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SGK1 INHIBITOREN BEI DER BEHANDLUNG VON LONG-QT-SYNDROM

INHIBITEURS DE LA SGK1 DANS LE TRAITEMENT DU SYNDROME DU QT LONG

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(56) References cited:

WO-A1-03/074497 **WO-A1-2013/041119**
WO-A2-01/93841 **US-A1- 2008 015 141**
US-A1- 2008 015 141 **US-A1- 2008 262 096**

- **G. SEEBOHM ET AL: "Long QT Syndrome-Associated Mutations in KCNQ1 and KCNE1 Subunits Disrupt Normal Endosomal Recycling of IKs Channels", CIRCULATION RESEARCH., vol. 103, no. 12, 5 December 2008 (2008-12-05), pages 1451-1457, XP055436405, US ISSN: 0009-7330, DOI: 10.1161/CIRCRESAHA.108.177360**
- **G. SEEBOHM ET AL: "Long QT Syndrome-Associated Mutations in KCNQ1 and KCNE1 Subunits Disrupt Normal Endosomal Recycling of IKs Channels", CIRCULATION RESEARCH., vol. 103, no. 12, 5 December 2008 (2008-12-05), pages 1451-1457, XP055436405, US ISSN: 0009-7330, DOI: 10.1161/CIRCRESAHA.108.177360**

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Description**Field of the Technology**

5 [0001] The present invention relates to treatment of heart conditions such as Long QT Syndrome and cardiovascular disease by inhibition of SGK1.

Statement of Government Interest

10 [0002] This invention was made with government support under Grant No. R01 HL094677-01A1 (PI: Rosenzweig; 1/15/10-12/31/13; Role of Serum- and Glucocorticoid-Regulated Kinase-1 in Electrical Remodeling) awarded by the NHLBPI and Grant No. R21 HL104370-01 awarded by the NHLBI (PI: Rosenzweig; 7/01/10-6/30/12; Computational discovery of SGK1 inhibitors for the treatment of heart disease). The government may have certain rights in this invention.

Background

15 [0003] The annual incidence of sudden cardiac death in the United States is in the range of 180,000 to 250,000 per year and unlikely to subside in the future with the increase in the age of the population as well an increase in cardiac risk factors such as diabetes mellitus and obesity. Alterations in ionic fluxes comprise the cellular basis for the arrhythmias that underlie sudden arrhythmic deaths. These alterations usually lead to an increase in myocardial cell repolarization time and an increase in the duration of the action potential. This corresponds to an increase in the QT interval on a surface electrocardiogram. Prolongation in the QT interval has been noted in diverse conditions including myocardial ischemia, in response to administration of commonly used drugs such as erythromycin and haloperidol (as an unintended side-effect), and as a congenital condition.

20 [0004] Many genetic causes for long QT syndrome (LQTS) have been identified, with the majority of mutations seen in genes that encode for three main cardiac ion channels (KCNQ1, KCNH2 and SCN5a). Unfortunately there are no current therapies for the treatment of LQTS that address the underlying mechanistic problem, namely prolongation of the action potential and myocardial repolarization. Additionally, most anti-arrhythmic medications prolong myocardial repolarization and drugs that shorten QT interval are rare and have very small effects. Thus medications that alter ion channel function that lead to shortening of the action potential and myocardial repolarization would be a novel class of agents in treating patients with genetic and acquired LQTS.

25 [0005] Zebrafish recapitulate several key aspects of human myocardial repolarization and have a long history of being used successfully for high-throughput screening of drugs. Recently it was shown that zebrafish models of acquired LQTS as well as genetic LQTS can be used for screening small molecules that can shorten myocardial repolarization, thereby rescuing the zebrafish LQTS. In particular the *breakdance* (*bkd*) mutant carries a mutation in the KCNH2 gene with observed 2:1 heart block due to prolonged action potential and phenocopies human LQT2. High throughput screening for small molecules that rescue the phenotype of *bkd* has been successfully conducted.

30 [0006] In addition, cardiovascular disease is responsible for 600,000 deaths per year in the United States, making it the leading cause of death in both men and women. The total annual estimated costs associated with coronary artery disease alone amount to \$109 billion.

35 [0007] Heart failure (HF) is a clinical syndrome of growing prevalence that can either be acquired, for example after myocardial infarction or pressure overload as in hypertension and valvular stenosis, or can be genetic due to *de novo* or inherited mutations. While there are evidence-based therapies for systolic HF, there has not been a new class of medications approved for HF in approximately a decade. Thus there is a substantial unmet clinical need for novel therapeutic approaches in HF.

40 [0008] Cardiomyopathies are a group of diseases affecting heart muscles, also known as the myocardium. Dilated cardiomyopathy weakens the heart and affects its ability to pump enough blood to the organs. This can lead to heart failure and death, as well as problems elsewhere in the body. Approximately 25-30% of heart failure may be due to genetic mutations that can occur *de novo* in affected individuals or can be familial and inherited. Although the genetic causes of dilated cardiomyopathy (DCM) are diverse, the most common genes affected encode structural components of the cytoskeleton and/or sarcomere. Treatment for dilated cardiomyopathy is limited, and includes drug treatment to alleviate symptoms, implantation of devices, and heart transplants. However these may have unwanted side effects or be ineffective. Accordingly, there is a need for further treatment options to deal with dilated cardiomyopathy.

Summary

45 [0009] The present invention is as defined in the claims.

50 [0010] It has now been found that SGK1 inhibition in a zebrafish model of genetic, congenital Long QT syndrome has

beneficial effects. The results given in Examples 2 and 3 show that SGK1 inhibition demonstrated significant rescue of the 2:1 AV block. Based on these discoveries, methods of treating Long QT Syndrome with SGK1 inhibitors are described herein.

[0011] In one aspect, the present invention is a SGK1 inhibitor for use in a method for treating a subject for Long QT syndrome, said method comprising administering to the subject a therapeutically effective amount of at least one SGK1 inhibitor. In some embodiments, the Long QT syndrome is genetic. In one or more embodiments, the genetic long QT syndrome is characterized by a mutation in the KCNQ1 gene, the KCNH2 gene, or the SCN5a gene. In some embodiments, the Long QT syndrome is acquired (for instance as a side effect of drugs such as erythromycin or haloperidol).

[0012] The present invention provides a number of advantages over current methods of treating Long QT syndrome (genetic and acquired). For instance, the present invention describes the unexpected observation that SGK1 inhibition can rescue the 2:1 phenotype in *bkd* (breakdancer) mutants. The present invention also presents a new therapeutic target for Long QT syndrome.

[0013] Furthermore, it has now been found that SGK1 is persistently activated in a mouse model of pressure overload induced heart failure. The results given in Examples 2 and 3 show that genetic SGK1 inhibition mitigated the development of heart failure and fibrosis (scarring) after aortic constriction and reduced heart failure-associated biochemical changes in the sodium channel that lead to arrhythmia. Based on these discoveries, methods of treating cardiovascular diseases with SGK1 inhibitors are described herein.

[0014] Suitable SGK1 inhibitors include: a small molecule compound; an anti-SGK1 antibody, or antigen-binding fragment thereof; a peptide that inhibits the function of SGK1; a miRNA; an siRNA; an shRNA; a dsRNA; an antisense RNA directed to SGK1 DNA or RNA; or a polynucleotide encoding any of the above inhibitors.

[0015] It has also now been found that SGK1 is activated in hearts of patients with DCM (see Example 6). It has also been found that SGK1 inhibition in a murine model of human genetic DCM due to a mutation in the myosin heavy chain (MHC-F764L) mitigates the development of genetic DCM. Based on these discoveries, methods for treating a subject with dilated cardiomyopathy with SGK1 inhibitors are disclosed.

[0016] The present invention provides a number of advantages over current methods of treating HF, including dilated cardiomyopathy. For instance, the present invention describes the unexpected observation that SGK1 inhibition can have a protective effect on cardiac cells. The present invention also describes the observation that SGK1 is activated in human dilated cardiomyopathy, and that inhibition of SGK1 mitigates the effects of dilated cardiomyopathy.

30 Brief Description of the Drawings

[0017]

Figure 1 shows the percent rescue of 2:1 AV block in *bkd*^{-/-} mutant zebrafish treated with SGK1 morpholino (500 mM) or SGK1-DN mRNA (500 mM) at the 1-2 cell stage and scored for the phenotype at 99 hours post fertilization. The total number of fish are indicated above the corresponding columns.

Figure 2 shows that SGK1 inhibitor 2 (also identified as Compound 1A or Compound 5377051) rescues 2:1 AV block phenotype of the *bkd*^{-/-} mutant zebrafish in a dose dependent manner. Peak rescue occurs at 45 mM with no significant increase in toxicity seen in the dosages tested (gray line, open circles, right). Percent rescue at this concentration is comparable to rescue by morpholino.

Figure 3 shows cell-based assays for SGK1 inhibition.

Figure 4A shows a graph of the heart weight to body weight (HW/BW) ratios in SGK1-inhibited and wild-type mice following TAC.

Figure 4B shows a graph of left ventricle (LV) wall thickness in SGK1-inhibited and wild-type mice following TAC.

Figure 4C shows a graph of fractional shortening (FS) in SGK1-inhibited and wild-type mice following TAC.

Figure 4D shows a graph of LV dilation (LVDD) in SGK1-inhibited and wild-type mice following TAC.

Figure 4E shows a graph of normalized fibrosis area in SGK1-inhibited and wild-type mice following TAC.

Figure 4F shows histology of cardiac cells in SGK1-inhibited and wild-type mice following TAC.

Figure 5 shows a graph of infarct size (normalized to AAR) in SGK1-inhibited and wild-type mice following ischemic

injury.

Figure 6 shows a graph demonstrating that SGK1 inhibition improves cardiac function in MHC-F764L mice (*p<0.05).

5 Figure 7 shows graphs regarding SGK1 regulates Nav 1.5 channel function: A) Expression of constitutively active SGK1 (SGK1-CA) in HEK-Nav1.5 cells leads to increase in I_{NA} density, while inhibition by expression of a dominant negative kinase dead form of SGK-1 (SGK1-DN) leads to decrease in I_{NA} density (*: p < 0.05 by one way ANOVA, n =7-11 for each condition); B) Expression of SGK1-CA in HEK-Nav 1.5 cells leads to a hyperpolarizing shift of steady state activation and inactivation curves for I_{NA} ; C) Mutation of a novel SGK1 putative site T1590A leads to abrogation of the increase in I_{NA} with SGK1 activation.

10 Figure 8 shows graphs regarding inhibition of SGK1 in cultured cardiomyocytes by lead compounds as follows: A) Inhibition of SGK1 activity assessed by GSK3-beta phosphorylation in CMs infected by Ad.SGK1-CA. Different concentrations of 5377051 were assessed at 48 hours after treatment with immunoblotting (left) and quantitated (right); B) Effect of SGK1 inhibitors on Akt-induced phosphorylation of GSK3-beta in CMs infected by Ad.-myr-Akt was assessed by immunoblotting; C) Comparison of potency of 10 micromolar compounds 5377051, 6410136 with the published inhibitor EMD63863.

15 Figure 9 shows inhibition of SGK1 activity in CMs transfected by Ad.SGK1-CA by inhibitor 6410136.

Detailed Description

20 [0018] The present invention relates to the treatment of Long QT syndrome (genetic or acquired), by the inhibition of SGK1. The present invention demonstrates that SGK1 inhibition *in vivo* has a protective effect and can alleviate symptoms associated with Long QT syndrome; can reduce and alleviate symptoms associated with heart failure, arrhythmia, ischemic injury, ischemic infarction, cardiac fibrosis, vascular proliferation, restenosis, genetic or acquired dilated cardiomyopathy, and stent failure.

Long QT Syndrome

25 [0019] Long QT syndrome can be genetic (e.g. caused by a mutation in the KCNQ1 gene, the KCNH2 gene, or the SCN5a gene). Alternatively, Long QT syndrome is not associated with a genetic mutation and is acquired as a result of exposure to an external stimulus. For instance, acquired Long QT syndrome can be a side effect of drugs such as erythromycin or haloperidol. Acquired Long QT syndrome is also associated with other heart conditions such as myocardial ischemia.

Dilated Cardiomyopathy

30 [0020] Genetic dilated cardiomyopathy is characterized by a mutation in any one of a number of genes. Below in Table 1 is given a list of genes that are known to be associated with genetic dilated cardiomyopathy. Table 1 is adapted from Hershberger et al. J. Am. Coll. Cardiol. 2011 April 19; 57(16) 1641-1649. Genetic cardiomyopathies treatable by the disclosed invention, i.e., by SGK1 inhibition, include those characterized by a mutation listed in Table 1.

45 Table 1. Genes Associated with Genetic Dilated Cardiomyopathy

Gene	Protein
LMNA	Lamin A/C
MYH6	α -miosin heavv chain
MYH7	β -miosin heavv chain
MYPN	Myopalladin
TNNT2	Cardiac troponin T
SCN5A	Sodium channel
MYBPC3	Myosin-binding, protein C
RBM20	RNA binding protein 20

(continued)

Gene	Protein
TMPO	Thymopoietin
LAMA4	Laminin a-4
VCL	Metavinculin
LDB3	Cypher/ZASP
TCAP	Titin-cap or telethonin
PSEN1/2	Presenilin 1/2
ACTN2	α -actinin-2
CRYAB	Alpha B crvstalin
TPM1	α -tropomyosin
ABCC9	SUR2A
ACTC	Cardiac actin
PDLIM3	PDZ LIM domain protein 3
ILK	Integrin-linked kinase
TNNC1	Cardiac troponin C
PLN	Phospholamban
DES	Desmin
SGCD	δ -sarcoglycan
CSRP3	Muscle LIM protein
TTN	Titin
EYA4	Eves-absent 4
ANKRD1	Ankvrin repeat domain containing protein 1
DMD	Dystrophin
TAZ/G4.5	tafazzin

Acquired dilated cardiomyopathy refers to dilated cardiomyopathy having external or environmental causes and is not caused by genetic mutations. It is also something referred to as "heart failure."

SGK1

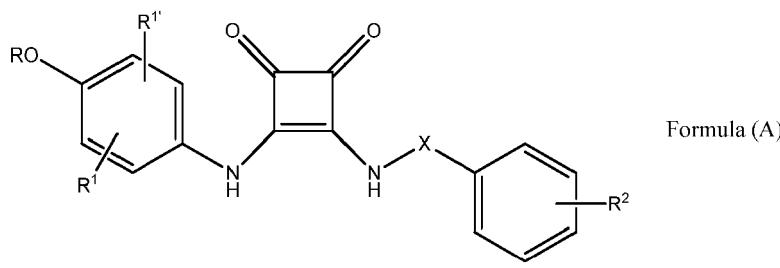
[0021] Serine/threonine-protein kinase (SGK1) (also known as serum/glucocorticoid-regulated kinase 1) is a protein kinase that plays a role in a cell's response to stress. *In vivo*, SGK1 activates certain potassium, sodium, and chloride channels. For instance, the protein is known to regulate the myo-inositol transporter during osmotic stress.

SGK1 Inhibitors

[0022] As used herein, the term "SGK1 inhibitor" refers to any agent that can block, arrest, interfere with, or reduce the biological activity of SGK1.

[0023] Inhibitors of SGK1 are known in the art, and can include small molecules. Although small molecules can have any molecular weight, they generally include molecules that are less than about 5,000 daltons.

[0024] For instance, small molecule SGK1 inhibitors are described in U.S. 2008/0262096 by Mederski et al. U.S. 2008/0262096 describes squaric acid compounds of the Formula (A), in which the variables R, R¹, R^{1'}, R² and X are defined in the specification as described below:



in which

R denotes H or A,

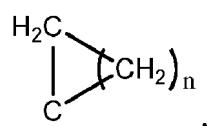
15 R¹, R^{1'} each, independently of one another, denote H, A, Hal, CN, NO₂, C(=O)A, CHO, CH(OH)A, NH₂, NH(C=O)A, COOH, COOA or SO₂NH₂, CONH₂ or CONA₂,

R² denotes OH, OA, Hal, CF₃, NO₂ or SO₂NH₂,

A denotes unbranched or branched alkyl having 1-10 C atoms, in which 1-7 H atoms may be replaced by F,

X is absent or denotes CH₂, CHA, CA₂ or

20



25

Hal denotes F, Cl, Br or I,

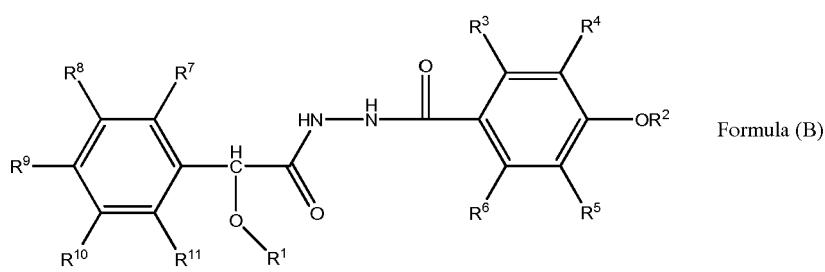
m denotes 0, 1 or 2, and

n denotes 1, 2, 3 or 4,

30 where bis(4-hydroxyphenylamino)cyclobut-3-ene-1,2-dione is excluded.

[0025] Small molecule SGK1 inhibitors have also been described in U.S. 2009/0221712 by Gericke et al. U.S. 2009/0221712 describes mandelic hydrazides of the Formula (B), in which the variables R¹- R¹¹ are defined in the specification as described below:

35



45 in which

R¹, R² each, independently of one another, denote H, CHO or acetyl,

50 R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, R¹¹ each, independently of one another, denote H, A, OSO₂A, Hal, NO₂, OR¹², N(R¹²)₂, CN, O-COA, -[C(R¹²)₂]_nCOOR¹², O-[C(R¹²)₂]_nCOOR¹², SO₃H, -[C(R¹²)₂]_nAr, -CO-Ar, O-[C(R²h)_nAr], -[C(R¹²)₂]_nHet, -[C(R¹²)₂]_nC≡CH, O-[C(R¹²)₂]_nC-CH, -[C(R¹²)₂]_nCON(R¹²)₂, -[C(R¹²)₂]_nCONR¹²N(R¹²)₂, O-[C(R¹²)₂]_nCON(R¹²)₂, O-[C(R²h)_nAr], NR¹²COA, NR¹²CON(R¹²)₂, NR¹²SO₂A, N(SO₂A)₂, COR¹², S(O)_mAr, SO₂NR² or S(O)_mA,

R³ and R⁴ together also denote CH=CH-CH=CH,

55 R³ and R⁴, R⁷ and R⁸ or R⁸ and R⁹ together also denote alkylene having 3, 4 or 5 C atoms, in which one or two CH₂ groups may be replaced by oxygen,

A denotes unbranched or branched alkyl having 1-6 C atoms, in which 1-7H atoms may be replaced by F, or cyclic alkyl having 3-7 C atoms,

Ar denotes phenyl, naphthyl or biphenyl, each of which is unsubstituted or mono-, di- or trisubstituted by Hal, A,

5 OR¹², N(R¹²)₂, NO₂, CN, phenyl, CON(R¹²)₂, NR¹²COA, NR¹²CON(R¹²)₂, NR¹²SO₂A, COR¹², SO₂N(R¹²)₂, S(O)_mA, -[C(R¹²)₂]_n-COOR¹² and/or -O[C(R¹²)₂]_n-COOR¹²,

Het denotes a mono- or bicyclic saturated, unsaturated or aromatic heterocycle having 1 to 4 N, O and/or S atoms, which may be mono-, di- or trisubstituted by Hal, A, OR¹², N(R¹²)₂, NO₂, CN, COOR¹²CON(R¹²)₂, NR¹²COA, NR¹²SO₂A, COR¹², SO₂NR¹², S(O)_mA, =S, =NR¹² and/or =O (carbonyl oxygen),

R¹² denotes H or A,

Hal denotes F, Cl, Br or I,

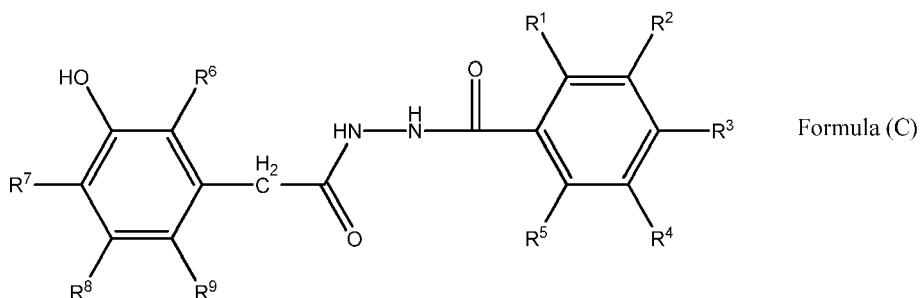
m denotes 0, 1 or 2,

n denotes 0, 1, 2 or 3, and

o denotes 1, 2 or 3.

10 [0026] Small molecule SGK1 inhibitors have further been described in U.S. 2008/0167380 by Gericke et al. U.S. 2008/0167380 describes acyl hydrazides of the Formula (C), in which R¹-R⁹ are defined in the specification as described below:

15



in which

30 R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, and R⁹ each, independently of one another, denote H, A, OSO₂A, Hal, NO₂, OR¹⁰, N(R¹⁰)₂, CN, -[C(R¹⁰)₂]_nCOOR¹⁰, O-[C(R¹⁰)₂]_oCOOR¹⁰, SO₃H, -[C(R¹⁰)₂]_nAr, -CO-Ar, O-[C(R¹⁰)₂]_nAr, -[C(R¹⁰)₂]_nHet, -[C(R¹⁰)₂]_nC≡CH, O-[C(R¹⁰)₂]_nC≡CH, -[C(R¹⁰)₂]_nCON(R¹⁰)₂, -[C(R¹⁰)₂]_nCONR¹⁰N(R¹⁰)₂, O-[C(R¹⁰)₂]_nCON(R¹⁰)₂, O-[C(R¹⁰)₂]_oCONR¹⁰N(R¹⁰)₂, NR¹⁰COA, NR¹⁰CON(R¹⁰)₂, NR¹⁰SO₂A, N(SO₂A)₂, COR¹⁰, S(O)_mAr, SO₂NR¹⁰ or S(O)_mA, or

35 R¹ and R², R² and R³, R³ and R⁴ or R⁴ and R⁵ together also denote CH=CH-CH=CH,

35 A denotes unbranched or branched alkyl having 1-6 C atoms, in which 1-7H atoms may be replaced by F, or cyclic alkyl having 3-7 C atoms,

40 Ar denotes phenyl, naphthyl or biphenyl, each of which is unsubstituted or mono-, di- or trisubstituted by Hal, A, OR¹⁰, N(R¹⁰)₂, NO₂, CN, phenyl, CON(R¹⁰)₂, NR¹⁰COA, NR¹⁰CON(R¹⁰)₂, NR¹⁰SO₂A, COR¹⁰, SO₂N(R¹⁰)₂, S(O)_mA, -[C(R¹⁰)₂]_n-COOR¹⁰ and/or -O[C(R¹⁰)₂]_o-COOR¹⁰,

45 Het denotes a mono- or bicyclic saturated, unsaturated or aromatic heterocycle having 1 to 4 N, O and/or S atoms, which may be mono-, di- or trisubstituted by Hal, A, OR¹⁰, N(R¹⁰)₂, NO₂, CN, COOR¹⁰, CON(R¹⁰)₂, NR¹⁰COA, NR¹⁰SO₂A, COR¹⁰, SO₂NR¹⁰ S(O)_mA, =S, NR¹⁰ and/or =O (carbonyl oxygen),

R¹⁰ denotes H or A,

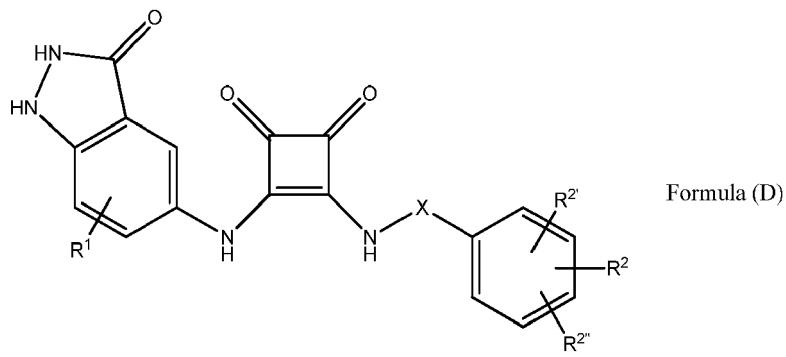
Hal denotes F, Cl, Br or I,

m denotes 0, 1 or 2,

n denotes 0, 1, 2 or 3, and

o denotes 1, 2 or 3.

55 [0027] Small molecule SGK1 inhibitors are also described in U.S. 2008/0234348 by Mederski et al. U.S. 2008/0234348 describes squaric acid compounds of the Formula (D) in which R1, R2, R2', R2" and X are defined in the specification as described below:



in which

15 R^1 denotes H, A, Hal, CN, NO_2 , $C(=O)A$, CHO , $CH(OH)A$, NH_2 , $NH(C=O)A$, $COOH$, $COOA$, SO_2NH_2 , $CONH_2$, $CONA_2$, $(CH_2)_mAr$ or Het,

20 R^2 denotes OH, OA, Hal, CF_3 , SO_2NH_2 , NHAc or $NHSO_2A$, $R^{2'}$, $R^{2''}$ each, independently of one another, denote H or Hal,

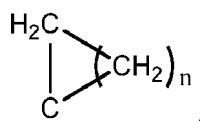
25 Ac denotes acetyl,

Ar denotes phenyl which is unsubstituted or mono-, di- or trisubstituted by Hal, A, OH, OA, NH_2 , NO_2 , CN, $COOH$, $COOA$, $CONH_2$, $NHCOA$, $NHCONH_2$, $NHSO_2A$, SO_2NH_2 and/or $S(O)_mA$,

25 Het denotes furyl, thienyl, pyrrolyl, imidazolyl, pyridyl, pyrimidinyl, pyrazolyl, thiazolyl or indolyl, each of which is unsubstituted or mono-, di- or trisubstituted by A, Hal, OH and/or OA,

25 A denotes unbranched or branched alkyl having 1-10 C atoms, in which 1-7H atoms may be replaced by F,

25 X is absent or denotes CH_2 , CHA , CA_2 or

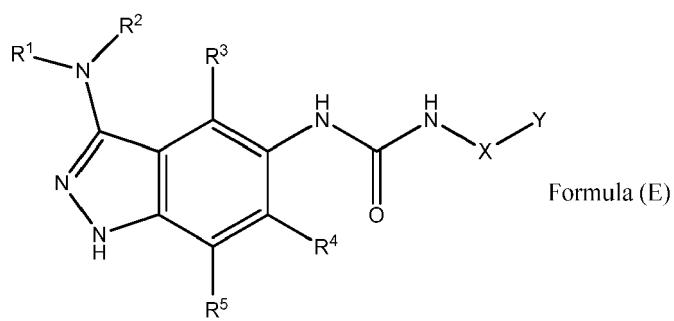


Hal denotes F, Cl, Br or I,

35 m denotes 0, 1 or 2, and

n denotes 1, 2, 3 or 4.

40 [0028] Small molecule SGK1 inhibitors are further described in U.S. 2009/0253767 by Klein et al. U.S. 2009/0253767 describes aminoindazolylurea derivatives of the Formula (E) in which R^1 , R^2 , R^3 , R^4 , R^5 , X and Y are defined in the specification as described below:



in which

55 R^1 , R^2 each, independently of one another, denote H, A, $-[C(R^7)_2]_nAr$, $-[C(R^7)_2]_nHet$, $-COHet$ or $-COAr$,

55 R^3 , R^4 , R^5 each, independently of one another, denote H, A, Hal, OH, OA, $-[C(R^7)_2]_nAr$, $-[C(R^7)_2]_nHet$, OA , $OHet$,

SH, SA, SAr, SHet, NH₂, NHA, NAA', NHAr, N(Ar)₂, NHHet, N(Het)₂, NAAr, NAHet, SOA, SOAr, SOHet, SO₂A, SO₂Ar, SO₂Het, NO₂, CN, COOH, COOA, CONH₂, CONHA, CONA₂, NHCOA, NACOA, NHCONH₂, NHCONHA, NHCONA₂, NSO₂A, NASO₂A, CHO, COA, COAr, COHet, SO₃H, SO₂NH₂, SO₂NHAr, SO₂N(Ar)₂, SO₂NHHet or SO₂N(Het)₂,

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X denotes -CR⁷R⁸-, -CR⁷R⁸CR⁹R¹- or -CR⁷R⁸C(OR⁹)R¹⁰-

Y denotes Ar or Het,

10 R⁷, R⁸, R⁹, R¹⁰ each, independently of one another, denote H or A,

R¹¹ denotes alkyl having 1-6 C atoms, in which 1-5H atoms may be replaced by F,

15 A, A' each, independently of one another, denote alkyl having 1-10 C atoms which is unsubstituted or mono-, di- or trisubstituted by R³, =S, =NR⁷ and/or =O (carbonyl oxygen) and in which one, two or three CH₂ groups may be replaced by O, S, SO, SO₂, NH, NR¹¹ and/or by -CH=CH- groups and/or, in addition, 1-7H atoms may be replaced by F and/or Cl, or cyclic alkyl having 3-7 C atoms,

20 Ar denotes phenyl, naphthyl or biphenyl, each of which is unsubstituted or mono-, di-, tri- or tetrasubstituted by A, Hal, OH, OA, Ar', OAr', Het, OHet, SH, SA, SAR', SHet, NH₂, NHA, NAA', NHAr, N(Ar)₂, NHHet, N(Het)₂, NAAr', NAHet, SOA, SOAr', SOHet, SO₂A, SO₂Ar', SO₂Het, NO₂, CN, COOH, COOA, CONH₂, CONHA, CONA₂, NHCOA, NACOA, NHCONH₂, NHCONHA, NHCONA₂, NSO₂A, NASO₂A, CHO, COA, COAr', COHet, SO₃H, SO₂NH₂, SO₂NHAr', SO₂N(Ar)₂, SO₂NHHet and/or SO₂N(Het)₂,

25 Het denotes a mono- or bicyclic saturated, unsaturated or aromatic heterocycle having 1 to 4 N, O and/or S atoms, which may be mono-, di- or trisubstituted by A, Hal, OH, OA, Ar, OAr, Het', OHet', SH, SA, SAR', SHet', NH₂, NHA, NAA', NHAr, N(Ar)₂, NHHet', N(Het)₂, NAAr', NAHet', SOA, SOAr', SOHet', SO₂A, SO₂Ar', SO₂Het', NO₂, CN, COOH, COOA, CONH₂, CONHA, CONA₂, NHCOA, NACOA, NHCONH₂, NHCONHA, NHCONA₂, NSO₂A, NASO₂A, CHO, COA, COAr', COHet', SO₃H, SO₂NH₂, SO₂NHAr', SO₂N(Ar)₂, SO₂NHHet' or SO₂N(Het)₂, =S, =NR⁷ and/or =O (carbonyl oxygen),

30 Ar' denotes phenyl which is unsubstituted or mono-, di-, tri- or tetrasubstituted by A, Hal, OH, OA, O-phenyl, SH, SA, NH₂, NHA, NAA', NH-phenyl, SOA, SO-phenyl, SO₂A, SO₂-phenyl, NO₂, CN, COOH, COOA, CONH₂, CONHA, CONA₂, NHCOA, NACOA, NHCONH₂, NHCONHA, NHCONA₂, NSO₂A, CHO, COA, CO-phenyl, SO₃H, SO₂NH₂, SO₂NH-phenyl and/or SO₂N(phenyl)₂,

35 Het' denotes a mono- or bicyclic saturated, unsaturated or aromatic heterocycle having 1 to 4 N, O and/or S atoms, which may be mono-, di- or trisubstituted by A, Hal, OH, OA, NH₂, NHA, NAA', SOA, SOAr', SO₂A, SO₂Ar', SO₂Het', NO₂, CN, COOH, COOA, CONH₂, CONHA, CONA₂, NHCOA, NACOA, NHCONH₂, NHCONHA, NHCONA₂, NSO₂A, CHO, COA, COAr', COHet', SO₃H, SO₂NH₂, SO₂NHAr', SO₂N(Ar)₂, =S, =NR⁷ and/or =O (carbonyl oxygen),

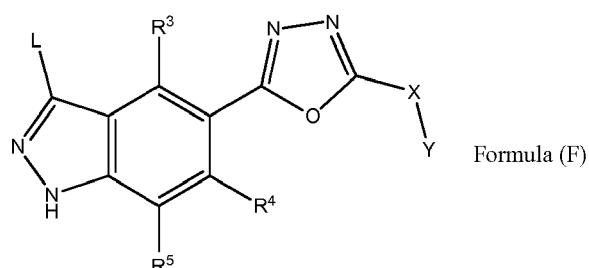
Hal denotes F, Cl, Br or I, and

40 n denotes 0, 1 or 2.

45

[0029] Small molecule SGK1 inhibitors are also described in U.S. 2010/0063115 by Klein et al. U.S. 2010/0063115 describes aminoindazolylurea derivatives of the Formula (F) in which L, X, Y, R³, R⁴ and R⁵ are described in the specification:

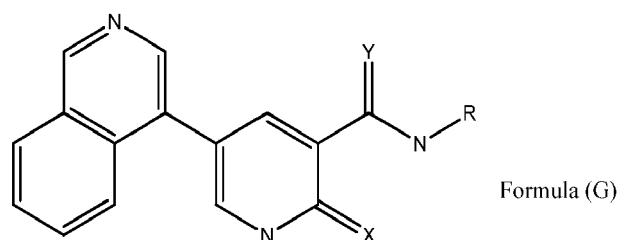
50



in which

L denotes R¹, R² or -(X)_mR³,
 X denotes CR⁷R⁸, CR⁷R⁸CR⁹R¹⁰, CR⁷R⁸C(OR⁹)R¹⁰, NR⁷, O, NR⁶CR⁷R⁸, CR⁷R⁸NR⁹, OCR⁷R⁸, OCR⁷R⁸CR⁹R¹⁰,
 5 CR⁷R⁸O, CR⁷R⁸CR⁹R¹⁰O, NR⁶CR⁷R⁸CR⁹R¹⁰, CR⁷R⁸SO₂, NR⁷CONR⁸, NR⁷CONR⁸CR⁹R¹⁰, COCR⁷R⁸, CONR⁷,
 CONR⁷CR⁸R⁹, NR⁷CR⁸R⁹CONR¹⁰, NR⁷CO or NR⁷COCR⁸R⁹,
 Y denotes H, A, Ar or Het,
 R¹ denotes CR⁹=CR⁹R¹⁰ or CR¹²=CR¹³R¹⁴,
 R² denotes C=CR¹² or C=C-Het,
 10 R³, R⁴, R⁵ each, independently of one another, denote H, A, Hal, OH, OA, -[C(R⁷)₂]_nAr, -[C(R⁷)₂]_nHet, OAr, OHet, S H, SA, SAR, S Het, NH₂, NHA, NAA', NHAr, N(Ar)₂, NHHet, N(Het)₂, NAAr, NAHet, SOA, SOAr, SOHet, SO₂A, SO₂Ar, SO₂Het, NO₂, CN, COOH, COOA, CONH₂, CONHA, CONA₂, NHCOA, NACOA, NHCONH₂, NHCONHA,
 15 NHCONA₂, NSO₂A, NASO₂A, CHO, COA, COAr, COHet, SO₃H, SO₂NH₂, SO₂NHAr, SO₂N(Ar)₂, SO₂NHHet or SO₂N(Het)₂,
 R⁶, R⁷, R⁸, R⁹, R¹⁰ each, independently of one another, denote H or A,
 R¹¹ denotes alkyl having 1-6 C atoms, in which 1-5 H atoms may be replaced by F,
 R¹², R¹³, R¹⁴ each, independently of one another, denote H or Ar,
 A, A' each, independently of one another, denote alkyl having 1-10 C atoms which is unsubstituted or mono-, di- or
 20 trisubstituted by R³, =S, =NR⁷ and/or =O (carbonyl oxygen) and in which one, two or three CH₂ groups may be replaced by O, S, SO, SO₂, NH, NR¹¹ and/or by -CH=CH-groups and/or, in addition, 1-7 H atoms may be replaced by F and/or Cl,
 or cyclic alkyl having 3-7 C atoms,
 Ar denotes phenyl, naphthyl or biphenyl, each of which is unsubstituted or mono-, di-, tri- or tetrasubstituted by A,
 25 Hal, OH, OA, Ar', OAr', Het, OHet, S H, SA, SAR', S Het, NH₂, NHA, NAA', NHAr', N(Ar')₂, NHHet, N(Het)₂, NAA', NAHet, SOA, SOAr', SOHet, SO₂A, SO₂Ar', SO₂Het, NO₂, CN, COOH, COOA, CONH₂, CONHA, CONA₂, NHCOA, NACOA, NHCONH₂, NHCONHA, NHCONA₂, NSO₂A, NASO₂A, CHO, COA, COAr', COHet, 50₃H, SO₂NH₂, SO₂NHAr', SO₂N(Ar')₂, SO₂NHHet and/or SO₂N(Het)₂,
 Het denotes a mono- or bicyclic saturated, unsaturated or aromatic heterocycle having 1 to 4 N, O and/or S atoms, which may be mono-, di- or trisubstituted by A, Hal, OH, OA, Ar, OAr, Het', OHet', SH, SA, SAR', SHet', NH₂, NHA, NAA', NHAr', N(Ar')₂, NHHet', N(Het')₂, NAA', NAHet', SOA, SOAr', SOHet', SO₂A, SO₂Ar', SO₂Het, NO₂, CN, COOH, COOA, CONH₂, CONHA, CONA₂, NHCOA, NACOA, NHCONH₂, NHCONHA, NHCONA₂, NSO₂A, NASO₂A, CHO, COA, COAr', COHet', SO₃H, SO₂NH₂, SO₂NHAr', SO₂N(Ar')₂, SO₂NHHet' or SO₂N(Het')₂, =S, =NR⁷ and/or =O (carbonyl oxygen),
 Ar' denotes phenyl which is unsubstituted or mono-, di-, tri- or tetrasubstituted by A, Hal, OH, OA, O-phenyl, SH, SA, NH₂, NHA, NAA', NH-phenyl, SOA, SO-phenyl, SO₂A, SO₂-phenyl, NO₂, CN, COOH, COOA, CONH₂, CONHA, CONA₂, NHCOA, NACOA, NHCONH₂, NHCONHA, NHCONA₂, NSO₂A, NASO₂A, CHO, COA, COAr', COHet', SO₃H, SO₂NH₂, SO₂NHAr', SO₂N(Ar')₂, SO₂NHHet' or SO₂N(Het')₂,
 35 Het' denotes a mono- or bicyclic saturated, unsaturated or aromatic heterocycle having 1 to 4 N, O and/or S atoms, which may be mono-, di- or trisubstituted by A, Hal, OH, OA, NH₂, NHA, NAA', SOA, SOAr', SO₂A, SO₂Ar', NO₂, CN, COOH, COOA, CONH₂, CONHA, CONA₂, NHCOA, NACOA, NHCONH₂, NHCONHA, NHCONA₂, NSO₂A, NASO₂A, CHO, COA, COAr', COHet', SO₃H, SO₂NH₂, SO₂NHAr', SO₂N(Ar')₂, =S, =NR⁷ and/or =O (carbonyl oxygen),
 Hal denotes F, Cl, Br or I,
 m denotes 0, 1,2 or 3, and
 n denotes 0, 1 or 2.
 45

[0030] Small molecule SGK1 inhibitors are yet further described in WO 2008/021725 by Drewry et al. WO 2008/021725 describes pyridine isoquinoline derivatives of the Formula (G) in which X, Y, and R are defined in the specification:



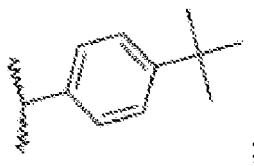
in which

5 X is O or S.

Y is O or S.

R is selected from:

10



;

(CH₂)_nR¹, wherein n is 1, 2, or 3 and R¹ is aryl;

heteroaryl optionally substituted with one or more groups independently selected from the group consisting of C₁-C₃ alkyl and oxo; and

15 aryl optionally substituted with one or more groups independently selected from the group consisting of d₃ alkyl, d₃ alkoxy, halo, C₁-C₃ haloalkyl, d₃ hydroxyalkyl, -C(O)NH₂, -OH, -CN, C₁-C₃ haloalkoxy, aryloxy, -(CH₂)_pS(O)₂NH₂, and -(CH₂)_qNR^aR^b; wherein:

20 p is 0, 1, 2, 3, 4, 5, or 6;

q is 1, 2, 3, 4, 5, or 6;

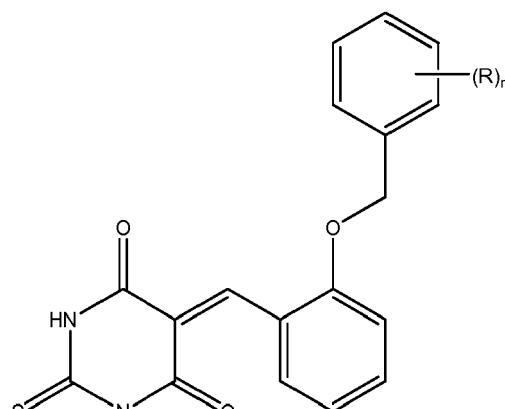
R^a is H or C₁-C₃ alkyl; and

R^b is H or C₁-C₃ alkyl.

[0031] Additional small molecule inhibitors of SGK1 can include compounds of the Formula 1:

25

30



40 (Formula 1),

wherein

n is 1, 2, 3, 4, or 5; each R is independently selected from the group consisting of hydrogen, a halogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, C₁-C₆ haloalkyl, C₁-C₆ haloalkoxy, CN, CO₂H, CO₂(C₁-C₆ alkyl), CO₂(C₁-C₆ haloalkyl), -OH, -NH₂,

45 -NH(C₁-C₆ alkyl), -N(C₁-C₆ alkyl)₂, -SH, and -S(C₁-C₆ alkyl); or a pharmaceutically acceptable salt thereof.

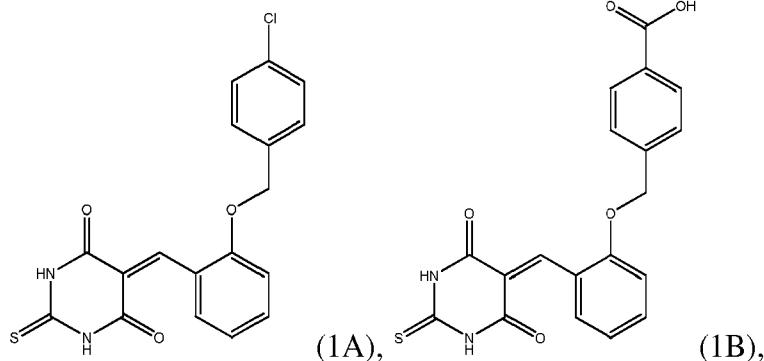
[0032] Exemplary small molecule inhibitors of SGK1 include:

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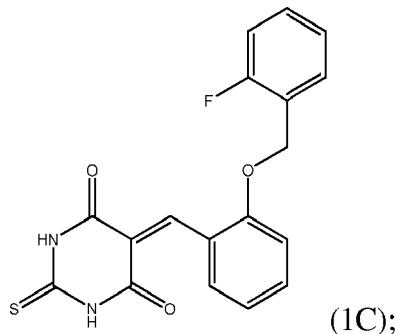


and/or

15

20

25



or a pharmaceutically acceptable salt thereof.

30 [0033] These three compounds were purchased from ChemBridge (San Diego, CA) (catalogue numbers 5377051 for 1A, 6333540 for 1B, and 6347949 for 1C).

35 [0034] Additional examples of SGK1 inhibitors can include antibodies or antigen-binding fragments thereof that selectively bind an SGK1 protein. The term "antibody" is intended to encompass all types of polyclonal and monoclonal antibodies (e.g., human, chimeric, humanized, primatized, veneered, single chain, domain antibodies (dAbs) and antigen-binding fragments of antibodies (e.g., Fv, Fc, Fd, Fab, Fab', F(ab'), dAb). (See e.g., Harlow et al. , Antibodies A Laboratory Manual, Cold Spring Harbor Laboratory, 1988). In a particular embodiment, an SGK1-specific antibody is a human antibody or humanized antibody. SGK1-specific antibodies can also be directly or indirectly linked to a cytotoxic agent.

40 [0035] Several antibodies that selectively bind SGK1 have been produced and are commercially available (e.g. from Abcam®, Cell Signaling Technology®, EMD Millipore, Sigma-Aldrich®, and Santa Cruz Biotechnology, Inc.).

45 [0036] Other antibodies or antibody fragments which can serve as SGK1 inhibitors can also be produced, constructed, engineered, and/or isolated by conventional methods or other suitable techniques. For example, antibodies which are specific for an SGK1 protein (and thus serve as SGK1 inhibitors) can be raised against an appropriate immunogen, such as a recombinant mammalian (e.g., human) SGK1 protein or portion thereof (including synthetic molecules, e.g., synthetic peptides). A variety of methods have been described (see e.g., Kohler et al., *Nature*, 256: 495-497 (1975) and *Eur. J. Immunol.* 6: 511-519 (1976); Milstein et al., *Nature* 266: 550-552 (1977); Koprowski et al., U.S. Patent No. 4,172,124; Harlow, E. and D. Lane, 1988, *Antibodies: A Laboratory Manual*, (Cold Spring Harbor Laboratory: Cold Spring Harbor, NY); *Current Protocols In Molecular Biology*, Vol. 2 (Supplement 27, Summer '94), Ausubel, F.M. et al. , Eds., (John Wiley & Sons: New York, NY), Chapter 11, (1991)).

50 [0037] Antibodies can also be raised by immunizing a suitable host (e.g., mouse) with cells that express SGK1 (e.g., cardiovascular cells/cell lines) or cells engineered to express SGK1 (e.g., transfected cells). (See e.g., Chuntharapai et al. , *J. Immunol.*, 152:1783-1789 (1994); Chuntharapai et al. U.S. Patent No. 5,440,021). For the production of monoclonal antibodies, a hybridoma can be produced by fusing a suitable immortal cell line (e.g., a myeloma cell line such as SP2/0 or P3X63Ag8.653) with antibody producing cells. The antibody-producing cells can be obtained from the peripheral blood, or preferably, the spleen or lymph nodes, of humans or other suitable animals immunized with the antigen of interest. The fused cells (hybridomas) can be isolated using selective culture conditions, and cloned by limited dilution. Cells which produce antibodies with the desired specificity can be selected by a suitable assay (e.g., ELISA).

55 [0038] Antibody fragments can be produced by enzymatic cleavage or by recombinant techniques. For example,

5 papain or pepsin cleavage can generate Fab or F(ab')₂ fragments, respectively. Other proteases with the requisite substrate specificity can also be used to generate Fab or F(ab')₂ fragments. Antibodies can also be produced in a variety of truncated forms using antibody genes in which one or more stop codons has been introduced upstream of the natural stop site. For example, a chimeric gene encoding a F(ab')₂ heavy chain portion can be designed to include DNA sequences encoding the CH₁ domain and hinge region of the heavy chain. Single chain antibodies, and human, chimeric, humanized or primatized (CDR-grafted), or veneered antibodies, as well as chimeric, CDR-grafted or veneered single chain antibodies, comprising portions derived from different species, and the like are also encompassed by the present invention and the term "antibody". The various portions of these antibodies can be joined together chemically by conventional techniques, or can be prepared as a contiguous protein using genetic engineering techniques. For example, nucleic acids encoding a chimeric or humanized chain can be expressed to produce a contiguous protein. See, e.g., Cabilly et al. , U.S. Patent No. 4,816,567; Cabilly et al. , European Patent No. 0,125,023 B1; Boss et al. , U.S. Patent No. 4,816,397; Boss et al. , European Patent No. 0,120,694 B1; Neuberger, M.S. et al. , WO 86/01533; Neuberger, M.S. et al. , European Patent No. 0,194,276 B1; Winter, U.S. Patent No. 5,225,539; Winter, European Patent No. 0,239,400 B1; Queen et al. , European Patent No. 0 451 216 B1; and Padlan, E.A. et al. , EP 0 519 596 A1. See also, Newman, R. et al. , BioTechnology, 10 10: 1455-1460 (1992), regarding primatized antibody, and Ladner et al., U.S. Patent No. 4,946,778 and Bird, R.E. et al. , Science, 242: 423-426 (1988) regarding single chain antibodies.

15 [0039] Humanized antibodies can be produced using synthetic or recombinant DNA technology using standard methods or other suitable techniques. Nucleic acid (e.g., cDNA) sequences coding for humanized variable regions can also be constructed using PCR mutagenesis methods to alter DNA sequences encoding a human or humanized chain, such as a DNA template from a previously humanized variable region (see e.g., Kamman, M., et al. , Nucl. Acids Res., 17: 20 5404 (1989)); Sato, K., et al. , Cancer Research, 53: 851-856 (1993); Daugherty, B.L. et al., Nucleic Acids Res., 19(9): 2471-2476 (1991); and Lewis, A.P. and J.S. Crowe, Gene, 101: 297-302 (1991)). Using these or other suitable methods, variants can also be readily produced. In one embodiment, cloned variable regions (e.g., dAbs) can be mutated, and sequences encoding variants with the desired specificity can be selected (e.g., from a phage library; see e.g., Krebber et al. U.S. 5,514,548; Hoogenboom et al. , WO 93/06213, published April 1, 1993).

25 [0040] Other suitable methods of producing or isolating antibodies of the requisite specificity can be used, including, for example, methods which select a recombinant antibody or antibody-binding fragment (e.g., dAbs) from a library (e.g., a phage display library), or which rely upon immunization of transgenic animals (e.g., mice). Transgenic animals capable of producing a repertoire of human antibodies are well-known in the art (e.g., Xenomouse® (Abgenix, Fremont, CA)) and can be produced using suitable methods (see e.g., Jakobovits et al. , Proc. Natl. Acad. Sci. USA, 90: 2551-2555 (1993); Jakobovits et al. , Nature, 362: 255-258 (1993); Lonberg et al. , U.S. Patent No. 5,545,806; Surani et al. , U.S. Patent No. 5,545,807; Lonberg et al. , WO 97/13852).

30 [0041] SGK1 inhibitors are also agents that inhibit (reduce, decrease, prevent, block, arrest or interfere with) the expression of an SGK1 protein. In one or more embodiments, the SGK1 inhibitor is a miRNA, a siRNA, a shRNA, a dsRNA or an antisense RNA directed to SGK1 DNA or mRNA; a polynucleotide encoding the miRNA, siRNA, shRNA, dsRNA or antisense RNA; or an equivalent of each thereof.

35 [0042] Agents (small molecules, peptides, nucleic acids, oligonucleotides) that inhibit SGK1 gene expression (e.g., transcription, mRNA processing, translation) can be effective SGK1 inhibitors. For example, small interfering ribonucleic acids (siRNAs) and, similarly, short hairpin ribonucleic acids (shRNAs) which are processed into short siRNA-like molecules in a cell, can prevent the expression (translation) of the SGK1 protein. siRNA molecules can be polynucleotides that are generally about 20 to about 25 nucleotides long and are designed to bind specific RNA sequence (e.g., SGK1 mRNA). siRNAs silence gene expression in a sequence-specific manner, binding to a target RNA (e.g., an RNA having the complementary sequence) and causing the RNA to be degraded by endoribonucleases. siRNA molecules able to inhibit the expression of the SGK1 gene product can be produced by suitable methods. There are several algorithms 40 that can be used to design siRNA molecules that bind the sequence of a gene of interest (see e.g., Mateeva O. et al. Nucleic Acids Res. 35(8):Epub, 2007; Huesken D. et al. , Nat. Biotechnol. 23:995-1001; Jagla B. et al. , RNA 11:864-872, 2005; Shabalina S.A. BMC Bioinformatics 7:65, 2005; Vert J.P. et al. BMC Bioinformatics 7:520, 2006). Expression vectors that can stably express siRNA or shRNA are available. (See e.g., Brummelkamp, T.R., Science 296: 550-553, 2002; Lee, N.S. et al. , Nature Biotechnol. 20:500-505, 2002; Miyagishi, M., and Taira, K. Nature Biotechnol. 20:497-500, 2002; Paddison, P.J., et al. , Genes & Dev. 16:948-958, 2002; Paul, C.P., et al. , Nature Biotechnol. 20:505-508; 2002; Sui, G., et al. , Proc. Natl. Acad. Sci. USA 99(6):5515-5520, 2002; Yu, J-Y., et al. , Proc. Natl. Acad. Sci. USA 99(9):6047-6052, 2002; Elbashir, S.M., et al. , Nature 411:494-498, 2001.).

45 [0043] "Short interfering RNAs" (siRNA) refer to double-stranded RNA molecules (dsRNA), generally, from about 10 to about 30 nucleotides in length that are capable of mediating RNA interference (RNAi). "RNA interference" (RNAi) refers to sequence-specific or gene specific suppression of gene expression (protein synthesis) that is mediated by short interfering RNA (siRNA). As used herein, the term siRNA includes short hairpin RNAs (shRNAs). A siRNA directed to a gene or the mRNA of a gene may be a siRNA that recognizes the mRNA of the gene and directs a RNA-induced silencing complex (RISC) to the mRNA, leading to degradation of the mRNA. A siRNA directed to a gene or the mRNA

of a gene may also be a siRNA that recognizes the mRNA and inhibits translation of the mRNA. A siRNA may be chemically modified to increase its stability and safety. See, e.g. Dykxhoorn and Lieberman (2006) *Annu. Rev. Biomed. Eng.* 8:377-402 and U.S. Patent Application Publication No.: 2008/0249055.

[0044] "Double stranded RNAs" (dsRNA) refer to double stranded RNA molecules that may be of any length and may be cleaved intracellularly into smaller RNA molecules, such as siRNA. In cells that have a competent interferon response, longer dsRNA, such as those longer than about 30 base pair in length, may trigger the interferon response. In other cells that do not have a competent interferon response, dsRNA may be used to trigger specific RNAi.

[0045] "MicroRNAs" (miRNA) refer to single-stranded RNA molecules of 21-23 nucleotides in length, which regulate gene expression. miRNAs are encoded by genes from whose DNA they are transcribed but miRNAs are not translated into protein (non-coding RNA); instead each primary transcript (a pri-miRNA) is processed into a short stem-loop structure called a pre-miRNA and finally into a functional miRNA. Mature miRNA molecules are partially complementary to one or more messenger RNA (mRNA) molecules, and their main function is to down-regulate gene expression.

[0046] siRNA, dsRNA, and miRNA to inhibit SGK1 gene expression can be designed following procedures known in the art. See, e.g., Dykxhoorn and Lieberman (2006) *Annu. Rev. Biomed. Eng.* 8:377-402; Dykxhoorn et al. (2006) *Gene Therapy* 13:541-52; Aagaard and Rossi (2007) *Adv. Drug Delivery Rev.* 59:75-86; de Fougerolles et al. (2007) *Nature Reviews Drug Discovery* 6:443-53; Krueger et al. (2007) *Oligonucleotides* 17:237-250; U.S. Patent Application Publication No.: 2008/0188430; and U.S. Patent Application Publication No.: 2008/0249055.

[0047] Delivery of siRNA, dsRNA or miRNA to a cell can be made with methods known in the art. See, e.g., Dykxhoorn and Lieberman (2006) *Annu. Rev. Biomed. Eng.* 8:377-402; Dykxhoorn et al. (2006) *Gene Therapy* 13:541-52; Aagaard and Rossi (2007) *Adv. Drug Delivery Rev.* 59:75-86; de Fougerolles et al. (2007) *Nature Reviews Drug Discovery* 6:443-53; Krueger et al. (2007) *Oligonucleotides* 17:237-250; U.S. Patent Application Publication No.: 2008/0188430; and U.S. Patent Application Publication No.: 2008/0249055.

[0048] Antisense oligonucleotides (e.g., RNA, DNA, riboprobes) can also be used as SGK1 inhibitors to inhibit SGK1 expression. Antisense oligonucleotides are generally short (~13 to ~25 nucleotides) single-stranded nucleic acids which specifically hybridize to a target nucleic acid sequence (e.g., mRNA or DNA) and induce the degradation of the target nucleic acid (e.g., degradation of the RNA through RNase H-dependent mechanisms) or sterically hinder the progression of splicing or translational machinery. (See e.g., Dias N. and Stein C.A., *Mol. Can. Ther.* 1:347-355, 2002). There are a number of different types of antisense oligonucleotides that can be used as SGK1 inhibitors including methylphosphonate oligonucleotides, phosphorothioate oligonucleotides, oligonucleotides having a hydrogen at the 2'-position of ribose replaced by an O-alkyl group (e.g., a methyl), polyamide nucleic acid (PNA), phosphorodiamidate morpholino oligomers (deoxyribose moiety is replaced by a morpholine ring), PN (N3'→P5' replacement of the oxygen at the 3' position on ribose by an amine group) and chimeric oligonucleotides (e.g., 2'-O-Methyl/phosphorothioate). Antisense oligonucleotides can be designed to be specific for an SGK1 protein using predictive algorithms. (See e.g., Ding, Y., and Lawrence, C. E., *Nucleic Acids Res.*, 29:1034-1046, 2001; Sczakiel, G., *Front. Biosci.*, 5:D194-D201, 2000; Scherr, M., et al. , *Nucleic Acids Res.*, 28:2455-2461, 2000; Patzel, V., et al. *Nucleic Acids Res.*, 27:4328-4334, 1999; Chiang, M.Y., et al. , *J. Biol. Chem.*, 266:18162-18171, 1991; Stull, R. A., et al. , *Nucleic Acids Res.*, 20:3501-3508, 1992; Ding, Y., and Lawrence, C. E., *Comput. Chem.*, 23:387-400, 1999; Lloyd, B. H., et al. , *Nucleic Acids Res.*, 29:3664-3673, 2001; Mir, K. U., and Southern, E. M., *Nat. Biotechnol.*, 17:788-792, 1999; Sohail, M., et al. , *Nucleic Acids Res.*, 29:2041 - 2051, 2001; Altman, R. K., et al. , *J. Comb. Chem.*, 1:493-508, 1999). The antisense oligonucleotides can be produced by suitable methods; for example, nucleic acid (e.g., DNA, RNA, PNA) synthesis using an automated nucleic acid synthesizer (from, e.g., Applied Biosystems) (see also Martin, P., *Helv. Chim. Acta* 78:486-504, 1995). Antisense oligonucleotides can also be stably expressed in a cell containing an appropriate expression vector.

[0049] Antisense oligonucleotides can be taken up by target cells (e.g., cardiac cells) via the process of adsorptive endocytosis. Thus, in the treatment of a subject (e.g., mammalian), antisense SGK1 oligonucleotides can be delivered to target cells (e.g., cardiac myocytes and/or other cardiac cells) by, for example, injection or infusion. For instance, purified oligonucleotides or siRNA/shRNA, can be administered alone or in a formulation with a suitable drug delivery vehicle (e.g., liposomes, cationic polymers, (e.g., poly-L-lysine' PAMAM dendrimers, polyalkylcyanoacrylate nanoparticles and polyethyleneimine) or coupled to a suitable carrier peptide (e.g., homeotic transcription factor, the *Antennapedia* peptide, Tat protein of HIV-1, E5CA peptide).

[0050] It will be appreciated that the oligonucleotides can be made using nonstandard bases (e.g., other than adenine, cytidine, guanine, thymine, and uridine) or nonstandard backbone structures to provide desirable properties (e.g., increased nuclease-resistance, tighter-binding, stability or a desired T_m). Techniques for rendering oligonucleotides nuclease-resistant include those described in PCT Publication WO 94/12633. A wide variety of useful modified oligonucleotides may be produced, including oligonucleotides having a peptide-nucleic acid (PNA) backbone (Nielsen et al. (1991) *Science* 254:1497) or incorporating 2'-O-methyl ribonucleotides, phosphorothioate nucleotides, methyl phosphonate nucleotides, phosphotriester nucleotides, phosphorothioate nucleotides, phosphoramidates. Another example of the modification is replacement of a non-bridging phosphoryl oxygen atom with a sulfur atom which increases resistance to nuclease digestion. Increased antisense polynucleotide stability can also be achieved using molecules with 2-meth-

yoxyethyl substituted backbones. See e.g., U.S. Patent Nos. 6,451,991 and 6,900,187.

[0051] Ribozymes can also be used as SGK1 inhibitors to inhibit SGK1 expression. Ribozymes are RNA molecules possessing enzymatic activity. One class of ribozymes is capable of repeatedly cleaving other separate RNA molecules into two or more pieces in a nucleotide base sequence specific manner. See Kim et al., Proc Natl Acad Sci USA, 84:8788 (1987); Haseloff & Gerlach, Nature, 334:585 (1988); and Jefferies et al., Nucleic Acid Res, 77:1371 (1989). Such ribozymes typically have two functional domains: a catalytic domain and a binding sequence that guides the binding of ribozymes to a target RNA through complementary base-pairing. Once a specifically-designed ribozyme is bound to a target mRNA, it enzymatically cleaves the target mRNA, typically reducing its stability and destroying its ability to directly translate an encoded protein. After a ribozyme has cleaved its RNA target, it is released from that target RNA and thereafter can bind and cleave another target. That is, a single ribozyme molecule can repeatedly bind and cleave new targets.

[0052] In accordance with the present invention, a ribozyme may target any portion of the mRNA encoding SGK1. Methods for selecting a ribozyme target sequence and designing and making ribozymes are generally known in the art. See e.g., U.S. Pat. Nos. 4,987,071; 5,496,698; 5,525,468; 5,631,359; 5,646,020; 5,672,511; and 6,140,491, For example, suitable ribozymes may be designed in various configurations such as hammerhead motifs, hairpin motifs, hepatitis delta virus motifs, group I intron motifs, or RNase P RNA motifs. See e.g., U.S. Pat. Nos. 4,987,071; 5,496,698; 5,525,468; 5,631,359; 5,646,020; 5,672,511; and 6,140,491; Rossi et al., AIDS Res Human Retroviruses 8:183 (1992); Hampel & Tritz, Biochemistry 28:4929 (1989); Hampel et al., Nucleic Acids Res, 18:299 (1990); Perrotta & Been, Biochemistry 31:16 (1992); and Guerrier-Takada et al., Cell, 35:849 (1983).

[0053] Ribozymes can be synthesized by the same methods used for normal RNA synthesis. For example, suitable methods are disclosed in Usman et al., JAm Chem Soc, 109:7845-7854 (1987) and Scaringe et al., Nucleic Acids Res, 18:5433-5441 (1990). Modified ribozymes may be synthesized by the methods disclosed in, e.g., U.S. Pat. No. 5,652,094; International Publication Nos. WO 91/03162; WO 92/07065 and WO 93/15187; European Patent Application No. 92110298.4; Perrault et al., Nature, 344:565 (1990); Pieken et al., Science, 253:314 (1991); and Usman & Cedergren, Trends Biochem Sci, 17:334 (1992).

[0054] "Triplex ribozymes" configurations allow for increased target cleavage relative to conventionally expressed ribozymes. Examples of triplex ribozymes include hairpin ribozymes and hammerhead ribozymes. Methods of making and using triplex ribozymes are found in, e.g., Aguino-Jarguin et al. (2008) Oligonucleotides 18(3):213-24 and U.S. Patent Application Publication No. 2005/0260163.

[0055] Proteins have been described that have the ability to translocate desired nucleic acids across a cell membrane. Typically, such proteins have amphiphilic or hydrophobic subsequences that have the ability to act as membrane-translocating carriers. For example, homeodomain proteins have the ability to translocate across cell membranes. The shortest internalizable peptide of a homeodomain protein, Antennapedia, was found to be the third helix of the protein, from amino acid position 43 to 58 (see, e.g., Prochiantz (1996) Current Opinion in Neurobiology 6:629-634. Another subsequence, the h (hydrophobic) domain of signal peptides, was found to have similar cell membrane translocation characteristics (see, e.g., Lin et al. (1995) J. Biol. Chem. 270:14255-14258). Such subsequences can be used to translocate oligonucleotides across a cell membrane. Oligonucleotides can be conveniently derivatized with such sequences. For example, a linker can be used to link the oligonucleotides and the translocation sequence. Any suitable linker can be used, e.g., a peptide linker or any other suitable chemical linker.

[0056] The present invention provides, in one embodiment, a polypeptide decoy that mimics a domain necessary for the action of SGK1. A polypeptide decoy of a protein for inhibiting the interaction between the protein and a substrate is a polypeptide that binds to the substrate but does not carry out the biological activity that such a binding would normally carry out. In one embodiment, a polypeptide decoy is a fragment of the SGK1 protein. For instance, a polypeptide decoy can comprise a 5 to 10, 10 to 15, 15 to 20, 20-25, 25-30, 30-40, 40-45 or more than 45 amino acid fragment thereof that regulates the myo-inositol transporter during osmotic stress. A polypeptide decoy can also arise from other modifications of an SGK1 protein, for instance through glycosylation or phosphorylation, or by designing similar systems that mimic peptides, such as using peptoids, D-amino acids or β -peptides. A peptide decoy that mimics the SGK1 protein can also arise from an alteration of the backbone of SGK1. SGK1 inhibitors can also include a cDNA or mRNA that encodes a portion or the full coding sequence of SGK1 that has been mutated in a way that will interfere with or inhibit the activity of the endogenous SGK1, a construct commonly called a "dominant negative".

[0057] The SGK1 inhibitors described herein for a therapeutic use may be administered with an acceptable pharmaceutical carrier. Acceptable "pharmaceutical carriers" are well known to those of skill in the art and can include, but not be limited to any of the standard pharmaceutical carriers, such as phosphate buffered saline, water and emulsions, such as oil/water emulsions and various types of wetting agents. Suitable pharmaceutical carriers can contain inert ingredients which do not interact with the SGK1 inhibitor. Standard pharmaceutical formulation techniques can be employed, such as those described in Remington's Pharmaceutical Sciences, Mack Publishing Company, Easton, PA. Suitable pharmaceutical carriers for parenteral administration include, for example, sterile water, physiological saline, bacteriostatic saline (saline containing about 0.9% mg/ml benzyl alcohol), phosphate-buffered saline, Hank's solution, Ringer's lactate

and the like. Formulations can also include small amounts of substances that enhance the effectiveness of the active ingredient (e.g., emulsifying, solubilizing, pH buffering, wetting agents). Methods of encapsulation compositions (such as in a coating of hard gelatin or cyclodextran) are known in the art. For inhalation, the agent can be solubilized and loaded into a suitable dispenser for administration (e.g., an atomizer or nebulizer or pressurized aerosol dispenser).

[0058] Agents having SGK1 binding specificity, including small molecules, or any other SGK1 inhibitor described herein can be identified in a screen, for example, a high-throughput screen of chemical compounds and/or libraries (e.g., chemical, peptide, nucleic acid libraries). Compounds or small molecules can be identified from numerous available libraries of chemical compounds from, for example, the Chemical Repository of the National Cancer Institute and the Molecular Libraries Small Molecules Repository (PubChem), as well as libraries of the Institute of Chemistry and Cell Biology at Harvard University and other libraries that are available from commercial sources (e.g., Chembridge, Peakdale, CEREP, MayBridge, Bionet). Such libraries or collections of molecules can also be prepared using well-known chemical methods, such as well-known methods of combinatorial chemistry. The libraries can be screened to identify compounds that bind and inhibit SGK1. Identified compounds can serve as lead compounds for further diversification using well-known methods of medicinal chemistry. For example, a collection of compounds that are structural variants of the lead can be prepared and screened for SGK1 binding and/or inhibitory activity. This can result in the development of a structure activity relationship that links the structure of the compounds to biological activity. Compounds that have suitable binding and inhibitory activity can be developed further for *in vivo* use. Suitable assays for SGK1 are described in, for instance, Example 3, US 2008/0234348 (Mederski et al.), WO 2008/021725 (Drewry et al.), and WO 2005/106491 (Golz et al.).

[0059] The term "treatment" is meant administering a pharmaceutical composition for the purpose of therapeutic treatment by reducing, alleviating or reversing at least one adverse effect or symptom.

[0060] The term "administering" means providing the subject with an effective amount of the SGK1 inhibitor effective to treat a disease or condition in the subject. Methods of administering pharmaceutical compositions are well known to those of skill in the art and include, but are not limited to, microinjection, intravenous or parenteral administration. The compositions are intended for systemic, topical, oral, or local administration as well as intravenously, subcutaneously, or intramuscularly. Administration can be effected continuously or intermittently throughout the course of treatment. Methods of determining the most effective means and dosage of administration are well known to those of skill in the art and will vary with the vector used for therapy, the polypeptide or protein used for therapy, the purpose of the therapy, the target cell being treated, and the subject being treated. Single or multiple administrations can be carried out with the dose level and pattern being selected by the treating physician. For example, the compositions can be administered prior to a subject already suffering from a disease or condition that is linked to apoptosis.

[0061] The term "therapeutically effective amount" refers to a quantity of compound (e.g., an SGK1 inhibitor) delivered with sufficient frequency to provide a medical benefit to the patient. In one embodiment, an effective amount of an SGK1 inhibitor is an amount sufficient to treat or ameliorate a symptom of Long QT syndrome or an immune complex mediated disease. Exemplary effective amounts of an SGK1 inhibitor range from 0.1 ug/kg body weight to 100 mg/kg body weight; alternatively 1.0 ug/kg body weight to 10mg/kg body weight.

[0062] A "subject" includes mammals, e.g., humans, companion animals (e.g., dogs, cats, birds and the like), farm animals (e.g., cows, sheep, pigs, horses, fowl and the like) and laboratory animals (e.g., rats, mice, guinea pigs and the like). In a preferred embodiment of the disclosed methods, the subject is human.

[0063] The present invention is further defined in the following Examples. It should be understood that these Examples, while indicating preferred embodiments of the invention, are given by way of illustration only.

Examples

45 Example 1

SGK1 activity regulates Nav1.5-mediated sodium flux

[0064] HEK 293 cells with a stably transfected SCN5a gene (referred to as HEK-Nav1.5) were transiently transfected with a plasmid expressing either a constitutively-active form of SGK1 (SGK1 S422D, referred to as SGK1-CA), a dominant negative form of SGK1 (SGK1 K127M, referred to as SGK1-DN) or the vector with an empty cassette. 48 hours after transfection, cells were studied by whole cell voltage clamping.

[0065] Expression of SGK1-CA led to an increase, while expression of SGK1-DN led to a decrease in I_{NA} current density (Fig. 7A), compared to empty plasmid. The fact that inhibition of SGK1 in cells transfected with SGK1-DN led to a decrease in I_{NA} current density suggested that there is a baseline SGK1 activity that drive sodium flux in the cell.

[0066] In addition, HEK-Nav1.5 cells transfected with SGK1-CA had left-ward (hyperpolarizing) shifts of their steady state activation and inactivation curves (Fig. 7B). This is also expected to increase the 'window' current, leading to an increase in I_{NA} .

[0067] Three consensus sites for phosphorylation on Nav 1.5 had been identified, including two sites in the linker between subdomains I and II, and a previously undescribed site on the C-terminal domain. The first two sites had been shown to influence channel biophysical properties in oocyte expression experiments. Mutation of the serines to alanines (to abolish SGK1 phosphorylation) also abolished the hyperpolarization shifts in the activation and inactivation curves described above. Mutation of the C-terminal residue (T1590A) to a non-phosphorylatable residue prevented the effects of SGK1 activation on I_{NA} current density (Fig. 7C). Taken together, the data suggests that SGK1 regulates I_{NA} in multiple different ways, and inhibition of SGK1 may be a novel manner to reduce I_{NA} in pathological conditions.

Example 2

Genetic Inhibition of SGK1 Rescues 2:1 Phenotype in Bkd Mutant Zebrafish

[0068] The zebrafish breakdance mutant (Bkd^{-/-}) fish is a unique model of action potential prolongation due to a mutation in the Zerg (zebrafish HERG) channel, that recapitulates human LQT2 syndrome and has been used in high throughput screens to identify molecules that may shorten the action potential duration. The phenotypic manifestation of the prolonged QT interval in the Bkd^{-/-} fish is the presence of functional 2:1 AV block, and previous screens have taken advantage of this easily quantifiable phenotype to screen for 'rescue' from 2:1 AV block.

[0069] Bkd^{-/-} (Breakdance) adults were crossed, and the resultant embryos were reared in E3 buffer at 25°C. 24 hours post-fertilization (hpf) animals were dechorionated with pronase and plated per well in 96-well plates in a final volume of 200 μ L. At 48 hpf, compounds were added to the appropriate concentration. Zebrafish were visually scored at 72 hpf for presence or absence of 2:1 AV block. For the dose response curve, the fish at 48 hpf were treated for 24 hours at the desired concentration (from 20 mM stock in DMSO) and scored for suppression of breakdance phenotype. Genetic inhibition of SGK1 was achieved either by injection of 500 μ M of SGK1 morpholino or SGK1-dominant negative mRNA at the 1-2 cell stage and the fish were scored for rescue of 2:1 AV block at the 99 hpf stage.

[0070] Bkd^{-/-} fish injected with 500 μ M of SGK1 morpholino or SGK1-DN mRNA (that have previously been shown to inhibit SGK1 activity in a dominant negative fashion) demonstrated significant rescue of the 2:1 AV block. SGK-1 morpholino showed rescue of 58% of mutants (compared with 5% spontaneous reversion in control scrambled morpholino injections). SGK1-DN was an even more potent in rescuing the phenotype (90% rescue) (Fig. 1). These results demonstrated that SGK1 inhibition could rescue the 2:1 AV block phenotype in this zebrafish model of LQTS. This may be due to shortening of the action potential duration and myocardial repolarization by modulation of ion channel function

Example 3

Compound 1A is Effective in Reversing the 2:1 Block in *bkd* Mutants

[0071] Bkd^{-/-} fish treated with compound 1A at the concentrations indicated in Fig. 2 was effective at rescuing the 2:1 AV block phenotype with 58% rescue noted at a concentration of 45 μ M with minimal and stable toxicity in concentrations up to 50 μ M.

[0072] For Examples 4 and 5, to test the benefits of SGK1 inhibition, a kinase-dead, dominant negative mutant form of SGK1 was genetically expressed in the hearts of mice, under control of the myosin-heavy-chain-alpha promoter. These mice were normal at baseline, and subsequently subjected to either transvers aortic constriction (TAC)-induced pressure overload or transient surgical ligation of a coronary artery, to induce a heart attack.

Example 4

SGK1 Inhibition Mitigates the Effect of Pressure Overload

[0073] For TAC, 12 week old male mice were anesthetized and underwent thoracotomy at which time the transverse was constricted to the diameter of a blunted 25G needle. Mice recovered and cardiac function was followed by echocardiography. Seven weeks after surgery, mice were euthanized for tissue studies including analysis of fibrosis.

[0074] Seven weeks after TAC, heart weight to body weight (HW/BW) ratios and left ventricular (LV) wall thickness increased less in SGK1-inhibited (SGK1-DN) than in wild-type (WT) littermates (**Figure 4A and 4B**). Echocardiography seven weeks after TAC revealed reduced cardiac function (fractional shortening, FS) and increased LV dilatation (LVDD) in WT but not SGK1-DN mice (**Figure 4C and 4D**). SGK1-DN mice also had significantly less fibrosis than WT mice after TAC (**Figure 4E**). Thus, SGK1 inhibition protects against the development of heart failure, cardiac dilation, and fibrosis following pressure overload, which simulates events that occur in humans in response to hypertension or valvular stenosis.

Example 5SGK1 Inhibition Reduces Injury and Improves Function After Ischemic Injury (Heart Attack)

5 [0075] For ischemia-reperfusion injury (IRI) that simulates a heart attack receiving reperfusion therapy, the current standard of care, 12-week old SGK1-inhibited (SGK1-DN) mice and WT controls were subjected to 30 minutes of ischemia (with reversible left coronary ligation) followed by 24 hours of reperfusion. At twenty-four hours after IRI, hearts were analyzed for infarct size and "area-at-risk" (AAR) was assessed by injection of fluorescent microspheres at the time of infarction and area of infarct was assessed by TTC staining. Quantitation showed that there was a significant reduction 10 (40%, p<0.04) in infarct size with chronic SGK1 inhibition compared with WT mice (**Figure 5**). This reduction in infarct size (in response to the same ischemic stimulus) was associated with improved cardiac function as measured by echocardiography (FS, p<0.05) for at least two weeks after IRI.

Example 6SGK1 is Activated in Human Dilated Cardiomyopathy (DCM)

15 [0076] Ventricular tissue from patients with hypertensive heart disease who died from non-cardiac causes (HHD) as well as explants from patients with heart failure and dilated cardiomyopathy (DCM) at the time of orthotopic transplantation 20 was examined. Healthy unused donor hearts served as controls. The cohorts were age and gender matched. Coronary angiograms had confirmed lack of significant epicardial coronary artery disease in the patients with DCM, and gravimetry confirmed a trend towards increased LV mass in patients with HHD (CTRL 363±133 gm, HHD 518±112 gm, p =0.07). In HHD hearts, there was an increase in total SGK1 (1.22±0.12, p<0.04) without a change in pSGK1 compared with 25 controls. DCM hearts had a 2.3-fold increase in pSGK1 (2.3±0.85-fold, p<0.005 vs controls) without alterations in total SGK1. This phosphorylation site is indicative of SGK1 activation and correlates well with kinase activity. Thus these studies demonstrate that SGK1 is activated in human DCM.

Example 7SGK1 Inhibition Mitigates Genetic DCM

30 [0077] To determine whether activation of SGK1 is functionally important in DCM, a well-characterized model of genetic DCM caused by a mutation in the myosin heavy chain (MHC-F764L) was selected. While this model is the result of a specific mutation, the gene involved is commonly mutated in cardiomyopathy and is a member of the class of structural 35 proteins (involved in cytoskeleton and sarcomere function) that is most commonly mutated in DCM. SGK1 was activated or inhibited in MHC-F764L mice by breeding two mice expressing either a constitutively active (SGK1-CA) or dominant negative (SGK1-DN) SGK1 mutant. MHC-F764L heterozygotes (hets) have significant LV dysfunction compared to WT (Figure 6, p<0.05). At 12 weeks, SGK1 activation (SGK1-CA) worsened fractional shortening (FS) in MHC-F764L hets, 40 while SGK1 inhibition (SGK1-DN) substantially improved FS (Figure 6, p<0.05).

Example 8Discovery of Small Molecule SGK1 Inhibitors

45 [0078] Using a computational drug discovery platform, three iterative rounds of testing predicted inhibitors were performed in a polarization-based in-solution kinase assay using purified recombinant SGK1 protein and the inhibitors to identify three inhibitor 'lead compounds' that perform well in solution. Briefly, this assay utilizes a phosphorylated peptide that is labeled with a green fluorescent dye: this has a low molecular weight, hence a minimal fluorescence polarization value. This tracer binds to a phospho-specific antibody forming a high molecular weight complex with a high polarization 50 value. In the kinase reaction, a competitive non-labeled peptide gets phosphorylated by the kinase and competes with the labeled tracer for binding with the antibody. With increased kinase activity, less labeled tracer is bound to the antibody, which in turn results in a decrease in fluorescence polarization. First, the reproducibility and dynamic range of the assay was validated using different concentrations of a recombinant GST-purified SGK-1 and determined the EC70 of the recombinant kinase to be 1 ng. This assay was then used in a high-throughput validation format to evaluate the 1st and 55 2nd generation small molecule inhibitors identified using our CADD platform. Initial activity screening was done at a concentration of 50 M to ensure finding all compounds with activity and several compounds had demonstrable SGK1 inhibitory potential.

[0079] These inhibitors were further characterized using a cell-based cardiomyocyte system (Figure 3). 2 x 10⁶ neonatal

rat ventricular myocytes (NRVMs) were plated on 60 mm plates in DMEM containing 10% horse serum, 5% fetal bovine serum, 1% L-glutamine and 1% penstrep antibiotic. 24 h after plating media was changed to DMEM containing 1% L-glutamine, and 24 h following cells were infected with SGK-CA virus and treated with various concentrations of SGK inhibitors 5377051 (Compound 1A) and 6410136 (Chembio compound catalogue #6410136). 12 h after infection and treatment, the media was changed and cells were treated again. 12 h following the second treatment NRVMs were harvested and lysed in 300 μ L of cell lysis buffer (Cell Signaling) containing protease and phosphatase inhibitors and PMSF at 4 degrees C for 1 h with shaking. Lysates were subjected to immunoblotting with an antibody against phospho (ser38)-glycogen synthase kinase beta (p-GSK3 β), a well-established SGK1 substrate. For gel electrophoresis, 50 ug of protein per well was loaded, transferred to nitrocellulose membrane, blocked, and incubated with GAPDH and P-GSK3 β primary antibodies. As expected, Ad. SGK1-CA caused an increase in p-GSK3 β , which was inhibited by both 5377051 and 6410136, suggesting inhibition of SGK1 in CMs (Fig. 8A, Fig. 9). The SGK1 inhibitors reproducibly and effectively reduced SGK1-driven phosphorylation of GSK-3 β at concentrations in the 10-100's nM range (Figure 3). In contrast, none of the inhibitors had a significant effect on Akt-mediated phosphorylation of GSK-3 β even at significantly higher concentrations in a similar cardiomyocyte-based assay (Figure 3).

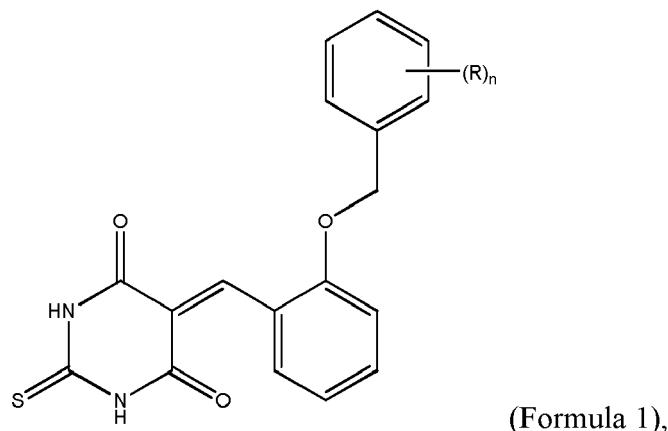
[0080] Because SGK1 and Akt are structurally related but have very different roles in the heart, with Akt deemed to be cardioprotective against stress, the inhibitors were assessed to determine if they have an off-target effect on Akt in CMs. NRVMs were infected with adenoviral-driven constitutively active Akt (myr-Akt) or control adenovirus, and treated with the inhibitors as above. The levels of p-GSK3 β , a common substrate for Akt and SGK1, were assessed again. While infection of NRVMs with Ad.myr-Akt caused an expected increase in p-GSK3 β , neither 5377051 nor 6410136 caused a significant inhibition of this activity (Fig. 8B).

[0081] The activity of 5377051 and 6410136 were compared with a previously reported SGK1 inhibitor EMD 638683 in this cellular assay and found that the lead compounds of the instant invention were better or at least equivalent in their ability to inhibit SGK1 (Fig. 8C).

Claims

1. A SGK1 inhibitor for use in a method for treating a subject for Long QT syndrome, said method comprising administering to the subject a therapeutically effective amount of at least one SGK1 inhibitor.
2. A SGK1 inhibitor for use as claimed in claim 1, wherein the Long QT syndrome is genetic.
3. A SGK1 inhibitor for use as claimed in claim 1, wherein the Long QT syndrome is acquired.
4. A SGK1 inhibitor for use as claimed in Claim 2, wherein the genetic long QT Syndrome is **characterized by** a mutation in the KCNQ1 gene.
5. A SGK1 inhibitor for use as claimed in Claim 2, wherein the genetic Long QT Syndrome is **characterized by** a mutation in the KCNH2 gene.
6. A SGK1 inhibitor for use as claimed in Claim 2 wherein the genetic Long QT Syndrome is **characterized by** a mutation in the SCN5a gene.
7. A SGK1 inhibitor for use as claimed in Claim 4, wherein the genetic Long QT syndrome is **characterized by** a loss of function in the KCNQ1 gene.
8. A SGK1 inhibitor for use as claimed in Claim 5, wherein the genetic Long QT syndrome is **characterized by** a loss of function in the KCNH2 gene.
9. A SGK1 inhibitor for use as claimed in Claim 6, wherein the genetic Long QT syndrome is **characterized by** a gain in function in the SCN5a gene.
10. A SGK1 inhibitor for use as claimed in any of claims 1-9, wherein the SGK1 inhibitor is a compound according to Formula 1:

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wherein:

n is 1, 2, 3, 4, or 5;

each R is independently selected from the group consisting of hydrogen, a halogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, C₁-C₆ haloalkyl, C₁-C₆ haloalkoxy, CN, CO₂H, CO₂(C₁-C₆ alkyl), CO₂(C₁-C₆ haloalkyl), -OH, -NH₂, -NH(C₁-C₆ alkyl), -N(C₁-C₆ alkyl)₂, -SH, and -S(C₁-C₆ alkyl);

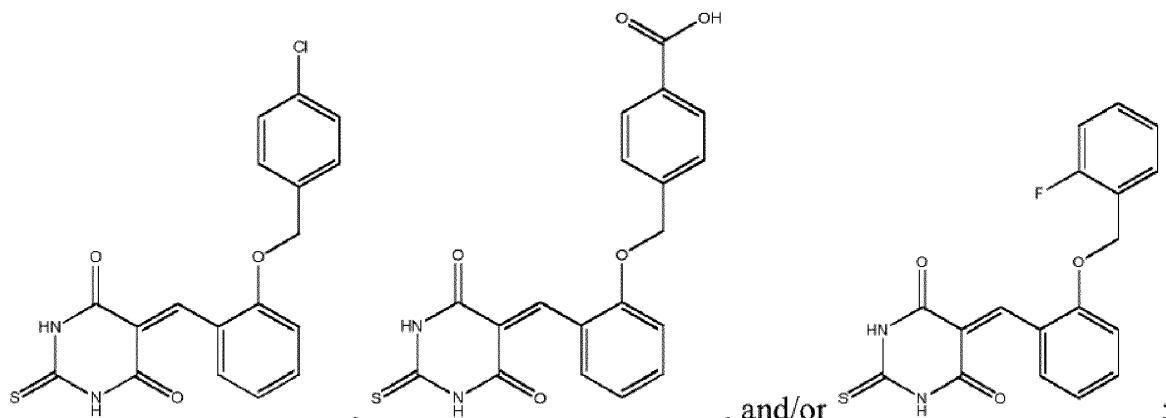
or a pharmaceutically acceptable salt thereof.

25 11. A SGK1 inhibitor for use as claimed in claim 10, wherein the SGK1 inhibitor is selected from:

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or a pharmaceutically acceptable salt thereof.

50 12. A SGK1 inhibitor for use as claimed in claims 1-9, wherein the SGK1 inhibitor is an anti-SGK1 antibody, or antigen-binding fragment thereof.

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13. A SGK1 inhibitor for use as claimed in claim 12, wherein the anti-SGK1 antibody, or antigen-binding fragment thereof is a human, chimeric, humanized, primatized, veneered, single chain, or domain (dAb) antibody, or antigen-binding fragment thereof.

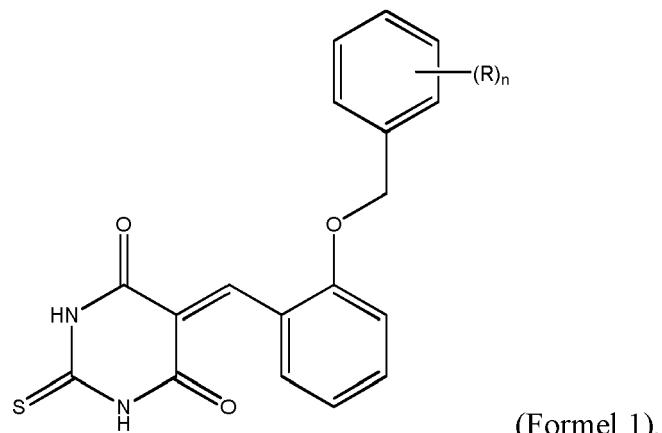
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Patentansprüche

1. SGK1-Inhibitor zur Verwendung bei einem Verfahren zur Behandlung eines Individuums gegen Long-QT-Syndrome,

wobei das Verfahren Verabreichen einer therapeutisch wirksamen Menge wenigstens eines SGK1-Inhibitors an das Individuum umfasst.

2. SGK1-Inhibitor zur Verwendung nach Anspruch 1, wobei das Long-QT-Syndrom erblich ist.
3. SGK1-Inhibitor zur Verwendung nach Anspruch 1, wobei das Long-QT-Syndrom erworben ist.
4. SGK1-Inhibitor zur Verwendung nach Anspruch 2, wobei das erbliche Long-QT-Syndrom durch eine Mutation im KCNQ1-Gen gekennzeichnet ist.
5. SGK1-Inhibitor zur Verwendung nach Anspruch 2, wobei das erbliche Long-QT-Syndrom durch eine Mutation im KCNH2-Gen gekennzeichnet ist.
6. SGK1-Inhibitor zur Verwendung nach Anspruch 2, wobei das erbliche Long-QT-Syndrom durch eine Mutation im SCN5a-Gen gekennzeichnet ist.
7. SGK1-Inhibitor zur Verwendung nach Anspruch 4, wobei das erbliche Long-QT-Syndrom durch einen Funktionsverlust beim KCNQ1-Gen gekennzeichnet ist.
8. SGK1-Inhibitor zur Verwendung nach Anspruch 5, wobei das erbliche Long-QT-Syndrom durch einen Funktionsverlust beim KCNH2-Gen gekennzeichnet ist.
9. SGK1-Inhibitor zur Verwendung nach Anspruch 6, wobei das erbliche Long-QT-Syndrom durch einen Funktionsverlust beim SCN5a-Gen gekennzeichnet ist.
10. SGK1-Inhibitor zur Verwendung nach einem der Ansprüche 1-9, wobei es sich bei dem SGK1-Inhibitor um eine Verbindung gemäß der Formel 1:



handelt, wobei:

n gleich 1, 2, 3, 4 oder 5 ist;

R jeweils unabhängig ausgewählt ist aus der Gruppe bestehend aus Wasserstoff, einem Halogen, C₁-C₆-Alkyl, C₁-C₆-Alkoxy, C₁-C₆-Halogenalkyl, C₁-C₆-Halogenalkoxy, CN, CO₂H, CO₂(C₁-C₆-Alkyl), CO₂(C₁-C₆-Halogenalkyl), -OH, -NH₂, -NH(C₁-C₆-Alkyl), -N(C₁-C₆-Alkyl)₂, -SH und -S(C₁-C₆-Alkyl);

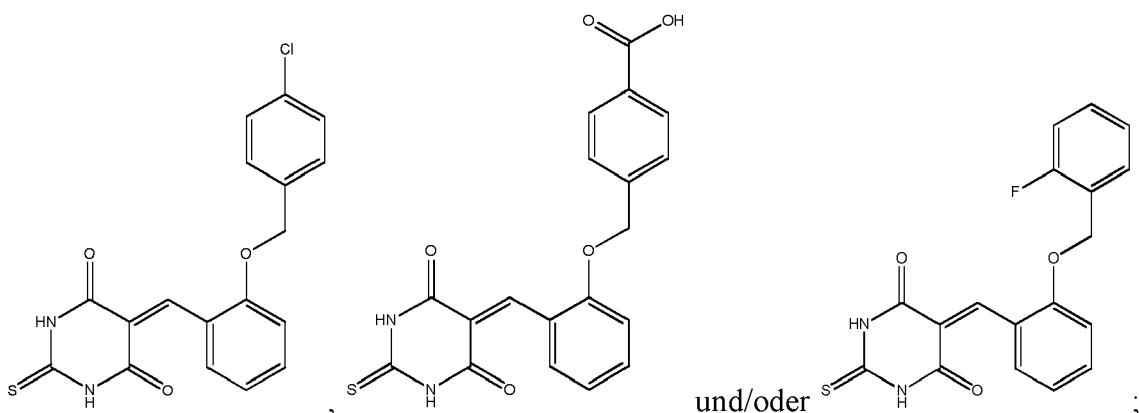
oder ein pharmazeutisch unbedenkliches Salz davon.

11. SGK1-Inhibitor zur Verwendung nach Anspruch 10, wobei der SGK1-Inhibitor ausgewählt ist aus:

5

10

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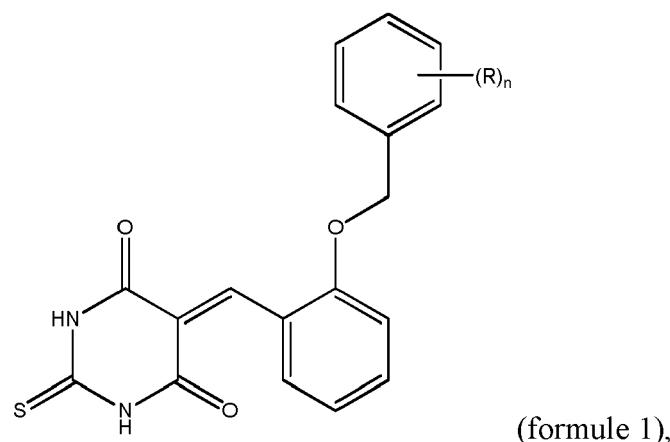
oder ein pharmazeutisch unbedenkliches Salz davon.

12. SGK1-Inhibitor zur Verwendung nach einem der Ansprüche 1-9, wobei es sich bei dem SGK1-Inhibitor um einen Anti-SGK1-Antikörper bzw. ein antigenbindendes Fragment davon handelt.
13. SGK1-Inhibitor zur Verwendung nach Anspruch 12, wobei es sich bei dem Anti-SGK1-Antikörper oder antigenbindenden Fragment davon um einen menschlichen, chimären, humanisierten, primatisierten, furnierten, Einzelkette- oder Domäne(dAb)-Antikörper bzw. ein antigenbindendes Fragment davon handelt.

25

Revendications

1. Inhibiteur de SGK1 pour utilisation dans un procédé de traitement d'un sujet pour le syndrome du QT long, ledit procédé comprenant l'administration au sujet d'une quantité thérapeutiquement efficace d'au moins un inhibiteur de SGK1.
2. Inhibiteur de SGK1 pour utilisation selon la revendication 1, où le syndrome du QT long est génétique.
3. Inhibiteur de SGK1 pour utilisation selon la revendication 1, où le syndrome du QT long est acquis.
4. Inhibiteur de SGK1 pour utilisation selon la revendication 2, où le syndrome du QT long génétique est **caractérisé par** une mutation dans le gène KCNQ1.
5. Inhibiteur de SGK1 pour utilisation selon la revendication 2, où le syndrome du QT long génétique est **caractérisé par** une mutation dans le gène KCNH2.
6. Inhibiteur de SGK1 pour utilisation selon la revendication 2 où le syndrome du QT long génétique est **caractérisé par** une mutation dans le gène SCN5a.
7. Inhibiteur de SGK1 pour utilisation selon la revendication 4, où le syndrome du QT long génétique est **caractérisé par** une perte de fonction dans le gène KCNQ1.
8. Inhibiteur de SGK1 pour utilisation selon la revendication 5, où le syndrome du QT long génétique est **caractérisé par** une perte de fonction dans le gène KCNH2.
9. Inhibiteur de SGK1 pour utilisation selon la revendication 6, où le syndrome du QT long génétique est **caractérisé par** un gain de fonction du gène SCN5a.
10. Inhibiteur de SGK1 pour utilisation selon l'une quelconque des revendications 1 à 9, où l'inhibiteur de SGK1 est un composé selon la formule 1 :



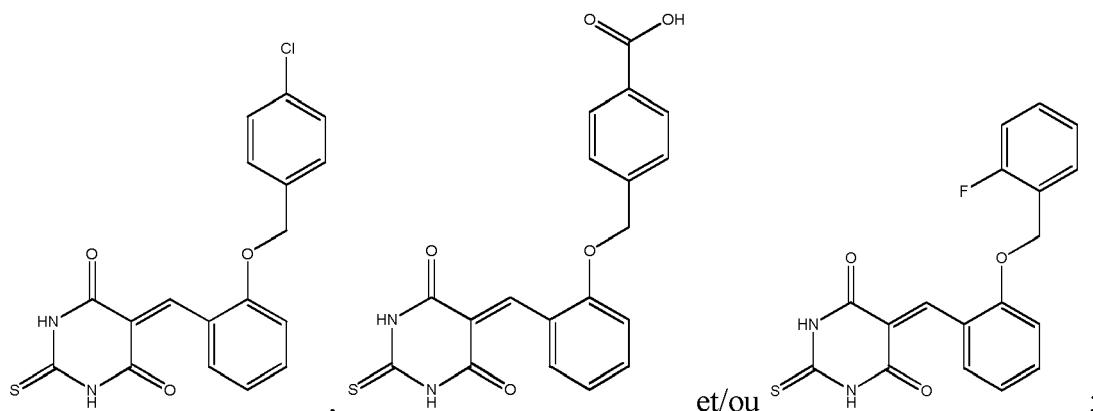
dans lequel :

n est 1, 2, 3, 4 ou 5 ;

20 chaque R est indépendamment choisi dans le groupe constitué d'hydrogène, un halogène, alkyle en C₁-C₆, alcoxy en C₁-C₆, halogénoalkyle en C₁-C₆, halogénoalcoxy en C₁-C₆, CN, CO₂H, CO₂(alkyle en C₁-C₆), CO₂(halogénoalkyle en C₁-C₆), -OH, -NH₂, -NH(alkyle en C₁-C₆), -N(alkyle en C₁-C₆)₂, -SH, et -S(alkyle en C₁-C₆) ;

25 ou un sel pharmaceutiquement acceptable de celui-ci.

11. Inhibiteur de SGK1 pour utilisation selon la revendication 10, où l'inhibiteur de SGK1 est choisi parmi :



45 ou un sel pharmaceutiquement acceptable de ceux-ci.

12. Inhibiteur de SGK1 pour utilisation selon les revendications 1 à 9, où l'inhibiteur de SGK1 est un anticorps anti-SGK1, ou un fragment de liaison d'antigène de celui-ci.

50 13. Inhibiteur de SGK1 pour utilisation selon la revendication 12, où l'anticorps anti-SGK1, ou un fragment de liaison d'antigène de celui-ci, est un anticorps humain, chimérique, humanisé, primatisé, plaqué, monocaténaire ou à domaine unique (dAb), ou un fragment de liaison d'antigène de ceux-ci.

Figure 1

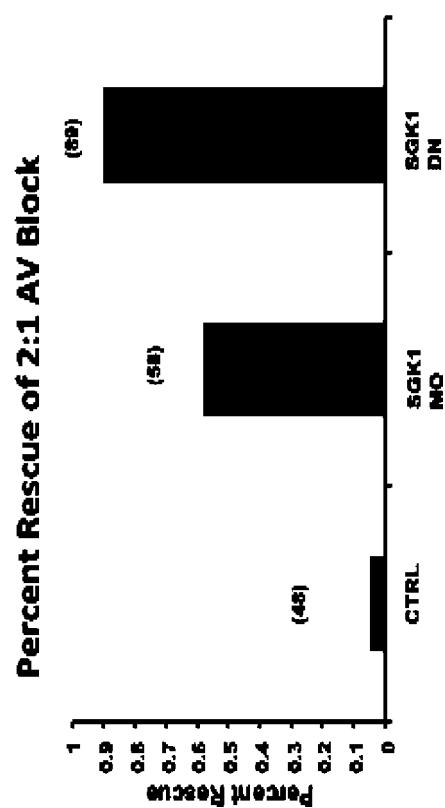


Figure 2

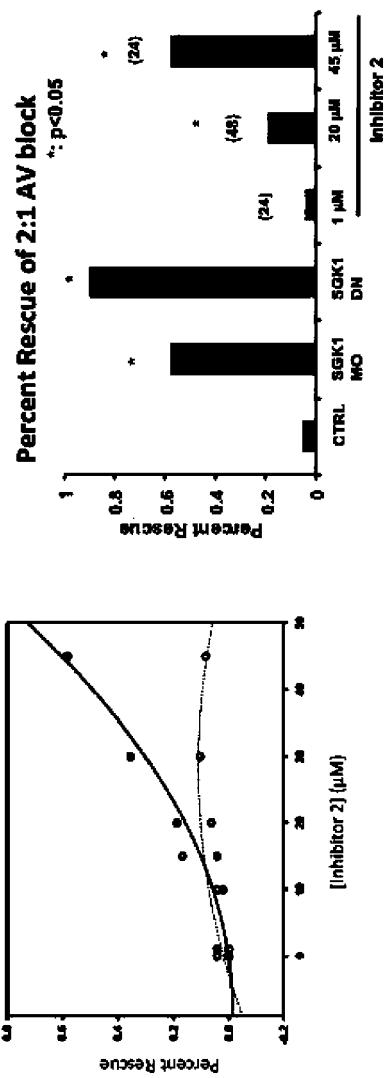


Figure 3

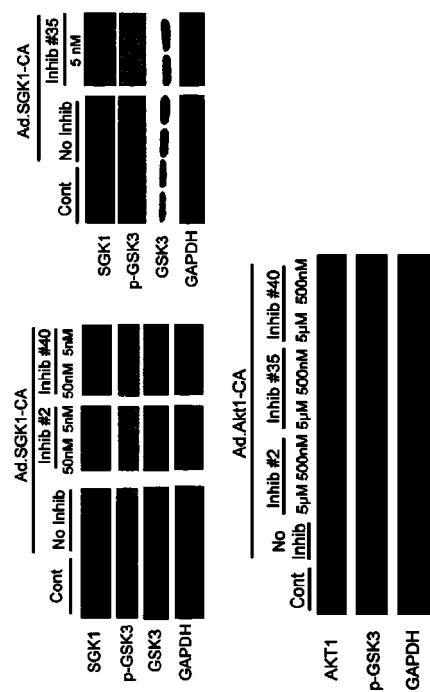
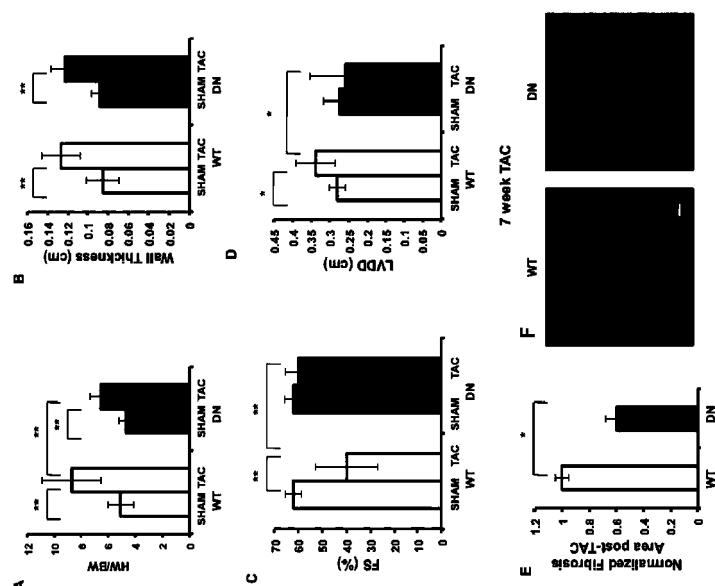


Figure 4



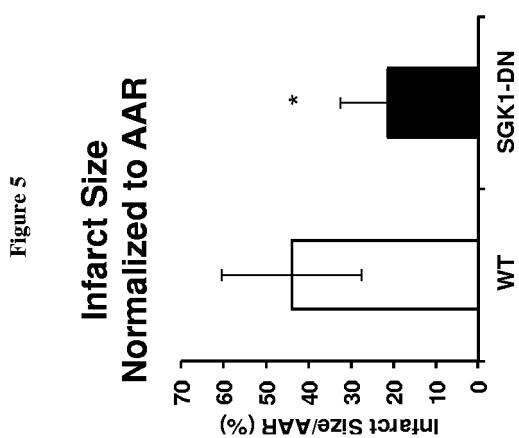
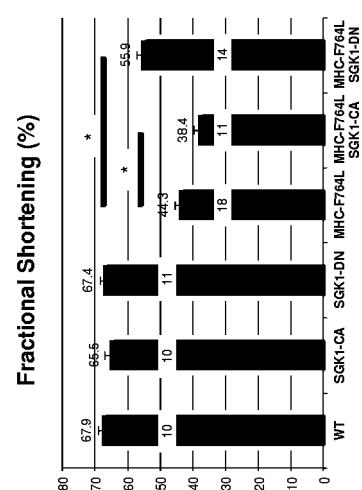


Figure 6



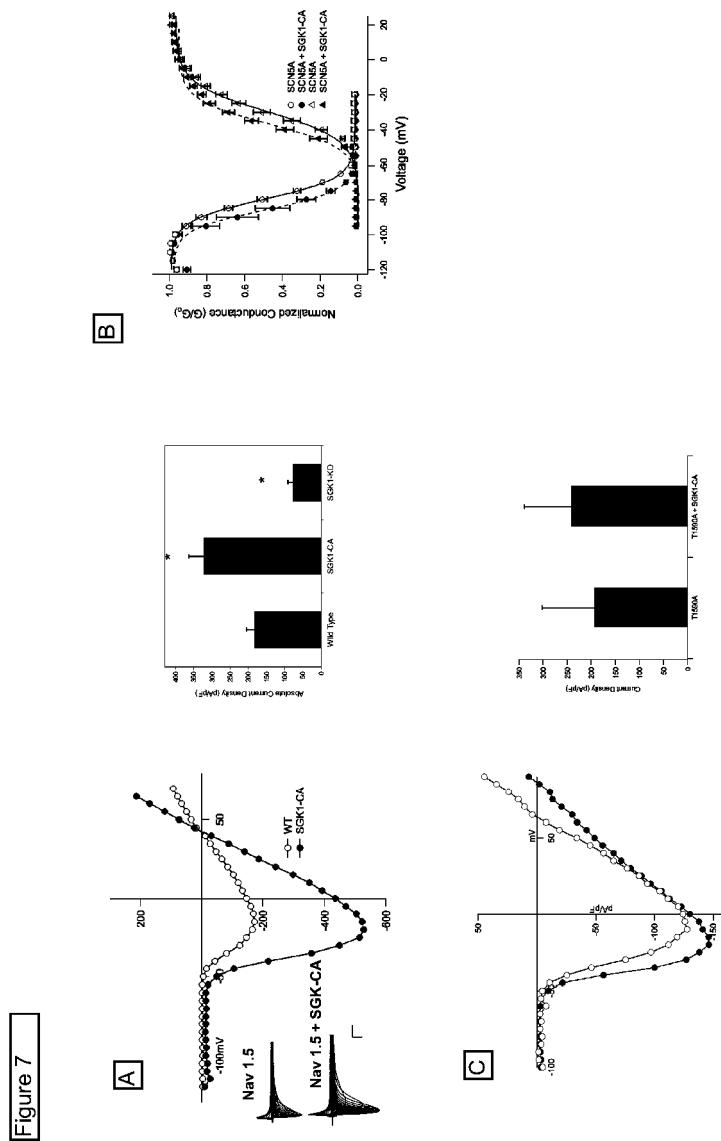
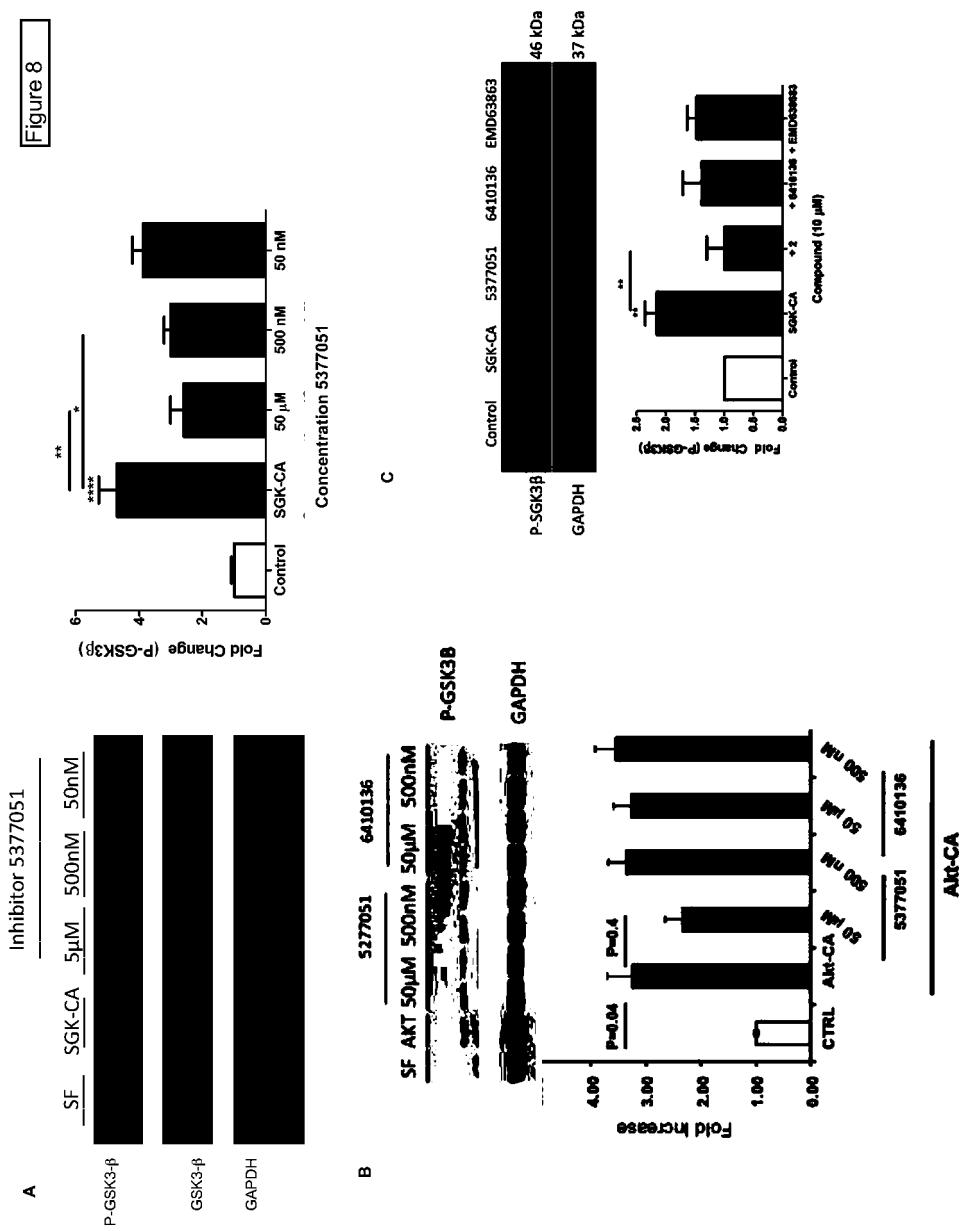
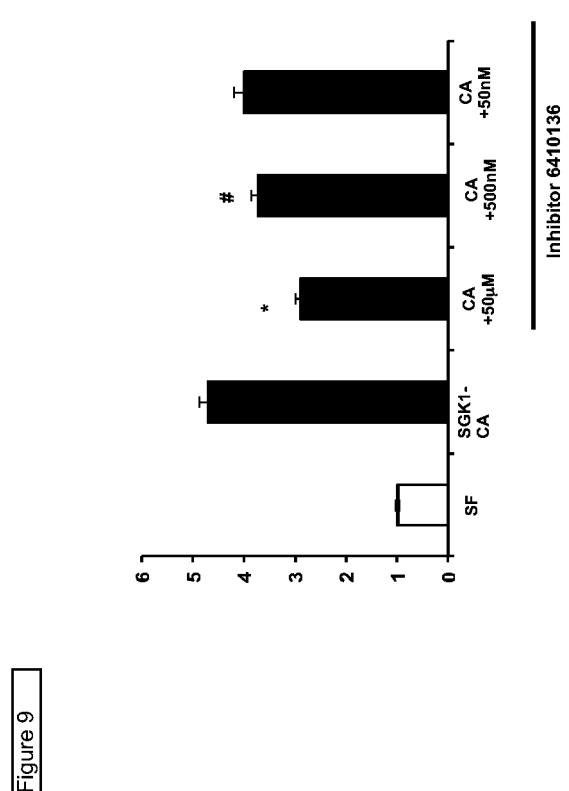


Figure 8





REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

- US 20080262096 A, Mederski [0024]
- US 20090221712 A, Gericke [0025]
- US 20080167380 A, Gericke [0026]
- US 20080234348 A, Mederski [0027] [0058]
- US 20090253767 A, Klein [0028]
- US 20100063115 A, Klein [0029]
- WO 2008021725 A, Drewry [0030] [0058]
- US 4172124 A, Koprowski [0036]
- US 5440021 A, Chuntharapai [0037]
- US 4816567 A, Cabilly [0038]
- EP 0125023 B1, Cabilly [0038]
- US 4816397 A, Boss [0038]
- EP 0120694 B1, Boss [0038]
- WO 8601533 A, Neuberger, M.S. [0038]
- EP 0194276 B1, Neuberger, M.S. [0038]
- US 5225539 A, Winter [0038]
- EP 0239400 B1, Winter [0038]
- EP 0451216 B1, Queen [0038]
- EP 0519596 A1, Padlan, E.A. [0038]
- US 4946778 A, Ladner [0038]
- US 5514548 A, Krebber [0039]
- WO 9306213 A, Hoogenboom [0039]
- US 5545806 A, Lonberg [0040]
- US 5545807 A, Surani [0040]
- WO 9713852 A, Lonberg [0040]
- US 20080249055 [0043] [0046] [0047]
- US 20080188430 [0046] [0047]
- WO 9412633 A [0050]
- US 6451991 B [0050]
- US 6900187 B [0050]
- US 4987071 A [0052]
- US 5496698 A [0052]
- US 5525468 A [0052]
- US 5631359 A [0052]
- US 5646020 A [0052]
- US 5672511 A [0052]
- US 6140491 A [0052]
- US 5652094 A [0053]
- WO 9103162 A [0053]
- WO 9207065 A [0053]
- WO 9315187 A [0053]
- EP 92110298 [0053]
- US 20050260163 [0054]
- WO 2005106491 A, Golz [0058]

Non-patent literature cited in the description

- **HERSHBERGER et al.** *J. Am. Coll. Cardiol.*, 19 April 2011, vol. 57 (16), 1641-1649 [0020]
- **HARLOW et al.** *Antibodies A Laboratory Manual*. Cold Spring Harbor Laboratory, 1988 [0034]
- **KOHLER et al.** *Nature*, 1975, vol. 256, 495-497 [0036]
- *Eur. J. Immunol.*, 1976, vol. 6, 511-519 [0036]
- **MILSTEIN et al.** *Nature*, 1977, vol. 266, 550-552 [0036]
- **HARLOW, E. ; D. LANE.** *Antibodies: A Laboratory Manual*. Cold Spring Harbor Laboratory, 1988 [0036]
- Current Protocols In Molecular Biology. John Wiley & Sons, 1991, vol. 2 [0036]
- **CHUNTHARAPAI et al.** *J. Immunol.*, 1994, vol. 152, 1783-1789 [0037]
- **NEWMAN, R. et al.** *BioTechnology*, 1992, vol. 10, 1455-1460 [0038]
- **BIRD, R.E. et al.** *Science*, 1988, vol. 242, 423-426 [0038]
- **KAMMAN, M. et al.** *Nucl. Acids Res.*, 1989, vol. 17, 5404 [0039]
- **SATO, K et al.** *Cancer Research*, 1993, vol. 53, 851-856 [0039]
- **DAUGHERTY, B.L. et al.** *Nucleic Acids Res.*, 1991, vol. 19 (9), 2471-2476 [0039]
- **LEWIS, A.P. ; J.S. CROWE.** *Gene*, 1991, vol. 101, 297-302 [0039]
- **JAKOBIVITS et al.** *Proc. Natl. Acad. Sci. USA*, 1993, vol. 90, 2551-2555 [0040]
- **JAKOBIVITS et al.** *Nature*, 1993, vol. 362, 255-258 [0040]
- **MATEEEVA O. et al.** *Nucleic Acids Res.*, 2007, vol. 35 (8 [0042]
- **HUESKEN D. et al.** *Nat. Biotechnol.*, vol. 23, 995-1001 [0042]
- **JAGLA B. et al.** *RNA*, 2005, vol. 11, 864-872 [0042]
- **SHABALINEA S.A.** *BMC Bioinformatics*, 2005, vol. 7, 65 [0042]
- **VERT J.P. et al.** *BMC Bioinformatics*, 2006, vol. 7, 520 [0042]
- **BRUMMELKAMP, T.R.** *Science*, 2002, vol. 296, 550-553 [0042]
- **LEE, NS et al.** *Nature Biotechnol.*, 2002, vol. 20, 500-505 [0042]
- **MIYAGISHI, M. ; TAIRA, K.** *Nature Biotechnol.*, 2002, vol. 20, 497-500 [0042]

- **PADDISON, P.J. et al.** *Genes & Dev.*, 2002, vol. 16, 948-958 **[0042]**
- **PAUL, C.P. et al.** *Nature Biotechnol.*, 2002, vol. 20, 505-508 **[0042]**
- **SUI, G. et al.** *Proc. Natl. Acad. Sci. USA*, 2002, vol. 99 (6), 5515-5520 **[0042]**
- **YU, J-Y et al.** *Proc. Natl. Acad. Sci. USA*, 2002, vol. 99 (9), 6047-6052 **[0042]**
- **ELBASHIR, SM et al.** *Nature*, 2001, vol. 411, 494-498 **[0042]**
- **DYKXHOORN ; LIEBERMAN.** *Annu. Rev. Biomed. Eng.*, 2006, vol. 8, 377-402 **[0043]** **[0046]** **[0047]**
- **DYKXHOORN et al.** *Gene Therapy*, 2006, vol. 13, 541-52 **[0046]** **[0047]**
- **AAGAARD ; ROSSI.** *Adv. Drug Delivery Rev.*, 2007, vol. 59, 75-86 **[0046]** **[0047]**
- **DE FOUGEROLLES et al.** *Nature Reviews Drug Discovery*, 2007, vol. 6, 443-53 **[0046]** **[0047]**
- **KRUEGER et al.** *Oligonucleotides*, 2007, vol. 17, 237-250 **[0046]** **[0047]**
- **DIAS N. ; STEIN C.A.** *Mol. Can. Ther.*, 2002, vol. 1, 347-355 **[0048]**
- **DING, Y. ; LAWRENCE, C. E.** *Nucleic Acids Res.*, 2001, vol. 29, 1034-1046 **[0048]**
- **SCZAKIEL, G.** *Front. Biosci.*, 2000, vol. 5, D194-D201 **[0048]**
- **SCHERR, M. et al.** *Nucleic Acids Res.*, 2000, vol. 28, 2455-2461 **[0048]**
- **PATZEL, V. et al.** *Nucleic Acids Res.*, 1999, vol. 27, 4328-4334 **[0048]**
- **CHIANG, M.Y. et al.** *J. Biol. Chem.*, 1991, vol. 266, 18162-18171 **[0048]**
- **STULL, R. A. et al.** *Nucleic Acids Res.*, 1992, vol. 20, 3501-3508 **[0048]**
- **DING, Y. ; LAWRENCE, C. E.** *Comput. Chem.*, 1999, vol. 23, 387-400 **[0048]**
- **LLOYD, B. H. et al.** *Nucleic Acids Res.*, 2001, vol. 29, 3664-3673 **[0048]**
- **MIR, K. U. ; SOUTHERN, E. M.** *Nat. Biotechnol.*, 1999, vol. 17, 788-792 **[0048]**
- **SOHAIL, M. et al.** *Nucleic Acids Res.*, 2001, vol. 29, 2041-2051 **[0048]**
- **ALTMAN, R. K. et al.** *J. Comb. Chem.*, 1999, vol. 1, 493-508 **[0048]**
- **MARTIN, P.** *Helv. Chim. Acta*, 1995, vol. 78, 486-504 **[0048]**
- **NIELSEN et al.** *Science*, 1991, vol. 254, 1497 **[0050]**
- **KIM et al.** *Proc Natl Acad Sci USA*, 1987, vol. 84, 8788 **[0051]**
- **HASELOFF ; GERLACH.** *Nature*, 1988, vol. 334, 585 **[0051]**
- **JEFFERIES et al.** *Nucleic Acid Res*, 1989, vol. 77, 1371 **[0051]**
- **ROSSI et al.** *AIDS Res Human Retroviruses*, 1992, vol. 8, 183 **[0052]**
- **HAMPEL ; TRITZ.** *Biochemistry*, 1989, vol. 28, 4929 **[0052]**
- **HAMPEL et al.** *Nucleic Acids Res*, 1990, vol. 18, 299 **[0052]**
- **PERROTTA ; BEEN.** *Biochemistry*, 1992, vol. 31, 16 **[0052]**
- **GUERRIER-TAKADA et al.** *Cell*, 1983, vol. 35, 849 **[0052]**
- **USMAN et al.** *JAm Chem Soc*, 1987, vol. 109, 7845-7854 **[0053]**
- **SCARINGE et al.** *Nucleic Acids Res*, 1990, vol. 18, 5433-5441 **[0053]**
- **PERRAULT et al.** *Nature*, 1990, vol. 344, 565 **[0053]**
- **PIEKEN et al.** *Science*, 1991, vol. 253, 314 **[0053]**
- **USMAN ; CEDERGREEN.** *Trends Biochem Sci*, 1992, vol. 17, 334 **[0053]**
- **AGUINO-JARGUIN et al.** *Oligonucleotides*, 2008, vol. 18 (3), 213-24 **[0054]**
- **PROCHIANTZ.** *Current Opinion in Neurobiology*, 1996, vol. 6, 629-634 **[0055]**
- **LIN et al.** *J. Biol. Chem.*, 1995, vol. 270, 14255-14258 **[0055]**
- Remington's Pharmaceutical Sciences. Mack Publishing Company **[0057]**