs as described in Example 1 (amino acid sequence of OCT6 is SEQ ID NO:2). The gene sequence proved to be identical to the recently submitted cDNA OKB1 (GenBank AF268892) submitted by M. Okabe and T. Abe, incorporated herein in its entirety. It is also contained within the submitted BAC clone CTA-331P3 (SEQ ID NO: 3) (GenBank AC002464) located at chromosome 6q21, incorporated herein in its entirety. The gene has a predicted protein structure typical of transport proteins with two groups of six transmembrane domains separated by a hydrophilic region (FIG. 1A). CLUSTALW alignment produced a dendrogram showing the phylogenetic relationship between OCT6 and other OAT and OCT proteins (FIG. 1B). This dendrogram suggests that the distinction between OAT and OCT genes, based on functional studies, obscures the common origin of both families of transporters. The actual CLUSTALW alignment of these genes is shown in FIG. 2 and demonstrates multiple regions of conservation among all of these genes.

Next, according to the methods described in Example 3, quantitative RT PCR analysis of the expression of OCT6 was performed, along with the expression of other OCT genes, in 50 cell lines. The results are shown in Table 3. The two highest expressing cell lines for OCT6 in this panel were two leukemia cell lines, HL60, a human promyelocytic leukemia cell line, and MOLT4, a human acute lymphoblastic leukemia (T-cell) cell line. There was only a low level of expression detected in most of the other cell lines.

TABLE 3

OCT expression in 50 cell lines of the NCI Drug Screen							
No.	Cell Line	source	OCT1	OCT2	OCT3	OCTN 2	OCT6
1	CCRF-CEM	Leukemia	0.7	0.7	0.2	0.1	5.7
2	HL-60	Leukemia	0.5	1.3	0.0	0.4	716
3	K-562	Leukemia	1.4	1.2	0.2	1.4	5.2
4	MOLT-4	Leukemia	0.1	1.1	0.5	0.6	46.8
5	RPMI-8226	Leukemia	2.8	2.0	0.1	3.7	6.02
6	SR	Leukemia	1.9	1.1	0.0	0.3	2.6
7	A549/ATCC	Lung cancer	1.7	1.2	161	4.3	1.2
8	HOP-62	Lung cancer	0.8	4.8	0.6	2.4	4.1
9	NCI-H226	Lung cancer	4.8	0.5	0.1	21.1	4.8
10	NCI-H23	Lung cancer	0.5	0.7	0.0	0.3	5.2
11	NCI-H460	Lung cancer	0.7	1.0	0.0	1.7	1.8
12	COLO205	Colon Ca.	4.9	5.3	30.9	2.2	3.6
13	HCC-2998	Colon Ca.	1.5	1.0	0.0	2.6	5.4
14	HCT-116	Colon Ca.	1.7	2.1	0.1	2.8	9.7
15	HCT-15	Colon Ca.	0.9	1.7	0.1	3.5	4.2
16	HT-29	Colon Ca.	1.9	1.2	18.1	1.5	1.5
17	KM-12	Colon Ca.	0.6	1.0	12.2	0.7	2.1
18	SW-620	Colon Ca.	1.0	2.6	40.4	1.9	3.7
19	SF-268	CNS Tumor	0.4	0.8	0.0	0.9	2
20	SF-295	CNS Tumor	0.5	1.2	0.2	1.1	2.5
21	SF-539	CNS Tumor	0.5	0.6	2.3	0.2	5.3
22	SNB-75	CNS Tumor	0.8	1.8	0.0	0.6	2.3
23	U251	CNS Tumor	0.8	0.9	0.0	0.6	7.4
24	LOCIMVI	Melanoma	2.9	2.1	0.1	0.4	3.6
25	MALME-3M	Melanoma	1.5	1.5	0.0	2.3	3
26	M14	Melanoma	1.9	1.4	0.0	1.9	4.7
27	SK-MEL-2	Melanoma	2.1	1.9	0.0	2.2	3.9
28	SK-MEL-5	Melanoma	2.6	1.5	0.0	1.9	2.7
29	UACC-257	Melanoma	3.2	3.6	0.0	1.1	5.4
30	IGROV1	Ovarian Ca.	4.9	5015	17.9	1.8	2.5
31	OVCAR-3	Ovarian Ca.	1.4	0.1	0.0	2.2	14
32	OVCAR-4	Ovarian Ca.	2.6	1.4	0.0	8.9	3.4
33	OVCAR-5	Ovarian Ca.	3.5	2.7	105	10.0	4.8
34	OVCAR-8	Ovarian Ca.	1.1	1.0	0.0	0.8	1.6
35	SK-OV-3	Ovarian Ca.	3.9	1995	9.2	8.5	9.8
36	A498	Renal Ca.	2.2	13.4	180	4.7	1.3
37	ACHN	Renal Ca.	1.1	1.1	0.7	1.2	1.1
38	CAKL_1	Renal Ca.	3.5	2.5	4.8	1.8	2.8



TABLE 3-continued

OCT expression in 50 cell lines of the NCI Drug Screen							
No.	Cell Line	source	OCT1	OCT2	OCT3	OCTN 2	OCT6
39	RXF-393	Renal Ca.	1.7	1.2	3.0	0.6	1.2
40	TK-10	Renal Ca.	3.6	5.0	16.8	2.5	8
41	UO-31	Renal Ca.	4.4	1.6	31.2	1.2	2.3
42	PC-3	Prostate Ca.	2.1	0.8	9.6	3.3	4.7
43	DU-145	Prostate Ca.	1.1	1.1	3.4	1.6	3
44	MCF-7	Breast Ca.	0.8	1.8	0.0	10.4	3.5
45	NCI/ADR-RES	Breast Ca.	1.4	1.3	1.1	2.0	2.1
46	MDA-MB-231	Breast Ca.	1.2	0.4	3.9	4.8	1.8
47	HS578T	Breast Ca.	1.0	1.5	0.0	1.2	8.3
48	MDA-MB-435	Breast Ca.	1.9	0.6	0.1	0.7	2.7
49	BT-549	Breast Ca.	1.2	0.8	0.1	0.3	2.6
50	T-47D	Breast Ca.	0.7	1.1	0.1	4.2	8.7

OCT6 is unique among the known members of OCT and OAT genes because of its pattern of tissue distribution. The pattern of expression of the OCT6 gene in the 50 cell lines suggested that its expression might be restricted to hematopoietic tissues. The restricted pattern of expression observed for OCT6 also suggests that therapies using OCT6-specific substrates are unlikely to have widespread toxicity to normal tissues. Therefore, we examined OCT6 expression in a cDNA panel representing a wide cross-section of normal tissues according to the methods of Example 4 (FIG. 3). This study revealed that OCT6 RNA levels are highest in testis and fetal liver, with lower but detectable levels in peripheral blood leukocytes and bone marrow. Since fetal hematopoiesis occurs in the liver, it is possible that the fetal liver sample may have included both hepatocytes and hematopoietic cells. OCT6 RNA levels were also barely detectable in pancreatic and adrenal tissue. Unlike other OCT genes, expression was not detectable in liver, kidney or placenta.

To determine whether OCT6 RNA expression in hematopoietic cells was lineage-specific, leukocytes were sorted from discarded buffy coat specimens by flow cytometry, and purified subpopulations were examined for OCT6 RNA expression according to the methods described in Example 5. OCT6 expression was also examined in a population of CD34+ cells. As can be seen in FIG. 4, the expression of OCT6 was highly enriched in CD34+ cells in comparison to the other cell populations. Also, significant levels of OCT6 expression (relative to MOLT4) were found in other hematopoietic cell lines: U937, a human histiocytic lymphoma cell line; THP-1, a human acute monocytic leukemia cell line; KG-1, a human erythroleukemia cell line; and MV-4-11, a human biphenotypic (B-cell and myelomonocytic) leukemia cell line.

The high levels of OCT6 RNA in some leukemia cell lines and CD34+ cells also raised the question as to whether this gene was highly expressed in actual leukemias. To address this issue, the RNA levels of OCT6 in 25 samples of peripheral leukemic cells were measured according to the methods set out in Example 6. The FAB classification of these samples are shown in Table 4. These results are shown in FIG. 5, and demonstrate that the majority of specimens contained RNA levels for OCT6 that exceeded the level found in MOLT4 cell line, the second highest expressing cell line among those examined, and exceed by orders of magnitude the levels found in placenta, kidney and liver.

TABLE 4

Phenotypes of leukemia specimens	
Sample Number	Description
1	CML, blast crisis
2	CML, blast crisis
3	CML, stable phase
4	CML, probably stable phase
5	CML, accelerated phase
6	ALL
7	ALL
8	AML
9	ALL
10	ALL
11	ALL
12	AML
13	AML
14	AML
15	AML
16	ALL, biphenotypic
17	ALL, biphenotypic
18	AML
19	AML, M2
20	AML, M2
21	AML, M4
22	AML, M4
23	AML, M1
24	AML
25	AML, M4

Due to the OCT6 protein's location on the cellular membrane and its function as an intracellular transporter, the OCT6 transporter protein has been identified as a therapeutic target. Basic principles of cellular pharmacology suggest that increase in intracellular accumulation will lead to increased intracellular effect. For anticancer drugs, this principle has been studied extensively in the context of lipophilic drugs, which require no specific mechanism for cellular uptake, and export pumps such as the product of the multidrug resistance gene, MDR1, whose overexpression of MDR1 leads to increased cellular resistance by decreasing intracellular concentrations of drug (Moscow, J. A., Schneider, E. S., Ivy, S. P., and Cowan, K. H. Multidrug resistance. In: H. M. Pinedo, D. L. Longo, and B. A. Chabner (eds.), Cancer chemotherapy and biological response modifiers. Annual 17. New York: Elsevier, 1997). The same principle applies to charged, hydrophilic drugs of the present invention, except that the determinants of sensitivity depend on uptake as opposed to efflux. As such, cells overexpressing an OCT6 transporter are likely to be highly sensitive to cytotoxic OCT6 substrates.

## Drug Screening

Accordingly, the present invention provides methods for screening potential substrates of, and potential therapeutic agents against hematological malignancies like leukemia that overexpress, the OCT6 transporter. In particular, potential therapeutic agents are screened for the ability to be a substrate recognized by an OCT6 transporter protein. Preferably, potential substrates are screened for the ability to confer cytotoxic effects on a cell overexpressing OCT6 transporter protein. More preferably, agents are screened for the ability to preferentially cause cellular uptake into, and cell death of, cells overexpressing the OCT6 transporter. Most preferably, the agents are screened for the ability to cause cell death of cancer cells such as leukemia overexpressing the OCT6 transporter as compared to normal cells.

A method for screening potential substrates of the OCT6 transporter protein comprises providing a cell or cell line which expresses OCT6 and a test compound, incubating the test compound and cell line and analyzing the cell or cell line to determine if there was a cellular influx of the test compound. Analysis of the cell line to determine whether cellular uptake of the test compound occurred can be accomplished by any means known in the art. For example, a test compound can be tagged with a detectable label prior to contact with a cell and then observed under microscopy or by other means for its location. Non-limiting examples of labels include green fluorescent protein, alkaline phosphatase, horseradish peroxidase, rease,  $\beta$ -galactosidase, CAT, luciferase, an immunogenic tag peptide sequence, an extrinsically activatable enzyme, an extrinsically activatable toxin, an extrinsically activatable fluor, an extrinsically activatable quenching agent, a radioactive element or an antibody.

A method for screening candidate anti-cancer agents comprises determining the viability of a mammalian cell which expresses OCT6 incubated in the presence and absence of a test compound and identifying the test compound as a potential anti-leukemia agent if there is a cellular uptake of the test compound and cell death. Analysis of cell viability can be accomplished by any means known in the art.

It is well known in the art that viability of a cell can be determined by contacting the cell with a dye and viewing it under a microscope. Viable cells can be observed to have an intact membrane and do not stain, whereas dying or dead cells having "leaky" membranes do stain. Incorporation of the dye by the cell indicates the death of the cell. The most common dye used in the art for determining viability is trypan blue. Viability of cells can also be determined by detecting DNA synthesis. Cells can be cultured in cell medium with labeled nucleotides (e.g.,  $^3\text{H}$  thymidine). The uptake or incorporation of the labeled nucleotides indicates DNA synthesis and cell viability. In addition, colonies formed by cells cultured in medium indicate cell growth and is another means to test viability of the cells.

Identification and/or observation of cells undergoing apoptosis can be another method of determining cell viability. Apoptosis is a specific mode of cell death recognized by a characteristic pattern of morphological, biochemical, and molecular changes. Cells going through apoptosis appear shrunken, and rounded; they also can be observed to become detached from culture dish. Thermophological changes involve a characteristic pattern of condensation of chromatin and cytoplasm which can be readily identified by microscopy. When stained with a DNA-binding dye, such as H33258, apoptotic cells display classic condensed and punctate nuclei instead of homogeneous and round nuclei.

The hallmark of apoptosis is the endonucleolysis, a molecular change in which nuclear DNA is initially degraded

at the linker sections of nucleosomes to give rise to fragments equivalent to single and multiple nucleosomes. When these DNA fragments are subjected to gel electrophoresis, they reveal a series of DNA bands which are positioned approximately equally distant from each other on the gel. The size difference between the two bands next to each other is about the length of one nucleosome (i.e., 20 base pairs). This characteristic display of the DNA bands is called a DNA ladder and it indicates apoptosis of the cell. Apoptotic cells can be identified by flow cytometric methods based on measurement of cellular DNA content, increased sensitivity of DNA to denaturation, or altered light scattering properties. These methods are well known in the art and are within the contemplation of the invention.

Abnormal DNA breaks are also characteristic of apoptosis and can be detected by any means known in the art. In one embodiment, DNA breaks are labeled with biotinylated dUTP (b-dUTP). Cells are fixed and incubated in the presence of biotinylated dUTP with either exogenous terminal transferase (terminal DNA transferase assay; TdT assay) or DNA polymerase (nick translation assay; NT assay). The biotinylated dUTP is incorporated into the chromosome at the places where abnormal DNA breaks are repaired, and are detected with fluorescein conjugated to avidin under fluorescence microscopy.

## Kits

The present invention provides kits that can be used in the above screening methods. In one embodiment, a kit comprises a substantially isolated polypeptide comprising an OCT6 epitope which is specifically immunoreactive with only test compound(s) that are substrates of the OCT6 transporter protein. Binding of a test compound to the OCT6 epitope is indicative that the test compound is a OCT6 substrate. In another embodiment, a kit comprises a cell line that overexpresses an OCT6 transporter protein. Binding and/or cellular uptake of a test compound via the OCT6 protein is indicative that the test compound is a OCT6 substrate. Preferably, the kits of the present invention further comprise a control compound or antibody which does not react with the OCT6 transporter protein. In another specific embodiment, the kits of the present invention contain a means for detecting the binding of a test compound to an OCT6 epitope and/or cellular uptake of a test compound. For example, the test compound may be conjugated to a detectable substrate such as a fluorescent compound, an enzymatic substrate, a radioactive compound or a luminescent compound, or a second antibody which recognizes the first antibody may be conjugated to a detectable substrate.

The detectable substance may be coupled or conjugated either directly to the test compound (or fragment thereof) or indirectly, through an intermediate (such as, for example, a linker known in the art) using techniques known in the art. See, for example, U.S. Pat. No. 4,741,900 for metal ions which can be conjugated to antibodies for use as diagnostics according to the present invention. Further non-limiting examples of detectable substances include various enzymes, prosthetic groups, fluorescent materials, luminescent materials, bioluminescent materials, radioactive materials, positron emitting metals using various positron emission tomographies, nonradioactive paramagnetic metal ions, immunogenic tag peptide sequences, extrinsically activatable toxins, extrinsically activatable quenching agents, or antibodies. Non-limiting examples of suitable materials include horseradish peroxidase, alkaline phosphatase, beta-galactosidase, or acetylcholinesterase; examples of suitable prosthetic group complexes include streptavidin/biotin and avidin/bi-

otin; examples of suitable fluorescent materials include umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride or phycoerythrin; an example of a luminescent material includes luminol; examples of bioluminescent materials include luciferase, luciferin, and aequorin; and examples of suitable radioactive material include  $^{125}\text{I}$ ,  $^{131}\text{I}$ ,  $^{111}\text{In}$  or  $^{99}\text{Tc}$ .

#### Immunogenic Compositions

The present invention also provides immunogenic compositions for the treatment of hematological malignancies. Non-limiting exemplary hematological malignancies include, but are not limited to, Hodgkin's disease, leukemia such as, acute lymphoid (lymphocytic or lymphoblastic) leukemia (ALL), acute myeloid (myelogenous or myeloblastic) leukemia (AML), acute lymphoid leukemia, biphenotypic (ALL, biphenotypic), acute undifferentiated leukemia (AUL), chronic myeloid (myelogenous or granulocytic) leukemia (CML), erythroleukemia, granulocytic leukemia, lymphoma, monocytic leukemia, myeloma, myelomonocytic leukemia, myelodysplastic syndromes, non-Hodgkin lymphoma, progranulocytic leukemia.

According to the invention immunogenic compositions for the treatment of hematological malignancies comprise a substrate recognized by an OCT6 transporter protein. Preferably, the substrate is a compound that binds selectively or specifically to a OCT6 transporter protein. In a preferred embodiment, the compound binds selectively to the OCT6 transporter protein encoded by a nucleotide sequence of SEQ ID NO:1. The compound may be a cytotoxin or coupled or conjugated with a cytotoxic agent. Preferably the cytotoxin or cytotoxic agent is a chemotherapeutic agent.

The present invention also provides pharmaceutical compositions. Such compositions comprise a therapeutically effective amount of a compound, and a pharmaceutically acceptable carrier.

Cell surface proteins like the OCT6 transporter can be utilized in antibody-based targeting strategies. In still another aspect of the invention, antibodies can be developed by known methods in the art against the external epitope of OCT6 transporter protein. In a preferred embodiment, antibodies are substrates of the OCT6 protein. The antibodies may be polyclonal antibodies or monoclonal antibodies.

Polyclonal antibodies to an antigen-of-interest can be produced by various procedures well known in the art. For example, a polypeptide of the invention can be administered to various host animals including, but not limited to, rabbits, mice, rats, etc. to induce the production of sera containing polyclonal antibodies specific for the antigen. Various adjuvants may be used to increase the immunological response, depending on the host species, and include but are not limited to, Freund's (complete and incomplete), mineral gels such as aluminum hydroxide, surface active substances such as lysollecithin, pluronic polyols, polyanions, peptides, oil emulsions, keyhole limpet hemocyanins, dinitrophenol, and potentially useful human adjuvants such as BCG (*Bacille Calmette-Guerin*) and *Corynebacterium parvum*. Such adjuvants are also well known in the art.

Monoclonal antibodies can be prepared using a wide variety of techniques known in the art including the use of hybridoma, recombinant, and phage display technologies, or a combination thereof. For example, monoclonal antibodies can be produced using hybridoma techniques including those known in the art and taught, for example, in Harlow et al., *Antibodies: A Laboratory Manual*, (Cold Spring Harbor Laboratory Press, 2nd ed. 1988); Hammerling, et al., in:

*Monoclonal Antibodies and T-Cell Hybridomas 563-681* (Elsevier, N.Y., 1981) (said references incorporated by reference in their entireties).

The present invention further encompasses antibodies or fragments thereof conjugated to a diagnostic or therapeutic agent. The antibodies can be used diagnostically to, for example, monitor the development or progression of a tumor as part of a clinical testing procedure to, e.g., determine the efficacy of a given treatment regimen. Detection can be facilitated by coupling the antibody to a detectable substance. Examples of detectable substances include various enzymes, prosthetic groups, fluorescent materials, luminescent materials, bioluminescent materials, radioactive materials, positron emitting metals using various positron emission tomographies, and nonradioactive paramagnetic metal ions. The detectable substance may be coupled or conjugated either directly to the antibody (or fragment thereof) or indirectly, through an intermediate, such as, for example, a linker known in the art, using techniques known in the art. (See, for example, U.S. Pat. No. 4,741,900 for metal ions which can be conjugated to antibodies for use as diagnostics according to the present invention.) Examples of suitable enzymes include horseradish peroxidase, alkaline phosphatase, beta-galactosidase, or acetylcholinesterase; examples of suitable prosthetic group complexes include streptavidin/biotin and avidin/biotin; examples of suitable fluorescent materials include umbelliferone, fluorescein, fluorescein isothiocyanate, rhodamine, dichlorotriazinylamine fluorescein, dansyl chloride or phycoerythrin; an example of a luminescent material includes luminol; examples of bioluminescent materials include luciferase, luciferin, and aequorin; and examples of suitable radioactive material include  $^{125}\text{I}$ ,  $^{131}\text{I}$ ,  $^{111}\text{In}$  or  $^{99}\text{Tc}$ .

Further, an antibody or fragment thereof may be conjugated to a therapeutic moiety such as a cytotoxin, e.g., a cytostatic or cytotoxic agent, a therapeutic agent or a radioactive metal ion, e.g., alpha-emitters such as, for example,  $^{213}\text{Bi}$ . Non-limiting examples include paclitaxol, cytochalasin B, gramicidin D, ethidium bromide, emetine, mitomycin, etoposide, teniposide, vincristine, vinblastine, colchicin, doxorubicin, daunorubicin, dihydroxy anthracin dione, mitoxantrone, mithramycin, actinomycin D, 1-dehydrotestosterone, glucocorticoids, procaine, tetracaine, lidocaine, propranolol, and puromycin and analogs or homologs thereof. Therapeutic agents include, but are not limited to, antimetabolites (e.g., methotrexate, 6-mercaptopurine, 6-thioguanine, cytarabine, 5-fluorouracil decarbazine), alkylating agents (e.g., mechlorethamine, thioepa chlorambucil, melphalan, carmustine (BSNU) and lomustine (CCNU), cyclophosphamide, busulfan, dibromomannitol, streptozotocin, mitomycin C, and cis-dichlorodiamine platinum (II) (DDP) cisplatin), anthracyclines (e.g., daunorubicin (formerly daunomycin) and doxorubicin), antibiotics (e.g., dactinomycin (formerly actinomycin), bleomycin, mithramycin, and anthramycin (AMC)), and anti-mitotic agents (e.g., vincristine and vinblastine).

The conjugates of the invention can be used for modifying a given biological response such as inducing cell death for the treatment and prevention of hematological malignancies like leukemia. The therapeutic agent or drug moiety is not to be construed as limited to classical chemical therapeutic agents. For example, the drug moiety may be a protein or polypeptide possessing a desired biological activity for inducing cell death. Such proteins may include, for example, a toxin such as abrin, ricin A, pseudomonas exotoxin, or diphtheria toxin; a protein such as tumor necrosis factor, alpha-interferon, beta-interferon, nerve growth factor, platelet derived growth factor, tissue plasminogen activator, an apoptotic agent, e.g.,

TNF-alpha, TNF-beta, AIM I (See, International Publication No. WO 97/33899), AIM II (See, International Publication No. WO 97/34911), Fas Ligand (Takahashi et al., *Int. Immunol.*, 6:1567-1574 (1994)), VEGF (See, International Publication No. WO 99/23105), a thrombotic agent or an anti-angiogenic agent, e.g., angiostatin or endostatin; or, biological response modifiers such as, for example, lymphokines, interleukin-1 ("IL-1"), interleukin-2 ("IL-2"), interleukin-6 ("IL-6"), granulocyte macrophage colony stimulating factor ("GM-CSF"), granulocyte colony stimulating factor ("G-CSF"), or other growth factors.

#### Therapeutic Treatment

The present invention is further directed to methods for preventing and treating hematological malignancies such as leukemia. According to the invention, hematological malignancies comprise without limitation, Hodgkin's disease, leukemia such as, acute lymphoid (lymphocytic or lymphoblastic) leukemia (ALL), acute myeloid (myelogenous or myeloblastic) leukemia (AML), acute lymphoid leukemia, biphenotypic (ALL, biphenotypic), acute undifferentiated leukemia (AUL), chronic myeloid (myelogenous or granulocytic) leukemia (CML), erythroleukemia, granulocytic leukemia, lymphoma, monocytic leukemia, myeloma, myelomonocytic leukemia, myelodysplastic syndromes, non-Hodgkin lymphoma, progranulocytic leukemia.

Methods of treatment of the present invention comprise administering to a subject in need thereof an immunogenic composition of the present invention. The compositions may be administered with a pharmaceutically acceptable carrier.

Such pharmaceutical carriers can be sterile liquids, such as water and oils, including those of petroleum, animal, vegetable or synthetic origin, such as peanut oil, soybean oil, mineral oil, sesame oil and the like. Water is a preferred carrier when the pharmaceutical composition is administered intravenously. Saline solutions and aqueous dextrose and glycerol solutions can also be employed as liquid carriers, particularly for injectable solutions. Suitable pharmaceutical excipients include starch, glucose, lactose, sucrose, gelatin, malt, rice, flour, chalk, silica gel, sodium stearate, glycerol monostearate, talc, sodium chloride, dried skim milk, glycerol, propylene, glycol, water, ethanol and the like. The composition, if desired, can also contain minor amounts of wetting or emulsifying agents, or pH buffering agents. These compositions can take the form of solutions, suspensions, emulsion, tablets, pills, capsules, powders, sustained-release formulations and the like. The composition can be formulated as a suppository, with traditional binders and carriers such as triglycerides. Oral formulation can include standard carriers such as pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, sodium saccharine, cellulose, magnesium carbonate, etc. Examples of suitable pharmaceutical carriers are described in "Remington's Pharmaceutical Sciences" by E. W. Martin. Such compositions will contain a therapeutically effective amount of the compound, preferably in purified form, together with a suitable amount of carrier so as to provide the form for proper administration to the patient. The formulation should suit the mode of administration.

In a preferred embodiment, the composition is formulated in accordance with routine procedures as a pharmaceutical composition adapted for intravenous administration to human beings. Typically, compositions for intravenous administration are solutions in sterile isotonic aqueous buffer. Where necessary, the composition may also include a solubilizing agent and a local anesthetic such as lignocaine to ease pain at the site of the injection. Generally, the ingredients are supplied either separately or mixed together in unit dosage form, for example, as a dry lyophilized powder or water free concentrate in a hermetically sealed container such as an ampoule or sachette indicating the quantity of active agent.

Where the composition is to be administered by infusion, it can be dispensed with an infusion bottle containing sterile pharmaceutical grade water or saline. Where the composition is administered by injection, an ampoule of sterile water for injection or saline can be provided so that the ingredients may be mixed prior to administration.

The compounds of the invention can be formulated as neutral or salt forms. Pharmaceutically acceptable salts include those formed with anions such as those derived from hydrochloric, phosphoric, acetic, oxalic, tartaric acids, etc., and those formed with cations such as those derived from sodium, potassium, ammonium, calcium, ferric hydroxides, isopropylamine, triethylamine, 2-ethylamino ethanol, histidine, procaine, etc.

The amount of the compound of the invention which will be effective in the treatment, inhibition and prevention of hematological malignancies can be determined by standard clinical techniques. In addition, in vitro assays may optionally be employed to help identify optimal dosage ranges. The precise dose to be employed in the formulation will also depend on the route of administration, and the seriousness of the disease or disorder, and should be decided according to the judgment of the practitioner and each patient's circumstances. Effective doses may be extrapolated from dose-response curves derived from in vitro or animal model test systems.

Various other delivery systems are known and can be used to administer a composition of the invention, e.g., encapsulation in liposomes, microparticles, microcapsules, recombinant cells capable of expressing the compound, receptor-mediated endocytosis (See, e.g., Wu and Wu, *J. Biol. Chem.* 262:4429-4432 (1987)), construction of a nucleic acid as part of a retroviral or other vector, etc. Methods of introduction include but are not limited to intradermal, intramuscular, intraperitoneal, intravenous, subcutaneous, intranasal, epidural, and oral routes. The compounds or compositions may be administered by any convenient route, for example by infusion or bolus injection, by absorption through epithelial or mucocutaneous linings (e.g., oral mucosa, rectal and intestinal mucosa, etc.) and may be administered together with other biologically active agents. Administration can be systemic or local. In addition, it may be desirable to introduce the pharmaceutical compounds or compositions of the invention into the central nervous system by any suitable route, including intraventricular and intrathecal injection; intraventricular injection may be facilitated by an intraventricular catheter, for example, attached to a reservoir, such as an Ommaya reservoir. Pulmonary administration can also be employed, e.g., by use of an inhaler or nebulizer, and formulation with an aerosolizing agent.

In a specific embodiment, it may be desirable to administer the pharmaceutical compounds or compositions of the invention locally to the area in need of treatment; this may be achieved by, for example, and not by way of limitation, local infusion during surgery, topical application, e.g., in conjunction with a wound dressing after surgery, by injection, by means of a catheter, by means of a suppository, or by means of an implant, said implant being of a porous, non-porous, or gelatinous material, including membranes, such as sialastic membranes, or fibers. Preferably, when administering a protein, including an antibody, of the invention, care must be taken to use materials to which the protein does not absorb.

In another embodiment, the compound or composition can be delivered in a vesicle, in particular a liposome (See Langer, *Science* 249:1527-1533 (1990); Treat et al., in *Liposomes in the Therapy of Infectious Disease and Cancer*, Lopez-Berstein and Fidler (eds.), Liss, New York, pp. 353-365 (1989); Lopez-Berstein, *ibid.*, pp. 317-327; see generally *ibid.*)

In yet another embodiment, the compound or composition can be delivered in a controlled release system. In one

embodiment, a pump may be used (see Langer, supra; Sefton, *CRC Crit. Ref. Biomed. Eng.* 14:201 (1987); Buchwald et al., *Surgery* 88:507 (1980); Saudek et al., *N. Engl. J. Med.* 321:574 (1989)). In another embodiment, polymeric materials can be used (see *Medical Applications of Controlled Release*, Langer and Wise (eds.), CRC Pres., Boca Raton, Fla. (1974); *Controlled Drug Bioavailability, Drug Product Design and Performance*, Smolen and Ball (eds.), Wiley, New York (1984); Ranger and Peppas, J., *Macromol. Sci. Rev. Macromol. Chem.* 23:61 (1983); see also Levy et al., *Science* 228:190 (1985); During et al., *Ann. Neurol.* 25:351 (1989); Howard et al., *J. Neurosurg.* 71:105 (1989)). In yet another embodiment, a controlled release system can be placed in proximity of the therapeutic target, i.e., the brain, thus requiring only a fraction of the systemic dose (see, e.g., Goodson, in *Medical Applications of Controlled Release*, supra, vol. 2, pp. 115-138 (1984)).

In a specific embodiment where the compound of the invention is a nucleic acid encoding a protein, the nucleic acid can be administered in vivo to promote expression of its encoded protein, by constructing it as part of an appropriate nucleic acid expression vector and administering it so that it becomes intracellular, e.g., by use of a retroviral vector (see U.S. Pat. No. 4,980,286), or by direct injection, or by use of microparticle bombardment (e.g., a gene gun; Biolistic, Dupont), or coating with lipids or cell-surface receptors or transfecting agents, or by administering it in linkage to a homeobox-like peptide which is known to enter the nucleus (see e.g., Joliot et al., *Proc. Natl. Acad. Sci. USA* 88:1864-1868 (1991)), etc. Alternatively, a nucleic acid can be introduced intracellularly and incorporated within host cell DNA for expression, by homologous recombination.

#### EXAMPLES

The following examples are presented for the illustrative purposes and it is to be understood that the present invention is not limited to those precise embodiments, and that various changes and modifications can be effected therein by one skilled in the art without departing from the scope and spirit of the invention as defined by the appended claims.

##### Example 1

##### OCT6 Nucleotide Sequence Identification and Analysis

OCT6 was first identified as a potential OCT gene by assembling and sequencing ESTs. BLAST searches of human ESTs in GenBank data base identified AI040384 (654 bp), AA033971 (714 bp) and H70190 (474 bp) sequences from three fetal liver IMAGE clones, 1656502, 429904 and 212935 respectively. IMAGE clone 1656502 (3', insert 1337 bp) ended the predicted 3' stop codon, whereas IMAGE clone 429904 (5', insert 996 bp) and IMAGE clone 212935 (5', insert 966 bp) aligned with the 5'-coding region. All clones were obtained from the IMAGE Consortium through the American Type Culture Collection (Manassas, Va.). Each clone was sequenced in both directions. The sequences were determined using ABI Prism™ 377 DNA sequencer (Perkin-Elmer). Our assemblage proved to be identical to the recently submitted cDNA OKB1 (AF268892) submitted by M. Okabe and T. Abe. We have dubbed this gene OCT6 as OCTN1 and OCTN2 may be considered as OCT4 and OCT5 respectively.

The OCT6 gene (SEQ ID NO:1) is also contained within BAC clone CTA-331P3 (SEQ ID NO:3) (GenBank AC002464) located at chromosome 6q21. It is divided into 6 exons that span 42 kb on the human genome, from nucleotide 79,570 to nucleotide 120490 on CTA-331P3.

The gene has a predicted protein structure typical of transport proteins with 2 groups of 6 transmembrane domains separated by a hydrophilic region (FIG. 1A). The large hydrophilic region between TMD1 and TMD2 is typical of OCT and OAT genes and is presumed to be located on the outside surface of the cell membrane. The OCT6 protein contains potential sites for N-glycosylation and phosphorylation, which will be described below in Methods. Of interest, the protein sequence also contains a 22 amino acid leucine zipper motif, starting at amino acid 146, suggesting that there may be a physical interaction between OCT6 and ion channels or other membrane-associated proteins.

CLUSTALW alignment produced a dendrogram showing the phylogenetic relationship between OCT6 and other OAT and OCT proteins (FIG. 1B). This dendrogram suggests that the distinction between OAT and OCT genes, based on functional studies, obscures the common origin of both families of transporters. The actual CLUSTALW alignment of these genes is shown in FIG. 2 and demonstrates multiple regions of conservation among all of these genes.

The hydropathy profile analysis, multiple sequence alignments of amino acid sequences using CLUSTALW and the phylogenetic tree were all produced with MacVector software.

##### Example 2

##### Molecular Cloning of OCT6

BLAST searches of human ESTs in GenBank data base identified AI040384 (654 bp), AA033971 (714 bp) and H70190 (474 bp) sequences from three fetal liver IMAGE clones, 1656502, 429904 and 212935 respectively. IMAGE clone 1656502 (3', insert 1337 bp) ended the predicted 3' stop codon, whereas IMAGE clone 429904 (5', insert 996 bp) and IMAGE clone 212935 (5', insert 966 bp) aligned with the 5'-coding region. All clones were obtained from the IMAGE Consortium through the American Type Culture Collection (Manassas, Va.). Each clone was sequenced in both directions. The sequences were determined using ABI Prism™ 377 DNA sequencer (Perkin-Elmer).

##### Example 3

##### Quantitative RT-PCR of OCT6 RNA Levels in Cancer Cell Lines

Total RNA isolated from 50 cell lines used in the NCI drug screen program was provided by the Developmental Therapeutics Program, NCI. Quantitative RT-PCR for detecting OAT-X transporter gene expression was performed by using a Roche LightCycler, which uses real time fluorescence detection for quantitative measurement of PCR products. A gene-specific primer pair was designed with Oligo 4.0 software and purchased from Integrated DNA Technologies, Inc. (Coralville, Iowa) (F: 5'-GGCACATTTATTTCACCAAGACCAG-3') (SEQ ID NO: 13) and (R: 5'-TGTGGACCTCAGCAG-CATTTGGAT-3') (SEQ ID NO:14). The specificity of the PCR reaction was confirmed by directly determining the DNA sequence of the PCR product. First, cDNA was synthesized from total RNA using SuperScript First-Strand Synthesis System (GIBCO/BRL) in a 20 µl volume following the instructions supplied by the manufacturer. The cDNA treated with RNase H for 20 minutes at 37° C. and stored at -20° C. Then, 2 µl of cDNA reaction was amplified in a standard PCR reaction condition, using 0.3 µM primer concentration, with the addition of SYBR Green I Dye. After 30 seconds denature at 95° C., the amplification reaction proceeded through 45-50 cycles of 95° C. denature for 0 second, 62-65° C. annealing

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for 10 seconds and a 72° C. extension for 40 seconds, with slopes of 20° C./s, 20° C./s and 2° C./s, respectively.

Fluorescence was acquired during each cycle after heating to a temperature just below the product melting temperature. Quantification was performed using the LightCycler analysis software. The log-linear portion of the standard amplification curve was identified, and the 'crossing point', a threshold of relative fluorescence, was determined as the best fit through the log-linear region above the background fluorescence (noise) band. The quantification of PCR product then was derived by plotting fluorescence data in the log linear region of each sample to determine a calculated number of cycles needed to reach the fluorescence crossing point. The calculated number of cycles required to reach the crossing point is proportional to the amount of target RNA in the sample. The relative amount of product was described in arbitrary units by interpolation of the data using a standard curve of a series of dilutions of a standard cell line RNA. The quantitative measurement of each gene in each cell line was normalized to the relative amount of actin RNA in each cell line, as a control for equivalent cDNA loading in each sample. The results represent the average of 3 independent determinations performed in duplicate.

A melting curve analysis was performed with positive control RNA prior analysis of the cell lines to enhance sensitivity and the specificity of the data. Amplified products usually melt quickly at a temperature characteristic for the products. The fluorescence signal was acquired at a temperature just below the T<sub>m</sub> of the specific PCR product and above the T<sub>m</sub> of the primer dimers. All specific PCR products displayed a single, sharply melting curve with a narrow peak. In addition, PCR products were confirmed for specificity and correct size by visualization of the LightCycler products on a 1% agarose gel.

## Example 4

## Tissue Distribution

First strand cDNAs derived from 24 adult and fetal tissues (RAPID-SCAN gene expression panel, OriGene Technologies, Rockville, Md.). The PCR primers used in this study

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were the same as used in the quantitative RT-PCR studies. The PCR reaction samples were denatured at 94° C. for 30 seconds, annealed and extended at 64° C. for 30 sec for 35 cycles. The PCR products were then visualized on 1% agarose gels.

## Example 5

## Cell Sorting

All human specimens were obtained in accordance with institutional IRB guidelines. Leukocytes from fresh discarded buffy coats were isolated after RBC lysis with ammonium chloride and labeled with lineage specific antibodies (CD14, monocytes; CD15, granulocytes; CD3, T-cells; and CD20, B-cells), and isolated using a FACSVantage flow cytometer. Each population was sorted twice to ensure purities of at least 99%. CD34 cells were obtained from discarded aliquots of G-CSF-mobilized peripheral blood stem cell collections from cancer patients. For each sample, the PCR results represent the pooled average of cells from 2 individuals performed in triplicate or quadruplicate.

## Example 6

## OCT6 RNA Levels in Leukemic Blasts

Total RNA was extracted from leukemia specimens using QIAGEN RNeasy midi kit. 150 ng of total RNA were used as a template for the first strand cDNA synthesis with the Oligo (dT) primer using the super script system (GIBCO BRL) according to the manufacturer's protocol. Quantitative real-time RT-PCR was performed using an iCycler thermal cycler with methods similar to those described above for the Roche LightCycler. The results represent the average of 3 independent determination performed in duplicate.

Although illustrative embodiments of the present invention have been described in detail, it is to be understood that the present invention is not limited to those precise embodiments, and that various changes and modifications can be effected therein by one skilled in the art without departing from the scope and spirit of the invention as defined by the appended claims.

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His His Val Cys Arg Pro Pro Gly Asn Val Ser Gln Val Val Phe His
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<210> SEQ ID NO 4
<211> LENGTH: 551
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens

<400> SEQUENCE: 4

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Met Arg Asp Tyr Asp Glu Val Ile Ala Phe Leu Gly Glu Trp Gly Pro
 1             5             10             15
Phe Gln Arg Leu Ile Phe Phe Leu Leu Ser Ala Ser Ile Ile Pro Asn
 20             25             30
Gly Phe Asn Gly Met Ser Val Val Phe Leu Ala Gly Thr Pro Glu His
 35             40             45
Arg Cys Arg Val Pro Asp Ala Ala Asn Leu Ser Ser Ala Trp Arg Asn
 50             55             60
Asn Ser Val Pro Leu Arg Leu Arg Asp Gly Arg Glu Val Pro His Ser
 65             70             75             80
Cys Ser Arg Tyr Arg Leu Ala Thr Ile Ala Asn Phe Ser Ala Leu Gly
 85             90             95
Leu Glu Pro Gly Arg Asp Val Asp Leu Gly Gln Leu Glu Gln Glu Ser
 100            105            110
Cys Leu Asp Gly Trp Glu Phe Ser Gln Asp Val Tyr Leu Ser Thr Val
 115            120            125
Val Thr Glu Trp Asn Leu Val Cys Glu Asp Asn Trp Lys Val Pro Leu

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130				135				140							
Thr	Thr	Ser	Leu	Phe	Phe	Val	Gly	Val	Leu	Leu	Gly	Ser	Phe	Val	Ser
145					150					155					160
Gly	Gln	Leu	Ser	Asp	Arg	Phe	Gly	Arg	Lys	Asn	Val	Leu	Phe	Ala	Thr
				165					170					175	
Met	Ala	Val	Gln	Thr	Gly	Phe	Ser	Phe	Leu	Gln	Ile	Phe	Ser	Ile	Ser
			180					185					190		
Trp	Glu	Met	Phe	Thr	Val	Leu	Phe	Val	Ile	Val	Gly	Met	Gly	Gln	Ile
		195					200					205			
Ser	Asn	Tyr	Val	Val	Ala	Phe	Ile	Leu	Gly	Thr	Glu	Ile	Leu	Gly	Lys
	210					215					220				
Ser	Val	Arg	Ile	Ile	Phe	Ser	Thr	Leu	Gly	Val	Cys	Thr	Phe	Phe	Ala
225					230					235					240
Val	Gly	Tyr	Met	Leu	Leu	Pro	Leu	Phe	Ala	Tyr	Phe	Ile	Arg	Asp	Trp
				245					250					255	
Arg	Met	Leu	Leu	Leu	Ala	Leu	Thr	Val	Pro	Gly	Val	Leu	Cys	Val	Pro
			260						265				270		
Leu	Trp	Trp	Phe	Ile	Pro	Glu	Ser	Pro	Arg	Trp	Leu	Ile	Ser	Gln	Arg
		275					280					285			
Arg	Phe	Arg	Glu	Ala	Glu	Asp	Ile	Ile	Gln	Lys	Ala	Ala	Lys	Met	Asn
	290					295					300				
Asn	Ile	Ala	Val	Pro	Ala	Val	Ile	Phe	Asp	Ser	Val	Glu	Glu	Leu	Asn
305					310					315					320
Pro	Leu	Lys	Gln	Gln	Lys	Ala	Phe	Ile	Leu	Asp	Leu	Phe	Arg	Thr	Arg
				325					330					335	
Asn	Ile	Ala	Ile	Met	Thr	Ile	Met	Ser	Leu	Leu	Leu	Trp	Met	Leu	Thr
			340					345					350		
Ser	Val	Gly	Tyr	Phe	Ala	Leu	Ser	Leu	Asp	Ala	Pro	Asn	Leu	His	Gly
		355					360					365			
Asp	Ala	Tyr	Leu	Asn	Cys	Phe	Leu	Ser	Ala	Leu	Ile	Glu	Ile	Pro	Ala
	370					375					380				
Tyr	Ile	Thr	Ala	Trp	Leu	Leu	Leu	Arg	Thr	Leu	Pro	Arg	Arg	Tyr	Ile
385					390					395					400
Ile	Ala	Ala	Val	Leu	Phe	Trp	Gly	Gly	Gly	Val	Leu	Leu	Phe	Ile	Gln
				405					410					415	
Leu	Val	Pro	Val	Asp	Tyr	Tyr	Phe	Leu	Ser	Ile	Gly	Leu	Val	Met	Leu
			420					425					430		
Gly	Lys	Phe	Gly	Ile	Thr	Ser	Ala	Phe	Ser	Met	Leu	Tyr	Val	Phe	Thr
		435					440					445			
Ala	Glu	Leu	Tyr	Pro	Thr	Leu	Val	Arg	Asn	Met	Ala	Val	Gly	Val	Thr
	450					455					460				
Ser	Thr	Ala	Ser	Arg	Val	Gly	Ser	Ile	Ile	Ala	Pro	Tyr	Phe	Val	Tyr
465					470					475					480
Leu	Gly	Ala	Tyr	Asn	Arg	Met	Leu	Pro	Tyr	Ile	Val	Met	Gly	Ser	Leu
				485					490					495	
Thr	Val	Leu	Ile	Gly	Ile	Leu	Thr	Leu	Phe	Phe	Pro	Glu	Ser	Leu	Gly
			500					505					510		
Met	Thr	Leu	Pro	Glu	Thr	Leu	Glu	Gln	Met	Gln	Lys	Val	Lys	Trp	Phe
		515					520					525			
Arg	Ser	Gly	Lys	Lys	Thr	Arg	Asp	Ser	Met	Glu	Thr	Glu	Glu	Asn	Pro
	530					535					540				
Lys	Val	Leu	Ile	Thr	Ala	Phe									
545					550										

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<210> SEQ ID NO 5
<211> LENGTH: 551
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens

<400> SEQUENCE: 5
Met Ala Gln Phe Val Gln Val Leu Ala Glu Ile Gly Asp Phe Gly Arg
 1           5           10          15
Phe Gln Ile Gln Leu Leu Ile Leu Leu Cys Val Leu Asn Phe Leu Ser
 20          25          30
Pro Phe Tyr Phe Phe Ala His Val Phe Met Val Leu Asp Glu Pro His
 35          40          45
His Cys Ala Val Ala Trp Val Lys Asn His Thr Phe Asn Leu Ser Ala
 50          55          60
Ala Glu Gln Leu Val Leu Ser Val Pro Leu Asp Thr Ala Gly His Pro
 65          70          75          80
Glu Pro Cys Leu Met Phe Arg Pro Pro Pro Ala Asn Ala Ser Leu Gln
 85          90          95
Asp Ile Leu Ser His Arg Phe Asn Glu Thr Gln Pro Cys Asp Met Gly
 100         105         110
Trp Glu Tyr Pro Glu Asn Arg Leu Pro Ser Leu Lys Asn Glu Phe Asn
 115         120         125
Leu Val Cys Asp Arg Lys His Leu Lys Asp Thr Thr Gln Ser Val Phe
 130         135         140
Met Gly Gly Leu Leu Val Gly Thr Leu Met Phe Gly Pro Leu Cys Asp
 145         150         155         160
Arg Ile Gly Arg Lys Ala Thr Ile Leu Ala Gln Leu Leu Leu Phe Thr
 165         170         175
Leu Ile Gly Leu Ala Thr Ala Phe Val Pro Ser Phe Glu Leu Tyr Met
 180         185         190
Ala Leu Arg Phe Ala Val Ala Thr Ala Val Ala Gly Leu Ser Phe Ser
 195         200         205
Asn Val Thr Leu Leu Thr Glu Trp Val Gly Pro Ser Trp Arg Thr Gln
 210         215         220
Ala Val Val Leu Ala Gln Cys Asn Phe Ser Leu Gly Gln Met Val Leu
 225         230         235         240
Ala Gly Leu Ala Tyr Gly Phe Arg Asn Trp Arg Leu Leu Gln Ile Thr
 245         250         255
Gly Thr Ala Pro Gly Leu Leu Leu Phe Phe Tyr Phe Trp Ala Leu Pro
 260         265         270
Glu Ser Ala Arg Trp Leu Leu Thr Arg Gly Arg Met Asp Glu Ala Ile
 275         280         285
Gln Leu Ile Gln Lys Ala Ala Ser Val Asn Arg Arg Lys Leu Ser Pro
 290         295         300
Glu Leu Met Asn Gln Leu Val Pro Glu Lys Thr Gly Pro Ser Gly Asn
 305         310         315         320
Ala Leu Asp Leu Phe Arg His Pro Gln Leu Arg Lys Val Thr Leu Ile
 325         330         335
Ile Phe Cys Val Trp Phe Val Asp Ser Leu Gly Tyr Tyr Gly Leu Ser
 340         345         350
Leu Gln Val Gly Asp Phe Gly Leu Asp Val Tyr Leu Thr Gln Leu Ile
 355         360         365
Phe Gly Ala Val Glu Val Pro Ala Arg Cys Ser Ser Ile Phe Met Met

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370				375				380							
Gln	Arg	Phe	Gly	Arg	Lys	Trp	Ser	Gln	Leu	Gly	Thr	Leu	Val	Leu	Gly
385					390					395					400
Gly	Leu	Met	Cys	Ile	Ile	Ile	Ile	Phe	Ile	Pro	Ala	Asp	Leu	Pro	Val
				405					410					415	
Val	Val	Thr	Met	Leu	Ala	Val	Val	Gly	Lys	Met	Ala	Thr	Ala	Ala	Ala
			420					425					430		
Phe	Thr	Ile	Ser	Tyr	Val	Tyr	Ser	Ala	Glu	Leu	Phe	Pro	Thr	Ile	Leu
		435					440					445			
Arg	Gln	Thr	Gly	Met	Gly	Leu	Val	Gly	Ile	Phe	Ser	Arg	Ile	Gly	Gly
	450					455					460				
Ile	Leu	Thr	Pro	Leu	Val	Ile	Leu	Leu	Gly	Glu	Tyr	His	Ala	Ala	Leu
465					470					475					480
Pro	Met	Leu	Ile	Tyr	Gly	Ser	Leu	Pro	Ile	Val	Ala	Gly	Leu	Leu	Cys
				485					490						495
Thr	Leu	Leu	Pro	Glu	Thr	His	Gly	Gln	Gly	Leu	Lys	Asp	Thr	Leu	Gln
			500					505					510		
Asp	Leu	Glu	Leu	Gly	Pro	His	Pro	Arg	Ser	Pro	Lys	Ser	Val	Pro	Ser
		515					520					525			
Glu	Lys	Glu	Thr	Glu	Ala	Lys	Gly	Arg	Thr	Ser	Ser	Pro	Gly	Val	Ala
	530					535					540				
Phe	Val	Ser	Ser	Thr	Tyr	Phe									
545					550										

&lt;210&gt; SEQ ID NO 6

&lt;211&gt; LENGTH: 557

&lt;212&gt; TYPE: PRT

&lt;213&gt; ORGANISM: Homo sapiens

&lt;400&gt; SEQUENCE: 6

Met	Arg	Asp	Tyr	Asp	Glu	Val	Thr	Ala	Phe	Leu	Gly	Glu	Trp	Gly	Pro
1				5					10					15	
Phe	Gln	Arg	Leu	Ile	Phe	Phe	Leu	Leu	Ser	Ala	Ser	Ile	Ile	Pro	Asn
			20					25					30		
Gly	Phe	Thr	Gly	Leu	Ser	Ser	Val	Phe	Leu	Ile	Ala	Thr	Pro	Glu	His
		35					40					45			
Arg	Cys	Arg	Val	Pro	Asp	Ala	Ala	Asn	Leu	Ser	Ser	Ala	Trp	Arg	Asn
	50					55					60				
His	Thr	Val	Pro	Leu	Arg	Leu	Arg	Asp	Gly	Arg	Glu	Val	Pro	His	Ser
65					70				75					80	
Cys	Arg	Arg	Tyr	Arg	Leu	Ala	Thr	Ile	Ala	Asn	Phe	Ser	Ala	Leu	Gly
			85					90						95	
Leu	Glu	Pro	Gly	Arg	Asp	Val	Asp	Leu	Gly	Gln	Leu	Glu	Gln	Glu	Ser
			100					105					110		
Cys	Leu	Asp	Gly	Trp	Glu	Phe	Ser	Gln	Asp	Val	Tyr	Leu	Ser	Thr	Ile
		115					120					125			
Val	Thr	Glu	Trp	Asn	Leu	Val	Cys	Glu	Asp	Asp	Trp	Lys	Ala	Pro	Leu
		130				135					140				
Thr	Ile	Ser	Leu	Phe	Phe	Val	Gly	Val	Leu	Leu	Gly	Ser	Phe	Ile	Ser
145					150					155				160	
Gly	Gln	Leu	Ser	Asp	Arg	Phe	Gly	Arg	Lys	Asn	Val	Leu	Phe	Val	Thr
				165					170					175	
Met	Gly	Met	Gln	Thr	Gly	Phe	Ser	Phe	Leu	Gln	Ile	Phe	Ser	Lys	Asn
			180					185						190	



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Phe Glu Met Phe Val Val Leu Phe Val Leu Val Gly Met Gly Gln Ile  
 195 200 205  
 Ser Asn Tyr Val Ala Ala Phe Val Leu Gly Thr Glu Ile Leu Gly Lys  
 210 215 220  
 Ser Val Arg Ile Ile Phe Ser Thr Leu Gly Val Cys Ile Phe Tyr Ala  
 225 230 235 240  
 Phe Gly Tyr Met Val Leu Pro Leu Phe Ala Tyr Phe Ile Arg Asp Trp  
 245 250 255  
 Arg Met Leu Leu Val Ala Leu Thr Met Pro Gly Val Leu Cys Val Ala  
 260 265 270  
 Leu Trp Trp Phe Ile Pro Glu Ser Pro Arg Trp Leu Ile Ser Gln Gly  
 275 280 285  
 Arg Phe Glu Glu Ala Glu Val Ile Ile Arg Lys Ala Ala Lys Ala Asn  
 290 295 300  
 Gly Ile Val Val Pro Ser Thr Ile Phe Asp Pro Ser Glu Leu Gln Asp  
 305 310 315 320  
 Leu Ser Ser Lys Lys Gln Gln Ser His Asn Ile Leu Asp Leu Leu Arg  
 325 330 335  
 Thr Trp Asn Ile Arg Met Val Thr Ile Met Ser Ile Met Leu Trp Met  
 340 345 350  
 Thr Ile Ser Val Gly Tyr Phe Gly Leu Ser Leu Asp Thr Pro Asn Leu  
 355 360 365  
 His Gly Asp Ile Phe Val Asn Cys Phe Leu Ser Ala Met Val Glu Val  
 370 375 380  
 Pro Ala Tyr Val Leu Ala Trp Leu Leu Leu Gln Tyr Leu Pro Arg Arg  
 385 390 395 400  
 Tyr Ser Met Ala Thr Ala Leu Phe Leu Gly Gly Ser Val Leu Leu Phe  
 405 410 415  
 Met Gln Leu Val Pro Pro Asp Leu Tyr Tyr Leu Ala Thr Val Leu Val  
 420 425 430  
 Met Val Gly Lys Phe Gly Val Thr Ala Ala Phe Ser Met Val Tyr Val  
 435 440 445  
 Tyr Thr Ala Glu Leu Tyr Pro Thr Val Val Arg Asn Met Gly Val Gly  
 450 455 460  
 Val Ser Ser Thr Ala Ser Arg Leu Gly Ser Ile Leu Ser Pro Tyr Phe  
 465 470 475 480  
 Val Tyr Leu Gly Ala Tyr Asp Arg Phe Leu Pro Tyr Ile Leu Met Gly  
 485 490 495  
 Ser Leu Thr Ile Leu Thr Ala Ile Leu Thr Leu Phe Leu Pro Glu Ser  
 500 505 510  
 Phe Gly Thr Pro Leu Pro Asp Thr Ile Asp Gln Met Leu Arg Val Lys  
 515 520 525  
 Gly Met Lys His Arg Lys Thr Pro Ser His Thr Arg Met Leu Lys Asp  
 530 535 540  
 Gly Gln Glu Arg Pro Thr Ile Leu Lys Ser Thr Ala Phe  
 545 550 555

<210> SEQ ID NO 7  
 <211> LENGTH: 555  
 <212> TYPE: PRT  
 <213> ORGANISM: Homo sapiens

<400> SEQUENCE: 7

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

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Phe Phe Gln Lys Gln Met Phe Phe Leu Leu Ala Leu Leu Ser Ala Thr  
 20 25 30  
 Phe Ala Pro Ile Tyr Val Gly Ile Val Phe Leu Gly Phe Thr Pro Asp  
 35 40 45  
 His Arg Cys Arg Ser Pro Gly Val Ala Glu Leu Ser Leu Arg Cys Gly  
 50 55 60  
 Trp Ser Pro Ala Glu Glu Leu Asn Tyr Thr Val Pro Gly Pro Gly Pro  
 65 70 75 80  
 Ala Gly Glu Ala Ser Pro Arg Gln Cys Arg Arg Tyr Glu Val Asp Trp  
 85 90 95  
 Asn Gln Ser Thr Phe Asp Cys Val Asp Pro Leu Ala Ser Leu Asp Thr  
 100 105 110  
 Asn Arg Ser Arg Leu Pro Leu Gly Pro Cys Arg Asp Gly Trp Val Tyr  
 115 120 125  
 Glu Thr Pro Gly Ser Ser Ile Val Thr Glu Phe Asn Leu Val Cys Ala  
 130 135 140  
 Asn Ser Trp Met Leu Asp Leu Phe Gln Ser Ser Val Asn Val Gly Phe  
 145 150 155 160  
 Phe Ile Gly Ser Met Ser Ile Gly Tyr Ile Ala Asp Arg Phe Gly Arg  
 165 170 175  
 Lys Leu Cys Leu Leu Thr Thr Val Leu Ile Asn Ala Ala Ala Gly Val  
 180 185 190  
 Leu Met Ala Ile Ser Pro Thr Tyr Thr Trp Met Leu Ile Phe Arg Leu  
 195 200 205  
 Ile Gln Gly Leu Val Ser Lys Ala Gly Trp Leu Ile Gly Tyr Ile Leu  
 210 215 220  
 Ile Thr Glu Phe Val Gly Arg Arg Tyr Arg Arg Thr Val Gly Ile Phe  
 225 230 235 240  
 Tyr Gln Val Ala Tyr Thr Val Gly Leu Leu Val Leu Ala Gly Val Ala  
 245 250 255  
 Tyr Ala Leu Pro His Trp Arg Trp Leu Gln Phe Thr Val Ala Leu Pro  
 260 265 270  
 Asn Phe Phe Phe Leu Leu Tyr Tyr Trp Cys Ile Pro Glu Ser Pro Arg  
 275 280 285  
 Trp Leu Ile Ser Gln Asn Lys Asn Ala Glu Ala Met Arg Ile Ile Lys  
 290 295 300  
 His Ile Ala Lys Lys Asn Gly Lys Ser Leu Pro Ala Ser Leu Gln Arg  
 305 310 315 320  
 Leu Arg Leu Glu Glu Thr Gly Lys Lys Leu Asn Pro Ser Phe Leu  
 325 330 335  
 Asp Leu Val Arg Thr Pro Gln Ile Arg Lys His Thr Met Ile Leu Met  
 340 345 350  
 Tyr Asn Trp Phe Thr Ser Ser Val Leu Tyr Gln Gly Leu Ile Met His  
 355 360 365  
 Met Gly Leu Ala Gly Asp Asn Ile Tyr Leu Asp Phe Phe Tyr Ser Ala  
 370 375 380  
 Leu Val Glu Phe Pro Ala Ala Phe Met Ile Ile Leu Thr Ile Asp Arg  
 385 390 395 400  
 Ile Gly Arg Arg Tyr Pro Trp Ala Ala Ser Asn Met Val Ala Gly Ala  
 405 410 415  
 Ala Cys Leu Ala Ser Val Phe Ile Pro Gly Asp Leu Gln Trp Leu Lys  
 420 425 430

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Ile Ile Ile Ser Cys Leu Gly Arg Met Gly Ile Thr Met Ala Tyr Glu
    435                440                445

Ile Val Cys Leu Val Asn Ala Glu Leu Tyr Pro Thr Phe Ile Arg Asn
    450                455                460

Leu Gly Val His Ile Cys Ser Ser Met Cys Asp Ile Gly Gly Ile Ile
465                470                475                480

Thr Pro Phe Leu Val Tyr Arg Leu Thr Asn Ile Trp Leu Glu Leu Pro
    485                490                495

Leu Met Val Phe Gly Val Leu Gly Leu Val Ala Gly Gly Leu Val Leu
    500                505                510

Leu Leu Pro Glu Thr Lys Gly Lys Ala Leu Pro Glu Thr Ile Glu Glu
    515                520                525

Ala Glu Asn Met Gln Arg Pro Arg Lys Asn Lys Glu Lys Met Ile Tyr
    530                535                540

Leu Gln Val Gln Lys Leu Asp Ile Pro Leu Asn
545                550                555

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<210> SEQ ID NO 8
<211> LENGTH: 554
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens

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<400> SEQUENCE: 8

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Met Pro Thr Val Asp Asp Ile Leu Glu Gln Val Gly Glu Ser Gly Trp
1                5                10                15

Phe Gln Lys Gln Ala Phe Leu Ile Leu Cys Leu Leu Ser Ala Ala Phe
    20                25                30

Ala Pro Ile Cys Val Gly Ile Val Phe Leu Gly Phe Thr Pro Asp His
    35                40                45

His Cys Gln Ser Pro Gly Val Ala Glu Leu Ser Gln Arg Cys Gly Trp
    50                55                60

Ser Pro Ala Glu Glu Leu Asn Tyr Thr Val Pro Gly Leu Gly Pro Ala
65                70                75                80

Gly Glu Ala Phe Leu Gly Gln Cys Arg Arg Tyr Glu Val Asp Trp Asn
    85                90                95

Gln Ser Ala Leu Ser Cys Val Asp Pro Leu Ala Ser Leu Ala Thr Asn
    100               105               110

Arg Ser His Leu Pro Leu Gly Pro Cys Gln Asp Gly Trp Val Tyr Asp
    115               120               125

Thr Pro Gly Ser Ser Ile Val Thr Glu Phe Asn Leu Val Cys Ala Asp
    130               135               140

Ser Trp Lys Leu Asp Leu Phe Gln Ser Cys Leu Asn Ala Gly Phe Leu
145               150               155               160

Phe Gly Ser Leu Gly Val Gly Tyr Phe Ala Asp Arg Phe Gly Arg Lys
    165               170               175

Leu Cys Leu Leu Gly Thr Val Leu Val Asn Ala Val Ser Gly Val Leu
    180               185               190

Met Ala Phe Ser Pro Asn Tyr Met Ser Met Leu Leu Phe Arg Leu Leu
    195               200               205

Gln Gly Leu Val Ser Lys Gly Asn Trp Met Ala Gly Tyr Thr Leu Ile
    210               215               220

Thr Glu Phe Val Gly Ser Gly Ser Arg Arg Thr Val Ala Ile Met Tyr
225               230               235               240

Gln Met Ala Phe Thr Val Gly Leu Val Ala Leu Thr Gly Leu Ala Tyr
    245               250               255

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Ala Leu Pro His Trp Arg Trp Leu Gln Leu Ala Val Ser Leu Pro Thr  
 260 265 270

Phe Leu Phe Leu Leu Tyr Tyr Trp Cys Val Pro Glu Ser Pro Arg Trp  
 275 280 285

Leu Leu Ser Gln Lys Arg Asn Thr Glu Ala Ile Lys Ile Met Asp His  
 290 295 300

Ile Ala Gln Lys Asn Gly Lys Leu Pro Pro Ala Asp Leu Lys Met Leu  
 305 310 315 320

Ser Leu Glu Glu Asp Val Thr Glu Lys Leu Ser Pro Ser Phe Ala Asp  
 325 330 335

Leu Phe Arg Thr Pro Arg Leu Arg Lys Arg Thr Phe Ile Leu Met Tyr  
 340 345 350

Leu Trp Phe Thr Asp Ser Val Leu Tyr Gln Gly Leu Ile Leu His Met  
 355 360 365

Gly Ala Thr Ser Gly Asn Leu Tyr Leu Asp Phe Leu Tyr Ser Ala Leu  
 370 375 380

Val Glu Ile Pro Gly Ala Phe Ile Ala Leu Ile Thr Ile Asp Arg Val  
 385 390 395 400

Gly Arg Ile Tyr Pro Met Ala Met Ser Asn Leu Leu Ala Gly Ala Ala  
 405 410 415

Cys Leu Val Met Ile Phe Ile Ser Pro Asp Leu His Trp Leu Asn Ile  
 420 425 430

Ile Ile Met Cys Val Gly Arg Met Gly Ile Thr Ile Ala Ile Gln Met  
 435 440 445

Ile Cys Leu Val Asn Ala Glu Leu Tyr Pro Thr Phe Val Arg Asn Leu  
 450 455 460

Gly Val Met Val Cys Ser Ser Leu Cys Asp Ile Gly Gly Ile Ile Thr  
 465 470 475 480

Pro Phe Ile Val Phe Arg Leu Arg Glu Val Trp Gln Ala Leu Pro Leu  
 485 490 495

Ile Leu Phe Ala Val Leu Gly Leu Leu Ala Ala Gly Val Thr Leu Leu  
 500 505 510

Leu Pro Glu Thr Lys Gly Val Ala Leu Pro Glu Thr Met Lys Asp Ala  
 515 520 525

Glu Asn Leu Gly Arg Lys Ala Lys Pro Lys Glu Asn Thr Ile Tyr Leu  
 530 535 540

Lys Val Gln Thr Ser Glu Pro Ser Gly Thr  
 545 550

<210> SEQ ID NO 9  
 <211> LENGTH: 539  
 <212> TYPE: PRT  
 <213> ORGANISM: Homo sapiens

<400> SEQUENCE: 9

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

Gln Met Leu His Leu Val Phe Ile Leu Pro Ser Leu Met Leu Leu Ile  
 20 25 30

Pro His Ile Leu Leu Glu Asn Phe Ala Ala Ala Ile Pro Gly His Arg  
 35 40 45

Cys Trp Val His Met Leu Asp Asn Asn Thr Gly Ser Gly Asn Glu Thr  
 50 55 60

Gly Ile Leu Ser Glu Asp Ala Leu Leu Arg Ile Ser Ile Pro Leu Asp

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65		70				75				80					
Ser	Asn	Leu	Arg	Pro	Glu	Lys	Cys	Arg	Phe	Phe	Val	His	Pro	Gln	Trp
				85					90					95	
Gln	Leu	Leu	His	Leu	Asn	Gly	Ile	His	Ser	Thr	Ser	Glu	Ala	Asp	Thr
			100					105					110		
Glu	Pro	Cys	Val	Asp	Gly	Trp	Val	Tyr	Asp	Gln	Ser	Tyr	Phe	Pro	Ser
		115					120					125			
Thr	Ile	Val	Thr	Lys	Trp	Asp	Leu	Val	Cys	Asp	Tyr	Gln	Ser	Leu	Lys
	130					135					140				
Ser	Val	Val	Gln	Phe	Leu	Leu	Leu	Thr	Gly	Met	Leu	Val	Gly	Gly	Ile
145					150					155					160
Ile	His	His	Gly	Val	Ser	Asp	Arg	Phe	Gly	Arg	Arg	Phe	Ile	Leu	Arg
			165						170						175
Trp	Cys	Leu	Leu	Gln	Leu	Ala	Ile	Thr	Asp	Thr	Cys	Ala	Ala	Phe	Ala
			180						185					190	
Pro	Thr	Phe	Pro	Val	Tyr	Cys	Val	Leu	Arg	Phe	Leu	Ala	Gly	Phe	Ser
		195					200					205			
Ser	Met	Ile	Ile	Ile	Ser	Asn	Asn	Ser	Leu	Pro	Ile	Thr	Glu	Trp	Ile
	210					215					220				
Arg	Pro	Asn	Ser	Lys	Ala	Leu	Val	Val	Ile	Leu	Ser	Ser	Gly	Ala	Leu
225					230					235					240
Ser	Ile	Gly	Gln	Ile	Ile	Leu	Gly	Gly	Leu	Ala	Tyr	Val	Phe	Arg	Asp
			245						250					255	
Trp	Gln	Thr	Leu	His	Val	Val	Ala	Ser	Val	Pro	Phe	Leu	Gly	Leu	Leu
			260						265				270		
Leu	Leu	Gln	Arg	Trp	Leu	Val	Glu	Ser	Ala	Arg	Trp	Leu	Ile	Ile	Thr
		275					280					285			
Asn	Lys	Leu	Asp	Glu	Gly	Leu	Lys	Ala	Leu	Arg	Lys	Val	Ala	Arg	Thr
	290					295					300				
Asn	Gly	Ile	Lys	Asn	Ala	Glu	Glu	Thr	Leu	Asn	Ile	Glu	Val	Val	Arg
305					310					315					320
Ser	Thr	Met	Gln	Glu	Glu	Leu	Asp	Ala	Ala	Gln	Thr	Lys	Thr	Thr	Val
			325						330					335	
Cys	Asp	Leu	Phe	Arg	Asn	Pro	Ser	Met	Arg	Lys	Arg	Ile	Cys	Ile	Leu
			340					345					350		
Val	Phe	Leu	Arg	Phe	Ala	Asn	Thr	Ile	Pro	Phe	Tyr	Gly	Thr	Met	Val
		355					360					365			
Asn	Leu	Gln	His	Val	Gly	Ser	Asn	Ile	Phe	Leu	Leu	Gln	Val	Leu	Tyr
	370					375						380			
Gly	Ala	Val	Ala	Leu	Ile	Val	Arg	Cys	Leu	Ala	Leu	Leu	Thr	Leu	Asn
385					390					395					400
His	Met	Gly	Arg	Arg	Ile	Ser	Gln	Ile	Leu	Phe	Met	Phe	Leu	Val	Gly
			405						410					415	
Leu	Ser	Ile	Leu	Ala	Asn	Thr	Phe	Val	Pro	Lys	Glu	Met	Gln	Thr	Leu
			420					425					430		
Arg	Val	Ala	Leu	Ala	Cys	Leu	Gly	Ile	Gly	Cys	Ser	Ala	Ala	Thr	Phe
		435					440					445			
Ser	Ser	Val	Ala	Val	His	Phe	Ile	Glu	Leu	Ile	Pro	Thr	Val	Leu	Arg
	450					455						460			
Ala	Arg	Ala	Ser	Gly	Ile	Asp	Leu	Thr	Ala	Ser	Arg	Ile	Gly	Ala	Ala
465					470					475					480
Leu	Pro	Leu	Leu	Met	Thr	Leu	Thr	Val	Phe	Phe	Thr	Thr	Leu	Pro	Trp
				485					490					495	

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Ile Ile Tyr Gly Ile Phe Pro Ile Ile Gly Gly Leu Ile Val Phe Leu
500 505 510

Leu Pro Glu Thr Lys Asn Leu Pro Leu Pro Asp Thr Ile Lys Asp Val
515 520 525

Glu Asn Gln Lys Lys Asn Leu Lys Glu Lys Ala
530 535

<210> SEQ ID NO 10
<211> LENGTH: 550
<212> TYPE: PRT
<213> ORGANISM: Homo sapiens

<400> SEQUENCE: 10

Met Ala Phe Ser Lys Leu Leu Glu Gln Ala Gly Gly Val Gly Leu Phe
1 5 10 15

Gln Thr Leu Gln Val Leu Thr Phe Ile Leu Pro Cys Leu Met Ile Pro
20 25 30

Ser Gln Met Leu Leu Glu Asn Phe Ser Ala Ala Ile Pro Gly His Arg
35 40 45

Cys Trp Thr His Met Leu Asp Asn Gly Ser Ala Val Ser Thr Asn Met
50 55 60

Thr Pro Lys Ala Leu Leu Thr Ile Ser Ile Pro Pro Gly Pro Asn Gln
65 70 75 80

Gly Pro His Gln Cys Arg Arg Phe Arg Gln Pro Gln Trp Gln Leu Leu
85 90 95

Asp Pro Asn Ala Thr Ala Thr Ser Trp Ser Glu Ala Asp Thr Glu Pro
100 105 110

Cys Val Asp Gly Trp Val Tyr Asp Arg Ser Val Phe Thr Ser Thr Ile
115 120 125

Val Ala Lys Trp Asp Leu Val Cys Ser Ser Gln Gly Leu Lys Pro Leu
130 135 140

Ser Gln Ser Ile Phe Met Ser Gly Ile Leu Val Gly Ser Phe Ile Trp
145 150 155 160

Gly Leu Leu Ser Tyr Arg Phe Gly Arg Lys Pro Met Leu Ser Trp Cys
165 170 175

Cys Leu Gln Leu Ala Val Ala Gly Thr Ser Thr Ile Phe Ala Pro Thr
180 185 190

Phe Val Ile Tyr Cys Gly Leu Arg Phe Val Ala Ala Phe Gly Met Ala
195 200 205

Gly Ile Phe Leu Ser Ser Leu Thr Leu Met Val Glu Trp Thr Thr Thr
210 215 220

Ser Arg Arg Ala Val Thr Met Thr Val Val Gly Cys Ala Phe Ser Ala
225 230 235 240

Gly Gln Ala Ala Leu Gly Gly Leu Ala Phe Ala Leu Arg Asp Trp Arg
245 250 255

Thr Leu Gln Leu Ala Ala Ser Val Pro Phe Phe Ala Ile Ser Leu Ile
260 265 270

Ser Trp Trp Leu Pro Glu Ser Ala Arg Trp Leu Ile Ile Lys Gly Lys
275 280 285

Pro Asp Gln Ala Leu Gln Glu Leu Arg Lys Val Ala Arg Ile Asn Gly
290 295 300

His Lys Glu Ala Lys Asn Leu Thr Ile Glu Val Leu Met Ser Ser Val
305 310 315 320

Lys Glu Glu Val Ala Ser Ala Lys Glu Pro Arg Ser Val Leu Asp Leu

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325					330					335					
Phe	Cys	Val	Pro	Val	Leu	Arg	Trp	Arg	Ser	Cys	Ala	Met	Leu	Val	Val
			340					345					350		
Asn	Phe	Ser	Leu	Leu	Ile	Ser	Tyr	Tyr	Gly	Leu	Val	Phe	Asp	Leu	Gln
		355					360					365			
Ser	Leu	Gly	Arg	Asp	Ile	Phe	Leu	Leu	Gln	Ala	Leu	Phe	Gly	Ala	Val
		370				375					380				
Asp	Phe	Leu	Gly	Arg	Ala	Thr	Thr	Ala	Leu	Leu	Leu	Ser	Phe	Leu	Gly
		385				390					395				400
Arg	Arg	Thr	Ile	Gln	Ala	Gly	Ser	Gln	Ala	Met	Ala	Gly	Leu	Ala	Ile
				405					410					415	
Leu	Ala	Asn	Met	Leu	Val	Pro	Gln	Asp	Leu	Gln	Thr	Leu	Arg	Val	Val
			420					425					430		
Phe	Ala	Val	Leu	Gly	Lys	Gly	Cys	Phe	Gly	Ile	Ser	Leu	Thr	Cys	Leu
		435					440					445			
Thr	Ile	Tyr	Lys	Ala	Glu	Leu	Phe	Pro	Thr	Pro	Val	Arg	Met	Thr	Ala
		450				455					460				
Asp	Gly	Ile	Leu	His	Thr	Val	Gly	Arg	Leu	Gly	Ala	Met	Met	Gly	Pro
		465				470					475				480
Leu	Ile	Leu	Met	Ser	Arg	Gln	Ala	Leu	Pro	Leu	Leu	Pro	Pro	Leu	Leu
				485					490					495	
Tyr	Gly	Val	Ile	Ser	Ile	Ala	Ser	Ser	Leu	Val	Val	Leu	Phe	Phe	Leu
			500					505					510		
Pro	Glu	Thr	Gln	Gly	Leu	Pro	Leu	Pro	Asp	Thr	Ile	Gln	Asp	Leu	Glu
		515					520					525			
Ser	Gln	Lys	Ser	Thr	Ala	Ala	Gln	Gly	Asn	Arg	Gln	Glu	Ala	Val	Thr
		530				535					540				
Val	Glu	Ser	Thr	Ser	Leu										
		545			550										

<210> SEQ ID NO 11  
 <211> LENGTH: 542  
 <212> TYPE: PRT  
 <213> ORGANISM: Homo sapiens

<400> SEQUENCE: 11

Met	Thr	Phe	Ser	Glu	Ile	Leu	Asp	Arg	Val	Gly	Ser	Met	Gly	His	Phe
1				5					10					15	
Gln	Phe	Leu	His	Val	Ala	Ile	Leu	Gly	Leu	Pro	Ile	Leu	Asn	Met	Ala
			20					25					30		
Asn	His	Asn	Leu	Leu	Gln	Ile	Phe	Thr	Ala	Ala	Thr	Pro	Val	His	His
		35				40						45			
Cys	Arg	Pro	Pro	His	Asn	Ala	Ser	Thr	Gly	Pro	Trp	Val	Leu	Pro	Met
		50				55					60				
Gly	Pro	Asn	Gly	Lys	Pro	Glu	Arg	Cys	Leu	Arg	Phe	Val	His	Pro	Pro
		65			70				75					80	
Asn	Ala	Ser	Leu	Pro	Asn	Asp	Thr	Gln	Arg	Ala	Met	Glu	Pro	Cys	Leu
				85				90						95	
Asp	Gly	Trp	Val	Tyr	Asn	Ser	Thr	Lys	Asp	Ser	Ile	Val	Thr	Glu	Trp
			100					105					110		
Asp	Leu	Val	Cys	Asn	Ser	Asn	Lys	Leu	Lys	Glu	Met	Ala	Gln	Ser	Ile
		115					120					125			
Phe	Met	Ala	Gly	Ile	Leu	Ile	Gly	Gly	Leu	Val	Leu	Gly	Asp	Leu	Ser
		130				135						140			

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Asp Arg Phe Gly Arg Arg Pro Ile Leu Thr Cys Ser Tyr Leu Leu Leu  
 145 150 155 160  
 Ala Ala Ser Gly Ser Gly Ala Ala Phe Ser Pro Thr Phe Pro Ile Tyr  
 165 170 175  
 Met Val Phe Arg Phe Leu Cys Gly Phe Gly Ile Ser Gly Ile Thr Leu  
 180 185 190  
 Ser Thr Val Ile Leu Asn Val Glu Trp Val Pro Thr Arg Met Arg Ala  
 195 200 205  
 Ile Met Ser Thr Ala Leu Gly Tyr Cys Tyr Thr Phe Gly Gln Phe Ile  
 210 215 220  
 Leu Pro Gly Leu Ala Tyr Ala Ile Pro Gln Trp Arg Trp Leu Gln Leu  
 225 230 235 240  
 Thr Val Ser Ile Pro Phe Phe Val Phe Phe Leu Ser Ser Trp Trp Thr  
 245 250 255  
 Pro Glu Ser Ile Arg Trp Leu Val Leu Ser Gly Lys Ser Ser Glu Ala  
 260 265 270  
 Leu Lys Ile Leu Arg Arg Val Ala Val Phe Asn Gly Lys Lys Glu Glu  
 275 280 285  
 Gly Glu Arg Leu Ser Leu Glu Glu Leu Lys Leu Asn Leu Gln Lys Glu  
 290 295 300  
 Ile Ser Leu Ala Lys Ala Lys Tyr Thr Ala Ser Asp Leu Phe Arg Ile  
 305 310 315 320  
 Pro Met Leu Arg Arg Met Thr Phe Cys Leu Ser Leu Ala Trp Phe Ala  
 325 330 335  
 Thr Gly Phe Ala Tyr Tyr Ser Leu Ala Met Gly Val Glu Glu Phe Gly  
 340 345 350  
 Val Asn Leu Tyr Ile Leu Gln Ile Ile Phe Gly Gly Val Asp Val Pro  
 355 360 365  
 Ala Lys Phe Ile Thr Ile Leu Ser Leu Ser Tyr Leu Gly Arg His Thr  
 370 375 380  
 Thr Gln Ala Ala Ala Leu Leu Leu Ala Gly Gly Ala Ile Leu Ala Leu  
 385 390 395 400  
 Thr Phe Val Pro Leu Asp Leu Gln Thr Val Arg Thr Val Leu Ala Val  
 405 410 415  
 Phe Gly Lys Gly Cys Leu Ser Ser Ser Phe Ser Cys Leu Phe Leu Tyr  
 420 425 430  
 Thr Ser Glu Leu Tyr Pro Thr Val Ile Arg Gln Thr Gly Met Gly Val  
 435 440 445  
 Ser Asn Leu Trp Thr Arg Val Gly Ser Met Val Ser Pro Leu Val Lys  
 450 455 460  
 Ile Thr Gly Glu Val Gln Pro Phe Ile Pro Asn Ile Ile Tyr Gly Ile  
 465 470 475 480  
 Thr Ala Leu Leu Gly Gly Ser Ala Ala Leu Phe Leu Pro Glu Thr Leu  
 485 490 495  
 Asn Gln Pro Leu Pro Glu Thr Ile Glu Asp Leu Glu Asn Trp Ser Leu  
 500 505 510  
 Arg Ala Lys Lys Pro Lys Gln Glu Pro Glu Val Glu Lys Ala Ser Gln  
 515 520 525  
 Arg Ile Pro Leu Gln Pro His Gly Pro Gly Leu Gly Ser Ser  
 530 535 540

&lt;210&gt; SEQ ID NO 12

&lt;211&gt; LENGTH: 550

&lt;212&gt; TYPE: PRT



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&lt;213&gt; ORGANISM: Homo sapiens

&lt;400&gt; SEQUENCE: 12

Met Ala Phe Asn Asp Leu Leu Gln Gln Val Gly Gly Val Gly Arg Phe  
 1 5 10 15  
 Gln Gln Ile Gln Val Thr Leu Val Val Leu Pro Leu Leu Leu Met Ala  
 20 25 30  
 Ser His Asn Thr Leu Gln Asn Phe Thr Ala Ala Ile Pro Thr His His  
 35 40 45  
 Cys Arg Pro Pro Ala Asp Ala Asn Leu Ser Lys Asn Gly Gly Leu Glu  
 50 55 60  
 Val Trp Leu Pro Arg Asp Arg Gln Gly Gln Pro Glu Ser Cys Leu Arg  
 65 70 75 80  
 Phe Thr Ser Pro Gln Trp Gly Leu Pro Phe Leu Asn Gly Thr Glu Ala  
 85 90 95  
 Asn Gly Thr Gly Ala Thr Glu Pro Cys Thr Asp Gly Trp Ile Tyr Asp  
 100 105 110  
 Asn Ser Thr Phe Pro Ser Thr Ile Val Thr Glu Trp Asp Leu Val Cys  
 115 120 125  
 Ser His Arg Ala Leu Arg Gln Leu Ala Gln Ser Leu Tyr Met Val Gly  
 130 135 140  
 Val Leu Leu Gly Ala Met Val Phe Gly Tyr Leu Ala Asp Arg Leu Gly  
 145 150 155 160  
 Arg Arg Lys Val Leu Ile Leu Asn Tyr Leu Gln Thr Ala Val Ser Gly  
 165 170 175  
 Thr Cys Ala Ala Phe Ala Pro Asn Phe Pro Ile Tyr Cys Ala Phe Arg  
 180 185 190  
 Leu Leu Ser Gly Met Ala Leu Ala Gly Ile Ser Leu Asn Cys Met Thr  
 195 200 205  
 Leu Asn Val Glu Trp Met Pro Ile His Thr Arg Ala Cys Val Gly Thr  
 210 215 220  
 Leu Ile Gly Tyr Val Tyr Ser Leu Gly Gln Phe Leu Leu Ala Gly Val  
 225 230 235 240  
 Ala Tyr Ala Val Pro His Trp Arg His Leu Gln Leu Leu Val Ser Ala  
 245 250 255  
 Pro Phe Phe Ala Phe Phe Ile Tyr Ser Trp Phe Phe Ile Glu Ser Ala  
 260 265 270  
 Arg Trp His Ser Ser Ser Gly Arg Leu Asp Leu Thr Leu Arg Ala Leu  
 275 280 285  
 Gln Arg Val Ala Arg Ile Asn Gly Lys Arg Glu Glu Gly Ala Lys Leu  
 290 295 300  
 Ser Met Glu Val Leu Arg Ala Ser Leu Gln Lys Glu Leu Thr Met Gly  
 305 310 315 320  
 Lys Gly Gln Ala Ser Ala Met Glu Leu Leu Arg Cys Pro Thr Leu Arg  
 325 330 335  
 His Leu Phe Leu Cys Leu Ser Met Leu Trp Phe Ala Thr Ser Phe Ala  
 340 345 350  
 Tyr Tyr Gly Leu Val Met Asp Leu Gln Gly Phe Gly Val Ser Ile Tyr  
 355 360 365  
 Leu Ile Gln Val Ile Phe Gly Ala Val Asp Leu Pro Ala Lys Leu Val  
 370 375 380  
 Gly Phe Leu Val Ile Asn Ser Leu Gly Arg Arg Pro Ala Gln Met Ala  
 385 390 395 400

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Ala Leu Leu Leu Ala Gly Ile Cys Ile Leu Leu Asn Gly Val Ile Pro  
 405 410 415

Gln Asp Gln Ser Ile Val Arg Thr Ser Leu Ala Val Leu Gly Lys Gly  
 420 425 430

Cys Leu Ala Ala Ser Phe Asn Cys Ile Phe Leu Tyr Thr Gly Glu Leu  
 435 440 445

Tyr Pro Thr Met Ile Arg Gln Thr Gly Met Gly Met Gly Ser Thr Met  
 450 455 460

Ala Arg Val Gly Ser Ile Val Ser Pro Leu Val Ser Met Thr Ala Glu  
 465 470 475 480

Leu Tyr Pro Ser Met Pro Leu Phe Ile Tyr Gly Ala Val Pro Val Ala  
 485 490 495

Ala Ser Ala Val Thr Val Leu Leu Pro Glu Thr Leu Gly Gln Pro Leu  
 500 505 510

Pro Asp Thr Val Gln Asp Leu Glu Ser Arg Lys Gly Lys Gln Thr Arg  
 515 520 525

Gln Gln Gln Glu His Gln Lys Tyr Met Val Pro Leu Gln Ala Ser Ala  
 530 535 540

Gln Glu Lys Asn Gly Leu  
 545 550

<210> SEQ ID NO 13  
 <211> LENGTH: 24  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Chemically synthesized

<400> SEQUENCE: 13

ggcacattta ttcaccaaga ccag

24

<210> SEQ ID NO 14  
 <211> LENGTH: 24  
 <212> TYPE: DNA  
 <213> ORGANISM: Artificial Sequence  
 <220> FEATURE:  
 <223> OTHER INFORMATION: Chemically synthesized

<400> SEQUENCE: 14

tgtggacctc agcagcattt ggat

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What is claimed is:

1. A method of screening candidate substrates of the organic cation transporter 6 (OCT6) comprising:

- a. providing a test agent;
- b. providing mammalian cells or a mammalian cell line which express OCT6;
- c. incubating the test agent with the cells or cell line; and
- d. determining whether the test agent is a substrate for OCT6,

wherein the mammalian cells or mammalian cell line provided in step b, are leukemia cells or a leukemia cell line, respectively.

2. The method of claim 1 wherein the test agent is coupled to a detectable substance.

3. The method of claim 2 wherein the detectable substance is selected from the group consisting of extrinsically activatable enzymes, prosthetic groups, fluorescent materials, lumi-

nescent materials, bioluminescent materials, radioactive materials, positron emitting metals using various positron emission tomographies, nonradioactive paramagnetic metal ions, immunogenic tag peptide sequences, extrinsically activatable toxins, extrinsically activatable quenching agents, and antibodies.

4. The method of claim 1 wherein the step of determining whether the test agent is a substrate for OCT6 comprises analyzing whether the test agent is located intracellularly.

5. The method of claim 1, wherein step (d) comprises determining the viability of the cells or cell line.

6. The method of claim 5, wherein the viability of the cells or cell line is determined by applying a dye to the cells or cell line, wherein incorporation of the dye by the cells is indicative of death of the cells or cell line.

7. The method of claim 6, wherein the dye is trypan blue.

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