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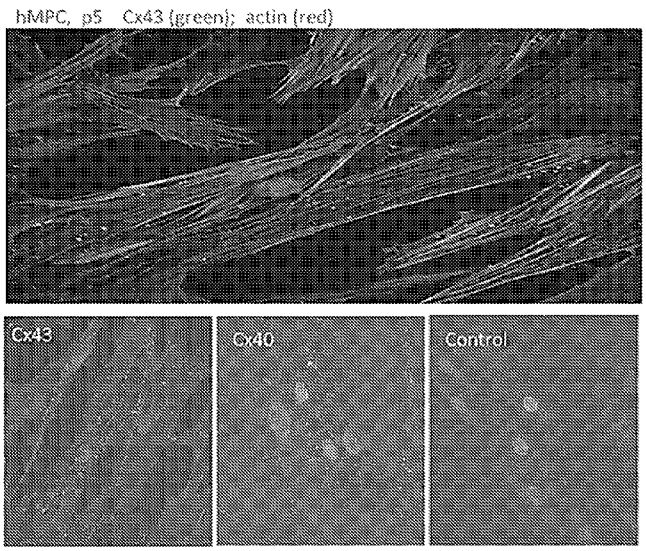


FIGURE 1A

(57) Abstract: The present disclosure relates to cellular compositions that are modified to introduce a nucleic acid or vector expressing the same. Such compositions may be used to deliver nucleic acid to a target cell and treat disease such as cancer.

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CELLULAR COMPOSITIONS AND METHODS OF TREATMENT I

FIELD OF THE INVENTION

The present disclosure relates to cellular compositions that are modified to introduce a nucleic acid or vector expressing the same. Such compositions may be used to deliver nucleic acid to a target cell and treat diseases such as cancer.

BACKGROUND OF THE INVENTION

Nucleic acids, either single or double stranded, or both, have been shown to pass through gap junctions formed by connexin proteins in HeLa cell pairs modified to express connexin proteins. Transfer was demonstrated by a single electrode delivery of fluorescent-tagged oligonucleotides to a donor cell and determining their transfer to the target cell via gap junction mediated communication.

Improved methods of delivering nucleic acids to target cells are required.

SUMMARY OF THE INVENTION

The present inventors have identified that mesenchymal lineage precursor or stem cells are able to direct a high rate of nucleic acid transfer to target cells. The present inventors have also identified that mesenchymal lineage precursor or stem cells can deliver nucleic acids to cancer cells to reduce cancer cell growth. Surprisingly, these capabilities of mesenchymal lineage precursor or stem cells are maintained despite knockout of a key protein involved in transfer of nucleic acids between cells.

Another advantage of using mesenchymal lineage precursor or stem cells for delivery of nucleic acids to target cells is the ability of the cells home to target tissues. The migration and adhesion capacity of mesenchymal lineage precursor or stem cells makes them particularly suitable for this purpose.

Accordingly, in an example, the present disclosure relates to a method of delivering an oligonucleotide into a target cell, the method comprising contacting a target cell with a mesenchymal lineage precursor or stem cell expressing one or more of the markers selected from the group consisting of α1, α2, α3, α4 and α5, αv, β1 and β3, wherein said cell has been modified to introduce an oligonucleotide or a vector expressing the oligonucleotide. In an example, the contacting occurs under conditions permitting the mesenchymal lineage precursor or stem cell to form a gap junction with the target cell, whereby the oligonucleotide is delivered to the target cell by traversing the gap junction. In an example, the gap junction is formed by Cx40 or Cx43. In another example, the gap junction is formed by Cx43. In another example, delivery of

the oligonucleotide is via a mechanism other than Cx43. For example, oligonucleotide may be delivered by traversing a gap junction formed by Cx40, Cx45, Cx30.3, Cx31 or Cx31.1. In another example, oligonucleotide may be delivered via formation of exosomes. In another example, delivery of the oligonucleotide is independent of cell to cell contact. In this example, delivery of the oligonucleotide does not occur via a connexin.

In an example, the target cell is a cancer cell or white blood cell. For example, the cancer cell can be a lung cancer, pancreatic cancer, colorectal cancer, liver cancer, cervical cancer, prostate cancer, osteosarcoma or melanoma cell. In another example, the target cell is a syncytial cell. For example, the syncytial cell can be a cardiac myocyte, smooth muscle cell, epithelial cell, connective tissue cell or syncytial cancer cell.

In another example, the present disclosure relates to a method of treating cancer in a subject, the method comprising administering to the subject a composition comprising mesenchymal lineage precursor or stem cells expressing one or more of the markers selected from the group consisting of α1, α2, α3, α4 and α5, αv, β1 and β3, wherein said cells are modified to introduce an oligonucleotide or a plasmid expressing the oligonucleotide.

In an example, the mesenchymal lineage precursor or stem cells express a connexin that is also expressed by cells comprising the subject's cancer. In an example, the connexin is Cx40 or Cx43. In an example, cells comprising the subject's cancer express Cx43.

In another example, the mesenchymal lineage precursor or stem cells express CD46.

In an example, the cancer is selected from the group consisting of lung cancer, pancreatic cancer, colorectal cancer, liver cancer, cervical cancer, prostate cancer, osteosarcoma and melanoma. In an example, the method comprises administering a composition according to the present disclosure.

In an example, the oligonucleotide is about 12 – 24 nucleotides in length. In another example, the oligonucleotide is RNA. In another example, the oligonucleotide is an antisense molecule. In another example, the oligonucleotide is a shRNA. In another example, the oligonucleotide is a siRNA. In an example, the siRNA is KIF11 siRNA or PLK1 siRNA. In another example, the siRNA is KIF11 siRNA.

In an example, the KIF11 siRNA binds towards the 5' end of KIF11 mRNA as shown in SEQ ID NO: 1 (NCBI reference number NM_004523.3). In an example, the KIF11 siRNA binds the KIF11 mRNA transcript between bp 800 and bp 3,600 from

the 5' end. In another example, the KIF11 siRNA binds the KIF11 mRNA transcript between bp 900 and bp 3,200 from the 5' end. In an example, the oligonucleotide comprises the nucleic acid sequence as shown in any one of SEQ ID NOs: 2, 3, 4, 5, 6 or 7. In an example, the oligonucleotide consists of the nucleic acid sequence as shown in any one of SEQ ID NOs: 2, 3, 4, 5, 6 or 7. In an example, the oligonucleotide comprises the nucleic acid sequence as shown in SEQ ID NO: 3. In another example, the oligonucleotide consists of the nucleic acid sequence as shown in SEQ ID NO: 3. In an example, the oligonucleotide comprises the nucleic acid sequence as shown in SEQ ID NO: 4. In another example, the oligonucleotide consists of the nucleic acid sequence as shown in SEQ ID NO: 4. In an example, the oligonucleotide comprises the nucleic acid sequence as shown in SEQ ID NO: 7. In another example, the oligonucleotide consists of the nucleic acid sequence as shown in SEQ ID NO: 7. In an example, the oligonucleotide comprises the nucleic acid sequence as shown in any one of SEQ ID NOs: 3, 4 or 7. In another example, the oligonucleotide consists of the nucleic acid sequence as shown in any one of SEQ ID NOs: 3, 4 or 7. In these examples, the target cell may be a bone cancer cell. In an example, the cancer may be sarcoma. In another example, the target cell may be a pancreatic cancer cell. In an example, the cancer may be pancreatic cancer. In another example, the target cell may be a prostate cancer cell. In an example, the cancer may be prostate cancer.

In another example, the oligonucleotide comprises the nucleic acid sequence as shown in any one of SEQ ID NOs: 2, 3, 4, 5, 6, 7, 8 or 9. In an example, the oligonucleotide consists of the nucleic acid sequence as shown in any one of SEQ ID NOs: 2, 3, 4, 5, 6, 7, 8 or 9. In an example, the oligonucleotide consists of the nucleic acid sequence as shown in SEQ ID NOs: 8 or 9.

In another example, the oligonucleotide is 18-22 nucleotides in length. In another example, the oligonucleotide is a miRNA. In an example, miRNAs can be selected from the group consisting of miR-155, miR-155-inh, miR-181-B1, miR-15a, miR-16-1, miR-21, miR-34a, miR-221, miR-29a, let-7b.

In another example, the mesenchymal lineage precursor or stem cells are modified to introduce an oligonucleotide or vector expressing the same that kills the target cell but does not substantially affect viability of the mesenchymal lineage precursor or stem cell. In another example, the mesenchymal lineage precursor or stem cells are modified to introduce an inhibitory oligonucleotide or vector expressing the same that does not kill the mesenchymal lineage precursor or stem cells before they can deliver the inhibitory oligonucleotide to a target cell such as a cancer cell.

The present inventors have also identified that mesenchymal lineage precursor or stem cells can be modified to introduce various inhibitory oligonucleotides known to kill cancer cells and/or reduce cancer cell growth. Surprisingly, these modified mesenchymal lineage precursor or stem cells remain viable and are able to deliver the inhibitory oligonucleotide to cancer cells in levels sufficient to kill cancer cells and/or reduce cell growth.

Accordingly, in another example, the present disclosure relates to a composition comprising mesenchymal lineage precursor or stem cells expressing one or more of the markers selected from the group consisting of α1, α2, α3, α4 and α5, αv, β1 and β3, wherein said cells are modified to introduce an inhibitory oligonucleotide or comprise a vector expressing the inhibitory oligonucleotide.

In an example, the inhibitory oligonucleotide is about 12 – 24 nucleotides in length. In another example, the inhibitory oligonucleotide is RNA. In another example, the inhibitory oligonucleotide is an antisense molecule. In another example, the inhibitory oligonucleotide is a shRNA. In another example, the inhibitory oligonucleotide is a siRNA. In another example, the inhibitory oligonucleotide is 18-22 nucleotides in length. In another example, the inhibitory oligonucleotide is a miRNA. In an example, miRNAs can be selected from the group consisting of miR-155, miR-155-inh, miR-181-B1, miR-15a, miR-16-1, miR-21, miR-34a, miR-221, miR-29a, let-7b.

In an example, the mesenchymal lineage precursor or stem cells also express STRO-1. Accordingly, in an example, the present disclosure relates to a method of delivering an oligonucleotide into a target cell, the method comprising contacting a target cell with a mesenchymal lineage precursor or stem cell expressing STRO-1 and one or more of the markers selected from the group consisting of α1, α2, α3, α4 and α5, αv, β1 and β3, wherein said cell has been modified to introduce an oligonucleotide or a vector expressing the oligonucleotide. In another example, the present disclosure relates to a method of treating cancer in a subject, the method comprising administering to the subject a composition comprising mesenchymal lineage precursor or stem cells expressing STRO-1 and one or more of the markers selected from the group consisting of α1, α2, α3, α4 and α5, αv, β1 and β3, wherein said cells are modified to introduce an oligonucleotide or a plasmid expressing the oligonucleotide. In another example, the present disclosure relates to a composition comprising mesenchymal lineage precursor or stem cells expressing STRO-1 and one or more of the markers selected from the group consisting of α1, α2, α3, α4 and α5, αv, β1 and β3, wherein said cells are modified to introduce an inhibitory oligonucleotide or comprise a vector expressing the

inhibitory oligonucleotide. In another example, the mesenchymal lineage precursor or stem cells are substantially STRO-1^{bri}. In another example, the mesenchymal lineage precursor or stem cell is not modified to express Cx40 or Cx43.

In an example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of oligonucleotide to reduce viability of a target cell such as, for example, a cancer cell. In an example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of between 0.25 to 5 nM of the inhibitory oligonucleotide to a target cell. In another example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of between 0.5 to 5 nM of the inhibitory oligonucleotide to a target cell. In another example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of between 0.5 to 4 nM of the inhibitory oligonucleotide to a target cell. In another example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of between 0.5 to 3.5 nM of the inhibitory oligonucleotide to a target cell.

In another example, the mesenchymal lineage precursor or stem cells are derived from pluripotent cells. For example, the mesenchymal lineage precursor or stem cells may be derived from induced pluripotent stem (iPS) cells.

In an example, mesenchymal lineage precursor or stem cells are modified to introduce the oligonucleotide using a viral vector. In an example, the viral vector is selected from the group consisting of Lentivirus, Baculovirus, Retrovirus, Adenovirus (AdV), Adeno-associated virus (AAV) or a recombinant form thereof. In another example, the viral vector is a AAV. In another example, the viral vector is a recombinant AAV.

In an example, modified mesenchymal lineage precursor or stem cells have been treated to effect modification of cell surface glycans on the mesenchymal lineage precursor or stem cells. In an example, the treatment involves exposure of the mesenchymal lineage precursor or stem cell to a glycosylstrasferase under conditions which result in modification of cell-surface glycans on the mesenchymal precursor lineage cell or stem cell.

In an example, mesenchymal lineage precursor or stem cell are exposed to an exogenous glycosyltranferase.

In another example, mesenchymal lineage precursor or stem cells have been modified to express a glycosyltranferase. In an example, the mesenchymal lineage precursor or stem cell have been modified to introduce a nucleic acid encoding a

glycosyltransferase, wherein expression of the glycosyltransferase in the cell results in enhanced retention of the cell at a site of inflammation *in vivo*.

In an example, the glycosyltransferase is a fucosyltransferase, a galactosyltransferase, or a sialyltransferase. In an example, the fucosyltransferase is an alpha 1,3 fucosyltransferase such as an alpha 1,3 fucosyltransferase III, alpha 1,3 fucosyltransferase IV, an alpha 1,3 fucosyltransferase VI, an alpha 1,3 fucosyltransferase VII or an alpha 1,3 fucosyltransferase IX.

Any example herein shall be taken to apply *mutatis mutandis* to any other example unless specifically stated otherwise.

The present disclosure is not to be limited in scope by the specific examples described herein, which are intended for the purpose of exemplification only. Functionally-equivalent products, compositions and methods are clearly within the scope of the disclosure, as described herein.

Throughout this specification, unless specifically stated otherwise or the context requires otherwise, reference to a single step, composition of matter, group of steps or group of compositions of matter shall be taken to encompass one and a plurality (i.e. one or more) of those steps, compositions of matter, groups of steps or group of compositions of matter.

The disclosure is hereinafter described by way of the following non-limiting Examples and with reference to the accompanying drawings.

BRIEF DESCRIPTION OF THE ACCOMPANYING DRAWINGS

Figure 1. MPCs (A) and MSCs (B) Express Gap Junction Proteins: Connexin Cx43 & Cx40.

Figure 2. Patch clamp technique – MPC expression of functional gap junctions.

Figure 3. Cx43 Expression in Tumour Lines in Screening Panel.

Figure 4. MPCs transfected with a Cyanine 3 Fluroescent Labeled siRNA using Lipofectamine: 1-1000nM extracellular concentration.

Figure 5. (i). MPCs Transfer of Cy-5-MiR16 via gap junctions to **(A)** GFP expressing HeLa-Cx43 cells and **(B)** GFP expressing PANC-1 cells. **(ii).** MPCs Transfer of KIF11 siRNA to GFP expressing PANC-1 cells.

Figure 6. (A) Co-Culture of KIF11 siRNA (100 nM) loaded MPCs & PANC-1 Tumour Cells. **(B)** Co-Culture of KIF11 siRNA (500 nM) loaded MPCs & PANC-1 Tumour Cells.

Figure 7. Co-Culture of KIF11 siRNA (500 nM) loaded MPCs & SNUC2A Tumour Cells.

Figure 8. (A) and (B) Co-Culture of KIF11 siRNA (500 nM) loaded MPCs & SAOS2 Tumour Cells.

Figure 9. (A) Co-Culture of Mir-16 (100 nM) loaded MPCs & PANC-1 Tumour Cells. **(B)** Co-Culture Mir-16 (100 nM) loaded MPCs & untransfected MPCs.

Figure 10. (A) Co-culture let7b (500 nM) loaded MPCs and PANC-1 (CRISP/Cas9 Cx43 KO) or PANC-1. Growth of PANC-1 cells transfected with let7b and untransfected PANC-1 cells also shown. **(B)** Growth of MPC transfected with let7b and untransfected MPC.

Figure 11. (A) Co-Culture of KIF11 siRNA (500 nM) loaded MPCs and PANC-1 (CRISP/Cas9 Cx43 KO) or PANC-1. Growth of PANC-1 (CRISP/Cas9 Cx43 KO) cells with untransfected MPC also shown. **(B)** Growth of KIF11 siRNA (500 nM) loaded MPCs and untransfected MPC.

Figure 12. Data combined: Co-culture KIF11 siRNA (500 nM) loaded MPCs and PANC-1 (CRISP/Cas9 Cx43 KO) or PANC-1. Growth of untransfected MPC co-cultured with PANC-1 (CRISP/Cas9 Cx43 KO) or PANC-1, and growth of PANC-1 (CRISP/Cas9 Cx43 KO) transfected with KIF11 also shown.

Figure 13. (A) Co-Culture of Mir-34a (500 nM) loaded MPCs and PANC-1 (CRISP/Cas9 Cx43 KO) or PANC-1. Growth of PANC-1 cells with untransfected MPC, and PANC-1 transfected cells also shown. **(B)** Growth of Mir-34a (500 nM) loaded MPCs and untransfected MPC.

Figure 14. (A) Death of MPCs transfected with mir-mimics, PLK1 siRNA or KIF11 siRNA, day 5; Fluorometric assay of cell number of MPCs transfected with non-targeting siRNA, death siRNA, PLK1 siRNA and KIF11 siRNA **(B)** WST assay of MPCs transfected with mir-mimics, PLK1 siRNA or KIF11 siRNA, day 5.

Figure 15. (A) Death of PANC-1 cells transfected with mir-mimics, PLK1 siRNA or KIF11 siRNA, day 5 **(B)** WST assay PANC-1 cells transfected with mir-mimics, PLK1 siRNA or KIF11 siRNA, day 5.

Figure 16. (A) Death of PC3 cells transfected with mir-mimics, PLK1 siRNA or KIF11 siRNA, day 5 (II) **(B)** WST assay of PC3 cells transfected with mir-mimics, PLK1 siRNA or KIF11 siRNA, day 5.

Figure 17. (A) Death of SAOS2 cells transfected with mir-mimics, PLK1 siRNA or KIF11 siRNA, day 5 (II) **(B)** WST assay of SAOS2 cells transfected with mir-mimics, PLK1 siRNA or KIF11 siRNA, day 5.

Figure 18. Comparative analysis of cell death and WST Assays (MPCs; PANC-1).

Figure 19. Comparative analysis of cell death and WST Assays (PC3; SAOS2).

Figure 20. PC3 prostate xenograft tumour model. Mice were orthotopically implanted in the prostate with 0.5 million PC3 cells expressing luciferase, with (treated) or without (control) 0.5 million MPC loaded with 500nM siRNA against KIF11. One week later, mice were injected with luciferin and imaged to visualise tumours. Panel **(A)** shows control mice and Panel **(B)** shows MPC treated mice. Panel **(C)** shows the survival curves for treated and untreated mice.

Figure 21. Measurement of Total (Panel A) and Live (Panel B) PANC-1 cells after Co-Culture with MPCs transfected with 500 nM KIF11 (1/1 ratio of MPCs/PANC-1 cells). X- Axis for Panel A and B:

- PANC-1, day 1: Number of PANC-1 cells after 1 day in culture (PANC-1 cells cultured alone)
    - PANC-1- KIF11 siRNA, day 1: Number of PANC-1 transfected with 500 nM KIF11 measured after 1 day in culture
- MPC- PANC-1, day 1: Number of PANC-1 cells after 1 day in co-culture with MPCs (1/1 ratio)
- MPC_KIF11- PANC-1: Number of PANC-1 cells after 1 day in co-culture with MPCs transfected with KIF11 (500 nM; 1/1 Ratio)
- PANC-1, day 6: Number of PANC-1 cells after 6 days in culture (PANC-1 cells cultured alone)
- PANC-1-KIF11 siRNA, day 6: Number of PANC-1 transfected with 500 nM KIF11 measured after 6 days in culture.
- MPC- PANC-1, day 1: Number of PANC-1 cells after 6 days in co-culture with MPCs (1/1 ratio)
- MPC_KIF11- PANC-1: Number of PANC-1 cells after 6 days in co-culture with MPCs transfected with KIF11 (500 nM; 1/1 Ratio).

Figure 22. Effect of KIF11 siRNA from MPCs to PANC-1 cells in co-culture. Panel A shows time course of PANC-1 cell growth in co-culture with MPCs loaded with KIF11-siRNA (0 – 500 nM). Panel B shows dose-response curve of PANC-1 cell growth in co-culture with MPCs loaded with KIF11-siRNA (0 – 500 nM) on day 5 of cell culture. Panel C shows time course of MPCs growth after transfection with KIF11 siRNA (0 – 500 nM) in co-culture with PANC-1. Results represent average of three independent measurements.

Figure 23. Dose dependent inhibition of PANC-1 cell viability by KIF11 (direct transfection).

Figure 24. Effect of KIF11 siRNA on PANC-1 cell growth. Panel A shows time course of PANC1 cell growth after transfection with KIF11 siRNA (0 – 500 nM). Panel B shows dose-response curve of PANC-1 cell growth after transfection with KIF11 siRNA (0 – 500 nM) on day 6 of cell culture.

Figure 25. Effect of KIF11 siRNA on MPC cell growth. Panel A shows time course of MPCs growth after transfection with KIF11 siRNA (0 – 500 nM). Panel B shows dose-response curve of MPCs cell growth after transfection with KIF11 siRNA (0 – 500 nM) on day 6 of cell culture.

Figure 26. Effect of KIF11 siRNA from MPCs to PC3 cells in co-culture. Panel A shows time course of PC3 cell growth in co-culture with MPCs loaded with KIF11 siRNA (0 – 500 nM). Panel B shows dose-response curve of PC3 cell growth in co-culture with MPCs loaded with KIF11-siRNA (0 – 500 nM) on day 5 of cell culture. Panel C shows time course of MPC growth after transfection with KIF11 siRNA (0 – 500 nM) in co-culture with PC3 cells. Results represent average of three independent measurements

Figure 27. Effect of KIF11 siRNA on PC3 cell growth. Panel A shows time course of PC3 cell growth after transfection with KIF11 siRNA (0 – 500 nM). Panel B shows dose-response curve of PC3 cell growth after transfection with KIF11 siRNA (0 – 500 nM) on day 6 of cell culture.

Figure 28. MPC donor cell KIF11 siRNA loading.

Figure 29. MPC donor cell KIF11 siRNA transfer to tumour cells.

Figure 30. KIF11 mRNA levels in KIF11 siRNA transfected cells.

Figure 31. KIF11 mRNA levels in PANC-1 cells following co-culture with KIF11 siRNA loaded MPCs. MPC = MPC directly transfected with KIF11 siRNA; MPC-Co = MPC directly transfected with KIF11 siRNA & co-cultured with PANC1; PANC1-Co = PANC1 cells co-cultured for 24 hours with MPCs transfected with KIF11 siRNA; KIF11 expression was assayed 48 hours after transfection. The Comparative Ct Method (ΔΔCt Method) for analyzing qRT-PCR data - 1. Sample KIF11 Average Ct - GAPDH Average Ct = ΔCt KIF11-GAPDH; 2. ΔΔCT=ΔCt treated-ΔCt untreated; 3. Fold difference in KIF11 relative to untreated = 2-ΔΔCt.

Figure 32. Alignment of siRNAs to human KIF11 mRNA sequence.

Figure 33. Effects of KIF11 siRNAs on MPC and tumour cell viability.

Figure 34. Gene expression analysis in MPCs and MSCs transfected with KIF11 siRNA compared with control.

KEY TO THE SEQUENCE LISTING

SEQ ID NO: 1: KIF11 mRNA (NCBI reference number NM_004523.3)
SEQ ID NO: 2: Hs_KIF11_8 siRNA
SEQ ID NO: 3: Hs_KIF11_9 siRNA
SEQ ID NO: 4: Hs_KIF11_6 siRNA
SEQ ID NO: 5: Hs_KIF11_12 siRNA
SEQ ID NO: 6: Hs_KIF11_7 siRNA
SEQ ID NO: 7: Hs_KIF11_4 siRNA

DETAILED DESCRIPTION OF THE INVENTION

General Techniques and Selected Definitions

Unless specifically defined otherwise, all technical and scientific terms used herein shall be taken to have the same meaning as commonly understood by one of ordinary skill in the art (e.g., molecular biology, cell culture, stem cell differentiation, cell therapy, genetic modification, biochemistry, physiology, and clinical studies).

Unless otherwise indicated, the molecular and statistical techniques utilized in the present disclosure are standard procedures, well known to those skilled in the art. Such techniques are described and explained throughout the literature in sources such as, J. Perbal, A Practical Guide to Molecular Cloning, John Wiley and Sons (1984), J. Sambrook et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbour Laboratory Press (1989), T.A. Brown (editor), Essential Molecular Biology: A Practical Approach, Volumes 1 and 2, IRL Press (1991), D.M. Glover and B.D. Hames (editors), DNA Cloning: A Practical Approach, Volumes 1-4, IRL Press (1995 and 1996), and F.M. Ausubel et al. (editors), Current Protocols in Molecular Biology, Greene Pub. Associates and Wiley-Interscience (1988, including all updates until present), Ed Harlow and David Lane (editors) Antibodies: A Laboratory Manual, Cold Spring Harbour Laboratory, (1988), and J.E. Coligan et al. (editors) Current Protocols in Immunology, John Wiley & Sons (including all updates until present).

As used in this specification and the appended claims, terms in the singular and the singular forms "a," "an" and "the," for example, optionally include plural referents unless the content clearly dictates otherwise. Thus, for example, reference to "an analyte" optionally includes one or more analytes.

As used herein, the term “about”, unless stated to the contrary, refers to +/- 10%, more preferably +/- 5%, more preferably +/- 1%, of the designated value.

The term “and/or”, e.g., “X and/or Y” shall be understood to mean either “X and Y” or “X or Y” and shall be taken to provide explicit support for both meanings or for either meaning.

Throughout this specification the word "comprise", or variations such as "comprises" or "comprising", will be understood to imply the inclusion of a stated element, integer or step, or group of elements, integers or steps, but not the exclusion of any other element, integer or step, or group of elements, integers or steps.

The term "connexin" as used herein means a large family of trans-membrane proteins that allow intercellular communication and the transfer of ions and small signalling molecules and assemble to form gap junctions. Connexins are four-pass transmembrane proteins with both C and N cytoplasmic termini, a cytoplasmic loop (CL) and two extra- cellular loops, (EL- I) and (EL-2). Connexins are assembled in groups of six to form hemichannels, or connexons, and two hemichannels , one on each cell, then combine to form a gap junction between the two cells. The term Connexin is abbreviated as Cx and the gene encoding for it *Cx*.

The term "gap junction" as used herein means a specialized intercellular connection between cell-types. A gap junction directly connects the cytoplasm of two cells, which allows various molecules such as nucleic acids, ions and electrical impulses to directly pass through a regulated gate between cells.

Various subjects can be administered cell compositions according to the present disclosure. In an example, the subject is a mammal. The mammal may be a companion animal such as a dog or cat, or a livestock animal such as a horse or cow. In another example, the subject is a human. Terms such as “subject”, “patient” or “individual” are terms that can, in context, be used interchangeably in the present disclosure.

As used herein, the term “treatment” refers to clinical intervention designed to alter the natural course of the individual or cell being treated during the course of clinical pathology. Desirable effects of treatment include decreasing the rate of disease progression, ameliorating or palliating the disease state, and remission or improved prognosis. An individual is successfully “treated”, for example, if one or more symptoms associated with a disease are mitigated or eliminated.

An “effective amount” refers to at least an amount effective, at dosages and for periods of time necessary, to achieve the desired therapeutic or prophylactic result. An effective amount can be provided in one or more administrations. In some examples of the present disclosure, the term “effective amount” is used to refer to an amount necessary to effect treatment of a disease or condition as hereinbefore described. The effective amount may vary according to the disease or condition to be treated and also according to the weight, age, racial background, sex, health and/or physical condition and other factors relevant to the mammal being treated. Typically,

the effective amount will fall within a relatively broad range (e.g. a “dosage” range) that can be determined through routine trial and experimentation by a medical practitioner. The effective amount can be administered in a single dose or in a dose repeated once or several times over a treatment period.

A “therapeutically effective amount” is at least the minimum concentration required to effect a measurable improvement of a particular disorder (e.g. cancer). A therapeutically effective amount herein may vary according to factors such as the disease state, age, sex, and weight of the patient, and the ability of the cellular composition to elicit a desired response in the individual. A therapeutically effective amount is also one in which any toxic or detrimental effects of the composition are outweighed by the therapeutically beneficial effects. In the case of cancer, a therapeutically effective amount can reduce the number of cancer cells; reduce the primary tumour size; inhibit (i.e., slow to some extent and, in some examples, stop) cancer cell infiltration into peripheral organs; inhibit (i.e., slow to some extent and, in some examples, stop) tumour metastasis; inhibit or delay, to some extent, tumour growth or tumour progression; and/or relieve to some extent one or more of the symptoms associated with the disorder. To the extent a composition according to the present disclosure may prevent growth and/or kill existing cancer cells, it may be cytostatic and/or cytotoxic. For cancer therapy, efficacy *in vivo* can, for example, be measured by assessing the duration of survival, time to disease progression (TTP), the response rates (RR), duration of response, and/or quality of life.

Mesenchymal lineage precursor (MPC) or stem cells

As used herein, the term "mesenchymal lineage precursor or stem cells" refers to undifferentiated multipotent cells that have the capacity to self renew while maintaining multipotentcy and the capacity to differentiate into a number of cell types either of mesenchymal origin, for example, osteoblasts, chondrocytes, adipocytes, stromal cells, fibroblasts and tendons, or non-mesodermal origin, for example, hepatocytes, neural cells and epithelial cells. .

The term "mesenchymal lineage precursor or stem cells" includes both parent cells and their undifferentiated progeny. The term also includes mesenchymal lineage precursor or stem cells (MPC), multipotent stromal cells, mesenchymal stem cells, perivascular mesenchymal lineage precursor or stem cells, and their undifferentiated progeny.

Mesenchymal lineage precursor or stem cells can be autologous, allogeneic, xenogeneic, syngeneic or isogeneic. Autologous cells are isolated from the same

individual to which they will be reimplanted. Allogeneic cells are isolated from a donor of the same species. Xenogeneic cells are isolated from a donor of another species. Syngeneic or isogeneic cells are isolated from genetically identical organisms, such as twins, clones, or highly inbred research animal models.

Mesenchymal lineage precursor or stem cells reside primarily in the bone marrow, but have also been shown to be present in diverse host tissues including, for example, cord blood and umbilical cord, adult peripheral blood, adipose tissue, trabecular bone and dental pulp.

In an example, mesenchymal lineage precursor or stem cells express STRO-1 and one or more integrins. Integrins are a class of cell adhesion receptors that mediate both cell-cell and cell-extracellular matrix adhesion events. Integrins consist of heterodimeric polypeptides where a single α chain polypeptide noncovalently associates with a single β chain. There are now about 16 distinct α chain polypeptides and at least about 8 different β chain polypeptides that constitute the integrin family of cell adhesion receptors. In general, different binding specificities and tissue distributions are derived from unique combinations of the α and β chain polypeptides or integrin subunits. The family to which a particular integrin is associated with is usually characterized by the β subunit. However, the ligand binding activity of the integrin is largely influenced by the α subunit.

In an example, mesenchymal lineage precursor or stem cells according to the present disclosure express STRO-1 and an integrin having a β1 (CD29) chain polypeptide.

In another example, mesenchymal lineage precursor or stem cells according to the present disclosure express STRO-1 and an integrin having an α chain polypeptide selected from the group consisting of α1 (CD49a), α2 (CD49b), α3 (CD49c), α4 (CD49d), α5 (CD49e) and αv (CD51). Accordingly, in an example, mesenchymal lineage precursor or stem cells according to the present disclosure express STRO-1 and α1. In another example, mesenchymal lineage precursor or stem cells express STRO-1 and α2. In another example, mesenchymal lineage precursor or stem cells express STRO-1 and α3. In another example, mesenchymal lineage precursor or stem cells express STRO-1 and α4. In another example, mesenchymal lineage precursor or stem cells express STRO-1 and α5. In another example, mesenchymal lineage precursor or stem cells express STRO-1 and αv. In another example, mesenchymal lineage precursor or stem cells express STRO-1, α2 and α3. In another example, mesenchymal lineage precursor or stem cells express STRO-1, α2 and α5. In another example,

mesenchymal lineage precursor or stem cells express STRO-1, α3 and α5. In another example, mesenchymal lineage precursor or stem cells express STRO-1, α2, α3 and α5.

In another example, the present disclosure encompasses a population of mesenchymal lineage precursor or stem cells enriched for STRO-1+ and α1+ cells. In this example, a population enriched for α1+ cells can comprise at least about 3% or 4% or 5% α1+ cells.

In another example, the present disclosure encompasses a population of mesenchymal lineage precursor or stem cells enriched for STRO-1+ and α2+ cells. In this example, a population enriched for α2+ cells can comprise at least about 30% or 40% or 50% α2+ cells.

In another example, the present disclosure encompasses a population of mesenchymal lineage precursor or stem cells enriched for STRO-1+ and α3+ cells. In this example, a population enriched for α3+ cells comprises at least about 40% or 45% or 50% α3+ cells.

In another example, the present disclosure encompasses a population of mesenchymal lineage precursor or stem cells enriched for STRO-1+ and α4+ cells. In this example, a population enriched for α4+ cells comprises at least about 5% or 6% or 7% α4+ cells.

In another example, the present disclosure encompasses a population of mesenchymal lineage precursor or stem cells enriched for STRO-1+ and α5+ cells. In this example, a population enriched for α5+ cells comprises at least about 45% or 50% or 55% α5+ cells.

In another example, the present disclosure encompasses a population of mesenchymal lineage precursor or stem cells enriched for STRO-1+ and αv+ cells. In this example, a population enriched for αv+ cells comprises at least about 5% or 6% or 7% αv+ cells.

In another example, the present disclosure encompasses a population of mesenchymal lineage precursor or stem cells enriched for STRO-1, α1+, α3+, α4+ and α5+ cells.

In another example, mesenchymal lineage precursor or stem cells according to the present disclosure can express integrins selected from the group consisting of α1β1, α2β1, α3β1, α4β1 and α5β1. Accordingly, in an example, mesenchymal lineage precursor or stem cells according to the present disclosure express STRO-1 and α1β1. In another example, mesenchymal lineage precursor or stem cells express STRO-1 and α2β1. In another example, mesenchymal lineage precursor or stem cells express

STRO-1 and α4β1. In another example, mesenchymal lineage precursor or stem cells express STRO-1 and α5β1.

In another example, mesenchymal lineage precursor or stem cells according to the present disclosure express STRO-1 and an integrin having a β3 (CD61) chain polypeptide. In an example, the present disclosure encompasses a population of mesenchymal lineage precursor or stem cells enriched for STRO-1+ and β3+ cells. In this example, a population enriched for β3+ cells comprises at least about 8% or 10% or 15% β3+ cells. In another example, mesenchymal lineage precursor or stem cells express STRO-1 and αvβ3. In another example, mesenchymal lineage precursor or stem cells according to the present disclosure express STRO-1 and an integrin having a β5 (ITGB5) chain polypeptide. In an example, mesenchymal lineage precursor or stem cells express STRO-1 and αvβ5. In another example, mesenchymal lineage precursor or stem cells express STRO-1 and αvβ6.

Identifying and/or enriching for mesenchymal lineage precursor or stem cells expressing above referenced integrins may be achieved using various methods known in the art. In one example, fluorescent activated cell sorting (FACS) can be employed using commercially available antibodies (e.g. Thermofisher; Pharmingen; Abcam) to identify and select for cells expressing a desired integrin polypeptide chain or combination thereof.

In an example, mesenchymal lineage precursor or stem cells express STRO-1 and coxsackievirus and adenovirus receptor. In another example, mesenchymal lineage precursor or stem cells express STRO-1, coxsackievirus and adenovirus receptor and one or more of the above referenced integrin's.

In another example, mesenchymal lineage precursor or stem cells express STRO-1, coxsackievirus and adenovirus receptor, αvβ3 and αvβ5.

In an example, mesenchymal lineage precursor or stem cells are genetically modified to express one or more of the above referenced integrin's or coxsackievirus and adenovirus receptor on their cell surface.

In an example, mesenchymal lineage precursor or stem cells express STRO-1 and a chimeric antigen receptor (CAR). In an example, mesenchymal lineage precursor or stem cells express STRO-1, CAR, αvβ3 and αvβ5.

In an example, mesenchymal lineage precursor or stem cells expressing CAR can trigger a T cell mediated immune response. In another example, the CAR acts as a means of attaching mesenchymal lineage precursor or stem cells to cancer cells. In another example, the CAR acts as a means of triggering enhanced adhesion of mesenchymal lineage precursor or stem cells to cancer cells.

In an example, the CAR is comprised of an extracellular antigen binding domain, a transmembrane domain, and an intracellular domain. In an example, the antigen binding domain possesses affinity for one or more tumour antigens. Exemplary tumour antigens include HER2, CLPP, 707-AP, AFP, ART-4, BAGE, MAGE, GAGE, SAGE, b-catenin/m, bcr-abl, CAMEL, CAP-1, CEA, CASP-8, CDK/4, CDC-27, Cyp-B, DAM-8, DAM-10, ELV-M2, ETV6, G250, Gp100, HAGE, HER-2/neu, EPV-E6, LAGE, hTERT, survivin, iCE, MART-1, tyrosinase, MUC-1, MC1-R, TEL/AML, and WT-1.

Exemplary intracellular domains include CD3-zeta, CD28 and 4- IBB. In some examples, the CAR can comprise a combination of CD3-zeta, CD28, 4- 1 BB, TLR-4.

Exemplary transmembrane domains can be derived from (i.e. comprise at least the transmembrane region(s) of) the alpha, beta or zeta chain of the T-cell receptor, CD28, CD3 epsilon, CD45, CD4, CD5, CDS, CD9, CD 16, CD22, CD33, CD37, CD64, CD80, CD86, CD 134, CD137, 35 CD 154. In another example, the transmembrane domain can be synthetic, in which case it will comprise predominantly hydrophobic residues such as leucine and valine.

Mesenchymal lineage precursor or stem cells can be isolated from host tissues such as those referred to above and enriched for by immunoselection. For example, a bone marrow aspirate from a subject may be further treated with an antibody to STRO-1 or TNAP to enable selection of mesenchymal lineage precursor or stem cells. In one example, the mesenchymal lineage precursor or stem cells can be enriched for by using the STRO-1 antibody described in Simmons & Torok-Storb, 1991.

STRO-1+ cells are cells found in bone marrow, blood, dental pulp cells, adipose tissue, skin, spleen, pancreas, brain, kidney, liver, heart, retina, brain, hair follicles, intestine, lung, lymph node, thymus, bone, ligament, tendon, skeletal muscle, dermis, and periosteum; and are capable of differentiating into germ lines such as mesoderm and/or endoderm and/or ectoderm. Thus, STRO-1+ cells are capable of differentiating into a large number of cell types including, but not limited to, adipose, osseous, cartilaginous, elastic, muscular, and fibrous connective tissues. The specific lineage-commitment and differentiation pathway which these cells enter depends upon various influences from mechanical influences and/or endogenous bioactive factors, such as growth factors, cytokines, and/or local microenvironmental conditions established by host tissues.

The term "enriched" as used herein describes a population of cells in which the proportion of one particular cell type or the proportion of a number of particular cell

types is increased when compared with an untreated population of the cells (e.g., cells in their native environment). In one example, a population enriched for STRO-1+ cells comprises at least about 0.1% or 0.5% or 1% or 2% or 5% or 10% or 15% or 20% or 25% or 30% or 50% or 75% STRO-1+ cells. In this regard, the term "population of cells enriched for STRO-1+ cells" will be taken to provide explicit support for the term "population of cells comprising X% STRO-1+ cells", wherein X% is a percentage as recited herein. The STRO-1+ cells can, in some examples, form clonogenic colonies, for example, CFU-F (fibroblasts) or a subset thereof (e.g., 50% or 60% or 70% or 70% or 90% or 95%) can have this activity.

In one example, the population of cells is enriched from a cell preparation comprising STRO-1+ cells in a selectable form. In this regard, the term "selectable form" will be understood to mean that the cells express a marker (e.g., a cell surface marker) permitting selection of the STRO-1+ cells. The marker can be STRO-1, but need not be. For example, as described and/or exemplified herein, cells (e.g., MPCs) expressing STRO-2 and/or STRO-3 (TNAP) and/or STRO-4 and/or VCAM-1 and/or CD146 and/or 3G5 also express STRO-1 (and can be STRO-1bright). Accordingly, an indication that cells are STRO-1+ does not mean that the cells are selected by STRO-1 expression. In one example, the cells are selected based on at least STRO-3 expression, e.g., they are STRO-3+ (TNAP+).

Reference to selection of a cell or population thereof does not necessarily require selection from a specific tissue source. As described herein, STRO-1+ cells can be selected from or isolated from or enriched from a large variety of sources. That said, in some examples, these terms provide support for selection from any tissue comprising STRO-1+ cells or vascularized tissue or tissue comprising pericytes (e.g., STRO-1+ or 3G5+ pericytes) or any one or more of the tissues recited herein.

In one example, the mesenchymal lineage precursor or stem cells of the disclosure express one or more markers individually or collectively selected from the group consisting of TNAP+, VCAM-1+, THY-1+, STRO-2+, STRO-4+ (HSP-90β), CD45+, CD146+, 3G5+.

By "individually" is meant that the disclosure encompasses the recited markers or groups of markers separately, and that, notwithstanding that individual markers or groups of markers may not be separately listed herein, the accompanying claims may define such marker or groups of markers separately and divisibly from each other.

By "collectively" is meant that the disclosure encompasses any number or combination of the recited markers or groups of markers, and that, notwithstanding that such numbers or combinations of markers or groups of markers may not be specifically

listed herein, the accompanying claims may define such combinations or sub-combinations separately and divisibly from any other combination of markers or groups of markers.

A cell that is referred to as being "positive" for a given marker may express either a low (lo or dim or dull), intermediate (median) or a high (bright, bri) level of that marker depending on the degree to which the marker is present on the cell surface, where the terms relate to intensity of fluorescence or other marker used in the sorting process of the cells or flow cytometric analysis of the cells. The distinction of low (lo or dim or dull), intermediate (median), or high (bright, bri) will be understood in the context of the marker used on a particular cell population being sorted or analysed. A cell that is referred to as being "negative" for a given marker is not necessarily completely absent from that cell. This term means that the marker is expressed at a relatively very low level by that cell, and that it generates a very low signal when detectably labeled or is undetectable above background levels, for example, levels detected using an isotype control antibody.

The term "bright" or bri as used herein, refers to a marker on a cell surface that generates a relatively high signal when detectably labeled. Whilst not wishing to be limited by theory, it is proposed that "bright" cells express more of the target marker protein (for example, the antigen recognized by a STRO-1 antibody) than other cells in the sample. For instance, STRO-1bri cells produce a greater fluorescent signal, when labeled with a FITC-conjugated STRO-1 antibody as determined by fluorescence activated cell sorting (FACS) analysis, than non-bright cells (STRO-1lo/dim/dull/intermediate/median). In one example, the mesenchymal lineage precursor or stem cells are isolated from bone marrow and enriched for by selection of STRO-1+ cells. In this example, "bright" cells constitute at least about 0.1% of the most brightly labeled bone marrow mononuclear cells contained in the starting sample. In other examples, "bright" cells constitute at least about 0.1%, at least about 0.5%, at least about 1%, at least about 1.5%, or at least about 2%, of the most brightly labeled bone marrow mononuclear cells contained in the starting sample. In an example, STRO-1bright cells have 2 log magnitude higher expression of STRO-1 surface expression relative to "background", namely cells that are STRO-1-. By comparison, STRO-1lo/dim/dull and/or STRO-1intermediate/median cells have less than 2 log magnitude higher expression of STRO-1 surface expression, typically about 1 log or less than "background".

In one example, the STRO-1+ cells are STRO-1bright. In one example, the STRO-1bright cells are preferentially enriched relative to STRO-1lo/dim/dull or STRO-1intermediate/median cells.

In one example, the STRO-1bright cells are additionally one or more of TNAP+, VCAM-1+, THY-1+, STRO-2+, STRO-4+ (HSP-90β) and/or CD146+. For example, the cells are selected for one or more of the foregoing markers and/or shown to express one or more of the foregoing markers. In this regard, a cell shown to express a marker need not be specifically tested, rather previously enriched or isolated cells can be tested and subsequently used, isolated or enriched cells can be reasonably assumed to also express the same marker.

In one example, the STRO-1bright cells are perivascular mesenchymal lineage precursor or stem cells as defined in WO 2004/85630, characterized by the presence of the perivascular marker 3G5.

As used herein the term "TNAP" is intended to encompass all isoforms of tissue non-specific alkaline phosphatase. For example, the term encompasses the liver isoform (LAP), the bone isoform (BAP) and the kidney isoform (KAP). In one example, the TNAP is BAP. In one example, TNAP refers to a molecule which can bind the STRO-3 antibody produced by the hybridoma cell line deposited with ATCC on 19 December 2005 under the provisions of the Budapest Treaty under deposit accession number PTA-7282.

Furthermore, in one example, the STRO-1+ cells are capable of giving rise to clonogenic CFU-F.

In one example, a significant proportion of the STRO-1+ cells are capable of differentiation into at least two different germ lines. Non-limiting examples of the lineages to which the cells may be committed include bone precursor cells; hepatocyte progenitors, which are multipotent for bile duct epithelial cells and hepatocytes; neural restricted cells, which can generate glial cell precursors that progress to oligodendrocytes and astrocytes; neuronal precursors that progress to neurons; precursors for cardiac muscle and cardiomyocytes, glucose-responsive insulin secreting pancreatic beta cell lines. Other lineages include, but are not limited to, odontoblasts, dentin-producing cells and chondrocytes, and precursor cells of the following: retinal pigment epithelial cells, fibroblasts, skin cells such as keratinocytes, dendritic cells, hair follicle cells, renal duct epithelial cells, smooth and skeletal muscle cells, testicular progenitors, vascular endothelial cells, tendon, ligament, cartilage, adipocyte, fibroblast, marrow stroma, cardiac muscle, smooth muscle, skeletal muscle, pericyte, vascular, epithelial, glial, neuronal, astrocyte and oligodendrocyte cells.

In one example, the mesenchymal lineage precursor or stem cells are MSCs. The MSCs may be a homogeneous composition or may be a mixed cell population enriched in MSCs. Homogeneous MSC compositions may be obtained by culturing adherent bone marrow or periosteal cells, and the MSCs may be identified by specific cell surface markers which are identified with unique monoclonal antibodies. A method for obtaining a cell population enriched in MSCs is described, for example, in US patent 5486359. MSC prepared by conventional plastic adherence isolation relies on the non-specific plastic adherent properties of CFU-F. Mesenchymal lineage precursor or stem cells isolated from bone marrow by immunoselection based on STRO-1 specifically isolates clonogenic mesenchymal precursors from bone marrow populations in the absence of other plastic adherent bone marrow populations. Alternative sources for MSCs include, but are not limited to, blood, skin, cord blood, muscle, fat, bone, and perichondrium.

In one example, the mesenchymal lineage precursor or stem cells are derived from pluripotent cells such as induced pluripotent stem cells (iPS cells). In one embodiment the pluripotent cells are human pluripotent cells. Suitable processes for generation of mesenchymal lineage precursor or stem cells from pluripotent cells are described, for example, in US 7,615,374 and US 2014273211, Barberi *et al*; Plos medicine, Vol 2(6):0554-0559 (2005), and Vodyanik *et al*. Cell Stem cell, Vol 7:718-728 (2010).

In another example, the mesenchymal lineage precursor or stem cells are immortalised. Exemplary processes for generation of immortalised mesenchymal lineage precursor or stem cells are described, for example, in Obinata M., Cell, Vol 2:235-244 (1997), US 9,453,203, Akimov et al. Stem Cells, Vol 23:1423-1433 and Kabara et al. Laboratory Investigation, Vol 94: 1340-1354 (2014).

In a preferred embodiment of the present disclosure, the mesenchymal lineage precursor or stem cells are obtained from a master cell bank derived from mesenchymal lineage precursor or stem cells enriched from the bone marrow of healthy volunteers. The use of mesenchymal lineage precursor or stem cells derived from such a source is particularly advantageous for subjects who do not have an appropriate family member available who can serve as the mesenchymal lineage precursor or stem cell donor, or are in need of immediate treatment and are at high risk of relapse, disease-related decline or death, during the time it takes to generate mesenchymal lineage precursor or stem cells.

In another example, mesenchymal lineage precursor cells express Cx43. In another example, mesenchymal lineage precursor cells express Cx40. In another

example, mesenchymal lineage precursor cells express Cx43 and Cx40. In another example, mesenchymal lineage precursor cells express Cx45, Cx32 and/or Cx37. In an example, mesenchymal lineage precursor cells are not modified to express a particular connexin.

In another example, mesenchymal lineage precursor cells express STRO-1 and CD46. In this example, mesenchymal lineage precursor cells can also express one or more integrin's referenced above.

Isolated or enriched mesenchymal lineage precursor cells can be expanded *in vitro* by culture. Isolated or enriched mesenchymal lineage precursor cells can be cryopreserved, thawed and subsequently expanded *in vitro* by culture.

In one example, isolated or enriched mesenchymal lineage precursor cells are seeded at 50,000 viable cells/cm^2 in culture medium (serum free or serum-supplemented), for example, alpha minimum essential media (αMEM) supplemented with 5% fetal bovine serum (FBS) and glutamine, and allowed to adhere to the culture vessel overnight at 37°C, 20% O^2. The culture medium is subsequently replaced and/or altered as required and the cells cultured for a further 68 to 72 hours at 37°C, 5% O^2.

As will be appreciated by those of skill in the art, cultured mesenchymal lineage precursor cells are phenotypically different to cells *in vivo*. For example, in one embodiment they express one or more of the following markers, CD44, CD46, NG2, DC146 and CD140b. Cultured mesenchymal lineage precursor cells are also biologically different to cells *in vivo*, having a higher rate of proliferation compared to the largely non-cycling (quiescent) cells *in vivo*.

Ang1 and VEGF levels

In an example, mesenchymal lineage precursor or stem cells express Ang1 in an amount of at least 0.1 μg/10^6 cells. However, in other examples, mesenchymal lineage precursor or stem cells express Ang1 in an amount of at least 0.2 μg/10^6 cells, 0.3 μg/10^6 cells, 0.4 μg/10^6 cells, 0.5 μg/10^6 cells, 0.6 μg/10^6 cells, 0.7 μg/10^6 cells, 0.8 μg/10^6 cells, 0.9 μg/10^6 cells, 1 μg/10^6 cells, 1.1 μg/10^6 cells, 1.2 μg/10^6 cells, 1.3 μg/10^6 cells, 1.4 μg/10^6 cells, 1.5 μg/10^6 cells.

In another example, mesenchymal lineage precursor or stem cells express VEGF in an amount less than about 0.05 μg/10^6 cells. However, in other examples, mesenchymal lineage precursor or stem cells express VEGF in an amount less than about 0.05 μg/10^6 cells, 0.04 μg/10^6 cells, 0.03 μg/10^6 cells, 0.02 μg/10^6 cells, 0.01 μg/10^6 cells, 0.009 μg/10^6 cells, 0.008 μg/10^6 cells, 0.007 μg/10^6 cells, 0.006 μg/10^6

cells, 0.005 μg/10^6 cells, 0.004 μg/10^6 cells, 0.003 μg/10^6 cells, 0.002 μg/10^6 cells, 0.001 μg/10^6 cells.

The amount of cellular Ang1 and/or VEGF that is expressed in a composition or culture of mesenchymal lineage precursor or stem cells may be determined by methods known to those skilled in the art. Such methods include, but are not limited to, quantitative assays such as quantitative ELISA assays, for example. In this example, a cell lysate from a culture of mesenchymal lineage precursor or stem cells is added to a well of an ELISA plate. The well may be coated with a primary antibody, either a monoclonal or a polyclonal antibody(ies), against the Ang1 or VEGF. The well then is washed, and then contacted with a secondary antibody, either a monoclonal or a polyclonal antibody(ies) , against the primary antibody. The secondary antibody is conjugated to an appropriate enzyme, such as horseradish peroxidase, for example. The well then may be incubated, and then is washed after the incubation period. The wells then are contacted with an appropriate substrate for the enzyme conjugated to the secondary antibody, such as one or more chromogens. Chromogens which may be employed include, but are not limited to, hydrogen peroxide and tetramethylbenzidine. After the substrate(s) is (are) added, the well is incubated for an appropriate period of time. Upon completion of the incubation, a "stop" solution is added to the well in order to stop the reaction of the enzyme with the substrate(s). The optical density (OD) of the sample is then measured. The optical density of the sample is correlated to the optical densities of samples containing known amounts of Ang1 or VEGF in order to determine the amount of Ang1 or VEGF expressed by the culture of stem cells being tested.

In another aspect, mesenchymal lineage precursor or stem cells express Ang1:VEGF at a ratio of at least about 2:1. However, in other examples, mesenchymal lineage precursor or stem cells express Ang1:VEGF at a ratio of at least about 10:1, 15:1, 20:1, 21:1, 22:1, 23:1, 24:1, 25:1, 26:1, 27:1, 28:1, 29:1, 30:1, 31:1, 32:1, 33:1, 34:1, 35:1, 50:1.

Methods for determining the Ang1:VEGF expression ratio will be apparent to one of skill in the art. For example Ang1 and VEGF expression levels can be quantitated via quantitative ELISA as discussed above. After quantifying the levels of Ang1 and VEGF, a ratio based on the quantitated levels of Ang1 and VEGF could be represented as: (level of Ang1 / level of VEGF) = Ang1:VEGF ratio.

In an example, the mesenchymal lineage precursor or stem cells of the present disclosure are not genetically modified to express Ang1 and/or VEGF at an above exemplified level or ratio. Cells that are not genetically modified to express Ang1

and/or VEGF have not been modified by transfection with a nucleic acid expressing or encoding Ang1 and/or VEGF. For the avoidance of doubt, in the context of the present disclosure a mesenchymal lineage precursor or stem cell transfected with a nucleic acid encoding Ang1 and/or VEGF would be considered genetically modified. In the context of the present disclosure cells not genetically modified to express Ang1 and/or VEGF naturally express Ang1 and/or VEGF to some extent without transfection with a nucleic acid encoding Ang1 and/or VEGF1.

Nucleic Acids

The present disclosure encompasses use of various nucleic acids. The terms "nucleic acid" and "nucleic acid molecule" may be used interchangeably throughout the disclosure. Exemplary nucleic acids include DNA (e.g., complementary DNA (cDNA), genomic DNA (gDNA)), RNA (e.g., message RNA (mRNA), short hairpin RNA (shRNA), short inhibitory RNA (siRNA), ribosomal RNA (rRNA), tRNA, microRNA, DNA or RNA analogues (e.g., containing base analogues, sugar analogues and/or a non-native backbone and the like), RNA/DNA hybrids and polyamide nucleic acids (PNAs), all of which can be in single- or double-stranded form. In an example, the nucleic acid is isolated. As used herein, the term "isolated nucleic acid" means a nucleic acid that is altered or removed from the natural state through human intervention.

Another exemplary nucleic acid is an oligonucleotide. The term "oligonucleotide" as used herein means a short DNA or RNA molecule. Oligonucleotides readily bind, in a sequence-specific manner, to their respective complementary oligonucleotides, DNA, or RNA to form duplexes.

In an example, the oligonucleotide can be a DNA molecule. In another example, the oligonucleotide can be an RNA molecule.

In various examples, oligonucleotides can be about five, six, seven, eight, nine, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30 nucleotides or larger in length.

In an example, oligonucleotides of the present disclosure are inhibitory oligonucleotides. In an example, the term "inhibitory oligonucleotide" refers to any oligonucleotide that reduces the production, expression or biological activity of one or more proteins. For example, an inhibitory oligonucleotide can interfere with translation of mRNA into protein in a ribosome. In another example, an inhibitory oligonucleotide can be sufficiently complementary to either a gene or a mRNA encoding one or more proteins to bind to (hybridize with) a targeted gene(s) or mRNA thereby reducing

expression or biological activity of the target protein. In another example, an inhibitory oligonucleotide inhibits the biological activity of an intracellular nucleic acid that does not code for a protein. For example, an inhibitory oligonucleotide can inhibit the biological activity of a non-coding RNA.

Exemplary inhibitory oligonucleotides include isolated or synthetic antisense RNA or DNA, siRNA or siDNA, miRNA, miRNA mimics, shRNA or DNA and Chimeric Antisense DNA or RNA.

The term "antisense" as used herein means a sequence of nucleotides complementary to and therefore capable of binding to a coding sequence, which may be either that of the strand of a DNA double helix that undergoes transcription, or that of a messenger RNA molecule. Antisense DNA is the non-coding strand complementary to the coding strand in double-stranded DNA. The antisense strand serves as the template for messenger RNA (mRNA) synthesis.

The terms "short hairpin RNA" or "shRNA" refer to an RNA structure having a duplex region and a loop region.

The term small interfering RNA (siRNA), sometimes known as short interfering RNA or silencing RNA, is a class of double-stranded RNA molecules, 20-25 base pairs in length. A siRNA that inhibits or prevents translation to a particular protein is indicated by the protein name coupled with the term siRNA. Thus a siRNA that interferes with the translation of KIF11 is indicated by the expression "KIF11 siRNA". Typically, a siRNA in various embodiments is a double-stranded nucleic acid molecule comprising two nucleotide strands, each strand having about 19 to about 28 nucleotides (i.e. about 19, 20, 21, 22, 23, 24, 25, 26, 27, or 28 nucleotides). Exemplary siRNAs encompassed by the present disclosure include KIF11 siRNA, KRAS siRNA and PLK1 siRNA.

The term “microRNA” (abbreviated miRNA) is a small non-coding RNA molecule (containing about 22 nucleotides) found in plants, animals and some viruses, that functions in RNA silencing and post-transcriptional regulation of gene expression. The prefix "miR" is followed by a dash and a number, the latter often indicating order of naming. Different miRNAs with nearly identical sequences except for one or two nucleotides are annotated with an additional lower case letter. Numerous miRNAs are known in the art (miRBase V.21 nomenclature; Kozomara et al. 2013; Griffiths-Jones, S. 2004). Examples of human miRNAs include, hsa-let-7d-3p, hsa-miR-101-5p, hsa-miR-106b-3p, hsa-miR-1179, hsa-miR-125a-5p, hsa-miR-141- 3p, hsa-miR-148a-3p, hsa-miR-16-5p, hsa-miR-192-5p, hsa-miR-195-5p, hsa-miR-19b-3p, hsa-miR-20a-5p, hsa-miR-200b-3p, hsa-miR-217, hsa-miR-223-3p, hsa-miR-371a-3p, hsa- miR-

487b, hsa-miR-515-3p, hsa-miR-605, hsa-let-7a-3p, hsa-let-7a-5p, hsa-let-7b-3p, hsa-let-7b-5p, hsa-let-7c, hsa-let-7d-5p, hsa-let-7e-3p, hsa-let-7e-5p, hsa-let-7f-l-3p, hsa-let-7f-2- 3p, hsa-let-7f-5p, hsa-let-7g-3p, hsa-let-7g-5p, hsa-let-7i-3p, hsa-let-7i-5p, hsa-miR-1, hsa- miR-100-3p, hsa-miR-100-5p, hsa-miR-101-3p, hsa-miR-103a-2-5p, hsa-miR-103a-3p, hsa- miR-103b, hsa-miR-105-5p, hsa-miR-106a-3p, hsa-miR-106a-5p, hsa-miR-106b-5p, hsa- miR-107, hsa-miR-10a-3p, hsa-miR-10a-5p, hsa-miR-10b-3p, hsa-miR-10b-5p, hsa-miR- 1178-3p, hsa-miR-1180, hsa-miR-1181, hsa-miR-1182, hsa-miR-1183, hsa-miR-1184, hsa- miR-1185-5p, hsa-miR-1204, hsa-miR-1207, hsa-miR-1208, hsa-miR-122-3p, hsa-miR-122- 5p, hsa-miR-1224, hsa-miR-1226, hsa-miR-1227-3p, hsa-miR-1228-5p, hsa-miR-1229-3p, hsa-miR-1231, hsa-miR-124-3p, hsa-miR-1245 a, hsa-miR-1246, hsa-miR-1249, hsa-miR- 125a-3p, hsa-miR- 125b- l-3p, hsa-miR- 125b-2-3p, hsa-miR- 125b-5p, hsa-miR-1251, hsa- miR-1252, hsa-miR-1253, hsa-miR- 1255 a, hsa-miR-1255b-5p, hsa-miR- 126-3p, hsa-miR- 1260a, hsa-miR-1260b, hsa-miR- 1262, hsa-miR- 1263, hsa-miR- 1265, hsa-miR- 1268a, hsa- miR-127-3p, hsa-miR-1270, hsa-miR-1272, hsa-miR- 1273 a, hsa-miR-1275, hsa-miR-1276, hsa-miR- 1277-3p, hsa-miR-1278, hsa-miR-128, hsa-miR-1285-3p, hsa-miR-1286, hsa-miR- 1287, hsa-miR- 129-2-3p, hsa-miR- 1292-5p, hsa-miR-1293, hsa-miR-1301, hsa-miR-1302, hsa-miR-1303, hsa-miR-1305, hsa-miR-130a-3p, hsa-miR- 130a-5p, hsa-miR-130b-3p, hsa- miR-130b-5p, hsa-miR- 132-3p, hsa-miR- 132-5p, hsa-miR-1321, hsa-miR-1323, hsa-miR- 1324, hsa-miR- 133a, hsa-miR- 133b, hsa-miR- 134, hsa-miR- 135a-3p, hsa-miR- 135a-5p, hsa- miR-135b-3p, hsa-miR- 135b-5p, hsa-miR-136-3p, hsa-miR-136-5p, hsa-miR-137, hsa-miR- 138-l-3p, hsa-miR- 138-2-3p, hsa-miR- 138-5p, hsa-miR- 139-3p, hsa-miR-139-5p, hsa-miR- 140-3p, hsa-miR- 140-5p, hsa-miR- 142-3p, hsa-miR- 142-5p, hsa-miR-143-3p, hsa-miR-143- 5p, hsa-miR- 144-3p, hsa-miR- 144-5p, hsa-miR-145-3p, hsa-miR-145-5p, hsa-miR-1468, hsa- miR-146a-5p, hsa-miR- 146b-3p, hsa-miR- 146b-5p, hsa-miR- 147a, hsa-miR- 147b, hsa-miR- 148a-5p, hsa-miR- 148b-3p, hsa-miR- 148b-5p, hsa-miR- 149-3p, hsa-miR- 149-5p, hsa-miR- 150-3p, hsa-miR- 150-5p, hsa-miR-151a-3p, hsa-miR-151a-5p, hsa-miR-152, hsa-miR-153, hsa-miR-1538, hsa-miR-1539, hsa-miR- 154-3p, hsa-miR- 154-5p, hsa-miR- 155-5p, hsa-miR- 15a-3p, hsa-miR- 15a-5p, hsa-miR- 15b-3p, hsa-miR- 15b-5p, hsa-miR- 16- l-3p, hsa-miR-16- 2-3p, hsa-miR- 17-3p, hsa-miR- 17-5p, hsa-miR-181a-2-3p, hsa-miR-181a-3p, hsa-miR-181a- 5p, hsa-miR-181b-5p, hsa-miR-181c-3p, hsa-miR-181c-5p, hsa-miR-181d, hsa-miR- 182-5p, hsa-miR-1827, hsa-miR- 183-3p, hsa-miR- 183-5p, hsa-miR-184, hsa-miR-185-3p, hsa-miR- 185-5p, hsa-miR- 186-3p, hsa-miR- 186-5p, hsa-miR- 187-3p, hsa-miR- 187-5p, hsa-miR-188- 5p, hsa-miR- 18a-3p, hsa-miR- 18a-5p, hsa-miR- 18b-3p, hsa-miR- 18b-5p, hsa-miR- 190a, hsa- miR-

190b, hsa-miR-191-3p, hsa-miR-191-5p, hsa-miR-1915-3p, hsa-miR- 192-3p, hsa-miR-193a-3p, hsa-miR- 193a-5p, hsa-miR- 193b-3p, hsa-miR- 193b-5p, hsa-miR- 194-5p, hsa-miR- 195-3p, hsa-miR- 196a-3p, hsa-miR- 196a-5p, hsa-miR- 196b-5p, hsa-miR-197-3p, hsa-miR- 198, hsa-miR- 199a-3p, hsa-miR- 199a-5p, hsa-miR- 199b-3p, hsa-miR- 199b-5p, hsa-miR-19a- 3p, hsa-miR- 19a-5p, hsa-miR- 1909-3p, hsa-miR-191 l-3p, hsa-miR-200a-3p, hsa-miR-200a- 5p, hsa-miR-200b-5p, hsa-miR-200c-3p, hsa-miR-200c-5p, hsa-miR-202-3p, hsa-miR-202- 5p, hsa-miR-203a, hsa-miR-203b-5p, hsa-miR-204-5p, hsa-miR-205-5p, hsa-miR-206, hsa- miR-208b, hsa-miR-20a-3p, hsa-miR-20b-3p, hsa-miR-20b-5p, hsa-miR-21-3p, hsa-miR-21 - 5p, hsa-miR-210, hsa-miR-21 l-5p, hsa-miR-2110, hsa-miR-212-3p, hsa-miR-214-3p, hsa- miR-214-5p, hsa-miR-215, hsa-miR-216a-5p, hsa-miR-218- l-3p, hsa-miR-218-5p, hsa-miR- 219-1 -3p, hsa-miR-219-5p, hsa-miR-22-3p, hsa-miR-22-5p, hsa-miR-221-3p, hsa-miR-221- 5p, hsa-miR-222-3p, hsa-miR-222-5p, hsa-miR-223-5p, hsa-miR-224-3p, hsa-miR-224-5p, hsa-miR-2355-3p, hsa-miR-23a-3p, hsa-miR-23a-5p, hsa-miR-23b-3p, hsa-miR-23b-5p, hsa- miR-24-l-5p, hsa-miR-24-2-5p, hsa-miR-24-3p, hsa-miR-25-3p, hsa-miR-25-5p, hsa-miR- 26a-2-3p, hsa-miR-26a-5p, hsa-miR-26b-3p, hsa-miR-26b-5p, hsa-miR-27a-3p, hsa-miR- 27a-5p, hsa-miR-27b-3p, hsa-miR-27b-5p, hsa-miR-28-3p, hsa-miR-28-5p, hsa-miR-296-3p, hsa-miR-296-5p, hsa-miR-297, hsa-miR-298, hsa-miR-299-3p, hsa-miR-299-5p, hsa-miR- 29a-3p, hsa-miR-29a-5p, hsa-miR-29b-l-5p, hsa-miR-29b-2-5p, hsa-miR-29b-3p, hsa-miR- 29c-3p, hsa-miR-29c-5p, hsa-miR-2909, hsa-miR-301a-3p, hsa-miR-301b, hsa-miR-302a-5p, hsa-miR-302b-5p, hsa-miR-302c-5p, hsa-miR-302d-3p, hsa-miR-302d-5p, hsa-miR-302f, hsa-miR-3064-5p, hsa-miR-3065-3p, hsa-miR-3065-5p, hsa-miR-3074-3p, hsa-miR-3074-5p, hsa-miR-30a-3p, hsa-miR-30a-5p, hsa-miR-30b-3p, hsa-miR-30b-5p, hsa-miR-30c-l-3p, hsa- miR-30c-2-3p, hsa-miR-30c-5p, hsa-miR-30d-3p, hsa-miR-30d-5p, hsa-miR-30e-3p, hsa- miR-30e-5p, hsa-miR-31-3p, hsa-miR-31-5p, hsa-miR-3120-3p, hsa-miR-3120-5p, hsa-miR- 3184-5p, hsa-miR-32-3p, hsa-miR-32-5p, hsa-miR-320a, hsa-miR-320b, hsa-miR-320c, hsa-miR-323a-3p, hsa-miR-323b-5p, hsa-miR-324-3p, hsa-miR-324-5p, hsa-miR-325, hsa-miR- 326, hsa-miR-328, hsa-miR-329, hsa-miR-330-3p, hsa-miR-330-5p, hsa-miR-331-3p, hsa- miR-331-5p, hsa-miR-335-3p, hsa-miR-335-5p, hsa-miR-337-3p, hsa-miR-337-5p, hsa-miR- 338-3p, hsa-miR-338-5p, hsa-miR-339-3p, hsa-miR-339-5p, hsa-miR-33a-5p, hsa-miR-33b- 3p, hsa-miR-33b-5p, hsa-miR-340-3p, hsa-miR-340-5p, hsa-miR-342-3p, hsa-miR-342-5p, hsa-miR-345-5p, hsa-miR-346, hsa-miR-34a-3p, hsa-miR-34a-5p, hsa-miR-34b-3p, hsa-miR- 34b-5p, hsa-miR-34c-3p, hsa-miR-34c-5p, hsa-miR-3591-3p, hsa-miR-361-3p, hsa-miR-361- 5p, hsa-miR-3613-5p, hsa-miR-3615, hsa-miR-3619, hsa-miR-362-3p, hsa-miR-362-5p, hsa- miR-363-3p, hsa-miR-

363-5p, hsa-mir-365a-3p, hsa-mir-365a-5p, hsa-mir-3653, hsa-miR- 3656, hsa-miR-367-3p, hsa-miR-367-5p, hsa-miR-3676-3p, hsa-miR-369-3p, hsa-miR-369- 5p, hsa-miR-370, hsa-miR-372, hsa-miR-373-3p, hsa-miR-373-5p, hsa-miR-374a-3p, hsa-miR-374a-5p, hsa-miR-374b-3p, hsa-miR-374b-5p, hsa-miR-375, hsa-miR-376a-2-5p, hsa- miR-376a-3p, hsa-miR-376a-5p, hsa-miR-376b-3p, hsa-miR-376c-3p, hsa-miR-377-3p, hsa- miR-377-5p, hsa-miR-378a-3p, hsa-miR-378a-5p, hsa-miR-378c, hsa-miR-378d, hsa-miR- 379-3p, hsa-miR-379-5p, hsa-miR-380-3p, hsa-miR-380-5p, hsa-miR-381-3p, hsa-miR-382- 3p, hsa-miR-382-5p, hsa-miR-383, hsa-miR-384, hsa-miR-3912, hsa-miR-3928, hsa-miR- 409-3p, hsa-miR-409-5p, hsa-miR-410, hsa-miR-41 l-3p, hsa-miR-41 l-5p, hsa-miR-412, hsa- miR-421, hsa-miR-422a, hsa-miR-423-3p, hsa-miR-423-5p, hsa-miR-424-3p, hsa-miR-424- 5p, hsa-miR-425-3p, hsa-miR-425-5p, hsa-miR-429, hsa-miR-4291, hsa-miR-431-5p, hsa- miR-432-5p, hsa-miR-433, hsa-miR-4421, hsa-miR-449a, hsa-miR-450a-5p, hsa-miR-450b- 3p, hsa-miR-450b-5p, hsa-miR-4505, hsa-miR-4510, hsa-miR-4516, hsa-miR-451a, hsa-miR- 452-3p, hsa-miR-452-5p, hsa-miR-4533, hsa-miR-4539, hsa-miR-454-3p, hsa-miR-454-5p, hsa-miR-455-3p, hsa-miR-455-5p, hsa-miR-4634, hsa-miR-4732-3p, hsa-miR-4732-5p, hsa- miR-4747-5p, hsa-miR-4792, hsa-miR-483-3p, hsa-miR-483-5p, hsa-miR-484, hsa-miR-485- 5p, hsa-miR-486-3p, hsa-miR-486-5p, hsa-miR-489, hsa-miR-490, hsa-miR-491-3p, hsa- miR-491-5p, hsa-miR-492, hsa-miR-493-3p, hsa-miR-493-5p, hsa-miR-494, hsa-miR-495- 3p, hsa-miR-496, hsa-miR-497-3p, hsa-miR-497-5p, hsa-miR-498, hsa-miR-499a-5p, hsa- miR-500a-3p, hsa-miR-501-3p, hsa-miR-501-5p, hsa-miR-502-3p, hsa-miR-502-5p, hsa- miR-503-5p, hsa-miR-504, hsa-miR-505-3p, hsa-miR-505-5p, hsa-miR-506-3p, hsa-miR- 508-3p, hsa-miR-508-5p, hsa-miR-509-3p, hsa-miR-511 , hsa-miR-512-5p, hsa-miR-513a-3p, hsa-miR-513a-5p, hsa-miR-513b, hsa-miR-514a-3p, hsa-miR-514a-5p, hsa-miR-515-5p, hsa- miR-516b-3p, hsa-miR-516b-5p, hsa-miR-517a-3p, hsa-miR-518a-3p, hsa-miR-518b, hsa- miR-518e-3p, hsa-miR-518e-5p, hsa-miR-518f-3p, hsa-miR-519a-5p, hsa-miR-519b-5p, hsa- miR-519c-3p, hsa-miR-519c-5p, hsa-miR-519d, hsa-miR-519e-5p, hsa-miR-520c-3p, hsa- miR-520e, hsa-miR-520f, hsa-miR-520g, hsa-miR-520h, hsa-miR-521, hsa-miR-522-5p, hsa-miR-523-5p, hsa-miR-525-3p, hsa-miR-532-3p, hsa-miR-532-5p, hsa-miR-539-5p, hsa-miR- 541, hsa-miR-542-3p, hsa-miR-542-5p, hsa-miR-543, hsa-miR-545-3p, hsa-miR-545-5p, hsa- miR-548a-3p, hsa-miR-548d-3p, hsa-miR-548e, hsa-miR-548i, hsa-miR-548m, hsa-miR-549, hsa-miR-550a-3p, hsa-miR-550a-5p, hsa-miR-551b-3p, hsa-miR-551b-5p, hsa-miR-552, hsa- miR-553, hsa-miR-554, hsa-miR-557, hsa-miR-563, hsa-miR-564, hsa-miR-567, hsa-miR- 569, hsa-miR-570-3p, hsa-miR-571, hsa-miR-572, hsa-miR-574-3p, hsa-miR-574-5p, hsa- miR-575, hsa-miR-576-3p, hsa-miR-576-

5p, hsa-miR-577, hsa-miR-578, hsa-miR-580, hsa- miR-582-3p, hsa-miR-582-5p, hsa-miR-583, hsa-miR-584-5p, hsa-miR-585, hsa-miR-586, hsa-miR-589-3p, hsa-miR-589-5p, hsa-miR-590-3p, hsa-miR-590-5p, hsa-miR-593-3p, hsa- miR-593-5p, hsa-miR-595, hsa-miR-598, hsa-miR-601, hsa-miR-602, hsa-miR-603, hsa- miR-606, hsa-miR-608, hsa-miR-609, hsa-miR-611, hsa-miR-612, hsa-miR-613, hsa-miR- 615-3p, hsa-miR-615-5p, hsa-miR-616-5p, hsa-miR-618, hsa-miR-619, hsa-miR-620, hsa- miR-623, hsa-miR-625-5p, hsa-miR-626, hsa-miR-627, hsa-miR-628-3p, hsa-miR-628-5p, hsa-miR-629-3p, hsa-miR-629-5p, hsa-miR-630, hsa-miR-631, hsa-miR-634, hsa-miR-635, hsa-miR-636, hsa-miR-638, hsa-miR-639, hsa-miR-641, hsa-miR-642a-3p, hsa-miR-642a- 5p, hsa-miR-643, hsa-miR-645, hsa-miR-646, hsa-miR-647, hsa-miR-649, hsa-miR-650, hsa- miR-651, hsa-miR-652-3p, hsa-miR-653, hsa-miR-654-3p, hsa-miR-655, hsa-miR-656, hsa- miR-657, hsa-miR-658, hsa-miR-659-3p, hsa-miR-660-5p, hsa-miR-663a, hsa-miR-663b, hsa-miR-664a-3p, hsa-miR-664a-5p, hsa-miR-668, hsa-miR-671-5p, hsa-miR-675-3p, hsa- miR-675-5p, hsa-miR-7-l-3p, hsa-miR-7-2-3p, hsa-miR-7-5p, hsa-miR-708-3p, hsa-miR- 708-5p, hsa-miR-718, hsa-miR-744-3p, hsa-miR-744-5p, hsa-miR-761, hsa-miR-765, hsa- miR-766, hsa-miR-767-3p, hsa-miR-769-3p, hsa-miR-769-5p, hsa-miR-770-5p, hsa-miR- 874, hsa-miR-875, hsa-miR-876, hsa-miR-877-3p, hsa-miR-877-5p, hsa-miR-885-3p, hsa- miR-887, hsa-miR-888-3p, hsa-miR-889, hsa-miR-890, hsa-miR-891a, hsa-miR-891b, hsa- miR-9-3p, hsa-miR-9-5p, hsa-miR-92a-3p, hsa-miR-92b-3p, hsa-miR-92b-5p, hsa-miR-921, hsa-miR-922, hsa-miR-93-3p, hsa-miR-93-5p, hsa-miR-935, hsa-miR-940, hsa-miR-941, hsa-miR-942, hsa-miR-95, hsa-miR-96-3p, hsa-miR-96-5p, hsa-miR-98-5p, hsa-miR-99a-3p, hsa-miR-99a-5p, hsa-miR-99b-3p, and/or hsa-miR-99b-5p. Sequences of these miRNAs are well known in the art and may be found, for example, on the world wide web at mirbase dot org.

Suitable example miRNAs from the above list include hsa-miR-155, hsa-miR-155-inh, hsa-miR-181-B1, hsa-miR-15a, hsa-miR-16-1, hsa-miR-21, hsa-miR-34a, hsa-miR-221 and hsa-miR-29a.

In an example, “inhibitory oligonucleotides” mimic the activity of one or more miRNA. The term “miRNA mimic”, as used herein, refers to small, double-stranded RNA molecules designed to mimic endogenous mature miRNA molecules when introduced into cells. miRNA mimics can be obtained from various suppliers such as Sigma Aldrich and Thermo Fisher Scientific.

In another example, “inhibitory oligonucleotides” inhibit the activity of one or more miRNA. Various miRNA species are suitable for this purpose. Examples include, without limitation, antagomirs, interfering RNA, ribozymes, miRNA sponges

and miR-masks. The term “antagomir” is used in the context of the present disclosure to refer to chemically modified antisense oligonucleotides that bind to a target miRNA and inhibit miRNA function by preventing binding of the miRNA to its cognate gene target. Antagomirs can include any base modification known in the art. In an example, the above referenced miRNA species are about 10 to 50 nucleotides in length. For example, antagomirs can have antisense portions of 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, or 50 nucleotides in length.

In an example, the miRNA species are chimeric oligonucleotides that contain two or more chemically distinct regions, each made up of at least one nucleotide. These oligonucleotides typically contain at least one region of modified nucleotides that confers one or more beneficial properties (such as, for example, increased nuclease resistance, increased uptake into cells, increased binding affinity for the target) and a region that is a substrate for enzymes capable of cleaving RNA:DNA or RNA:RNA hybrids.

In an example, nucleic acids encompassed by the present disclosure are synthetic. The term "synthetic nucleic acid" means that the nucleic acid does not have a chemical structure or sequence of a naturally occurring nucleic acid. Synthetic nucleotides include an engineered nucleic acid such as a DNA or RNA molecule. It is contemplated, however, that a synthetic nucleic acid administered to a cell may subsequently be modified or altered in the cell such that its structure or sequence is the same as non-synthetic or naturally occurring nucleic acid, such as a mature miRNA sequence. For example, a synthetic nucleic acid may have a sequence that differs from the sequence of a precursor miRNA, but that sequence may be altered once in a cell to be the same as an endogenous, processed miRNA. Consequently, it will be understood that the term "synthetic miRNA" refers to a "synthetic nucleic acid" that functions in a cell or under physiological conditions as a naturally occurring miRNA. In another example, the nucleic acid structure can also be modified into a locked nucleic acid (LNA) with a methylene bridge between the 2' Oxygen and the 4' carbon to lock the ribose in the 3'-endo (North) conformation in the A- type conformation of nucleic acids (Lennox et al 2011; Bader et al 2011). In the context of miRNAs, this modification can significantly increase both target specificity and hybridization properties of the molecule.

Nucleic acids for use in the methods disclosed herein can be designed using routine methods as required. For example, in the context of inhibitory oligonucleotides, target segments of 5, 6, 7, 8, 9, 10 or more nucleotides in length

comprising a stretch of at least five (5) consecutive nucleotides within the seed sequence, or immediately adjacent thereto, are considered to be suitable for targeting a gene. Exemplary target segments can include sequences that comprise at least the 5 consecutive nucleotides from the 5'-terminus of one of the seed sequence (the remaining nucleotides being a consecutive stretch of the same RNA beginning immediately upstream of the 5'-terminus of the seed sequence and continuing until the nucleic acid contains about 5 to about 30 nucleotides). In another example, target segments are represented by RNA sequences that comprise at least the 5 consecutive nucleotides from the 3'-terminus of one of the seed sequence (the remaining nucleotides being a consecutive stretch of the same RNA beginning immediately downstream of the 3 '-terminus of the target segment and continuing until the nucleic acid contains about 5 to about 30 nucleotides). The term "seed sequence" is used in the context of the present disclosure to refer to a 6-8 nucleotide (nt) long substring within the first 8 nt at the 5 -end of the miRNA (i.e., seed sequence) that is an important determinant of target specificity. Once one or more target regions, segments or sites have been identified, inhibitory nucleic acid compounds are chosen that are sufficiently complementary to the target, i.e., that hybridize sufficiently well and with sufficient specificity (i.e., do not substantially bind to other non- target nucleic acid sequences), to give the desired effect.

Modification

Mesenchymal lineage precursor or stem cells of the present disclosure can be modified to introduce an above referenced nucleic acid. The term “introduced” is used in the context of the present disclosure to refer to the introduction of a nucleic acid into the nucleus or cytoplasm of a mesenchymal lineage precursor or stem cell according to the present disclosure.

Mesenchymal lineage precursor or stem cells are considered “modified” when a nucleic acid has been transferred into the cell by any suitable means of artificial manipulation, or where the cell is a progeny of an originally altered cell that has inherited a nucleic acid.

Terms such as “genetically altered”, “transfected”, “transduced” or “genetically transformed” can also be used interchangeably in the context of the present disclosure to refer to modified mesenchymal lineage precursor or stem cells. Mesenchymal lineage precursor or stem cells can be modified in a stable or transient fashion.

In an example, mesenchymal lineage precursor or stem cells of the present disclosure can be modified to introduce at least one, at least two, at least three, at least four, at least five, at least six, at least seven, at least eight, at least nine, a least 10, at least 15, at least 20 nucleic acids such as siRNA or miRNA. For example, mesenchymal lineage precursor or stem cells of the present disclosure can be modified to introduce KIF11 siRNA, KRAS siRNA, PLK1 siRNA or a combination thereof. For example, mesenchymal lineage precursor or stem cells of the present disclosure can be modified to introduce KIF11 siRNA. For example, the mesenchymal lineage precursor or stem cells can be modified to introduce KIF11 siRNA that binds towards the 5' end of KIF11 mRNA as shown in SEQ ID NO: 1 (NCBI reference number NM_004523.3). In an example, the KIF11 siRNA binds the KIF11 mRNA transcript between bp 800 and bp 3,600 from the 5' end. In another example, the KIF11 siRNA binds the KIF11 mRNA transcript between bp 900 and bp 3,200 from the 5' end. In another example, the KIF11 siRNA binds the KIF11 mRNA transcript between bp 900 and bp 2,500 from the 5' end. In another example, the KIF11 siRNA binds the KIF11 mRNA transcript before bp 3,200 from the 5' end. In another example, the KIF11 siRNA binds the KIF11 mRNA transcript after bp 3,600 from the 5' end. In another example, the KIF11 siRNA binds the KIF11 mRNA transcript after bp 4,000 from the 5' end. In another example, the KIF11 siRNA binds the KIF11 mRNA transcript between bp 3,600 and bp 5101 from the 5' end. In another example, the KIF11 siRNA binds the KIF11 mRNA transcript between bp 4,600 and bp 5101 from the 5' end.

In another example, mesenchymal lineage precursor or stem cells of the present disclosure can be modified to introduce KIF11_4 siRNA. In another example, mesenchymal lineage precursor or stem cells can be modified to introduce KIF11_6. In another example, mesenchymal lineage precursor or stem cells can be modified to introduce KIF11_9. In another example, mesenchymal lineage precursor or stem cells can be modified to introduce one or more of KIF11_8, KIF11_9, KIF11_6, KIF11_12, KIF11_7 or KIF11_4.

In another example, the the KIF11 siRNA binds the KIF11 mRNA transcript between bp 800 and bp 4,800 from the 5' end. In another example, the the KIF11 siRNA binds the KIF11 mRNA transcript between bp 900 and bp 4,800 from the 5' end. In another example, mesenchymal lineage precursor or stem cells can be modified to introduce one or more of KIF11_8, KIF11_9, KIF11_6, KIF11_12, KIF11_7, KIF11_4, KIF11_15 or KIF11_13. In another example, mesenchymal lineage precursor or stem cells can be modified to introduce KIF11_15 or KIF11_13.

The sequence details for the above referenced KIF11 siRNA are shown below in Table 1.

Table 1. KIF11 siRNA sequences

SEQ ID NO	KIF11 siRNA reference (#Qiagen product number)	Sequence
SEQ ID NO: 2	Hs_KIF11_8 (#SI03019793)	CTCGGGAAGCTGGAAATATAA
SEQ ID NO: 3	Hs_KIF11_9 (#SI03104038)	GAGGGCGTACAAGAACATCTA
SEQ ID NO: 4	Hs_KIF11_6 (#SI02653693)	ACGGAGGAGATAGAACGTTTA
SEQ ID NO: 5	Hs_KIF11_12 (#SI04376358)	CAGGAATTGATTAATGTACTC
SEQ ID NO: 6	Hs_KIF11_7 (#SI02653770)	GCCGATAAGATAGAAGATCAA
SEQ ID NO: 7	Hs_KIF11_4 (#SI00064855)	CTAGATGGCTTTCTCAGTATA
SEQ ID NO: 8	Hs_KIF11_15 (#SI05064353)	CAGCTTGAGCTTACATAGGTA
SEQ ID NO: 9	Hs_KIF11_13 (#SI04435655)	TAAGCGATGGATAATACCTAA

In another example, mesenchymal lineage precursor or stem cells of the present disclosure can be modified to introduce has-miR-let7b, hsa-miR-155, hsa-miR-155-inh, hsa-miR-181-B1, hsa-miR-15a, hsa-miR-16-1, hsa-miR-21, hsa-miR-34a, hsa-miR-221, hsa-miR-29a or a combination thereof. In another example, mesenchymal lineage precursor or stem cells are modified to introduce an oligonucleotide or vector expressing the same that preferentially kills the target cell compared with the mesenchymal lineage precursor or stem cell.

In another example, mesenchymal lineage precursor or stem cells are modified to introduce an oligonucleotide or vector expressing the same that kills the target cell but does not substantially affect viability of the mesenchymal lineage precursor or stem cell.

In another example, mesenchymal lineage precursor or stem cells are modified to introduce an oligonucleotide or vector expressing the same that does not kill the mesenchymal lineage precursor or stem cells before they can deliver the oligonucleotide to a target cell such as a cancer cell.

In an example, mesenchymal lineage precursor or stem cells of the present disclosure can be modified to introduce a vector expressing an above referenced nucleic acid. Numerous vectors for expression in cells are known in the art. Vector components generally include, but are not limited to, one or more of the following: a signal sequence, a sequence encoding a nucleic acid such as an oligonucleotide, an enhancer element, a promoter, and a transcription termination sequence.

Exemplary expression vectors include plasmid, phage, autonomously replicating sequence (ARS), viral, centromere, artificial chromosome, chromosome, or other structure able to express a nucleic acid in a mesenchymal lineage precursor or stem cell according to the present disclosure.

Suitable vector plasmids for transfecting into mesenchymal lineage precursor or stem cells include lipid/DNA complexes, such as those described in U.S. Pat. Nos. 5,578,475; 6,020,202; and 6,051,429. Suitable reagents for making DNA-lipid complexes include lipofectamine (Gibco/Life Technologies #11668019) and FuGENE™ 6 (Roche Diagnostics Corp. #1814443); and LipoTAXI™ (Invitrogen Corp., #204110).

In another example, mesenchymal lineage precursor or stem cells are modified to introduce an oligonucleotide using a viral expression vector. Exemplary viral expression vectors include Lentivirus, Baculovirus, Retrovirus, Adenovirus (AdV), Adeno-associated virus (AAV) including recombinant forms such as recombinant adeno-associated virus (rAAV) and derivatives thereof such as self-complementary AAV (scAAV) and non-integrating AV.

In an example, the viral vector is replication-defective. In this example, replication genes are deleted or replaced with an expression cassette with a high activity promoter. For example, in the context of AV, E1/E3 genes can be deleted or replaced. In the context of AAV, E1A and E1B genes can be deleted or replaced. Exemplary high activity promoters include CMV, EF1a, SV40, PGK1, Ubc, human beta actin, CAG, TRE, UAS and Ac5.

In an example, mesenchymal lineage precursor or stem cells are modified to introduce an oligonucleotide using an AV vector or a recombinant form thereof. Various AV serotypes may be suitable for modifying mesenchymal lineage precursor or stem cells to introduce an oligonucleotide. In an example, AV serotype 1 (AV1) is used to modify mesenchymal lineage precursor or stem cells. In another example, AV2 is used to modify mesenchymal lineage precursor or stem cells. In other examples, AV3, AV4, AV7, AV8, AV9, AV10, AV11, AV12 or AV13 is used to modify mesenchymal lineage precursor or stem cells. In another example, AV5 is used to modify mesenchymal lineage precursor or stem cells. In another example, AV6 is used to modify mesenchymal lineage precursor or stem cells.

In an example, mesenchymal lineage precursor or stem cells are modified to introduce an oligonucleotide using an AAV vector or a recombinant form thereof. Various AAV serotypes may also be suitable for modifying mesenchymal lineage precursor or stem cells to introduce an oligonucleotide.

In an example, AAV serotype 1 (AAV1) is used to modify mesenchymal lineage precursor or stem cells. In another example, AAV2 is used to modify mesenchymal lineage precursor or stem cells. In other examples, AAV3, AAV4, AAV7, AAV8, AAV9, AAV10, AAV11, AAV12 or AAV13 is used to modify mesenchymal lineage precursor or stem cells. In another example, AAV5 is used to modify mesenchymal lineage precursor or stem cells. In another example, AAV6 is used to modify mesenchymal lineage precursor or stem cells.

The optimal vector can be identified using various techniques known in the art. In an example, mesenchymal lineage precursor or stem cells can be contacted/transfected with various vectors expressing green fluorescent protein (GFP). In this example, optimal vectors can be identified based on transfection/transduction efficiency, GFP expression level, cellular tropism, and/or persistence of GFP expression.

Methods of viral transduction are known in the art (e.g. U.S. Pat. Nos. 6,723,561; 6,627,442). Various viral expression vector systems are also available from commercial suppliers such as Miltenyi Biotech (MACSductin), Sigma-Aldrich (ExpressMag) and Thermo Fisher Scientific (ViraPower).

Efficiencies of modification are rarely 100%, and it is usually desirable to enrich the population for cells that have been successfully modified. In an example, modified cells can be enriched by taking advantage of a functional feature of the new genotype. One exemplary method of enriching modified cells is positive selection using resistance to a drug such as neomycin.

In an example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of oligonucleotide to reduce viability of a target cell. In an example, the target cell is a cancer cell. Accordingly, in an example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of oligonucleotide to reduce viability of a cancer cell. In an example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of at least 0.25 nM of the inhibitory oligonucleotide to a target cell. In an example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of at least 0.3 nM of the inhibitory oligonucleotide to a target cell. In an example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of at least 0.4 nM of the inhibitory oligonucleotide to a target cell. In an example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of at least 0.5

nM of the inhibitory oligonucleotide to a target cell. In an example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of at least 1 nM of the inhibitory oligonucleotide to a target cell. In an example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of at least 3 nM of the inhibitory oligonucleotide to a target cell. In another example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of between 0.25 to 5 nM of the inhibitory oligonucleotide to a target cell. In another example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of between 0.5 to 5 nM of the inhibitory oligonucleotide to a target cell. In another example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of between 0.5 to 4 nM of the inhibitory oligonucleotide to a target cell. In another example, modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of between 0.5 to 3.5 nM of the inhibitory oligonucleotide to a target cell.

In another example, modified mesenchymal lineage precursor or stem cells comprise at least 100 nM of inhibitory oligonucleotide. In another example, modified mesenchymal lineage precursor or stem cells comprise at least 200 nM of inhibitory oligonucleotide. In another example, modified mesenchymal lineage precursor or stem cells comprise at least 300 nM of inhibitory oligonucleotide. In another example, modified mesenchymal lineage precursor or stem cells comprise at least 400 nM of inhibitory oligonucleotide. In another example, modified mesenchymal lineage precursor or stem cells comprise at least 500 nM of inhibitory oligonucleotide. In another example, modified mesenchymal lineage precursor or stem cells comprise at least 600 nM of inhibitory oligonucleotide. In another example, modified mesenchymal lineage precursor or stem cells comprise at least 700 nM of inhibitory oligonucleotide. In another example, modified mesenchymal lineage precursor or stem cells comprise at least 1 μM of inhibitory oligonucleotide.

Delivery to target cells

The present inventors have identified that mesenchymal lineage precursor or stem cells can transfer nucleic acids to target cells. Accordingly, in an example, the present disclosure encompasses methods of delivering an above referenced nucleic acid to target cells by contacting them with mesenchymal lineage precursor or stem cells

that have been modified to introduce an above referenced nucleic acid or vector expressing the same. For the avoidance of doubt the nucleic acid being delivered to a target cell is the nucleic acid introduced to the modified mesenchymal lineage precursor or stem cell.

The term "contacting" is used in the context of the present disclosure to refer to "direct" or "indirect" contact. "Direct contact" is used in the context of the present disclosure to refer to physical contact between the target cell and a modified mesenchymal lineage precursor or stem cell that facilitates transfer of a nucleic acid. For example, a target cell and a modified mesenchymal lineage precursor or stem cell can be in direct contact via a common connexin (i.e. a connexin that is expressed by both the target cell and the modified mesenchymal lineage precursor or stem cell). In this example, the common connexin facilitates transfer of the nucleic acid from the mesenchymal lineage precursor or stem cell to the target cell via a gap junction. Accordingly, in an example, contacting occurs under conditions permitting the mesenchymal lineage precursor or stem cell to form a gap junction with the target cell, whereby a nucleic acid is delivered to the target cell by traversing the gap junction. In an example, the gap junction is formed by Cx40. In another example, the gap junction is formed by Cx43. In another example, the gap junction is formed Cx45, Cx32 and/or Cx37.

"Indirect contact" is used in the context of the present disclosure to refer to delivery of a nucleic acid from a modified mesenchymal lineage precursor or stem cell to a target cell without direct contact. For example, a modified mesenchymal lineage precursor or stem cell in close proximity to a target cell may be in indirect contact with the target cell. In an example, a modified mesenchymal lineage precursor or stem cell in indirect contact with a target cell can deliver a nucleic acid to the target cell via exosomes.

In another example, a modified mesenchymal lineage precursor or stem cell in direct contact with a target cell can deliver a nucleic acid to the target cell via a common connexin and indirectly via exosomes.

Target cells receiving a nucleic acid from a modified mesenchymal lineage precursor or stem cell are not particularly limited so long as they can directly or indirectly contacted by the modified mesenchymal lineage precursor or stem cell to facilitate transfer of a nucleic acid. Exemplary target cells include cells in syncytial tissue. The term "syncytial" is used in the context of the present disclosure to refer to tissue that is made up of cells interconnected by specialized membrane with gap junctions, which are synchronized electrically in an action potential. Exemplary

syncytial cells include cardiac myocytes, smooth muscle cells, epithelial cells, connective tissue cells and syncytial cancer cells. In an example, the target cell is an immune cell. For example, the target cell can be a white blood cell.

In an example, the target cell is a cancer cell. For example, the target cell can be a pancreatic cancer cell. In another example, the target cell can be a lung cancer cell. In another example, the target cell can be a cervical cancer cell. In another example, the target cell can be a colorectal cancer cell. In another example, the target cell can be a liver cancer cell. In another example, the target cell can be an bone cancer cell. In another example, the target cell can be an osteosarcoma cell. In another example, the target cell can be a prostate cancer cell. In another example, the target cell can be a melanoma cell.

In another example, a target cell has a common connexin with the modified mesenchymal lineage precursor or stem cell. In an example, a target cell expresses Cx40. In another example, a target cell expresses Cx43. In another example, a target cell expresses Cx45, Cx32 and/or Cx37.

Delivery of nucleic acids from a modified mesenchymal lineage precursor or stem cell to a target cell can be facilitated *in vitro* or *in vivo*. In an example, delivery of nucleic acids from a modified mesenchymal lineage precursor or stem cell to a target cell can be facilitated *in vitro* by co-culturing the modified mesenchymal lineage precursor or stem cell with the target cell. In an example, delivery of nucleic acids from a modified mesenchymal lineage precursor or stem cell to a target cell can be facilitated *in vivo* by administering the modified mesenchymal lineage precursor or stem cell to a subject. For example, mesenchymal lineage precursor or stem cells may be administered systemically, such as, for example, by intravenous, intraarterial, or intraperitoneal administration. The mesenchymal lineage precursor or stem cells may also be administered by intranasal, intramuscular or intracardiac administration. In an example, the mesenchymal lineage precursor or stem cells are administered to a site in close proximity to a target cell such as surrounding tissue. In another example, the mesenchymal lineage precursor or stem cells are administered directly into tissue comprising a target cell.

Method of Treatment

In one example, compositions according to the present disclosure can be administered for the treatment of a cancer. The term "cancer" refers to or describes the physiological condition in mammals that is typically characterized by unregulated cell growth. Examples of cancer include but are not limited to, carcinoma, lymphoma, blastoma, sarcoma, and leukemia or lymphoid malignancies. More particular examples of such cancers include, but are not limited to, squamous cell cancer (e.g. , epithelial squamous cell cancer), lung cancer including small-cell lung cancer, non-small cell lung cancer, adenocarcinoma of the lung and squamous carcinoma of the lung, cancer of the peritoneum, hepatocellular cancer, gastric or stomach cancer including gastrointestinal cancer and gastrointestinal stromal cancer, pancreatic cancer, glioblastoma, cervical cancer, ovarian cancer, liver cancer, bladder cancer, cancer of the urinary tract, hepatoma, breast cancer, colon cancer, rectal cancer, colorectal cancer, endometrial or uterine carcinoma, salivary gland carcinoma, kidney or renal cancer, prostate cancer, vulval cancer, thyroid cancer, hepatic carcinoma, anal carcinoma, penile carcinoma, melanoma, superficial spreading melanoma, lentigo maligna melanoma, acral lentiginous melanomas, nodular melanomas, multiple myeloma and B-cell lymphoma (including low grade/follicular non-Hodgkin's lymphoma (NHL); small lymphocytic (SL) NHL; intermediate grade/follicular NHL; intermediate grade diffuse NHL; high grade immunoblastic NHL; high grade lymphoblastic NHL; high grade small non-cleaved cell NHL; bulky disease NHL; mantle cell lymphoma; AIDS-related lymphoma; and Waldenstrom's Macroglobulinemia); chronic lymphocytic leukemia (CLL); acute lymphoblastic leukemia (ALL); hairy cell leukemia; chronic myeloblastic leukemia; and post-transplant lymphoproliferative disorder (PTLD), as well as abnormal vascular proliferation associated with phakomatoses, edema (such as that associated with brain tumors), Meigs' syndrome, brain, as well as head and neck cancer, and associated metastases.

In an example, the cancer has the oncogenic G12D mutation in KRAS. Those of skill in the art will appreciate that the G12D mutation is a single nucleotide point mutation which is present in codon 12. For example, the cancer can be selected from the group consisting of colon cancer, lung cancer, melanoma, liver cancer and osteosarcoma, wherein the cancer, has the oncogenic G12D mutation in KRAS.

In an example, the cancer is pancreatic cancer. In another example, the cancer is lung cancer. In another example, the cancer is cervical cancer. In another example, the cancer is colorectal cancer. In another example, the cancer is liver cancer. In

another example, the cancer is osteosarcoma. In another example, the cancer is prostate cancer. In another example, the cancer is melanoma.

In another example, cancer treated according to the present disclosure comprises cells that share a common connexin with a mesenchymal lineage precursor or stem cell according to the present disclosure. In this example, the common connexin facilitates transfer of the nucleic acid from the mesenchymal lineage precursor or stem cell to cancer cells.

In an example, the cancer comprises cells expressing Cx40. In another example, the cancer comprises cells expressing Cx43. In another example, the cancer comprises cells expressing Cx40 and Cx43.

Improved preservation and/or homing of modified Mesenchymal lineage precursor or stem cells

In one aspect, the mesenchymal lineage precursor or stem cells defined herein are treated in order to modify their cell-surface glycans. Modification of glycans on cell surface proteins such as CD44 has been shown to create E-selectin ligands which can bind to the E-selectin molecules expressed *in vivo* on microvessels at sites of inflammation. In this way, modification of cell-surface glycans on mesenchymal lineage precursor or stem cells improves homing of the mesenchymal lineage precursor or stem cells to sites of tissue damage *in vivo*.

The present inventors have also identified that glycosyltransferase mediated modification of cell-surface glycans improves cell viability post-cryopreservation (i.e. more cells are viable following a freeze-thaw cycle). Accordingly, in an example, the present disclosure encompasses a cryopreserved population of mesenchymal lineage precursor or stem cells that have been treated with a glycosyltransferase (E.C 2.4) under conditions that modified cell-surface glycans on the cells. In another example, the present disclosure encompasses a method of cryopreserving mesenchymal lineage precursor or stem cells, the method comprising: treating a population of mesenchymal lineage precursor or stem cells with a glycosyltransferase under conditions that result in modification of cell-surface glycans on the cells, and cryopreserving the cells in a composition. In another example, the present disclosure encompasses a method of producing therapeutic cells, the method comprising: treating a population of mesenchymal lineage precursor or stem cells with a glycosyltransferase under conditions that result in modification of cell-surface glycans on the cells, and cryopreserving the cells in a composition.

In an example, mesenchymal lineage precursor or stem cell "treatment" includes contacting the cells with a glycosyltransferase under conditions in which the glycosyltransferase has enzymatic activity. In this example, the glycosyltransferase modifies cell surface glycans on mesenchymal lineage precursor or stem cells. An example of cell surface glycan modification is fucosylation. In an example, CD44 is modified. In another example, CD14 is modified. In another example, one or more of CD44, CD14, CD3 and CD19 are modified.

In an example, surface glycan modification is identified using flow cytometry. In this example, modified mesenchymal lineage precursor or stem cells have a 1 log magnitude higher expression of a fucosylated cell surface glycan(s) than untreated mesenchymal lineage precursor cells. In another example, modified mesenchymal lineage precursor or stem cells have a 2 log magnitude higher expression of a fucosylated cell surface glycan(s) than untreated mesenchymal lineage precursor cells. In another example, modified mesenchymal lineage precursor or stem cells have a 3 log magnitude higher expression of a fucosylated cell surface glycan(s) than untreated mesenchymal lineage precursor cells. For example, modified mesenchymal lineage precursor or stem cells can have a 1 log magnitude higher expression of fucosylated CD14 than untreated mesenchymal lineage precursor cells. In another example, modified mesenchymal lineage precursor or stem cells have a 2 log magnitude higher expression of fucosylated CD14 than untreated mesenchymal lineage precursor cells. In another example, modified mesenchymal lineage precursor or stem cells have a 3 log magnitude higher expression of fucosylated CD14 than untreated mesenchymal lineage precursor cells.

In an example, the "treatment" includes contacting the mesenchymal lineage precursor or stem cells with a glycosyltransferase in the presence of a nucleotide sugar donor substrate. Suitable donor substrates include fucose, galactose, sialic acid, or N-acetyl glucosamine. For example, the substrate can be GDP-fucose.

For example, treatment can involve contacting a population of mesenchymal lineage precursor or stem cells with an exogenous glycosyltransferase such as a fucosyltransferase. In this example, a glycosyltransferase can be added to cell culture media or other physiologically acceptable solution comprising mesenchymal lineage precursor or stem cells. For example, mesenchymal lineage precursor or stem cells can be cultured in medium comprising a glycosyltransferase. In another example, mesenchymal lineage precursor or stem cells are suspended in culture medium comprising a glycosyltransferase. For example, mesenchymal linage precursor or stem cells can be dissociated from culture and resuspending in a suitable medium comprising

a glycosyltransferase. In an example, cells can be dissociated using Ethylenediaminetetraacetic acid (EDTA). In another example, cells can be dissociated using a protease such as trypsin alone oFr in combination with EDTA.

In an example, the cell culture medium comprises at least 1.8 μg of glycosyltransferase. In another example, the cell culture medium comprises at least 2.0 μg of glycosyltransferase. In another example, the cell culture medium comprises at least 2.5 μg of glycosyltransferase. In another example, the cell culture medium comprises between 2 and 15 μg of glycosyltransferase. In another example, the cell culture medium comprises between 2 and 10 μg of glycosyltransferase. In another example, the cell culture medium comprises between 2 and 5 μg of glycosyltransferase. In an example, the cell culture medium comprises at least 1.8 μg of fucosyltransferase. In another example, the cell culture medium comprises at least 2.0 μg of fucosyltransferase. In another example, the cell culture medium comprises at least 2.5 μg of fucosyltransferase. In another example, the cell culture medium comprises between 2 and 15 μg of fucosyltransferase. In another example, the cell culture medium comprises between 2 and 10 μg of fucosyltransferase. In another example, between 2 and 5 μg of fucosyltransferase is added to the cell culture media. In these examples, the glycosyltransferase can be provided in 30 μl reaction volume to around $5x10^5$ mesenchymal lineage precursor or stem cells.

For example, mesenchymal lineage precursor or stem cells can be treated with exogenous glycosyltransferase in a process known as exofucosylation. In this embodiment the glycosyltransferase may be provided in a physiologically acceptable solution that has low levels of divalent metal co-factors. In various embodiments, the physiologically acceptable solution is buffered. The physiologically acceptable solution may be, for example, Hank's Balanced Salt Solution, Dulbecco's Modified Eagle Medium, a Good's buffer (see N. E. Good, G. D. Winget, W. Winter, T N. Conolly, S. Izawa and R. M. M. Singh, Biochemistry 5, 467 (1966); N. E. Good, S. Izawa, Methods Enzymol. 24, 62 (1972) such as a HEPES buffer, a 2-Morpholinoethanesulfonic acid (MES) buffer, phosphate buffered saline (PBS).

In an example, the physiologically acceptable solution is substantially free of glycerol.

In another example, mesenchymal lineage precursor or stem cells are treated with a glycosyltransferase by modifying the cells to express a glycosyltransferase. For example, the glycosyltransferase can be generated intracellularly by the mesenchymal lineage precursor or stem cell. In this embodiment, a nucleic acid molecule(s) which encodes a glycosyltransferase is introduced into the mesenchymal lineage precursor or

stem cell. The glycosyltransferase is then expressed by the mesenchymal lineage precursor or stem cells to effect modification of its surface glycans.

Mesenchymal lineage precursor or stem cells are considered "genetically modified to express a glycosyltransferase" when nucleic acid encoding a glycosyltransferase has been transferred into the cell by any suitable means of artificial manipulation, or where the cell is a progeny of an originally altered cell that carries the nucleic acid encoding the glycosyltransferase. Cells can be stably or transiently modified to express a glycosyltransferase.

In an example, expression of the glycosyltransferase in genetically modified mesenchymal lineage precursor or stem cells results in enhanced retention of the cells at a site of inflammation *in vivo*. For example, genetically modified mesenchymal lineage precursor or stem cells may be retained at a tumour or metastasis thereof. In another example, genetically modified mesenchymal lineage precursor or stem cells may be retained at a site of organ transplant rejection. In another example, genetically modified mesenchymal lineage precursor or stem cells may be retained at a site of injury such as an infarcted heart. Various methods are available for determining whether a genetically modified mesenchymal lineage precursor or stem cell is retained at a site of inflammation *in vivo*. In an example, cells are imaged *in vivo* using a radiotracer or other suitable label.

Mesenchymal lineage precursor or stem cells can be genetically modified using various methods known in the art. In an example, mesenchymal lineage precursor or stem cells are treated with a viral vector *in vitro*. Genetically modified viruses have been widely applied for the delivery of nucleic acids into cells. Exemplary viral vectors for genetic modification of the cells described herein include retroviral vectors such as gamma retroviral vectors, lentivirus, murine leukemia virus (MLV or MuLV), and adenovirus. For example, virus can be added to mesenchymal lineage precursor or stem cell culture medium. Non-viral methods may also be employed. Examples include plasmid transfer and the application of targeted gene integration through the use of integrase or transposase technologies, liposome or protein transduction domain mediated delivery and physical methods such as electroporation.

Efficiencies of genetic modification are rarely 100%, and it is usually desirable to enrich the population for cells that have been successfully modified. In an example, modified cells can be enriched by taking advantage of a functional feature of the new genotype. One exemplary method of enriching modified cells is positive selection

using resistance to a drug such as neomycin or colorimetric selection based on expression of lacZ.

In various embodiments, the mesenchymal lineage precursor or stem cell is contacted with more than one glycosyltransferase and its appropriate donor substrate (e.g. sugar). For example, the cell is contacted with two glycosyltransferases simultaneously, or sequentially, each adding a distinct monosaccharide in appropriate linkage to the (extending) core glycan structure. In another example, genetically modified cells express two glycosyltransferases.

In one embodiment, treated mesenchymal lineage precursor or stem cells expresses CD44, e.g., alpha(2,3)sialyated CD44. In another embodiment, the mesenchymal lineage precursor or stem cell does not express CD34 or PSGL-1. In an example, treated mesenchymal lineage precursor or stem cell binds E-selectin and or L-selectin. In an example, the modified mesenchymal lineage precursor or stem cell does not bind P-selectin.

In another example, CD14 is fucosylated on treated mesenchymal lineage precursor or stem cells. In another example, CD14 and CD3 are fucosylated on treated mesenchymal lineage precursor or stem cells.

In one embodiment, the glycosyltransferase is capable of transferring 1.0 mmole of sugar per minute at pH 6.5 at 37°C.

In an example, the glycosyltransferase is a fucosyltransferase (catalyses transfer of L-fucose sugar). In another example, the glycosyltransferase is an alpha 1,3 fucosyltransferase, e.g., an alpha 1,3 fucosyltransferase III, alpha 1,3 fucosyltransferase IV, an alpha 1,3 fucosyltransferase VI, an alpha 1,3 fucosyltransferase VII, an alpha 1,3 fucosyltransferase IX, an alpha 1,3 fucosyltransferase X, an alpha 1,3 fucosyltransferase XI). For example, cells can be treated with alpha 1,3 fucosyltransferase VII. In another example, cells can be treated with alpha 1,3 fucosyltransferase VI. In these examples, fucosylation of mesenchymal lineage precursor or stem cells can be identified by detecting an increase in the ability of treated cells to bind to a selectin such as E-selectin and/or an increase in the reactivity of treated cells with an antibody known in the art to bind to sLeX including, but not limited to, the HECA-452.

In another example, the glycosyltransferase is a galactosyltransferase (catalyses the transfer of galactose). In another example, the glycosyltransferase is a sialyltransferase (catalyses the transfer of sialic acid).

Cellular Compositions

In performing the methods of the present disclosure mesenchymal lineage precursor or stem cells can be administered in the form of a composition.

Exemplary compositions according to the present disclosure can comprise mesenchymal lineage precursor or stem cells that have been modified to introduce an siRNA or a miRNA or a vector expressing the same. For example, a composition according to the present disclosure can comprise mesenchymal lineage precursor or stem cells that have been modified to introduce KIF11 siRNA, KRAS siRNA, PLK1 siRNA, a combination thereof or a vector expressing the same.

In an example, the composition comprises stem cells that have been modified to introduce KIF11 siRNA, wherein the KIF11 siRNA binds the KIF11 mRNA transcript between bp 800 and bp 3,600 from the 5' end. In another example, the composition comprises stem cells that have been modified to introduce KIF11 siRNA, wherein the siRNA binds the KIF11 mRNA transcript between bp 900 and bp 3,200 from the 5' end. In another example, the composition comprises stem cells that have been modified to introduce KIF11 siRNA, wherein the siRNA binds the KIF11 mRNA transcript between bp 900 and bp 2,500 from the 5' end. In another example, the composition comprises stem cells that have been modified to introduce KIF11 siRNA, wherein the siRNA binds the KIF11 mRNA transcript before bp 3,200 from the 5' end. In these examples, the KIF11 mRNA transcript is as shown in SEQ ID NO: 1 (NCBI reference number NM_004523.3).

In an example, the composition comprises stem cells that have been modified to introduce KIF11_4. In another example, the composition comprises stem cells that have been modified to introduce KIF11_6. In another example, the composition comprises stem cells that have been modified to introduce KIF11_9. In another example, the composition comprises stem cells that have been modified to introduce one or more of KIF11_8, KIF11_9, KIF11_6, KIF11_12, KIF11_7 or KIF11_4.

In another example, the composition comprises stem cells that have been modified to introduce one or more KIF11 siRNA, wherein the siRNA binds the KIF11 mRNA transcript between bp 800 and bp 4,800 from the 5' end. In another example, the composition comprises stem cells that have been modified to introduce one or more KIF11 siRNA, wherein the siRNA binds the KIF11 mRNA transcript between bp 900 and bp 4,800 from the 5' end. In another example, the composition comprises stem cells that have been modified to introduce one or more KIF11 siRNA, wherein the siRNA binds the KIF11 mRNA transcript before bp 3,600 from the 5' end. In another example, the composition comprises stem cells that have been modified to introduce one or more KIF11 siRNA, wherein the siRNA binds the KIF11 mRNA transcript after

bp 3,600 from the 5' end. In another example, the composition comprises stem cells that have been modified to introduce one or more KIF11 siRNA, wherein the siRNA binds the KIF11 mRNA transcript after bp 4,000 from the 5' end. In another example, the KIF11 siRNA binds the KIF11 mRNA transcript between bp 3,600 and bp 5101 from the 5' end. In another example, the composition comprises stem cells that have been modified to introduce one or more KIF11 siRNA, wherein the siRNA binds the KIF11 mRNA transcript between bp 4,600 and bp 5101 from the 5' end.

In another example, the composition comprises stem cells that have been modified to introduce one or more of KIF11_8, KIF11_9, KIF11_6, KIF11_12, KIF11_7, KIF11_4, KIF11_15 or KIF11_13. In another example, the composition comprises stem cells that have been modified to introduce one or both of KIF11_15 or KIF11_13.

The sequence details for the above referenced KIF11 siRNA are shown above in Table 1.

In other examples, compositions according to the present disclosure can comprise mesenchymal lineage precursor or stem cells modified to introduce an above referenced miRNA, a combination thereof or a vector expressing the same. For example, compositions according to the present disclosure can comprise mesenchymal lineage precursor or stem cells modified to introduce hsa-miR-let7b, hsa-miR-155, hsa-miR-155-inh, hsa-miR-181-B1, hsa-miR-15a, hsa-miR-16-1, hsa-miR-21, hsa-miR-34a, hsa-miR-221, hsa-miR-29, a combination thereof or a vector expressing the same.

In another example, compositions according to the present disclosure can comprise mesenchymal lineage precursor or stem cells modified to introduce an inhibitory oligonucleotide or vector expressing the same that does not kill the mesenchymal lineage precursor or stem cells before they can deliver the at least one type of inhibitory oligonucleotide to a target cell such as a cancer cell.

In one example, such a composition comprises a pharmaceutically acceptable carrier and/or excipient.

The terms "carrier" and "excipient" refer to compositions of matter that are conventionally used in the art to facilitate the storage, administration, and/or the biological activity of an active compound (see, e.g., Remington's Pharmaceutical Sciences, 16th Ed., Mac Publishing Company (1980). A carrier may also reduce any undesirable side effects of the active compound. A suitable carrier is, for example, stable, e.g., incapable of reacting with other ingredients in the carrier. In one example, the carrier does not produce significant local or systemic adverse effect in recipients at the dosages and concentrations employed for treatment.

Suitable carriers for the present disclosure include those conventionally used, e.g., water, saline, aqueous dextrose, lactose, Ringer's solution, a buffered solution, hyaluronan and glycols are exemplary liquid carriers, particularly (when isotonic) for solutions. Suitable pharmaceutical carriers and excipients include starch, cellulose, glucose, lactose, sucrose, gelatin, malt, rice, flour, chalk, silica gel, magnesium stearate, sodium stearate, glycerol monostearate, sodium chloride, glycerol, propylene glycol, water, ethanol, and the like.

In another example, a carrier is a media composition, e.g., in which a cell is grown or suspended. Such a media composition does not induce any adverse effects in a subject to whom it is administered.

Exemplary carriers and excipients do not adversely affect the viability of a cell and/or the ability of a cell to treat or prevent disease.

In one example, the carrier or excipient provides a buffering activity to maintain the cells and/or soluble factors at a suitable pH to thereby exert a biological activity, e.g., the carrier or excipient is phosphate buffered saline (PBS). PBS represents an attractive carrier or excipient because it interacts with cells and factors minimally and permits rapid release of the cells and factors, in such a case, the composition of the disclosure may be produced as a liquid for direct application to the blood stream or into a tissue or a region surrounding or adjacent to a tissue, e.g., by injection.

The cellular compositions described herein may be administered alone or as admixtures with other cells. The cells of different types may be admixed with a composition of the disclosure immediately or shortly prior to administration, or they may be co-cultured together for a period of time prior to administration.

In one example, the composition comprises an effective amount or a therapeutically effective amount of cells. For example, the composition comprises about $1x10^5$ cells to about $1x10^9$ cells or about $1.25x10^3$ cells to about $1.25x10^7$ cells. The exact amount of cells to be administered is dependent upon a variety of factors, including the age, weight, and sex of the subject, and the extent and severity of the disorder being treated.

Exemplary dosages include at least about 1.2×10^8 to about 8×10^{10} cells, such as between about 1.3×10^8 to about 8×10^9 cells, about 1.4×10^8 to about 8×10^8 cells, about 1.5×10^8 to about 7.2×10^8 cells, about 1.6×10^8 to about 6.4×10^8 cells, about 1.7×10^8 to about 5.6×10^8 cells, about 1.8×10^8 to about 4.8×10^8 cells, about 1.9×10^8 to about 4.0×10^8 cells, about 2.0×10^8 to about 3.2×10^8 cells, about 2.1×10^8 to about 2.4×10^8 cells. For example, a dose can include at least about 1.5×10^8 cells. For example, a dose can include at least about 2.0×10^8 cells.

Put another way, exemplary doses include at least about 1.5 x 10^6 cells/kg (80kg subject). In an example, a dose can include at least about 2.5 x 10^6 cells/kg. In other examples, a dose can comprise between about 1.5 x 10^6 to about 1x10^9 cells/kg, about 1.6 x 10^6 to about 1 x 10^8 cells/kg, about 1.8 x 10^6 to about 1 x 10^7 cells/kg, about 1.9 x 10^6 to about 9 x 10^6 cells/kg, about 2.0 x 10^6 to about 8 x 10^6 cells/kg, about 2.1 x 10^6 to about 7 x 10^6 cells/kg, about 2.3 x 10^6 to about 6 x 10^6 cells/kg, about 2.4 x 10^6 to about 5 x 10^6 cells/kg, about 2.5 x 10^6 to about 4 x 10^6 cells/kg, about 2.6 x 10^6 to about 3 x 10^6 cells/kg.

In an example, modified mesenchymal lineage precursor or stem cells comprise at least about 5%, at least about 10%, at least about 15%, at least about 20%, at least about 25%, at least about 30%, at least about 35%, at least about 40%, at least about 45%, at least about 50%, at least about 55%, at least about 60%, at least about 65%, at least about 70%, at least about 75%, at least about 80%, at least about 85%, at least about 90%, at least about 95%, at least about 99% of the cell population of the composition.

Compositions of the disclosure may be cryopreserved. Cryopreservation of mesenchymal lineage precursor or stem cells can be carried out using slow-rate cooling methods or 'fast' freezing protocols known in the art. Preferably, the method of cryopreservation maintains similar phenotypes, cell surface markers and growth rates of cryopreserved cells in comparison with unfrozen cells.

The cryopreserved composition may comprise a cryopreservation solution. The pH of the cryopreservation solution is typically 6.5 to 8, preferably 7.4.

The cyropreservation solution may comprise a sterile, non-pyrogenic isotonic solution such as, for example, PlasmaLyte ATM. 100 mL of PlasmaLyte ATM contains 526 mg of sodium chloride, USP (NaCl); 502 mg of sodium gluconate (C6H11NaO7); 368 mg of sodium acetate trihydrate, USP (C2H3NaO2•3H2O); 37 mg of potassium chloride, USP (KCl); and 30 mg of magnesium chloride, USP (MgCl2•6H2O). It contains no antimicrobial agents. The pH is adjusted with sodium hydroxide. The pH is 7.4 (6.5 to 8.0).

The cryopreservation solution may comprise Profreeze™. The cryopreservation solution may additionally or alternatively comprise culture medium, for example, αMEM.

To facilitate freezing, a cryoprotectant such as, for example, dimethylsulfoxide (DMSO), is usually added to the cryopreservation solution. Ideally, the cryoprotectant should be nontoxic for cells and patients, nonantigenic, chemically inert, provide high survival rate after thawing and allow transplantation without washing. However, the

most commonly used cryoprotector, DMSO, shows some cytotoxicity . Hydroxylethyl starch (HES) may be used as a substitute or in combination with DMSO to reduce cytotoxicity of the cryopreservation solution.

The cryopreservation solution may comprise one or more of DMSO, hydroxyethyl starch, human serum components and other protein bulking agents. In one example, the cryopreserved solution comprises about 5% human serum albumin (HSA) and about 10% DMSO. The cryopreservation solution may further comprise one or more of methycellulose, polyvinyl pyrrolidone (PVP) and trehalose.

In one embodiment, cells are suspended in 42.5% ProfreezeTM/50% αMEM/7.5% DMSO and cooled in a controlled-rate freezer.

The cryopreserved composition may be thawed and administered directly to the subject or added to another solution, for example, comprising hyaluronic acid. Alternatively, the cryopreserved composition may be thawed and the mesenchymal lineage precursor or stem cells resuspended in an alternate carrier prior to administration.

In an example, the cellular compositions described herein may be administered as a single dose. In another example, cellular compositions are administered over multiple doses. For example, at least 2, at least 3, at least 4, at least 5, at least 6, at least 7, at least 8, at least 9, at least 10 doses.

In one example, the mesenchymal lineage precursor or stem cells are isolated, or isolated and enriched, and culture expanded *ex vivo or in vitro* prior to cryopreservation. In another example, the mesenchymal lineage precursor or stem cells are isolated, or isolated and enriched, cryopreserved, thawed and subsequently culture expanded *ex vivo or in vitro*. In yet another example, the mesenchymal lineage precursor or stem cells are culture expanded prior to and after cryopreservation. For example, mesenchymal lineage precursor or stem cells can be defrosted and then culture expanded. Various methods of mesenchymal lineage precursor or stem cell culture are known in the art. In an example, mesenchymal lineage precursor or stem cells are culture expanded in a serum free medium prior to administration. For example, mesenchymal lineage precursor or stem cells can be passaged at least once, twice, three, four, five, six, seven, eight, nine, 10 or more times prior to administration.

Mesenchymal precursor cells may be administered systemically, such as, for example, by intravenous, intraarterial, or intraperitoneal administration. The mesenchymal precursor cells may also be administered by intranasal, intramuscular or intracardiac administration. In an example, the mesenchymal precursor cells are administered directly into a subject's tumour.

EXAMPLES

EXAMPLE 1 – Cellular Transfer of MiRs/siRNAs by Mesenchymal Precursor Cells

Human mesenchymal precursor cell (MPC) expression of connexins was assessed via Immunohistochemistry and Western Blot. Data generated demonstrate that MPCs express functional Cx43 and Cx40 and make functional gap junctions with target cell types that also express a common connexin (Figure 1A). Human mesenchymal stem cell (MSC) expression of connexins was also assessed via Immunohistochemistry and Western Blot. Punctate Cx43 and Cx40 staining was observed on hMSCs (Figure 1B). Functional gap junctions were also demonstrated in MPCs using patch clamp technique (Figure 2). Various tumour cells also express Cx43 (Figure 3) and Cx40 (data not shown).

MPC mediated delivery of nucleic acids to target cells via gap junctional transfer was investigated. Lipid based loading of MPCs was demonstrated with fluorescently labelled MiRs. High level intracellular loading of miRs and siRNAs was achieved using lipid based transection (Lipofectamine™; Thermo Fisher Scientific; Figure 4).

Experiments were also conducted to show transfer of Mir16 to HeLa-Cx43 and PANC-1 tumour cells. MPCs transfected with Cy-5 Mir16 (Red; 100nM) were co-cultured for 24 hours with GFP expressing HeLa-Cx43 and GFP-expressing PANC-1 cells. Figure 5A shows transfer of the Cy-5-Mir16 to both the HeLa cells (A) and the PANC-1 cells (B). Figure 5B shows transfer of KIF11 siRNA to GFP expressing PANC-1 cells.

A co-culture assay was subsequently established to measure anti-proliferative and cytotoxic effects of MPC-loaded miRs and siRNAs on tumour cell lines. Results of co-culture of KIF11 siRNA loaded MPCs (100 nM) with PANC-1 are shown in Figure 6A. Results of co-culture of KIF11 siRNA loaded MPCs (500 nM) with PANC-1 are shown in Figure 6B. These results show that PANC-1 cell death is induced in the presence of KIF11 siRNA loaded MPCs (100 nM) and increased in the presence of a higher dose (500 nM) of KIF11 siRNA.

Results of co-culture of KIF11 siRNA loaded MPCs (500 nM) with SNUC2A cells are shown in Figure 7. These results show that MPC transfer of KIF11 induces SNUC2A tumor cell death.

Results of co-culture of KIF11 siRNA loaded MPCs (500 nM) with SAOS2 cells are shown in Figure 8 (A) and (B). These results show that MPC transfer of KIF11 induces SAOS2 tumor cell death.

Co-culture of Mir-16 loaded MPCs (100 nM) with PANC-1 tumour cell lines also induced tumour cell death (Figure 9A). Co-culture of Mir-16 loaded MPCs with unloaded MPCs inhibited MPC proliferation but did not significantly affect viability of unloaded MPCs (Figure 9B).

CRISPR/CAS9 mediated knockout of Cx43 was subsequently performed in PANC-1 cells to assess the role of Gap junctions in the transfer of nucleic acids from MPCs to target cells.

Co-culture of let7b loaded MPCs (500 nM) with PANC-1 cells inhibited PANC-1 cell growth. However, growth of Cx43 knockout PANC-1 cells was not significantly affected when co-cultured with the let7b loaded MPCs, suggesting transfer of let7b is occurring through Cx43 (Figure 10A). Figure 10B shows that direct transfection of let7b in MPCs resulted in decreased cell proliferation but not increased cell death as is observed with the PANC-1 cells.

Co-culture of KIF11 siRNA loaded MPCs with PANC-1 cells induced cell death. Reduced levels of cell death were observed in Cx43 knockout PANC-1 cells when co-cultured with KIF11 siRNA loaded MPCs. However, MPC directed inhibition of cell growth was surprisingly maintained in Cx43 knockout PANC-1 cells (Figures 11A and 12). This suggests that transfer of the siRNA is occurring not only via connexin 43, but also through another mechanism such as, for example, a different connexin (e.g. Cx40, Cx45, Cx30.3, Cx31 or Cx31.1) or via formation of exosomes. Figure 11B indicates that direct transfection of KIF11 siRNA in MPCs resulted in decreased cell proliferation but not increased cell death as is observed with the PANC-1 cells.

Co-culture of Mir-34a loaded MPCs with PANC-1 cells also induced cell death. Reduced levels of cell death were observed in Cx43 knockout PANC-1 cells when co-cultured with Mir-34a loaded MPCs. However, MPC directed inhibition of cell growth was surprisingly maintained in Cx43 knockout PANC-1 cells (Figure 13A). Again, this suggests that transfer of the Mir-34a is occurring not only via connexin 43, but also through another mechanism such as, for example, a different connexin or via formation of exosomes. Figure 13B indicates that direct transfection of Mir-34a in MPCs resulted in decreased cell proliferation but not increased cell death as is observed with the PANC-1 cells.

Effects of miRNAs and siRNAs on cell death and cell proliferation were assessed in MPCs, PANC-1, PC3 and SOAS2 cells. Cells were plated at ~25% confluency on 96-well or 3.5 cm cell culture plates and transfected with 100 nM siRNA or miR-mimic using Lipofectamine RNAiMax as a transfection reagent. Effects of hsa-miR-15A, hsa-miR-16-1, hsa-miR-34a, hsa-miR-155, hsa-let7b, hs_KIF11_4 siRNA and hs_PLK1_2 siRNA were assessed. Effects of siRNAs and miRNAs on cell death and cell growth were measured 5 days after transfection (Figure 14 - 17). Comparative analysis of cell death and WST Assays shown in Figures 18 and 19. These data show that transfected miRs and siRNAs can kill and/or inhibit growth of various cancer cells.

EXAMPLE 2: *In vivo* efficacy of MPC based delivery of siRNA

Six mice were orthotopically implanted in the prostate with 0.5 million PC3 cells stably expressing luciferase, with or without 0.5 million MPC loaded with 500 nM siRNA against KIF 11 (24 hours prior). Volume of the injections was 60ul in PBS. One week later, mice were injected with lucifern and then imaged with the IVIS imaging system to visualise the tumors (Figure 20 A and B). Thirty five days later, MPC-treated mice were injected with 1 million MPC loaded with 500 nM siRNA against KIF11. Control mice were injected with saline. As shown in Figure 20C, median survival times were 38 days for the control group and 57 days for the MPC-treated group, P==0.0067 (Log-Rank test).

These results show that KIF11 loaded MPCs significantly increased survival time in a mouse PC3 prostate xenograft tumour model.

EXAMPLE 3: MPC based delivery of siRNA and dose dependent reduction in cell viability

PANC-1 cells were co-cultured with KIF11 siRNA loaded MPCs (500 nM KIF11). Co-culture for 6 days reduced viability of PANC-1 cells by 77% (Figure 21). About an 80% decrease in cell growth rate was also observed (Figure 22).

PANC-1 cells were then directly transfected with increasing amounts of KIF11 siRNA (0 – 3.3 nM). Viability of PANC-1 cells was measured 6 days after transfection. A dose dependent reduction in cell viability was observed following direct transfection (Figure 23). Notably, maximum reduction in PANC-1 viability at 3.3 nM was 76%, similar to that seen after co-culture with 500nM-transfected MPC. As shown in Figures 6(a) and 6(b), MPCs transfected with 500nM KIF11 siRNA inhibited PANC-1 proliferation by more than 30% relative to MPCs transfected with 100nM KIF11 siRNA. These results are consistent with a step-wise inhibition of PANC-1

viability by MPCs transfected with concentrations of KIF11 siRNA from 100 to 500nM. Since the effect of direct oligonucleotide inhibition of PANC-1 viability was reduced by approximately 30% from 3.3 to 0.5nM, we conclude that MPC mediated transfer of the oligonucleotide to target tumour cells occurs at sufficient levels to result in a concentration of oligonucleotide inside the tumour cell of 0.5-3.3nM.

In a further experiment, cultures of MPC or PANC-1 cells were directly transfected with increasing amounts of KIF11 siRNA (0 – 500 nM). Dose response curves for PANC-1 cells are shown in Figure 24 and MPCs in Figure 25.

PC3 cells were also co-cultured with KIF11 siRNA loaded MPCs (500 nM KIF11). Co-culture for 6 days reduced cell growth rate by around 55% (Figure 26). PC3 cells were then directly transfected with increasing amounts of KIF11 siRNA (0 – 500 nM). Dose response curves are shown in Figure 27.

EXAMPLE 4: MPC loading and viability

MPCs from three donors were transfected with KIF11-FITC-siRNA (0–1000nM) using lipofectamine RNAiMax transfection reagent (15 µl/well). Accumulation of florescent label in MPCs 24 hours post transfection was then assayed by flow cytometry (Figure 28). The optimal concentration for siRNA loading into MPCs was determined to be ~500nM.

MPCs loaded with KIF11-FITC-siRNA were then co-cultured with PANC-1 or PC3 cells. Transfer of siRNA from MPCs loaded with KIF11-FITC-siRNA to cancer cells was measured 24 hours after co-culture. Within 24 hours, approximately 10% of siRNA was transferred to cancer cells with transfer showing little variability with respect to MPC donor (Figure 29).

MPCs and PANC-1 cells were then cultured in 6 well plates at 1.3 x 10^5 cells/well. Cells were transfected with siRNA at different concentrations (0-500 nM final) using lipofectamine RNAiMax transfection reagent (15 µl/well). RNA was harvested from cells 2 days post-transfection and gene expression was assessed using qRT-PCR (Taqman gene expression assays; Applied Biosystems; s03929097_g1 for housekeeping gene glyceraldehyde-3-phosphate dehydrogenase (GAPDH) and Hs00189698_m1 for KIF11). Figure 30 and 31 shows that KIF11 siRNA species retain their function in MPC and tumour cells (i.e. siRNA that is transfected into and transferred from MPCs to tumour cells is able to reduce KIF11 mRNA).

EXAMPLE 5: Effects of KIF11 siRNAs on MPC and Tumor cell viability

MPC, SJSA1, PANC-1, PC3, SAOS2 and SNUC2a cells were plated at 25 – 50% confluency in 96 well plates (n = 3 wells per treatment group). Cells were transfected with 100 nM of either KIF11 siRNA or control siRNA (Lipofectamine; RNAiMax; siRNA sequence details shown above in Table 1; KIF11 sequence target summarized in Figure 32). Cell viability was measured 5 days post transfection and viability data was normalized using control cell data (Figure 33). Surprisingly, KIF11 siRNA targeted towards 5' end of the KIF11 mRNA transcript had greater effect of reducing tumour cell viability, in particular in PANC-1 and SAOS2 lines. For example, KIF11_13 and KIF11_15 which target the 3' end of KIF11 mRNA transcript had relatively low effect on tumor cell viability compared to other KIF11 siRNA tested (Figure 33). It was also surprising to note that KIF11_4, KIF11_6 and KIF11_9 siRNAs consistently reduced tumor cell viability while having minimal effect on MPC viability (Figure 33).

EXAMPLE 6: Effects of KIF11 siRNAs on gene expression in MPCs and MSCs

MPCs and MSCs were plated in 6 well plates at 1.25 x 10^6 cells/well and transfected with KIF11 siRNA (100 nM) using lipofectamine. Control cells were treated with lipofectamine alone. RNA was harvested 48 hours after cell treatment and global gene expression was analysed using human Clariom S Assay (Affymetrix). Gene expression analysis indicated that MPCs and MSCs are similar in regards to genes that are downregulated by the KIF-11 siRNA (Figure 34; top 10 down regulated genes). KIF11 siRNA reduced KIF11 mRNA levels around 18 fold in MPCs and around 20 fold in MSCs. Looking at the top 10 downregulated genes, FSTL1:MIR198, ALDH9A1, ELK3 & SYPL1 mRNAs were also reduced in both MPCs and MSCs. Taken together with the findings outlined above, in particular Example 1 which notes that both MPCs and MSCs express Cx43 and Cx40, the gene expression analysis further supports that both MPCs and MSCs can be used as vehicles for oligonucleotide transfer to cancer cells.

It will be appreciated by persons skilled in the art that numerous variations and/or modifications may be made to the disclosure as shown in the specific embodiments without departing from the spirit or scope of the disclosure as broadly described. The present embodiments are, therefore, to be considered in all respects as illustrative and not restrictive.

The present application claims priority from 62/589,764 filed 22 November 2017, the disclosures of which are incorporated herein by reference.

All publications discussed above are incorporated herein in their entirety.

Any discussion of documents, acts, materials, devices, articles or the like which has been included in the present specification is solely for the purpose of providing a context for the present disclosure. It is not to be taken as an admission that any or all of these matters form part of the prior art base or were common general knowledge in the field relevant to the present disclosure as it existed before the priority date of each claim of this application.

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CLAIMS

1. A method of delivering an oligonucleotide into a target cell, the method comprising contacting a target cell with a mesenchymal lineage precursor or stem cell expressing one or more of the markers selected from the group consisting of α1, α2, α3, α4 and α5, αv, β1 and β3, wherein said cell has been modified to introduce an oligonucleotide or a vector expressing the oligonucleotide.

2. A method of treating cancer in a subject, the method comprising administering to the subject a composition comprising mesenchymal lineage precursor or stem cells expressing one or more of the markers selected from the group consisting of α1, α2, α3, α4 and α5, αv, β1 and β3, wherein said cells are modified to introduce an oligonucleotide or a plasmid expressing the oligonucleotide.

3. A composition comprising mesenchymal lineage precursor or stem cells expressing one or more of the markers selected from the group consisting of α1, α2, α3, α4 and α5, αv, β1 and β3, wherein said cells are modified to introduce an inhibitory oligonucleotide or comprise a vector expressing the inhibitory oligonucleotide.

4. The method or composition according to any one of claims 1 to 3, wherein modified mesenchymal lineage precursor or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of between 0.5 to 5 nM of the inhibitory oligonucleotide to a target cell.

5. The method or composition according to any one of claims 1 to 4, wherein the oligonucleotide is about 12 – 24 nucleotides in length.

6. The method or composition according to any one of claims 1 to 5, wherein the oligonucleotide is RNA.

7. The method or composition according to any one of claims 1 to 5, wherein the oligonucleotide is an antisense molecule.

8. The method or composition according to any one of claims 1 to 5, wherein the oligonucleotide is a shRNA, siRNA or miRNA.

9. The method or composition of claim 8, wherein the siRNA is KIF11 siRNA or PLK1 siRNA.

10. The method or composition according to any one of claims 7 to 9, wherein the oligonucleotide is a KIF11 siRNA which binds the KIF11 mRNA transcript as shown in SEQ ID NO: 1 between bp 800 and bp 3,600 from the 5' end.

11. The method or composition according to any one of claims 7 to 10, wherein the oligonucleotide comprises the nucleic acid sequence as shown in any one of SEQ ID NOs: 2, 3, 4, 5, 6, 7, 8 or 9.

12. The method or composition according to any one of claims 1 to 11, wherein the oligonucleotide is 18-22 nucleotides in length.

13. The method or composition of claim 8, wherein the miRNA is selected from the group consisting of miR-155, miR-155-inh, miR-181-B1, miR-15a, miR-16-1, miR-21, miR-34a, miR-221, miR-29a, let-7b.

14. The method according to any one of claims 1, 2 or 4 to 13, wherein the mesenchymal lineage precursor or stem cells are modified to introduce an oligonucleotide or vector expressing the same that kills the target cell but does not substantially affect viability of the mesenchymal lineage precursor or stem cell.

15. The method according to any one of claims 1, 2 or 4 to 13, wherein the mesenchymal lineage precursor or stem cells are modified to introduce an inhibitory oligonucleotide or vector expressing the same that does not kill the mesenchymal lineage precursor or stem cells before they can deliver the inhibitory oligonucleotide to a target cell such as a cancer cell.

16. The composition or method according to any one of claims 1 to 15, wherein the mesenchymal lineage precursor or stem cells also express STRO-1.

17. The composition or method according to any one of claims 1 to 16, wherein the mesenchymal lineage precursor or stem cells are substantially STRO-1^{bri}.

18. The composition or method according to any one of claims 1 to 15, wherein the mesenchymal lineage precursor or stem cells are derived from pluripotent cells.

19. The composition or method of claim 18, wherein the pluripotent cells are induced pluripotent stem (iPS) cells.

20. The method of claim 1, wherein the contacting occurs under conditions permitting the mesenchymal lineage precursor or stem cell to form a gap junction with the target cell, whereby the oligonucleotide is delivered to the target cell by traversing the gap junction.

21. The method of claim 20, wherein the gap junction is formed by Cx40 or Cx43.

22. The method or composition according to any one of claims 1 to 21, wherein the delivery of the oligonucleotide is via a mechanism other than Cx43.

23. The method according to any one of claims 1 or 4 to 22, wherein the target cell is a cancer cell or white blood cell.

24. The method of claim 23, wherein the cancer cell is a lung cancer, pancreatic cancer, colorectal cancer, liver cancer, cervical cancer, prostate cancer, osteosarcoma or melanoma cell.

25. The method according to any one of claims 1 or 4 to 22, wherein the target cell is a syncytial cell.

26. The method of claim 25, wherein the syncytial cell is a cardiac myocyte, smooth muscle cell, epithelial cell, connective tissue cell or syncytial cancer cell.

27. A method of treating cancer in a subject, the method comprising administering a composition according to any one of claims 3 to 16 or 22.

28. The method of claim 27, wherein the mesenchymal lineage precursor or stem cells express a connexin that is also expressed by a cancer cell comprising the subject's cancer.

29. The method of claim 28, wherein the connexin is Cx40 or Cx43.

30. The method of claim 28, wherein a cancer cell comprising the subject's cancer expresses Cx43.

31. The method according to any one of claims 2, 4 to 16 or 27 to 30, wherein the cancer is selected from the group consisting of lung cancer, pancreatic cancer, colorectal cancer, liver cancer, cervical cancer, prostate cancer, osteosarcoma and melanoma.

32. The method or composition according to any one of claims 1 to 31, wherein the mesenchymal lineage precursor or stem cell is modified to introduce the oligonucleotide using a viral vector.

33. The method or composition of claim 32, wherein the viral vector is selected from the group consisting of Lentivirus, Baculovirus, Retrovirus, Adenovirus (AdV), Adeno-associated virus (AAV) or a recombinant form thereof.

34. The method or composition of claim 32, wherein the viral vector is a AAV.

35. The method or composition of claim 32, wherein the viral vector is a recombinant AAV.

36. The method or composition according to any one of claims 1 to 35, wherein the mesenchymal lineage precursor or stem cells also expresses CD46.

37. The method or composition according to any one of claims 1 to 36, wherein the modified mesenchymal lineage precursor or stem cell has been treated effect modification of cell surface glycans on the mesenchymal lineage precursor or stem cell.

38. The method or composition according to claim 37, wherein the treatment involves exposure of the mesenchymal lineage precursor or stem cell to a glycosylstrasferase under conditions which result in modification of cell-surface glycans on the mesenchymal precursor lineage cell or stem cell.

39. The method or composition according to claims 37 or 38 wherein the mesenchymal lineage precursor or stem cell is exposed to an exogenous glycosyltranferase.

40. The method or composition according to claims 37 or 38 wherein the mesenchymal lineage precursor or stem cell has been modified to introduce a nucleic acid encoding a glycosyltransferase and wherein expression of the glycosyltransferase in the cell results in enhanced retention of the cell at a site of inflammation *in vivo*.

41. The method or composition according to any one of claims 38 to 40 wherein the glycosyltransferase is a fucosyltransferase, a galactosyltransferase, or a sialyltransferase.

42. The method or composition according to claim 41 wherein the fucosyltransferase is an alpha 1,3 fucosyltransferase such as an alpha 1,3 fucosyltransferase III, alpha 1,3 fucosyltransferase IV, an alpha 1,3 fucosyltransferase VI, an alpha 1,3 fucosyltransferase VII or an alpha 1,3 fucosyltransferase IX.

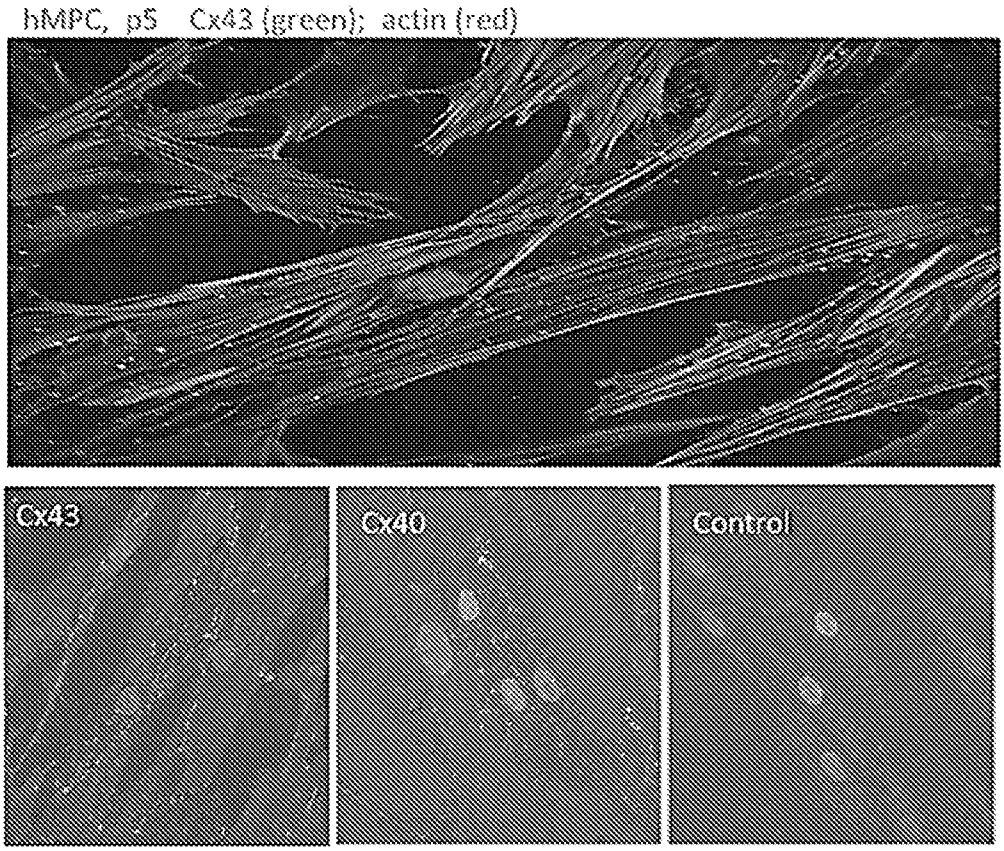


FIGURE 1A

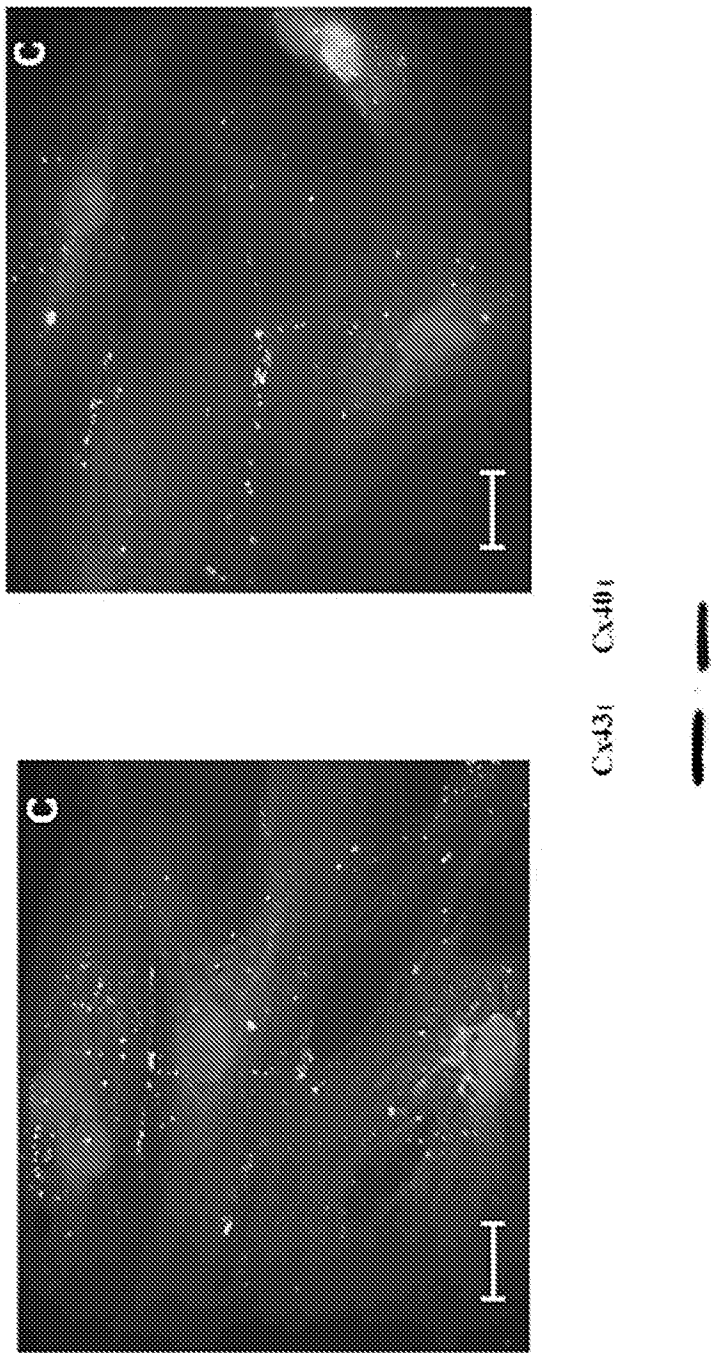
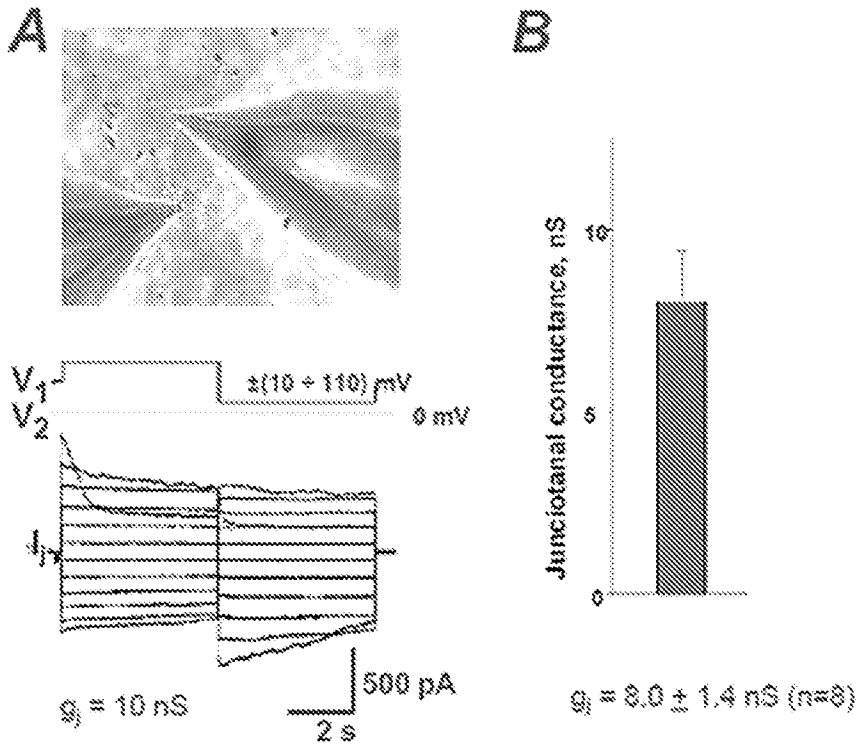


FIGURE 1B


A
B
V1
V2
±(10 ÷ 110) mV
0 mV
Ij
500 pA
2 s
gj = 10 nS
Junctional conductance, nS
10
5
0
gj = 8.0 ± 1.4 nS (n=8)

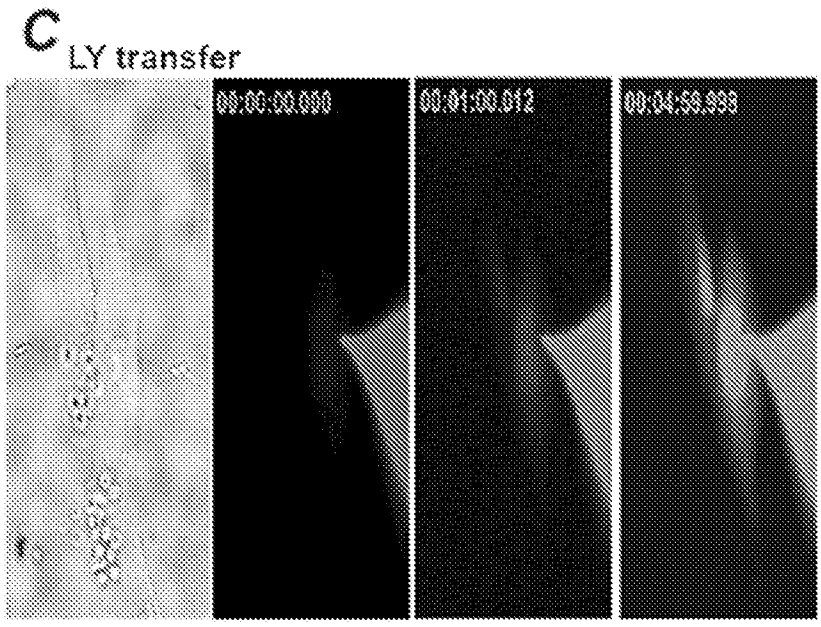

C
LY transfer
00:00:00.000
00:01:00.012
00:04:59.988

FIGURE 2

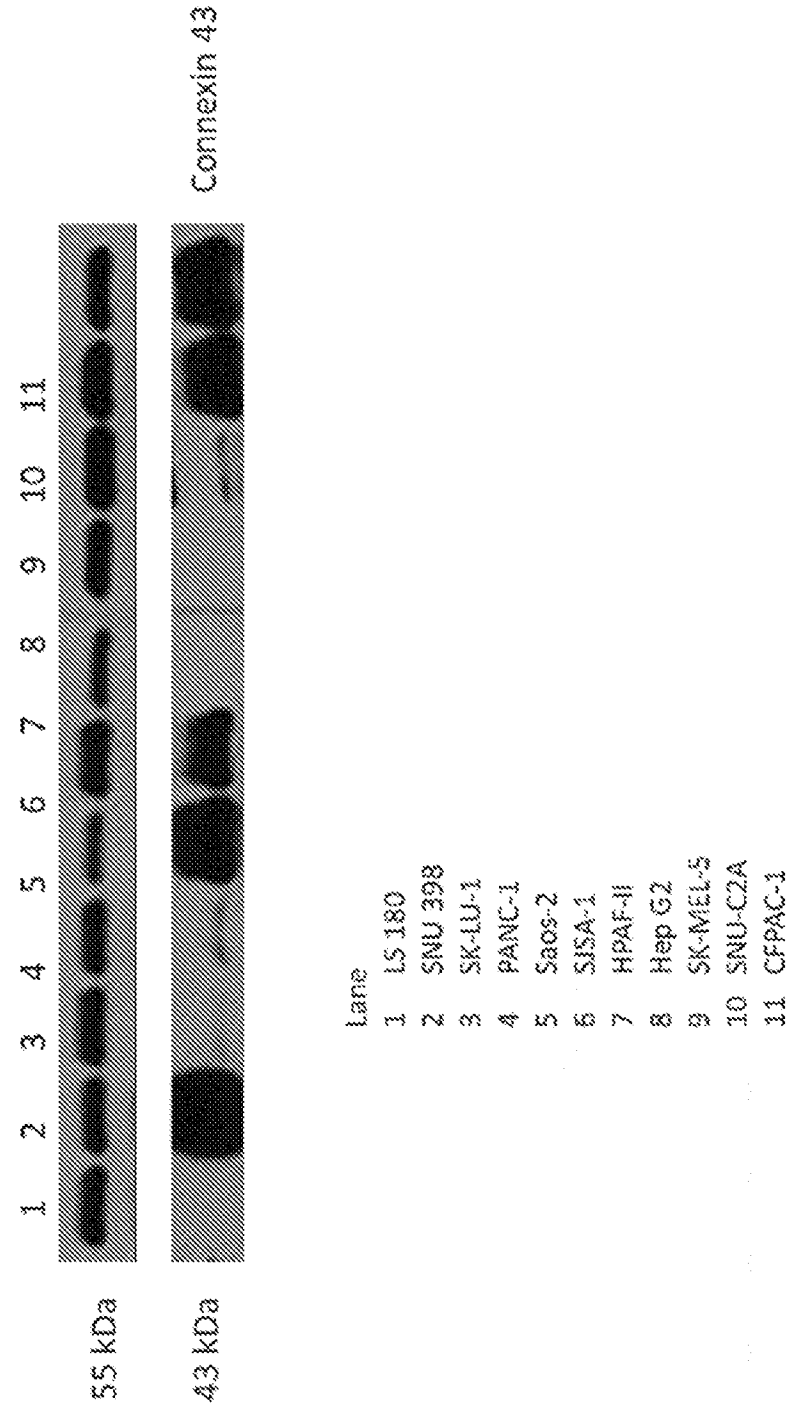

Connexin 43
1 2 3 4 5 6 7 8 9 10 11
55 kDa
43 kDa
Lane
1 LS 180
2 SNU 398
3 SK-LU-1
4 PANC-1
5 Saos-2
6 SISA-1
7 HPAF-II
8 Hep G2
9 SK-MEL-5
10 SNU-C2A
11 CFPAC-1

FIGURE 3-1

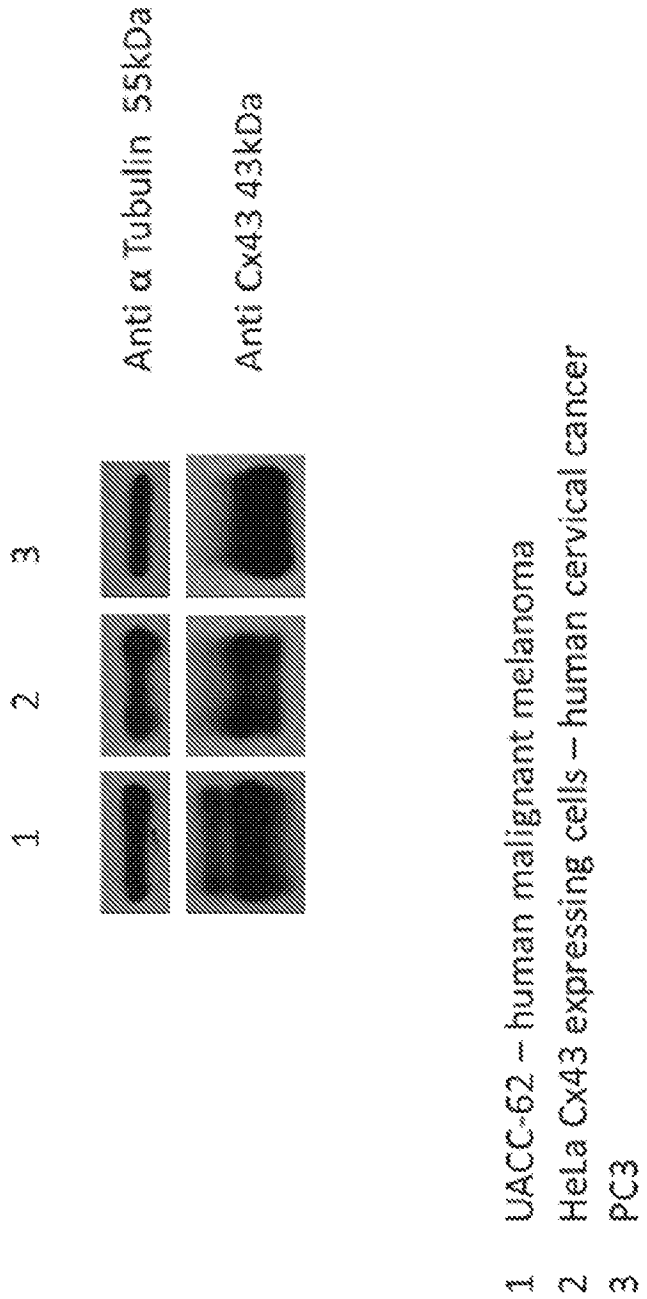

Anti α Tubulin 55kDa
Anti Cx43 43kDa
1 2 3
1 UACC-62 – human malignant melanoma
2 HeLa Cx43 expressing cells – human cervical cancer
3 PC3

FIGURE 3-2

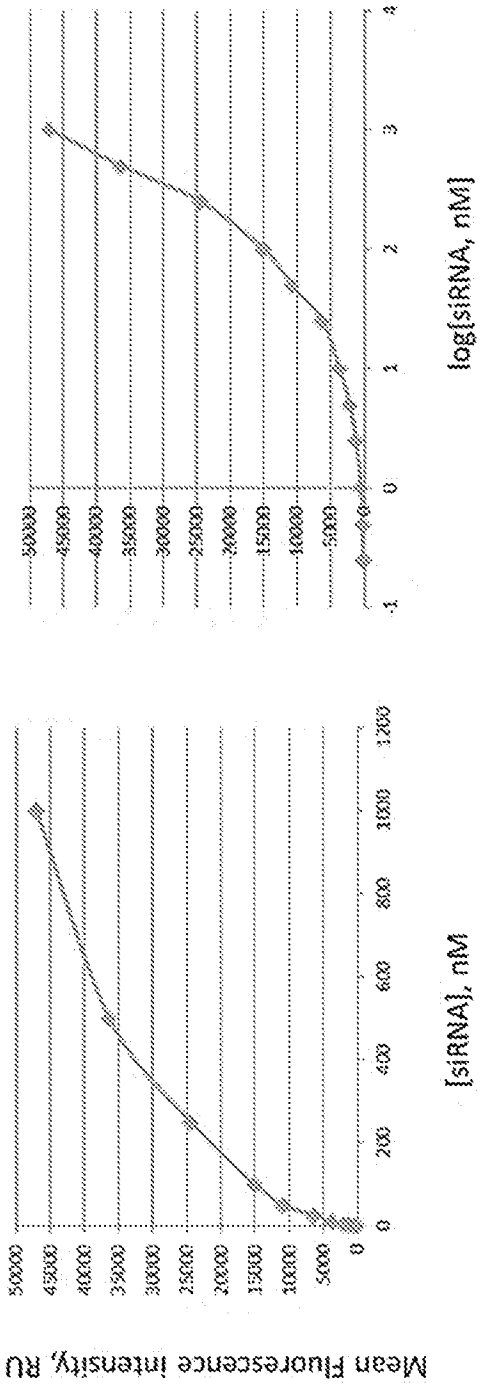

Mean Fluorescence Intensity, RU
50000
45000
40000
35000
30000
25000
20000
15000
10000
5000
0
0
200
400
600
800
1000
1200
[siRNA], nM
log[siRNA, nM]
-1
0
1
2
3
4

FIGURE 4

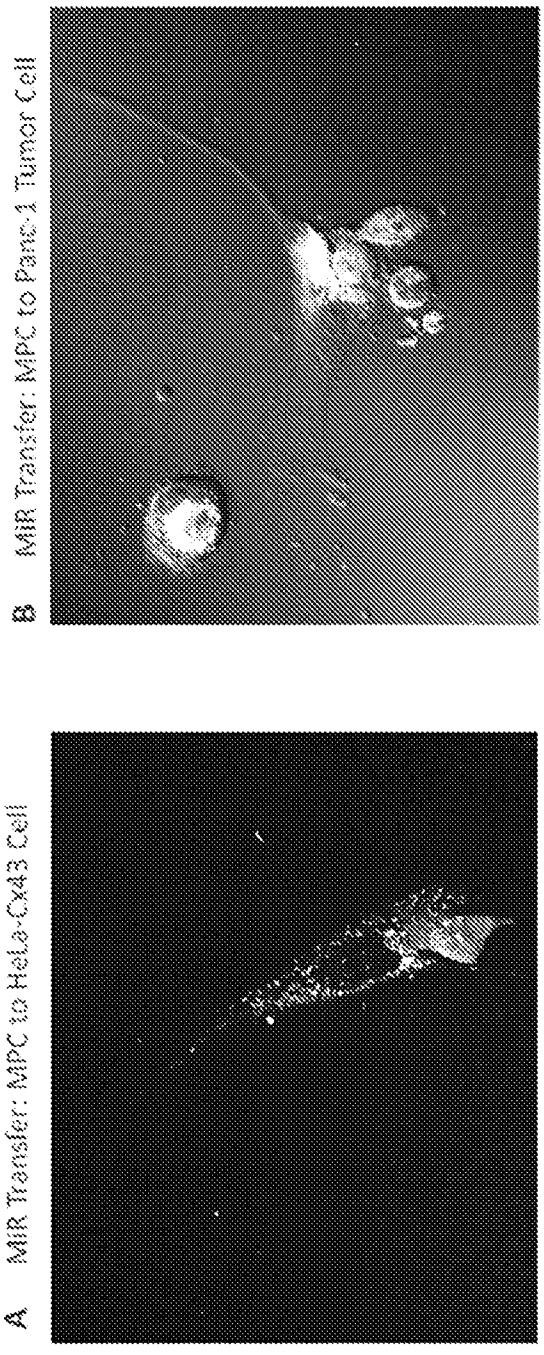

B MiR Transfer: MPC to Panc-1 Tumor Cell
A MiR Transfer: MPC to HeLa-Cx43 Cell

FIGURE 5(i)

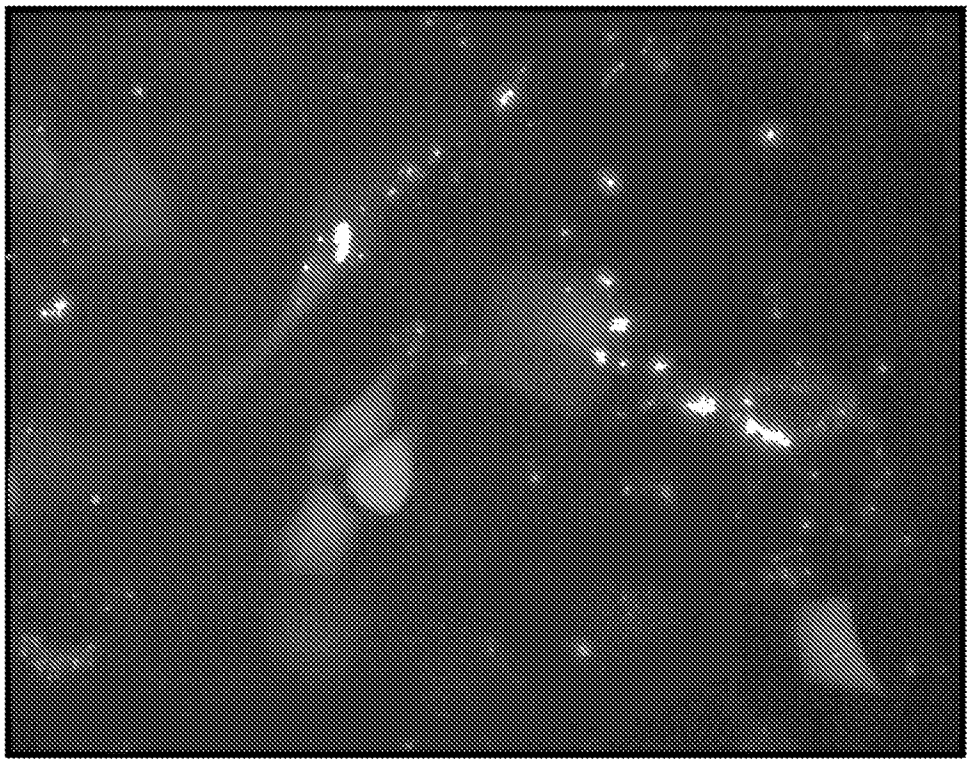


FIGURE 5(ii)

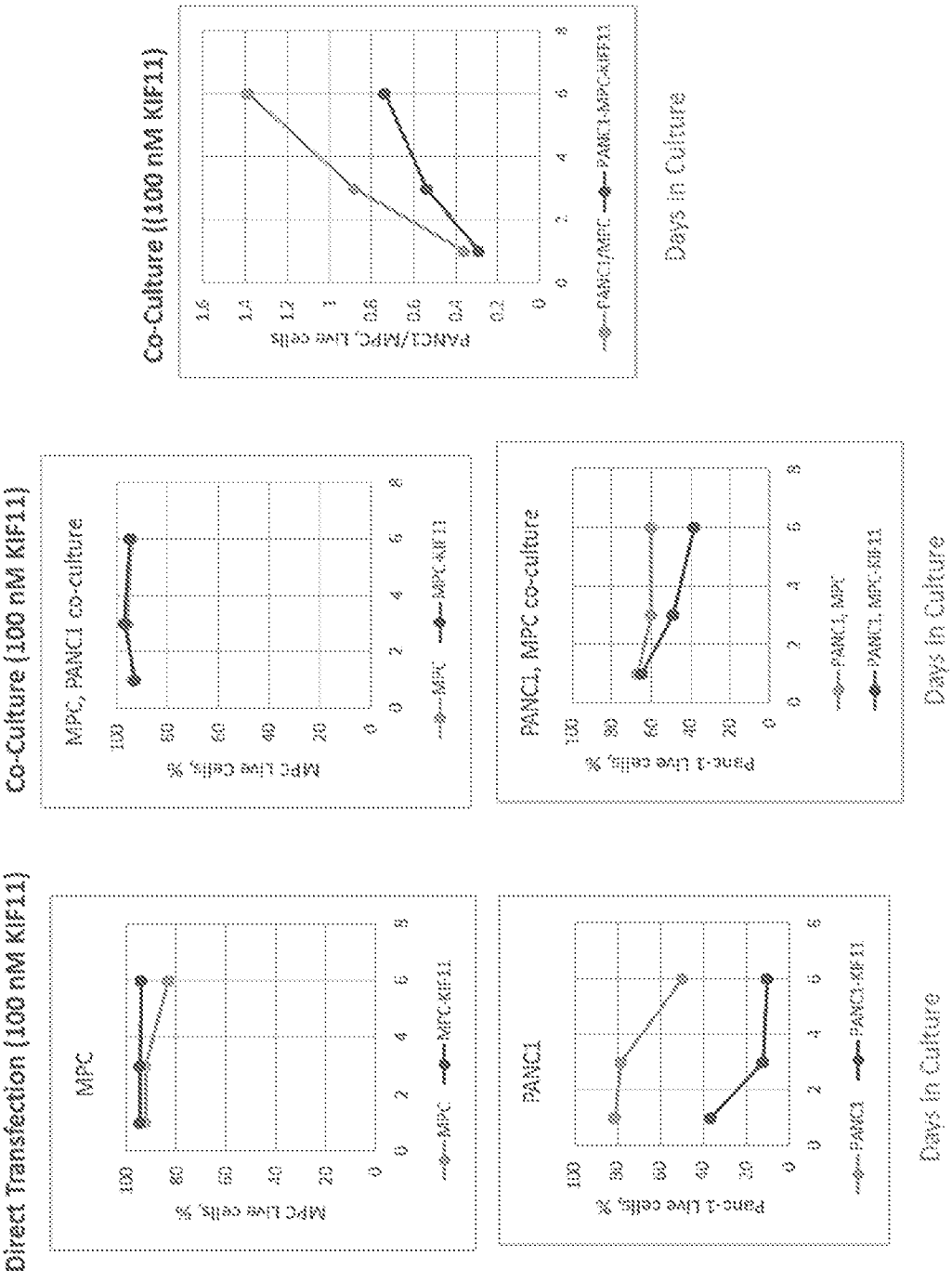

Direct Transfection (100 nM KIF11)
MPC
MPC Live cells, %
MPC
MPC-KIF11
PANC1
Panc-1 live cells, %
PANC1
PANC1-KIF11
Days in Culture
Co-Culture (100 nM KIF11)
MPC, PANC1 co-culture
MPC Live Cells, %
MPC
MPC-KIF11
PANC1, MPC co-culture
Panc-1 live cells, %
PANC1, MPC
PANC1, MPC-KIF11
Days in Culture
Co-Culture ((100 nM KIF11)
PANC1/MPC Live cells
PANC1/MPC
PANC1-MPC-KIF11
Days in Culture

FIGURE 6(A)

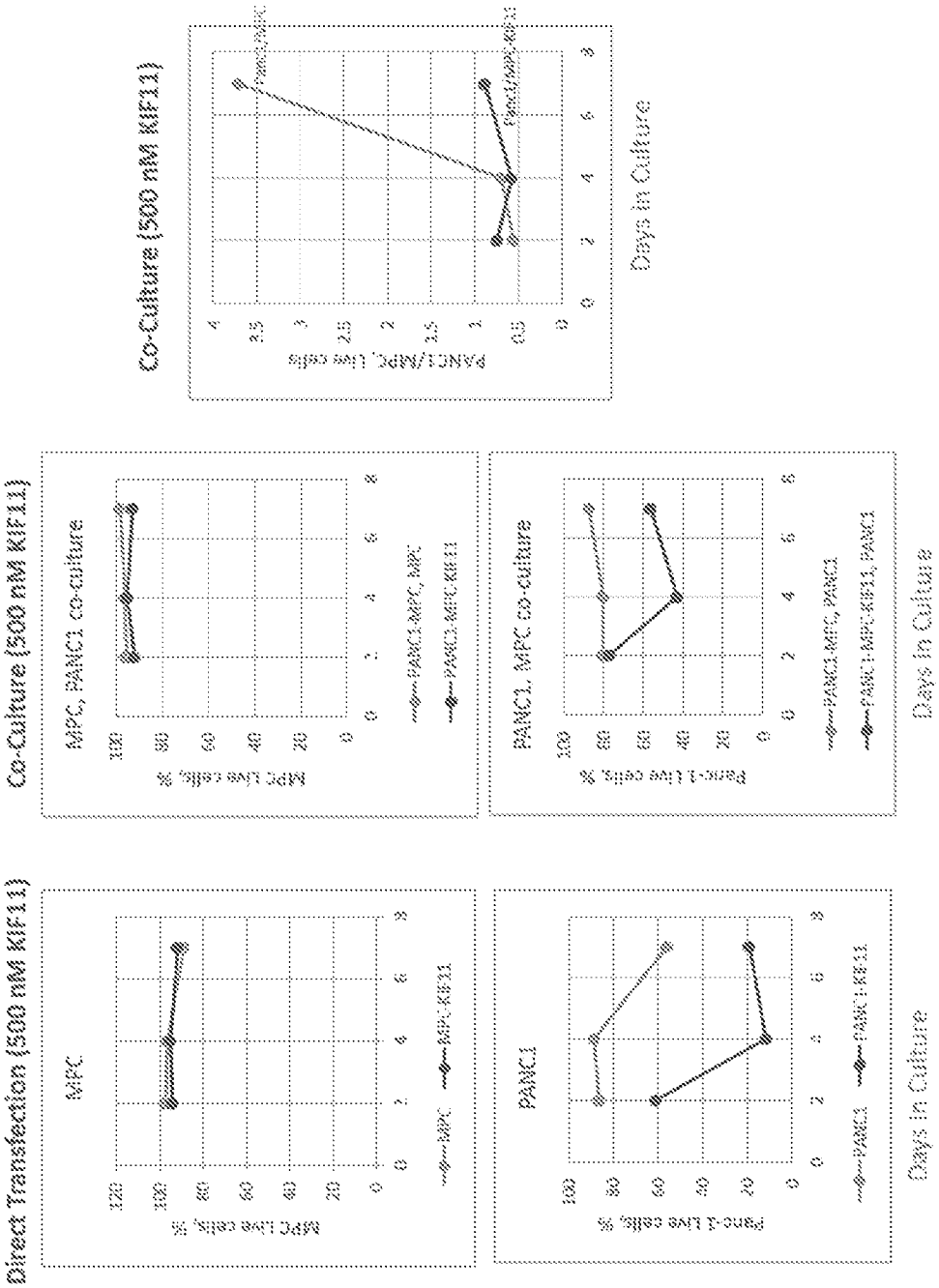

Co-Culture (500 nM KIF11)
PANC1/MPC, Live cells
Days in Culture
Co-Culture (500 nM KIF11)
MPC, PANC1 co-culture
MPC Live cells, %
PANC1 MPC, MPC
PANC1 MPC KIF11
PANC1, MPC co-culture
Panc-1 Live cells, %
PANC1 MPC, PANC1
PANC1 MPC KIF11, PANC1
Days in Culture
Direct Transfection (500 nM KIF11)
MPC
MPC Live cells, %
MPC
MPC KIF11
PANC1
Panc-1 Live cells, %
PANC1
PANC1 KIF11
Days in Culture

FIGURE 6(B)

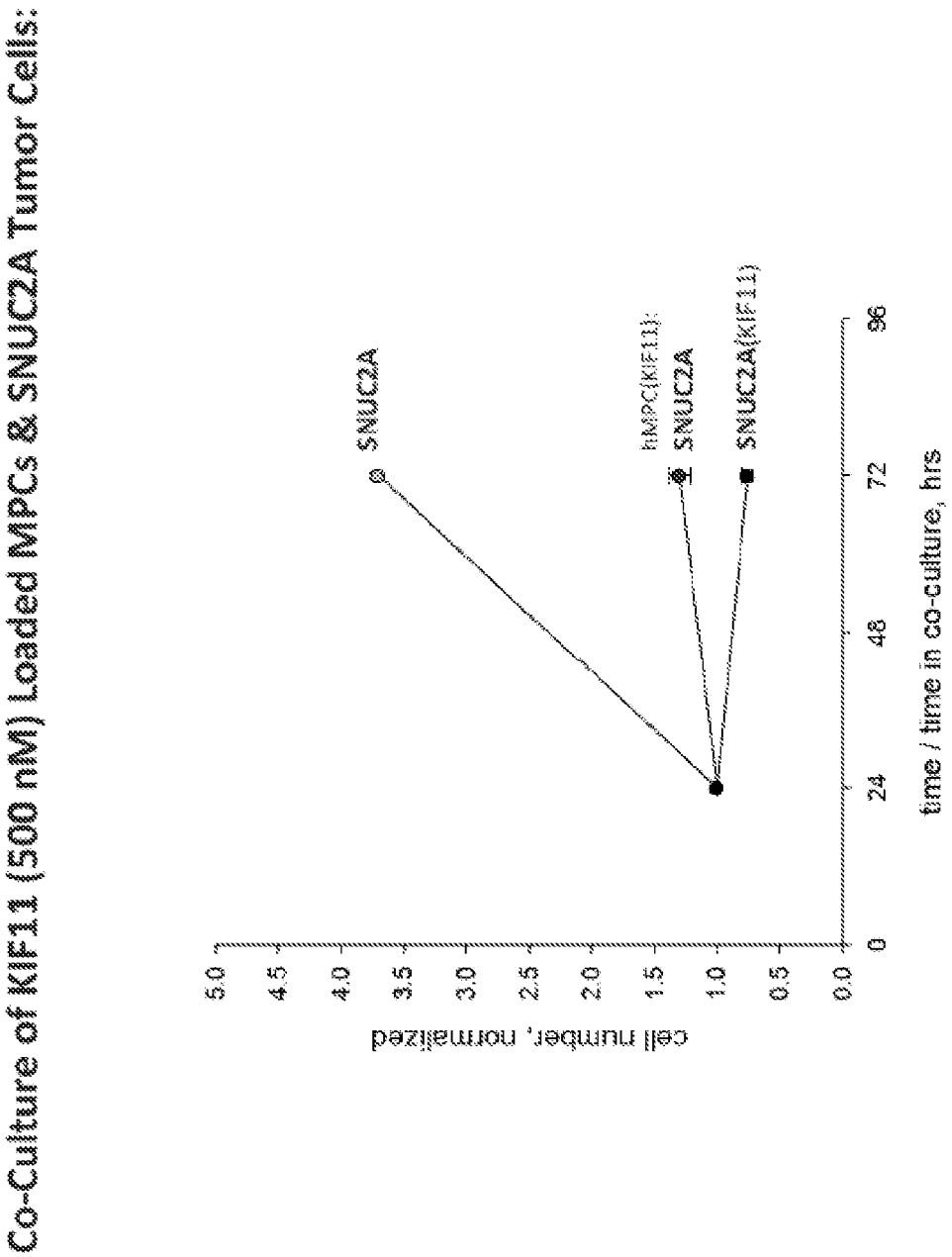

Co-Culture of KIF11 (500 nM) Loaded MPCs & SNUC2A Tumor Cells:
SNUC2A
hMPC(KIF11):
SNUC2A
SNUC2A(KIF11)
cell number, normalized
5.0
4.5
4.0
3.5
3.0
2.5
2.0
1.5
1.0
0.5
0.0
0
24
48
72
96
time / time in co-culture, hrs

FIGURE 7

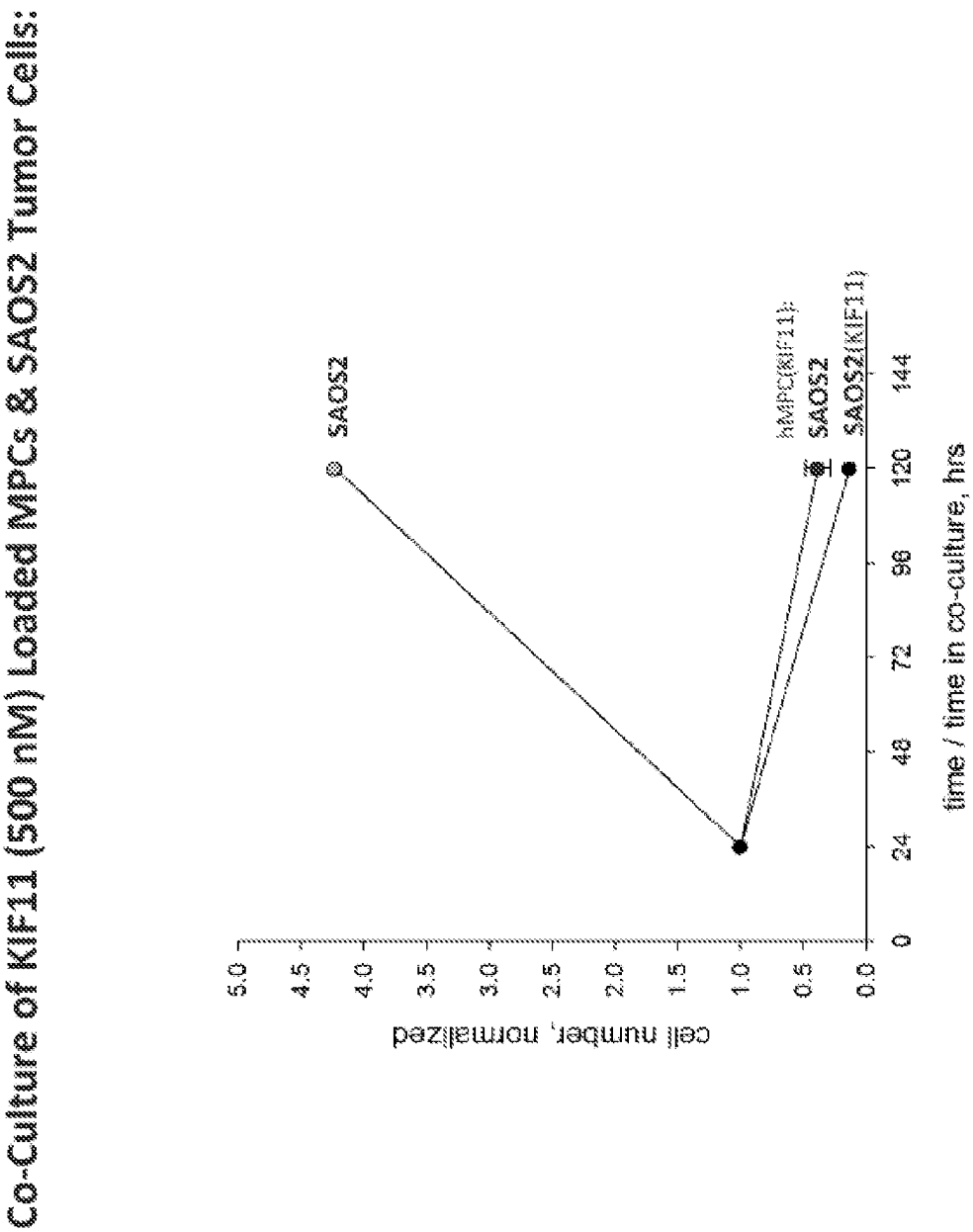

Co-Culture of KIF11 (500 nM) Loaded MPCs & SAOS2 Tumor Cells:
cell number, normalized
5.0
4.5
4.0
3.5
3.0
2.5
2.0
1.5
1.0
0.5
0.0
0
24
48
72
96
120
144
time / time in co-culture, hrs
SAOS2
hMPC(KIF11)+
SAOS2
SAOS2(KIF11)

FIGURE 8(A)

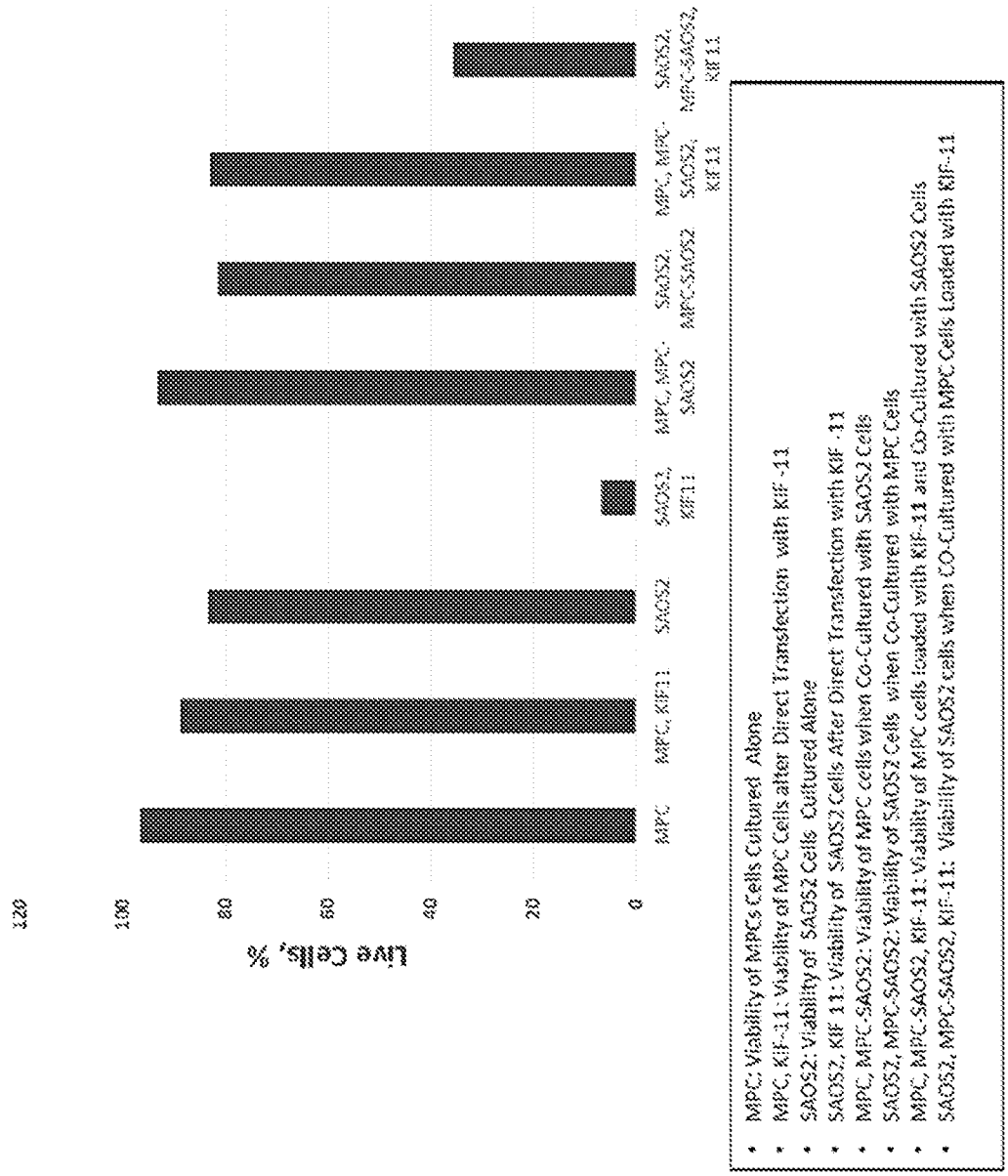

Live Cells, %
120
100
80
60
40
20
0
MPC
MPC, KIF11
SAOS2
SAOS2, KIF11
MPC, MPC-SAOS2
SAOS2, MPC-SAOS2
MPC, MPC-SAOS2, KIF11
SAOS2, MPC-SAOS2, KIF11
MPC: Viability of MPCs Cells Cultured Alone
MPC, KIF-11: Viability of MPC Cells after Direct Transfection with Kif - 11
SAOS2: Viability of SAOS2 Cells Cultured Alone
SAOS2, Kif 11: Viability of SAOS2 Cells After Direct Transfection with Kif - 11
MPC, MPC-SAOS2: Viability of MPC cells when Co-Cultured with SAOS2 Cells
SAOS2, MPC-SAOS2: Viability of SAOS2 Cells when Co-Cultured with MPC Cells
MPC, MPC-SAOS2, KIF-11: Viability of MPC cells loaded with KIF-11 and Co-Cultured with SAOS2 Cells
SAOS2, MPC-SAOS2, KIF-11: Viability of SAOS2 cells when CO-Cultured with MPC Cells Loaded with KIF-11

FIGURE 8(B)

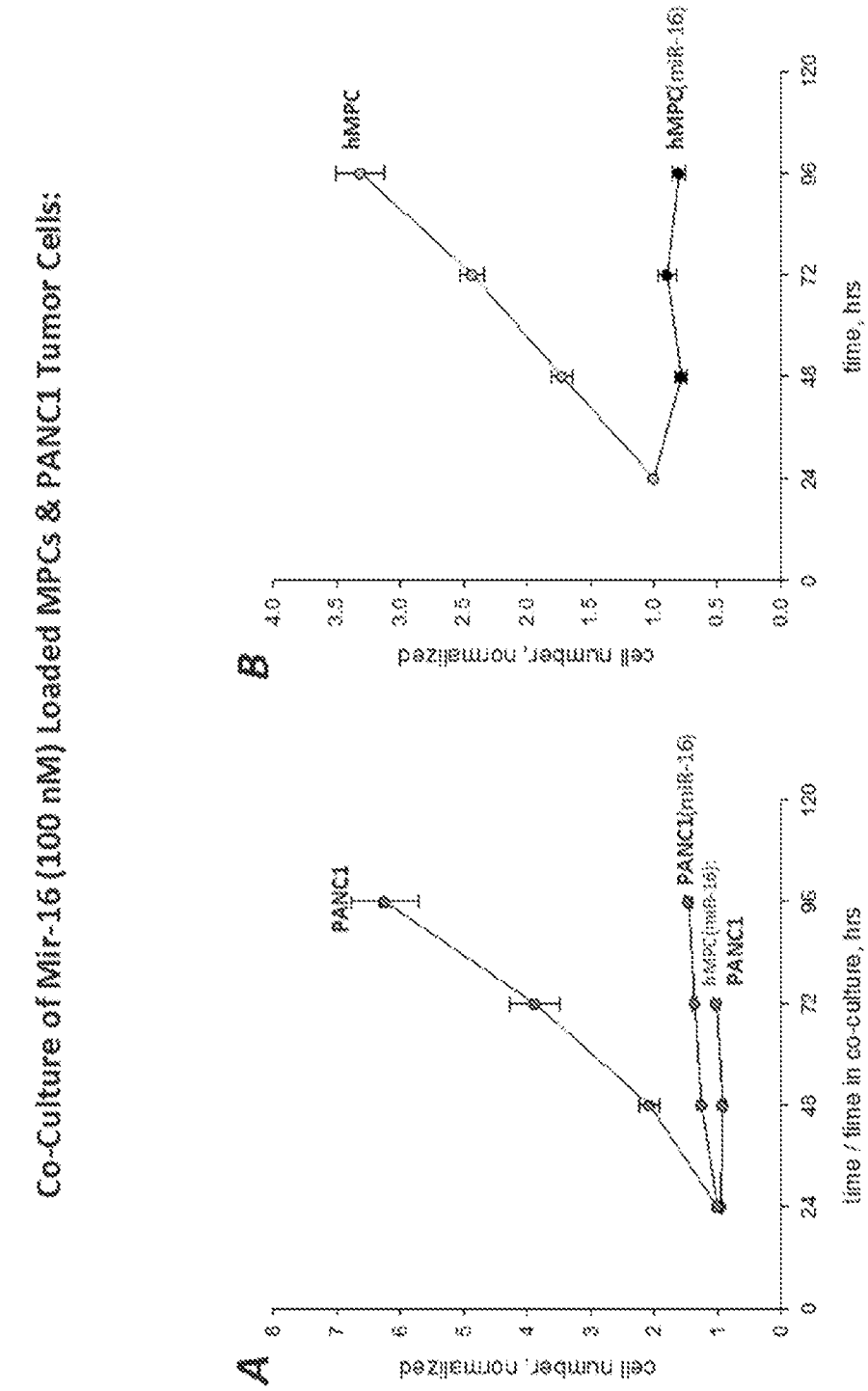

Co-Culture of Mir-16 (100 nM) Loaded MPCs & PANC1 Tumor Cells:
A
cell number, normalized
time / time in co-culture, hrs
PANC1
PANC1(miR-16)
hMPC(miR-16)
PANC1
B
cell number, normalized
time, hrs
hMPC
hMPC(miR-16)

FIGURE 9

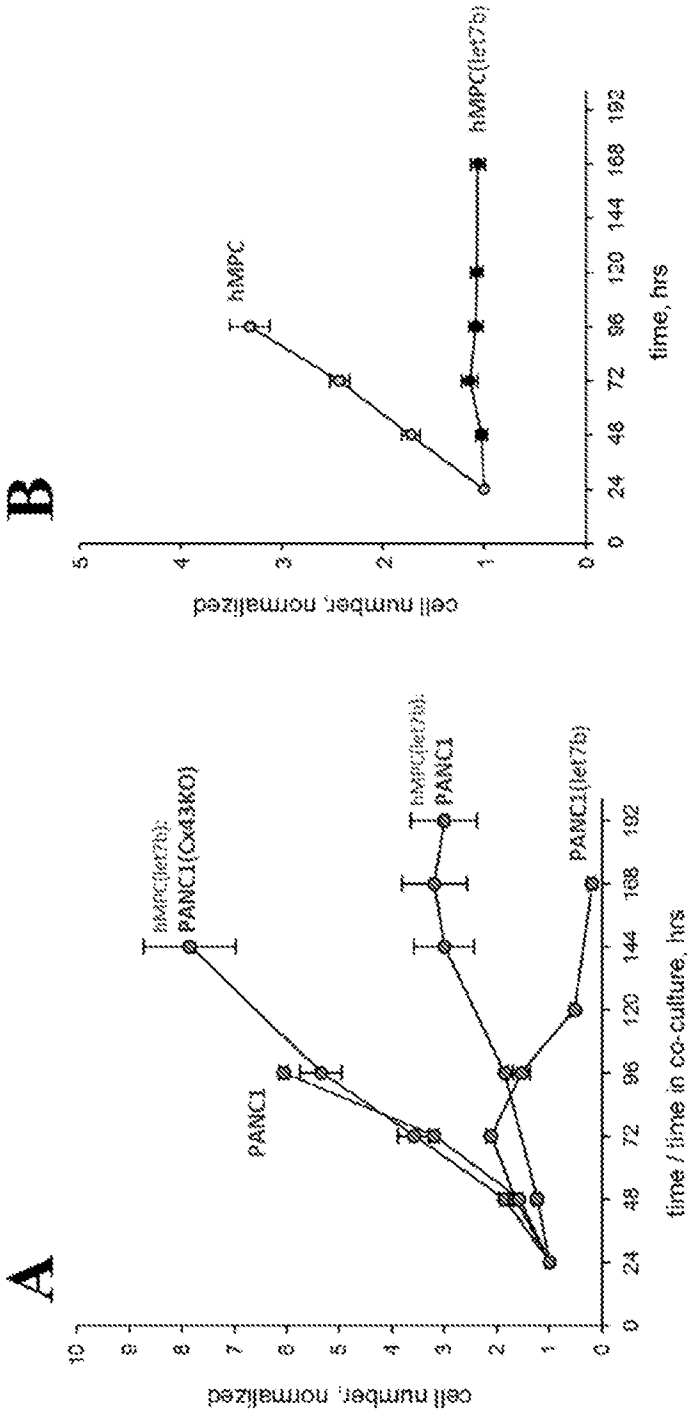

A
cell number, normalized
time / time in co-culture, hrs
PANC1
hMPC(let7b)
PANC1(Cx43KO)
hMPC(let7b)
PANC1
PANC1(let7b)
B
cell number, normalized
hMPC
hMPC(let7b)
time, hrs

FIGURE 10

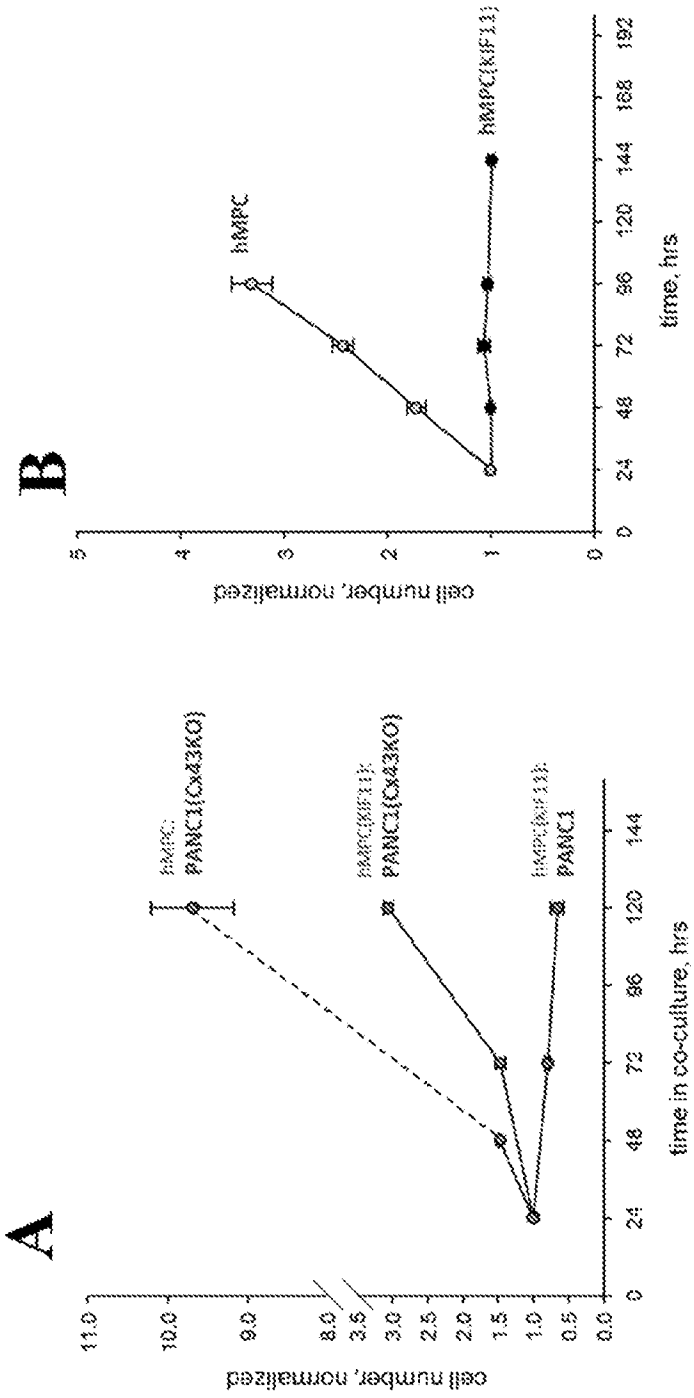

A
cell number, normalized
time in co-culture, hrs
PANC1(Cx43KO)
PANC1
B
cell number, normalized
time, hrs
hMPC
hMPC(KIF11i)

FIGURE 11

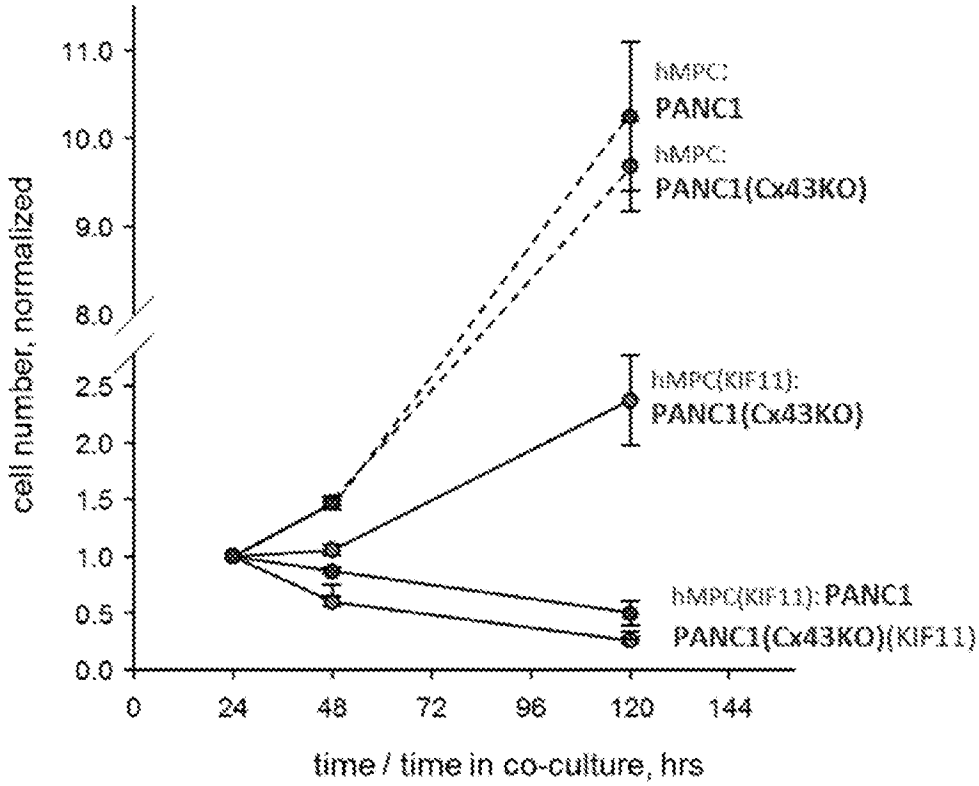

cell number, normalized
11.0
10.0
9.0
8.0
2.5
2.0
1.5
1.0
0.5
0.0
0
24
48
72
96
120
144
time / time in co-culture, hrs
hMPC:
PANC1
hMPC:
PANC1(Cx43KO)
hMPC(KIF11):
PANC1(Cx43KO)
hMPC(KIF11): PANC1
PANC1(Cx43KO)(KIF11)

FIGURE 12

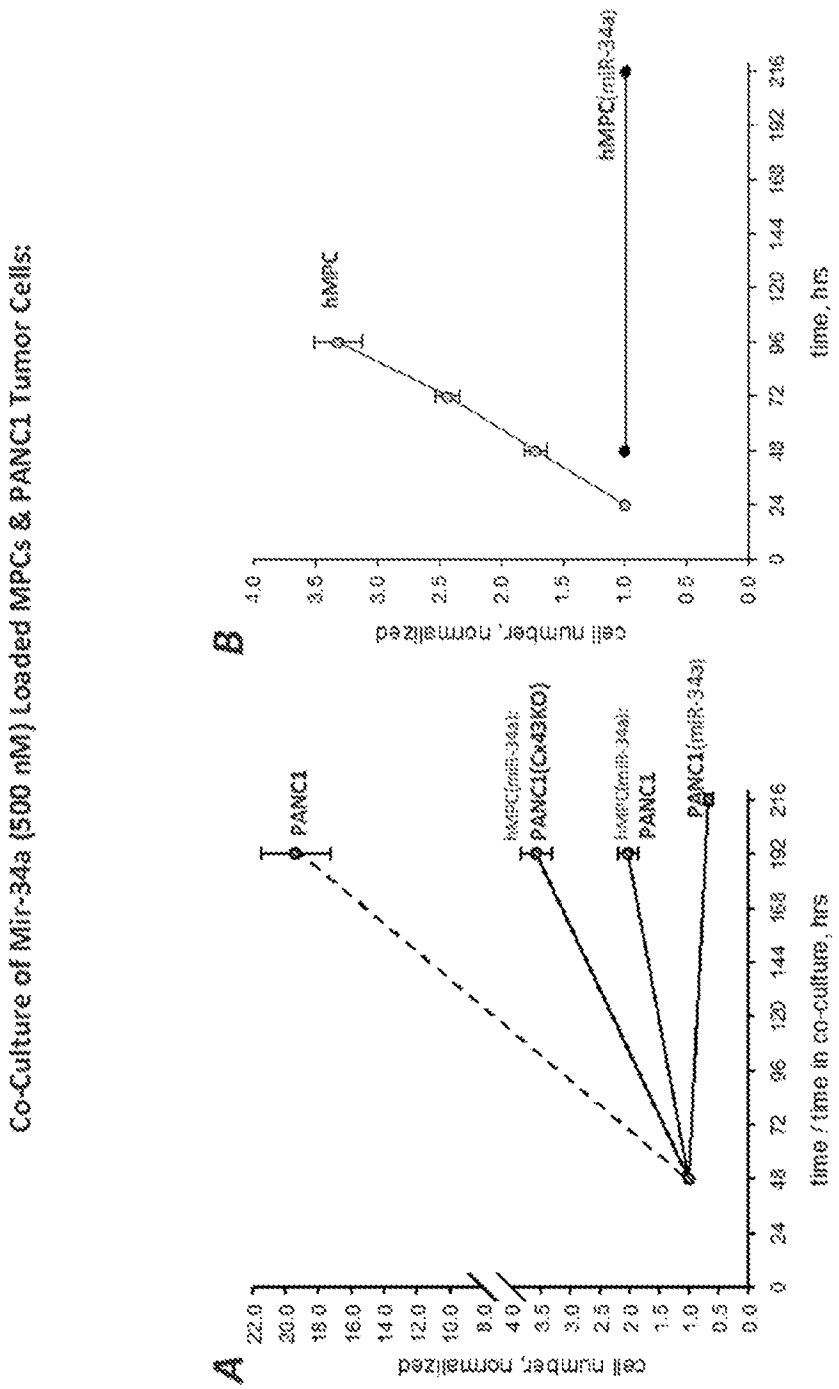

Co-Culture of Mir-34a (500 nM) Loaded MPCs & PANC1 Tumor Cells:
A
B
cell number, normalized
time / time in co-culture, hrs
PANC1
PANC1(Cx43KO)
PANC1
hMPC
hMPC(miR-34a)
time, hrs

FIGURE 13

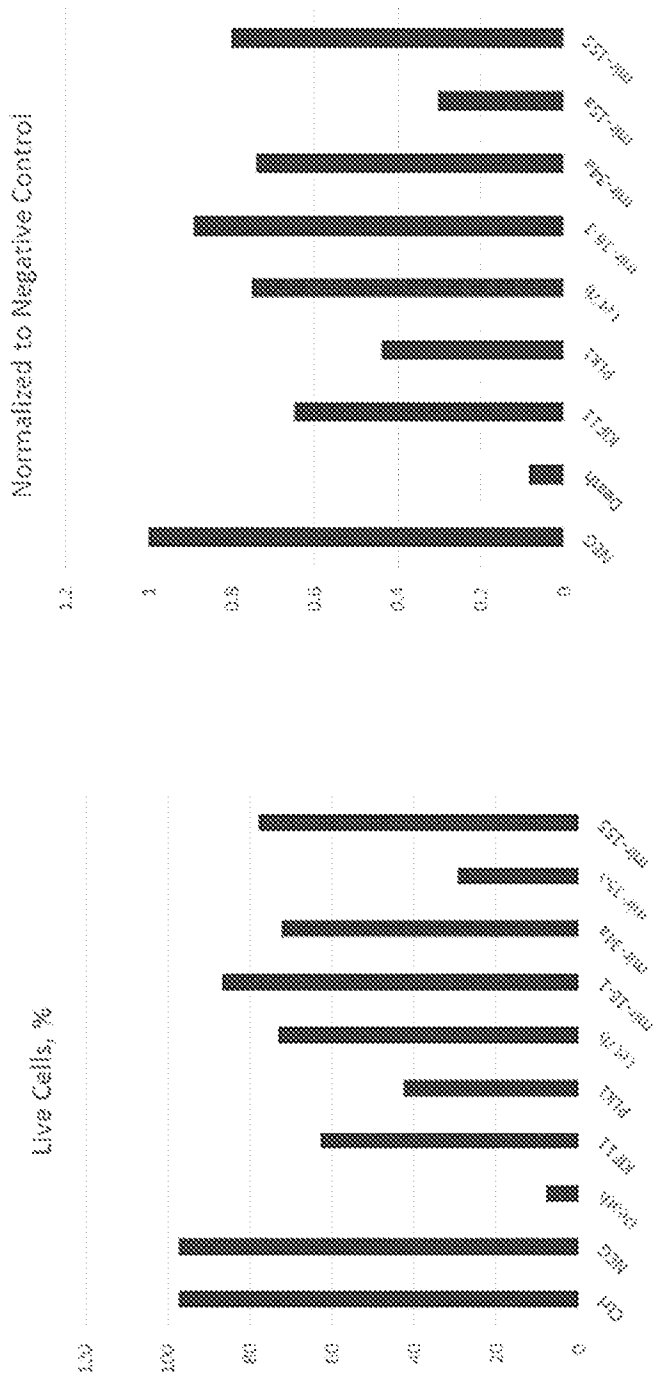
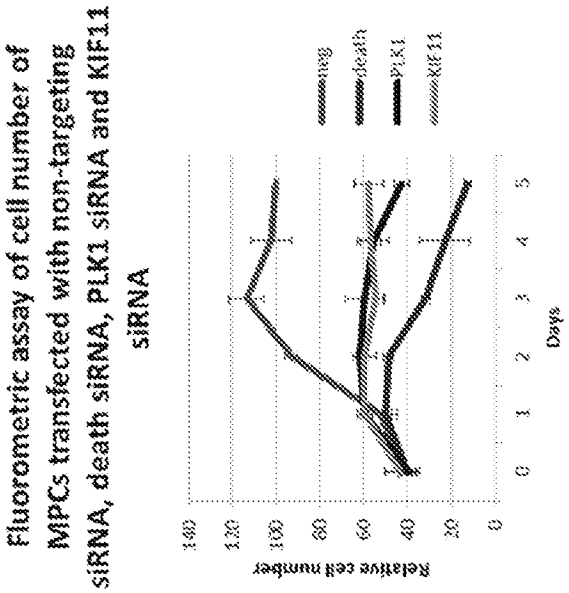
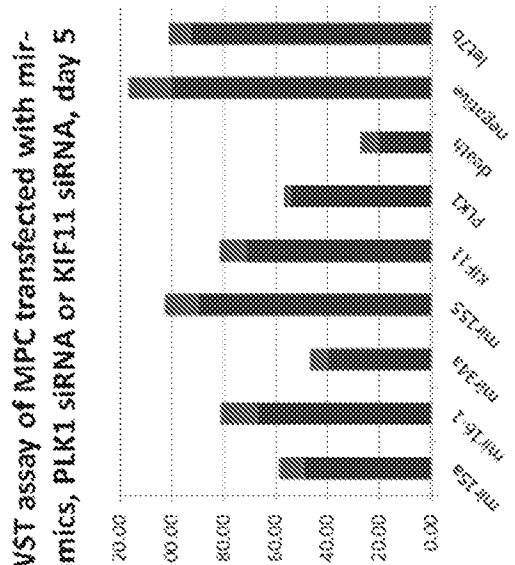


FIGURE 14(A)





	exp1	exp2	exp3		mean	STD
none	0.39	0.44	0.45	none	0.41	0.04
mir15a	0.27	0.35	0.31	mir15a	0.31	0.06
mir16-1	0.36	0.48	0.35	mir16-1	0.42	0.09
mir34a	0.22	0.28	0.21	mir34a	0.29	0.04
mir155	0.62	0.50	0.63	mir155	0.56	0.09
KIF11	0.49	0.40	0.64	KIF11	0.45	0.07
PLK1	0.33	0.35	0.39	PLK1	0.34	0.02
death	0.16	0.09	0.15	death	0.13	0.05
negative	0.70	0.55	0.84	negative	0.63	0.10
let7b	0.54	0.62	0.61	let7b	0.58	0.05

	mean	STD
none	85.94	5.83
mir15a	49.02	8.47
mir16-1	67.06	14.22
mir34a	39.84	5.69
mir155	89.34	13.86
KIF11	71.02	10.43
PLK1	53.39	7.73
death	20.05	7.27
negative	100.00	16.64
let7b	92.37	0.88

FIGURE 14(B)

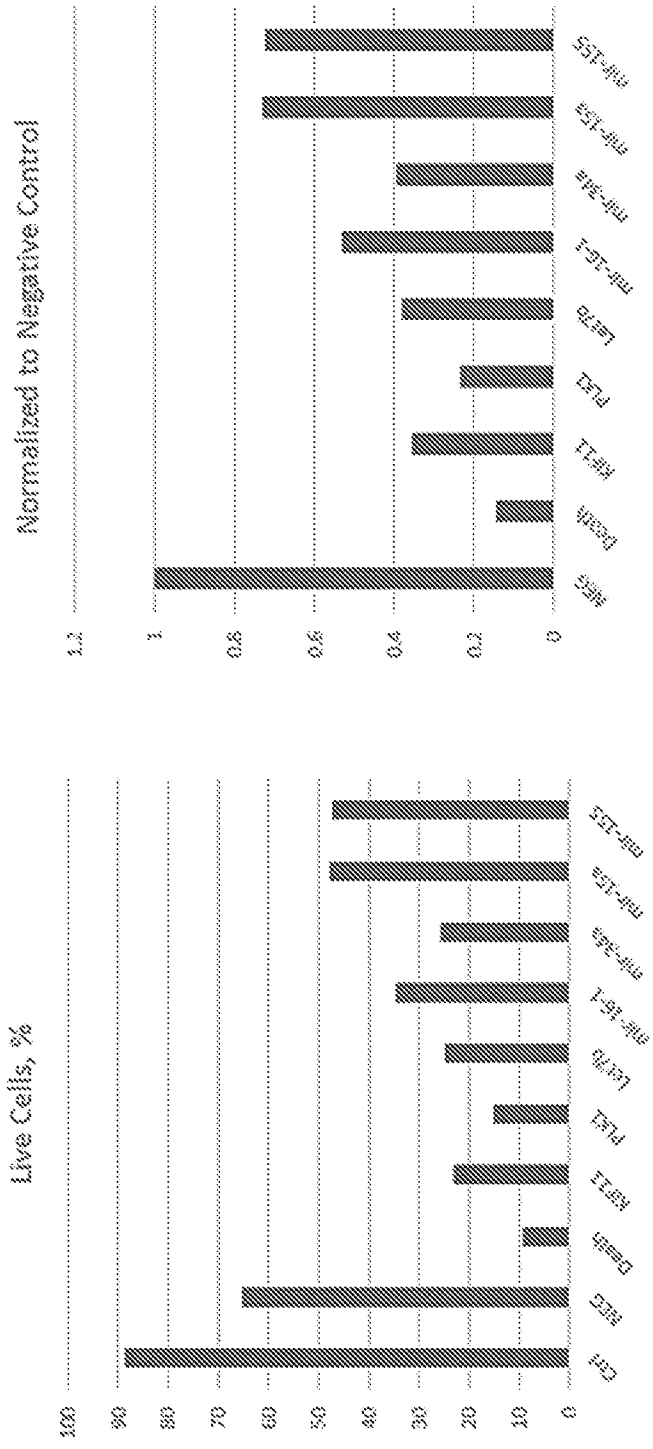
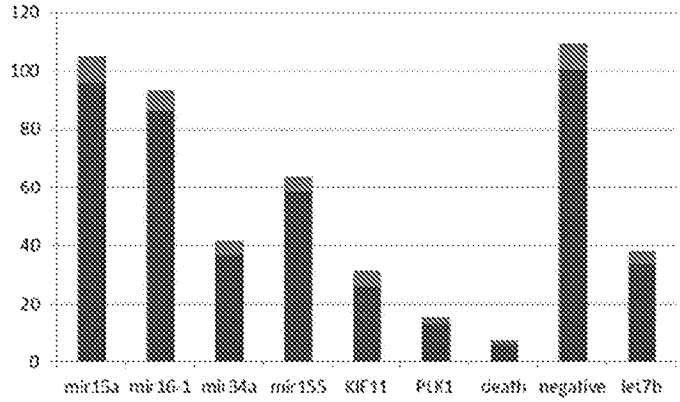

Normalized to Negative Control
1.2
1
0.8
0.6
0.4
0.2
0
Live Cells, %
100
90
80
70
60
50
40
30
20
10
0

FIGURE 15(A)

WST assay of PANC1 transfected with mir-mimics, PLK1 siRNA or KIF11 siRNA, day 5



	exp1	exp2	exp3		mean	STD		mean	STD
none	1.69	1.65	1.70	none	1.68	0.03	none	128.59	2.17
mir15a	1.29	1.35	1.11	mir15a	1.25	0.12	mir15a	95.74	9.27
mir16-1	1.06	1.23	1.09	mir16-1	1.12	0.09	mir16-1	86.09	7.04
mir34a	0.42	0.47	0.54	mir34a	0.48	0.06	mir34a	36.67	4.78
mir155	0.69	0.82	0.77	mir155	0.76	0.07	mir155	58.45	5.13
KIF11	0.26	0.37	0.39	KIF11	0.34	0.07	KIF11	25.91	5.50
PLK1	0.15	0.15	0.20	PLK1	0.17	0.03	PLK1	12.96	2.22
death	0.09	0.06	0.09	death	0.08	0.02	death	5.93	1.34
negative	1.17	1.36	1.33	negative	1.31	0.12	negative	100.00	9.17
let7b	0.38	0.49	0.45	let7b	0.44	0.06	let7b	33.54	4.58

FIGURE 15(B)

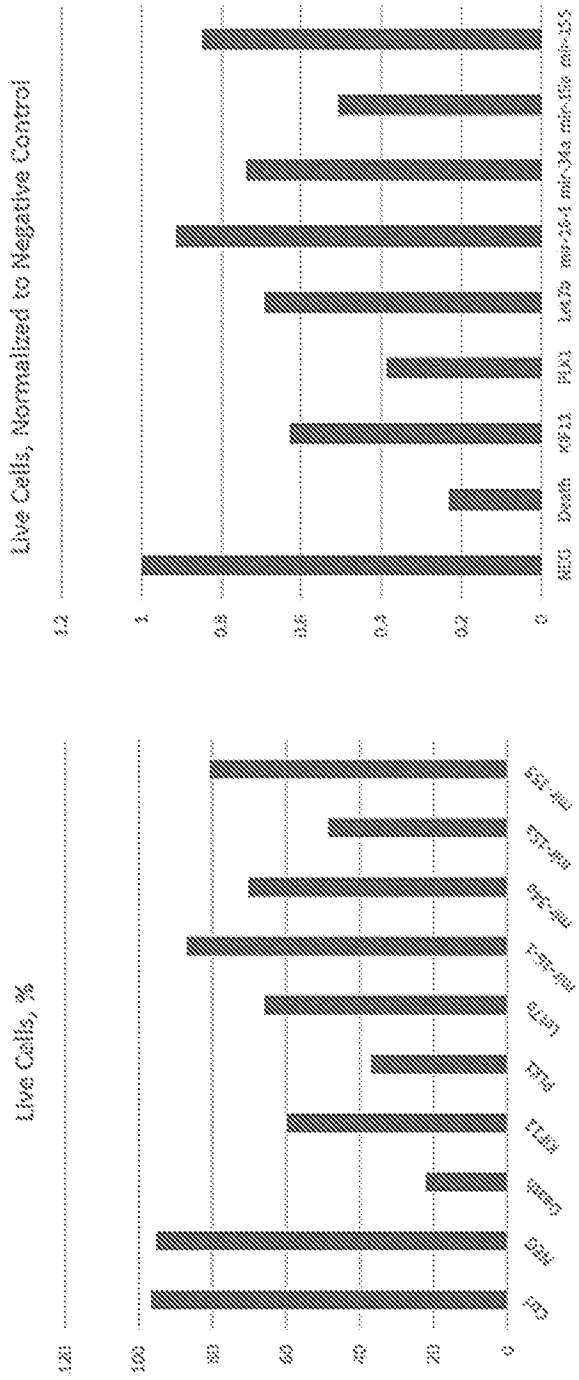
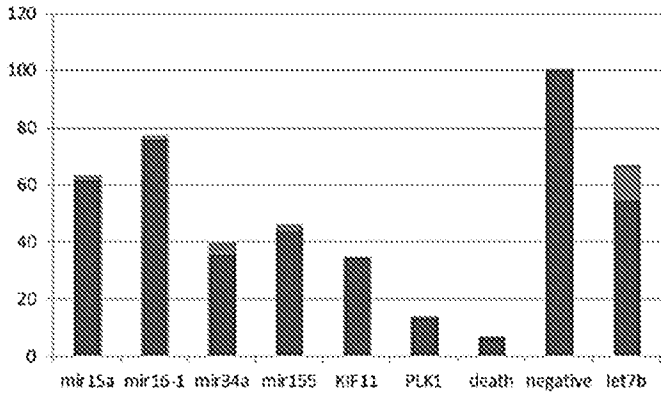

Live Cells, Normalized to Negative Control
1.2
1
0.8
0.6
0.4
0.2
0
NEG
Death
miR-21
miR-23
miR-155
miR-16-1 miR-34a miR-15a miR-155
Live Cells, %
120
100
80
60
40
20
0

FIGURE 16(A)

WST assay of PC3 cells transfected with mir-mimics, PLK1 siRNA or KIF11 siRNA, day 5



	exp1	exp2	exp3		mean	STD		mean	STD
								101.2	
none	1.83	1.90	1.82	none	1.87	0.05	none	1	2.54
mir15a	1.16	1.11	0.76	mir15a	1.14	0.03	mir15a	61.57	1.83
mir16-1	1.42	1.38	1.07	mir16-1	1.40	0.03	mir16-1	75.66	1.61
mir34a	0.71	0.60	0.69	mir34a	0.65	0.07	mir34a	35.49	4.06
mir155	0.79	0.84	0.61	mir155	0.81	0.04	mir155	44.12	1.95
KIF11	0.64	0.63	0.50	KIF11	0.64	0.01	KIF11	34.55	0.75
PLK1	0.24	0.25	0.17	PLK1	0.25	0.01	PLK1	13.30	0.49
death	0.12	0.11	0.03	death	0.11	0.01	death	6.17	0.35
								100.0	
negative	1.86	1.83	1.75	negative	1.85	0.02	negative	2	0.67
let7b	0.85	1.17	0.62	let7b	1.01	0.22	let7b	54.69	12.07

FIGURE 16(B)

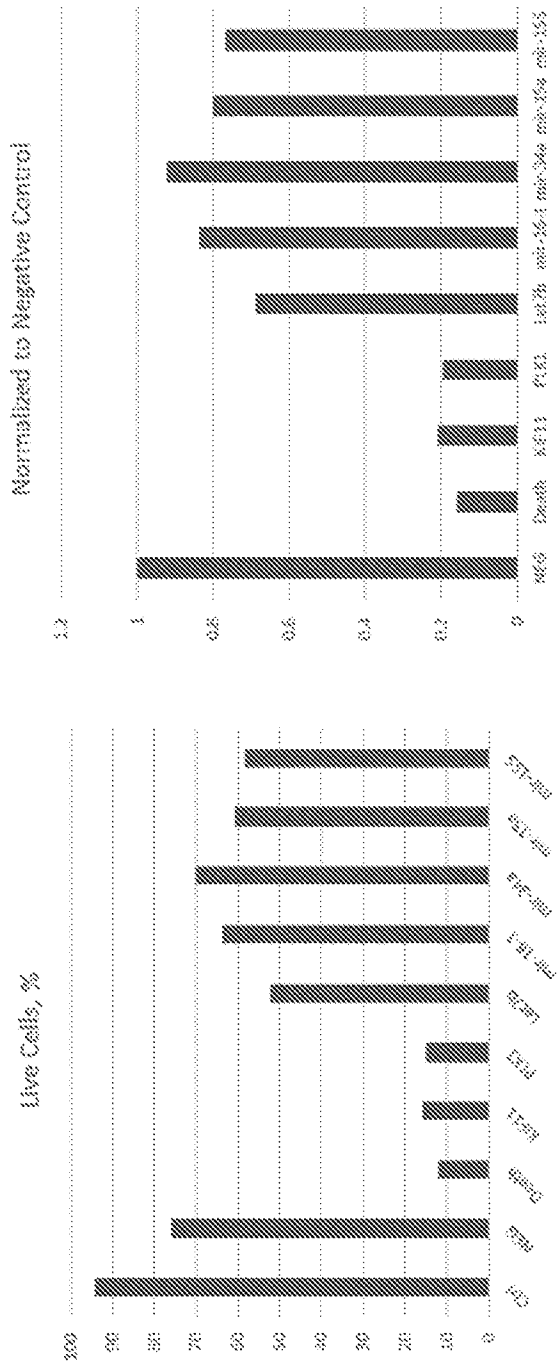
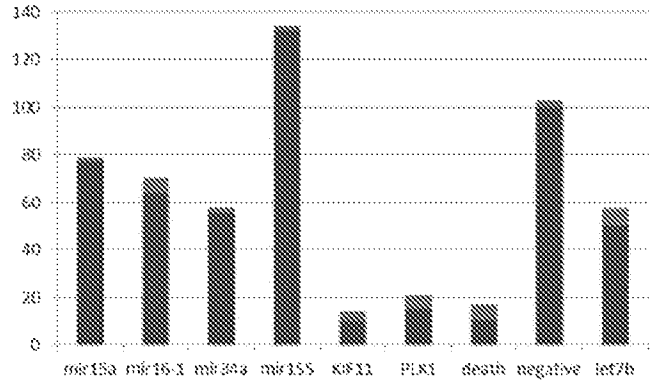

Normalized to Negative Control
1.2
1
0.8
0.6
0.4
0.2
0
miR-0
Death
miR-211
miR-211
miR-34a
miR-34a
miR-34a
miR-34a
Live Cells, %
100
90
80
70
60
50
40
30
20
10
0
-ve
+ve
miR-0
Death
miR-211
miR-211
miR-34a
miR-34a
miR-34a
miR-34a

FIGURE 17(A)

WST assay of SAOS2 cells transfected with mir-mimics, PLK1 siRNA or KIF11 siRNA, day 5



	exp1	exp2	exp3		mean	STD		mean	STD
none	1.20	1.19	0.98	none	1.19	0.01	none	129.00	1.26
mir15a	0.72	0.71	0.46	mir15a	0.72	0.01	mir15a	77.41	1.22
mir16-1	0.55	0.63	0.47	mir16-1	0.59	0.06	mir16-1	63.50	6.51
mir34a	0.50	0.53	0.53	mir34a	0.52	0.02	mir34a	56.04	1.99
mir155	1.21	1.24	0.96	mir155	1.22	0.02	mir155	132.05	1.99
KIF11	0.12	0.07	0.14	KIF11	0.09	0.04	KIF11	9.84	4.65
PLK1	0.18	0.11	0.10	PLK1	0.14	0.05	PLK1	15.23	5.43
death	0.14	0.06	0.07	death	0.10	0.06	death	10.60	6.58
negative	0.94	0.91	0.80	negative	0.93	0.02	negative	100.01	2.63
let7b	0.42	0.51	0.27	let7b	0.47	0.07	let7b	50.42	7.04

FIGURE 17(B)

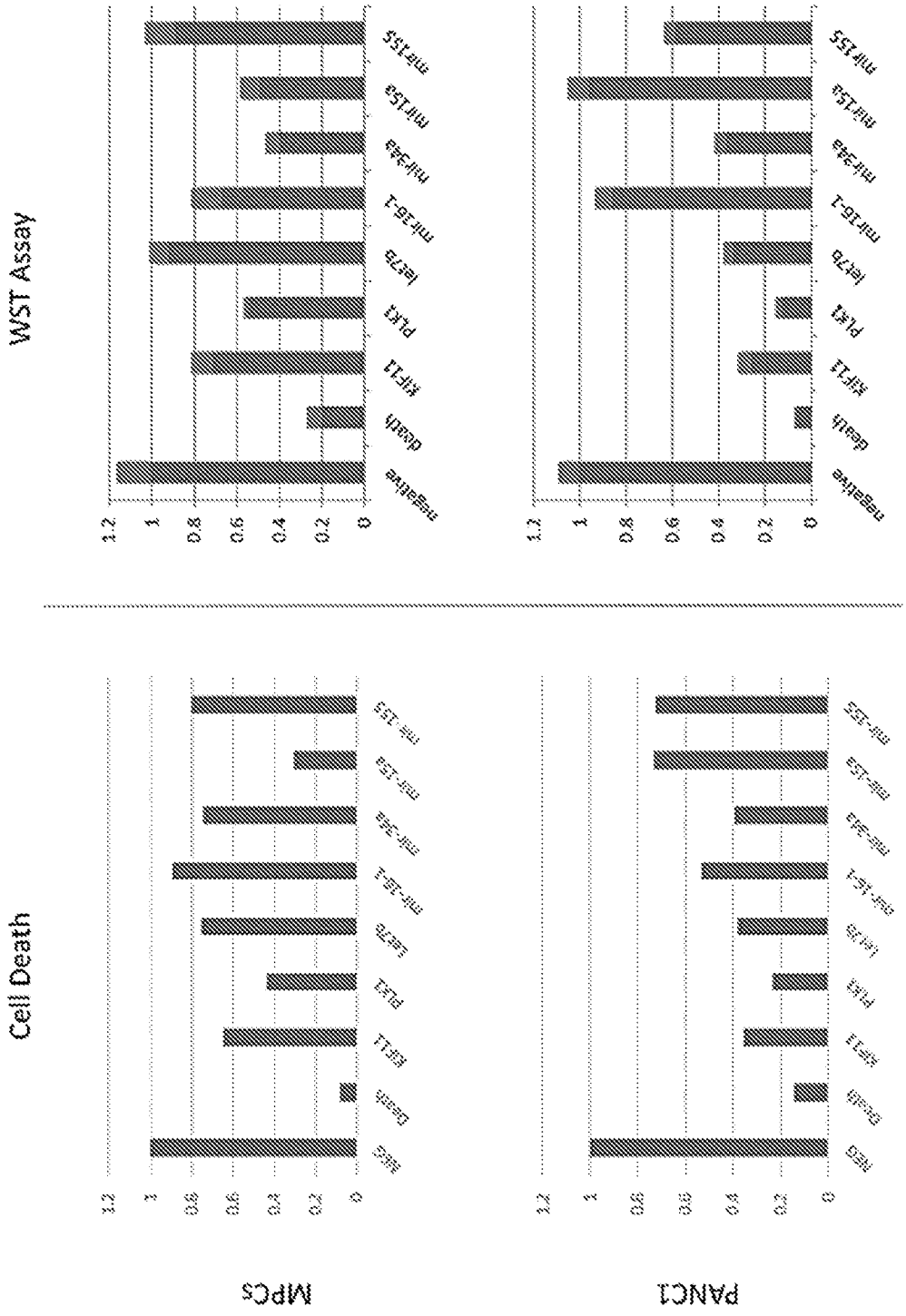

WST Assay
Cell Death
PANC1
MPCS
negative
death
KIF11
PLK1
tet7b
mir-16-1
mir-34a
mir-15a
mir-15b

FIGURE 18

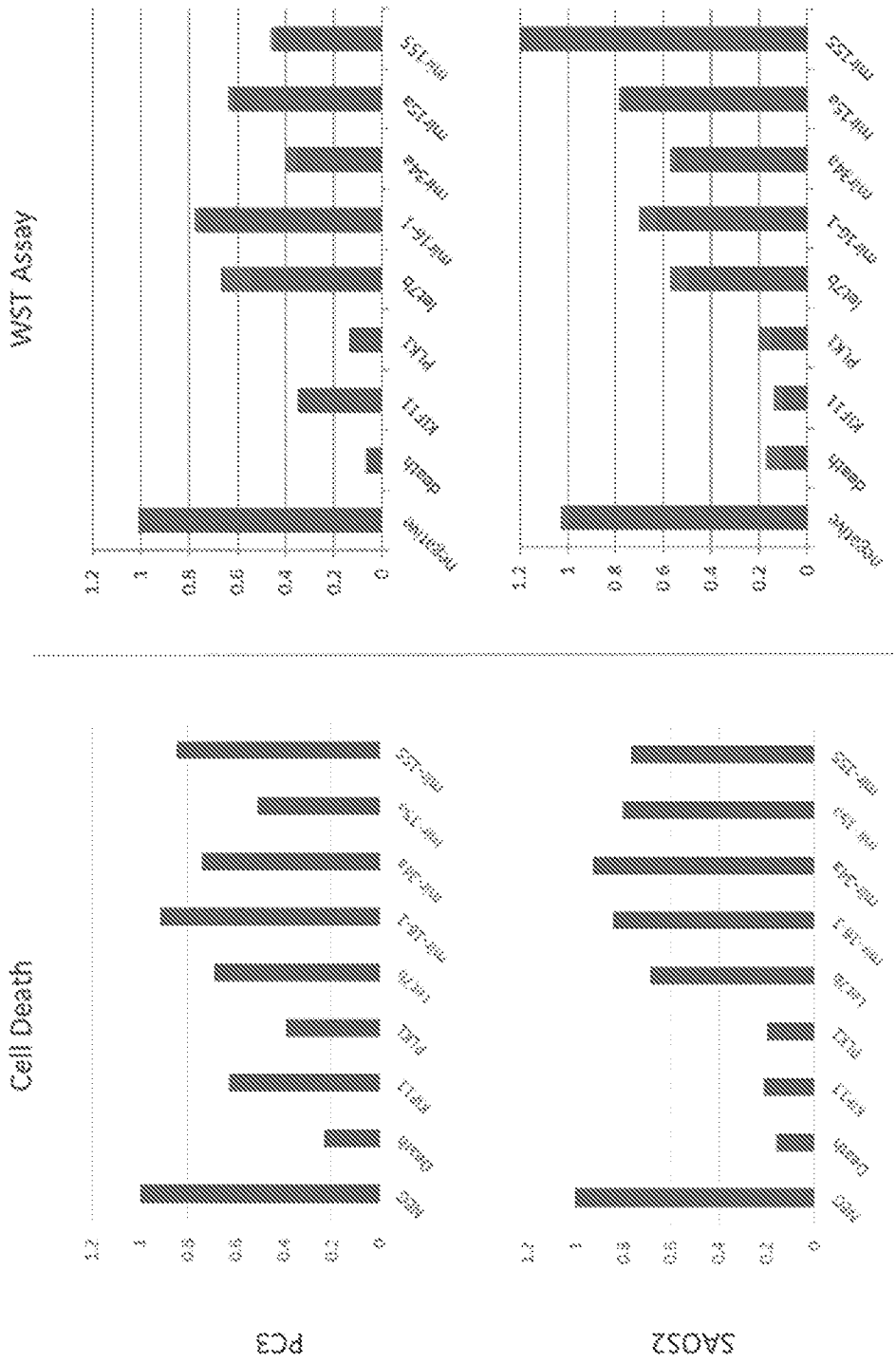


FIGURE 19

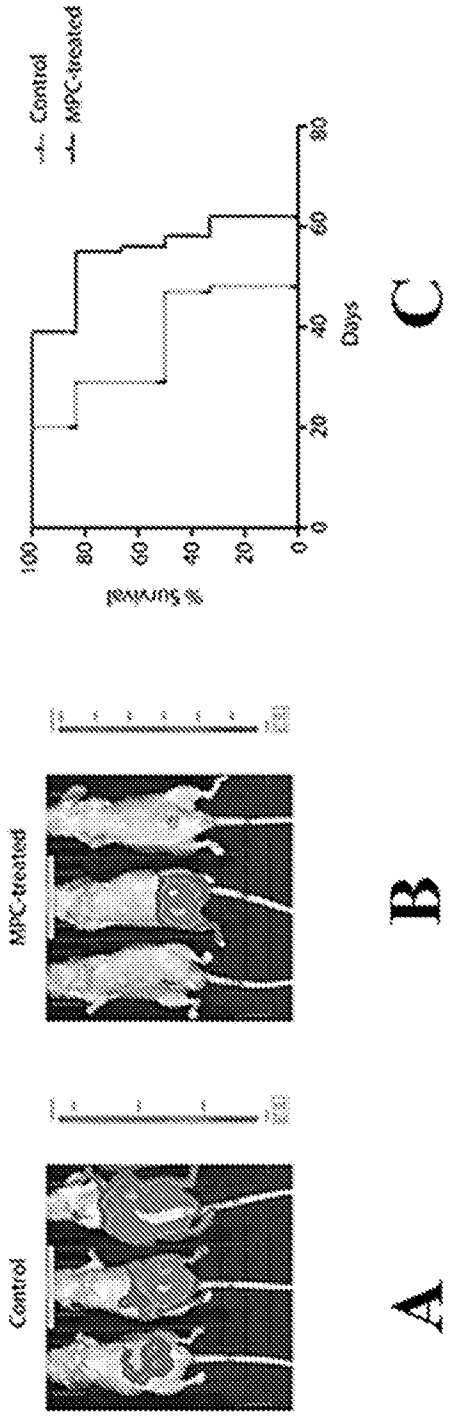


FIGURE 20

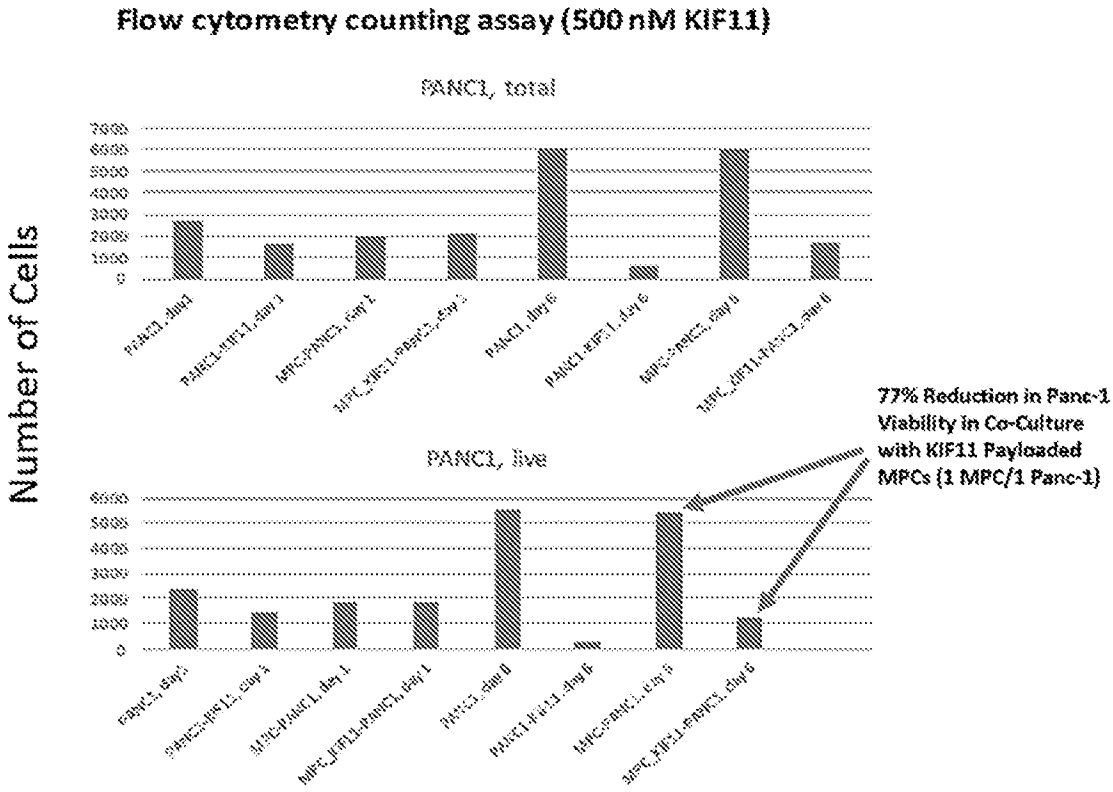

Flow cytometry counting assay (500 nM KIF11)
PANC1, total
Number of Cells
PANC1, live
77% Reduction in Panc-1 Viability in Co-Culture with KIF11 Payloaded MPCs (1 MPC/1 Panc-1)

FIGURE 21

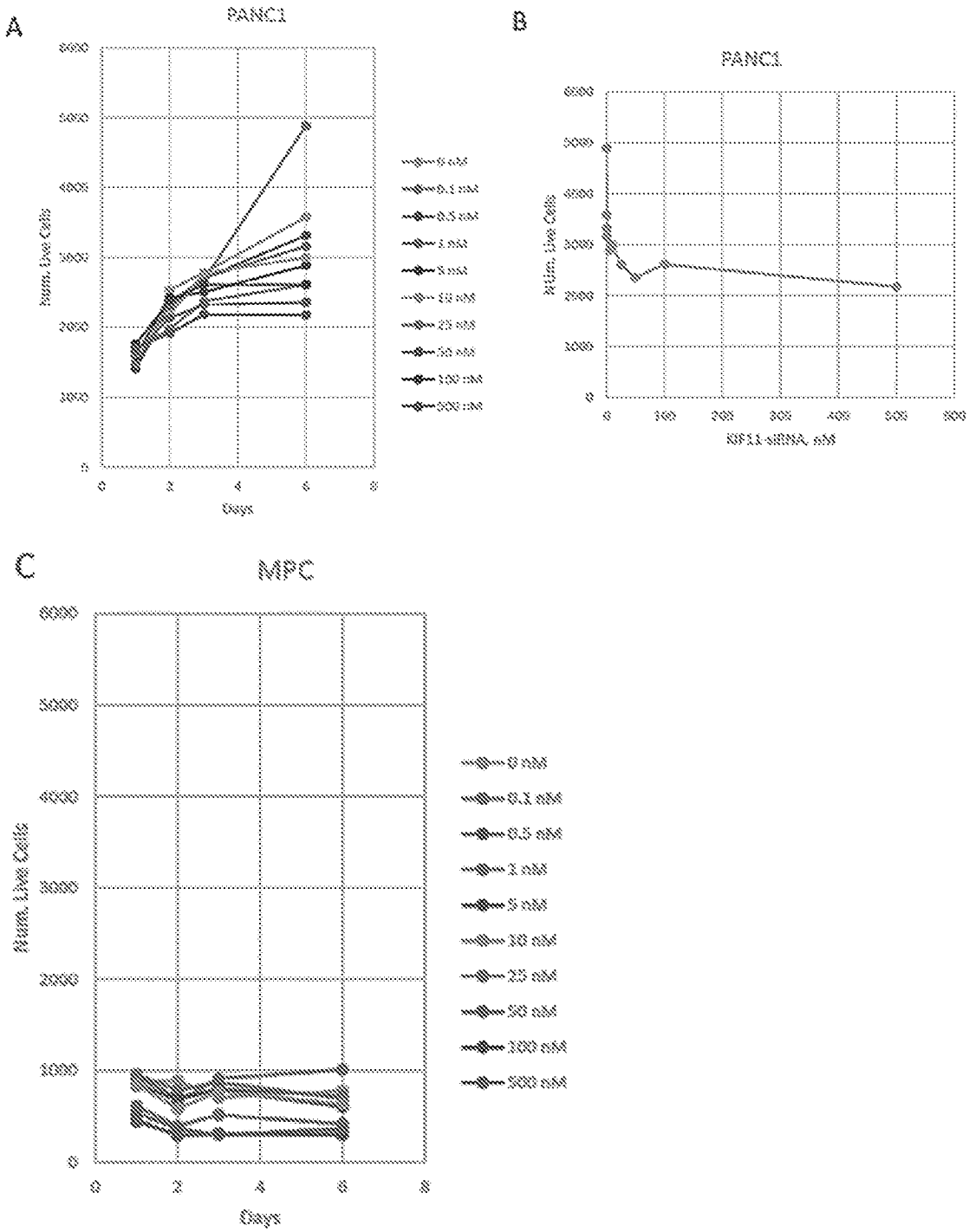

A
PANC1
Num. Live Cells
Days
B
PANC1
Num. Live Cells
KIF11 siRNA, nM
C
MPC
Num. Live Cells
Days
0 nM
0.1 nM
0.5 nM
1 nM
5 nM
10 nM
25 nM
50 nM
100 nM
500 nM

FIGURE 22

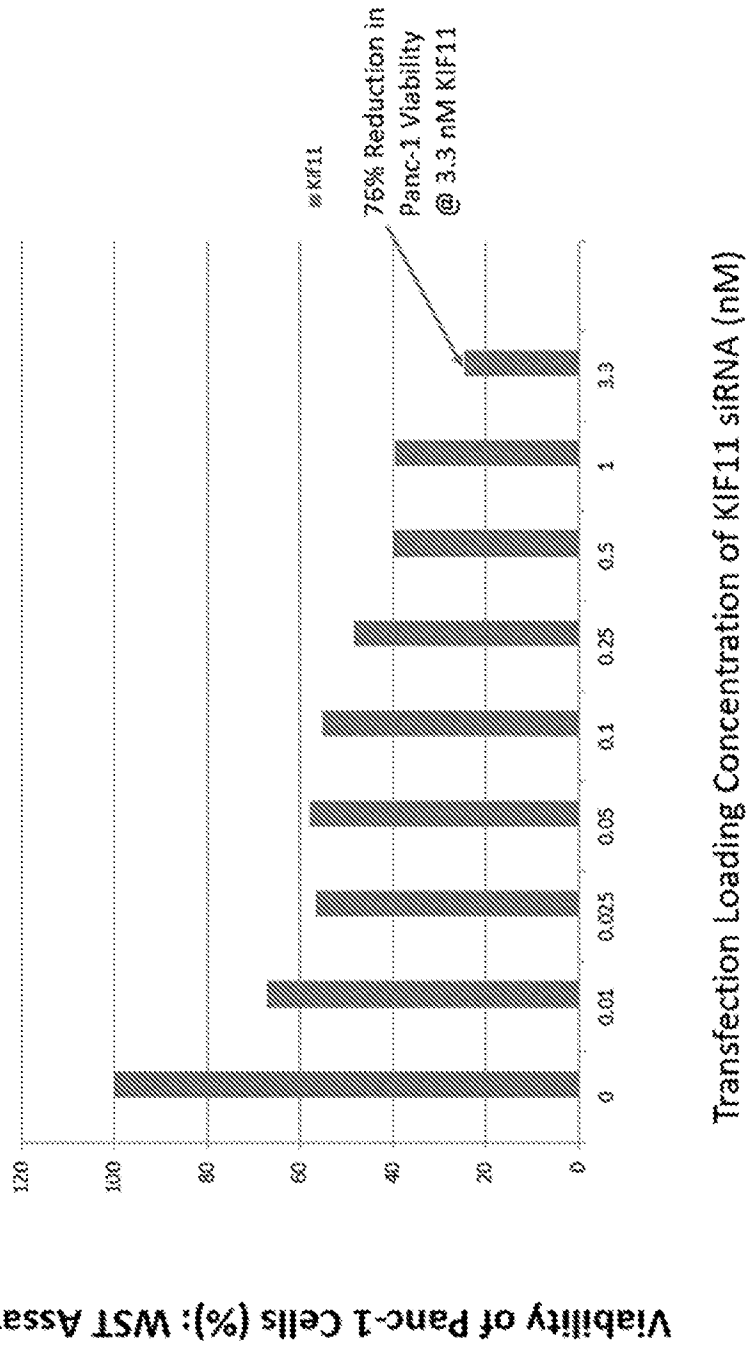

Viability of Panc-1 Cells (%): WST Assay
120
100
80
60
40
20
0
0
0.01
0.025
0.05
0.1
0.25
0.5
1
3.3
Transfection Loading Concentration of KIF11 siRNA (nM)
KIF11
76% Reduction in Panc-1 Viability @ 3.3 nM KIF11

FIGURE 23

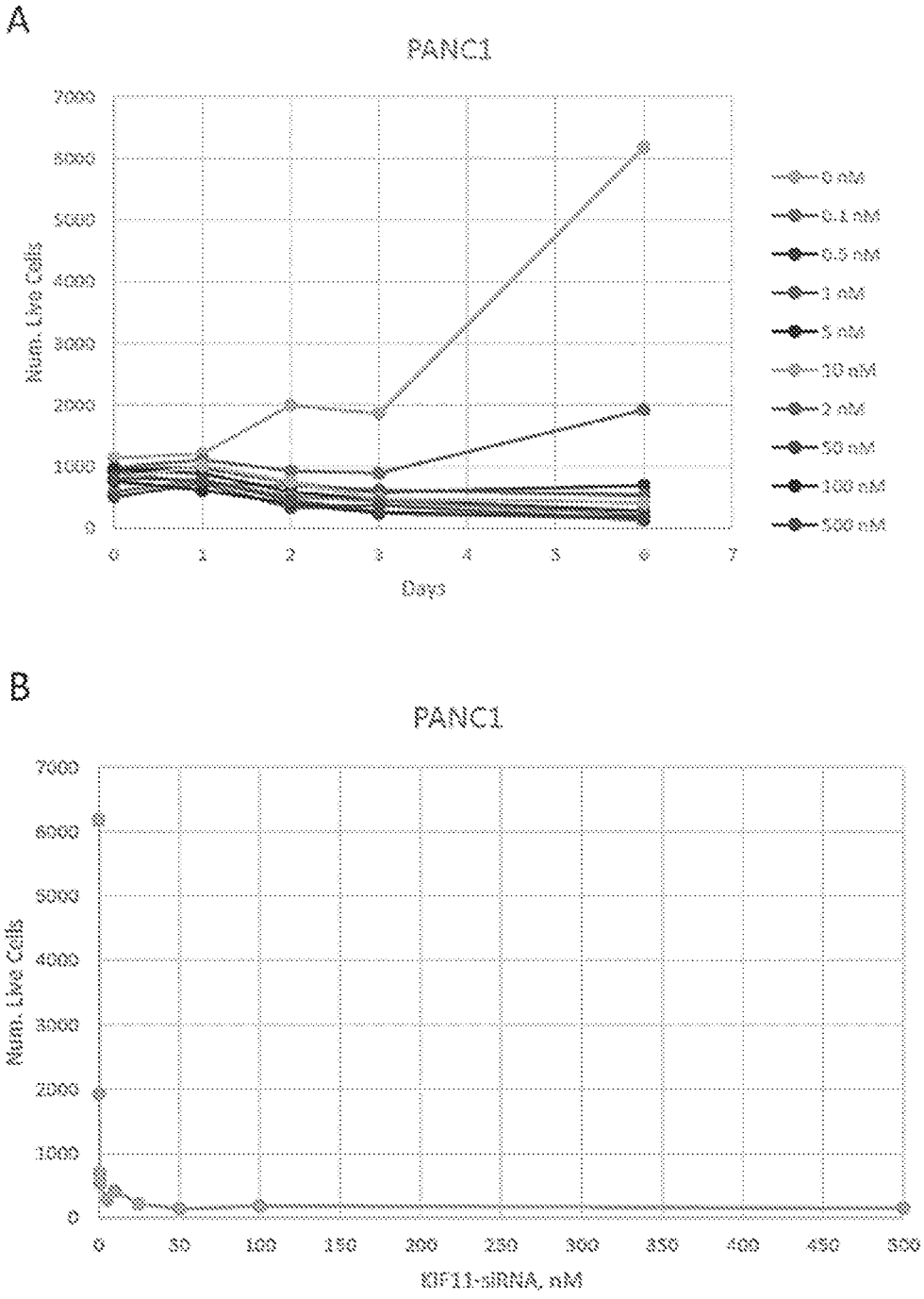
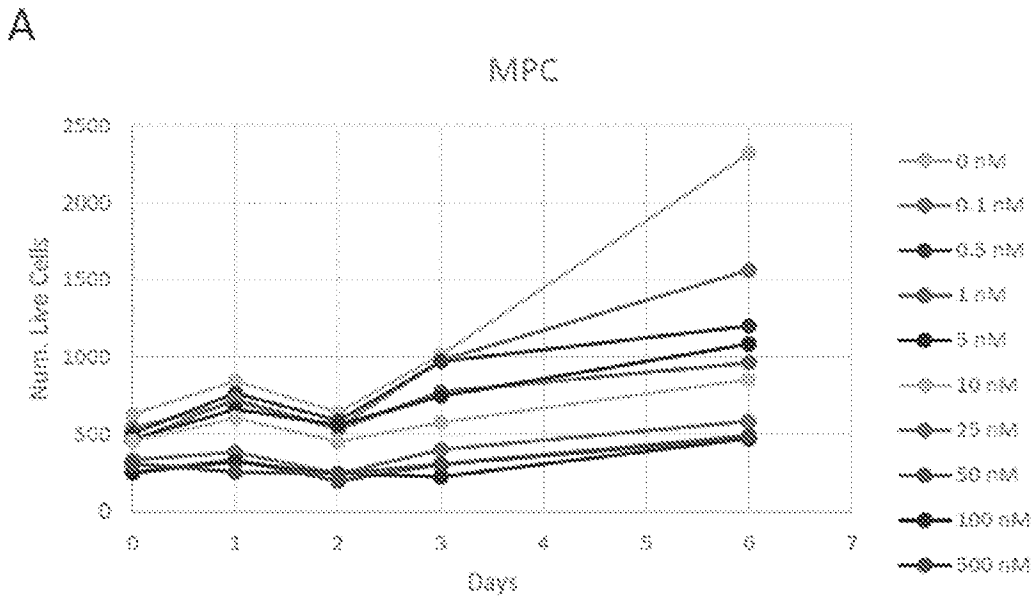

A
PANC1
Num. Live Cells
7000
6000
5000
4000
3000
2000
1000
0
0
1
2
3
4
5
6
7
Days
0 nM
0.1 nM
0.5 nM
1 nM
5 nM
10 nM
2 nM
50 nM
100 nM
500 nM
B
PANC1
Num. Live Cells
7000
6000
5000
4000
3000
2000
1000
0
0
50
100
150
200
250
300
350
400
450
500
KIF11-siRNA, nM

FIGURE 24


A
MPC
Num. Live Cells
2500
2000
1500
1000
500
0
0
1
2
3
4
5
6
7
Days
0 nM
0.1 nM
0.3 nM
1 nM
3 nM
10 nM
25 nM
50 nM
100 nM
300 nM

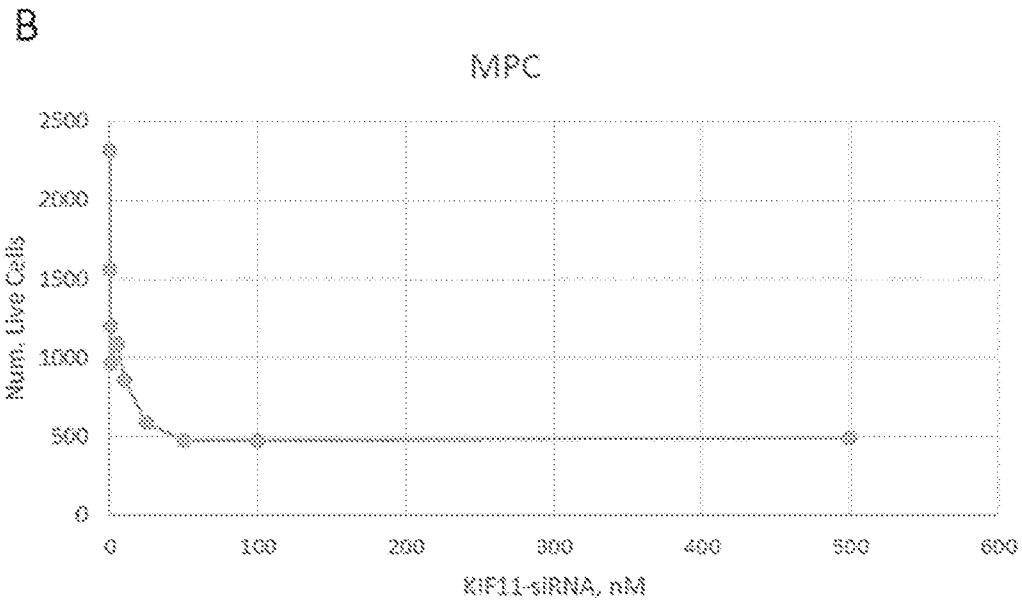

B
MPC
Num. Live Cells
2500
2000
1500
1000
500
0
0
100
200
300
400
500
600
KIF11-siRNA, nM

FIGURE 25

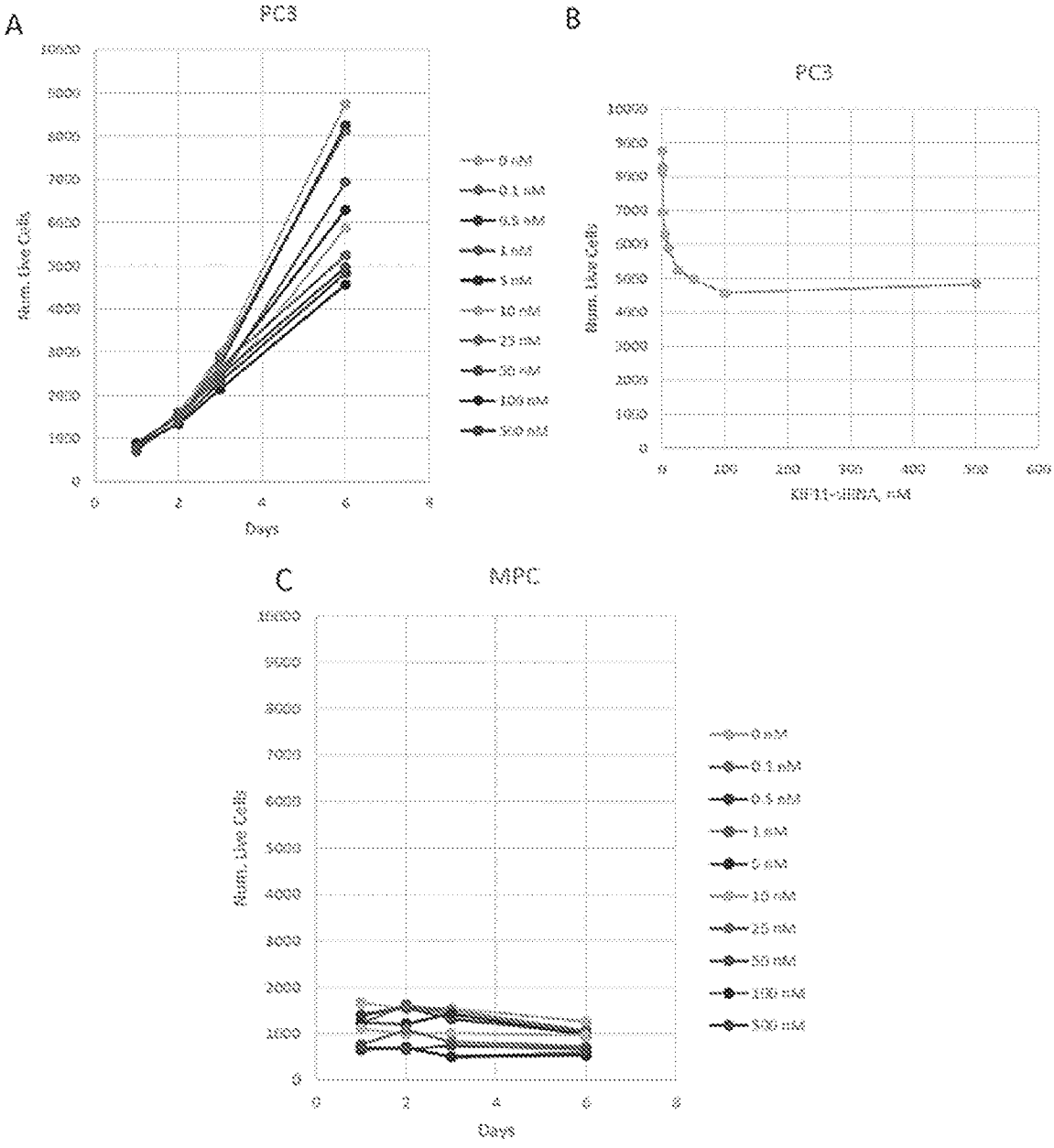

A
PC3
Num. Live Cells
Days
0 nM
0.1 nM
0.5 nM
1 nM
5 nM
10 nM
25 nM
50 nM
100 nM
500 nM
B
PC3
Num. Live Cells
KIF11-IN8A, nM
C
MPC
Num. Live Cells
Days

FIGURE 26

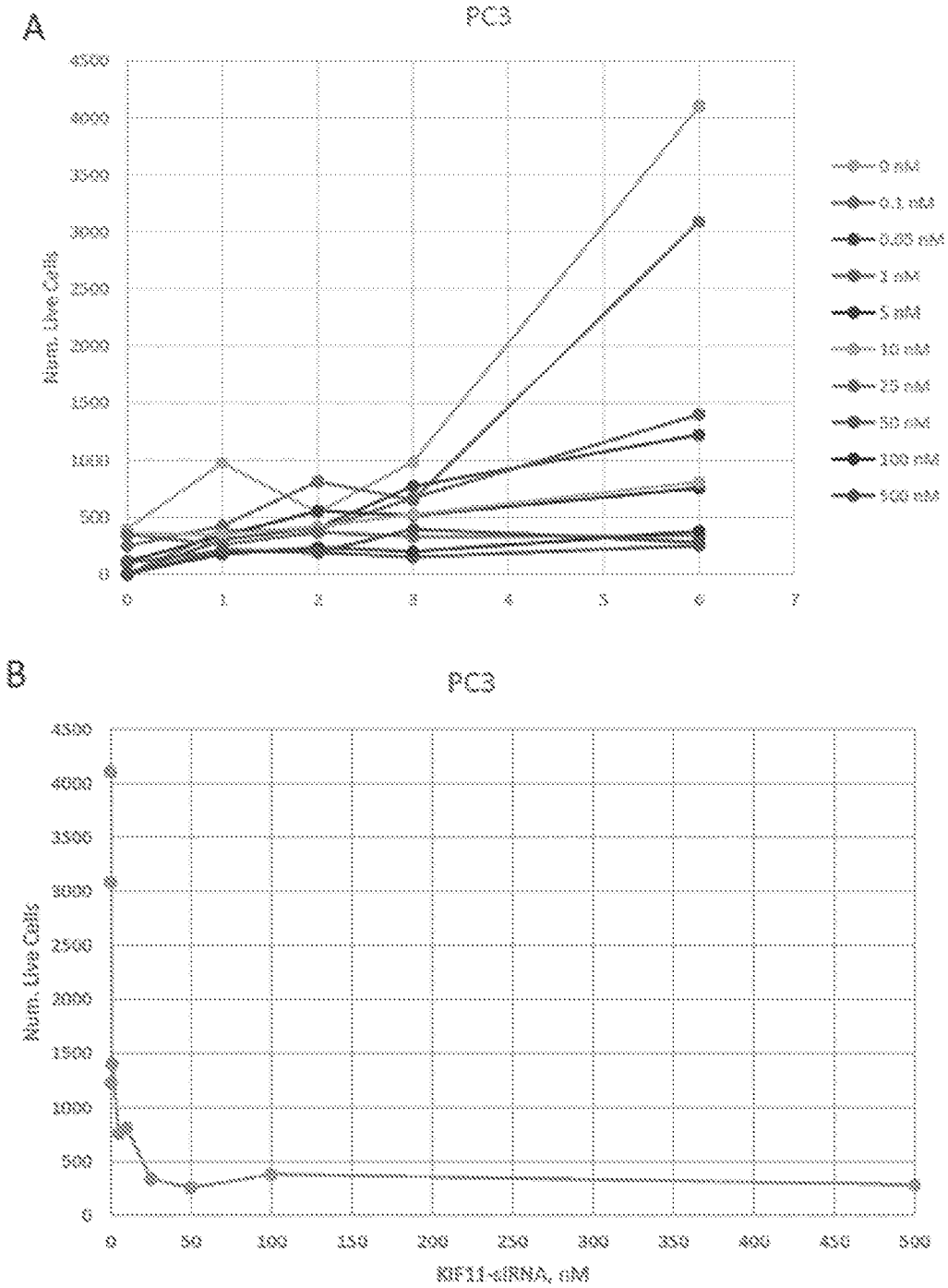
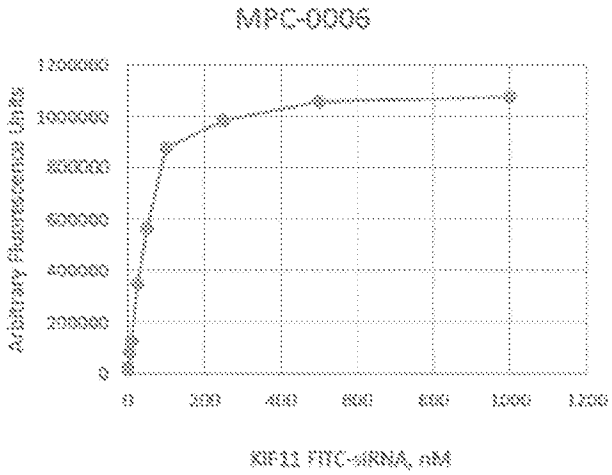
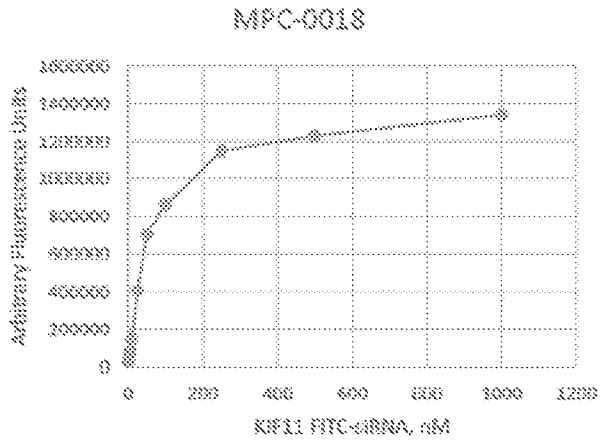

A
PC3
Num. Live Cells
0 nM
0.1 nM
0.05 nM
1 nM
5 nM
10 nM
25 nM
50 nM
100 nM
500 nM
B
PC3
Num. Live Cells
0
50
100
150
200
250
300
350
400
450
500
RF11-siRNA, nM

FIGURE 27

A



B



C

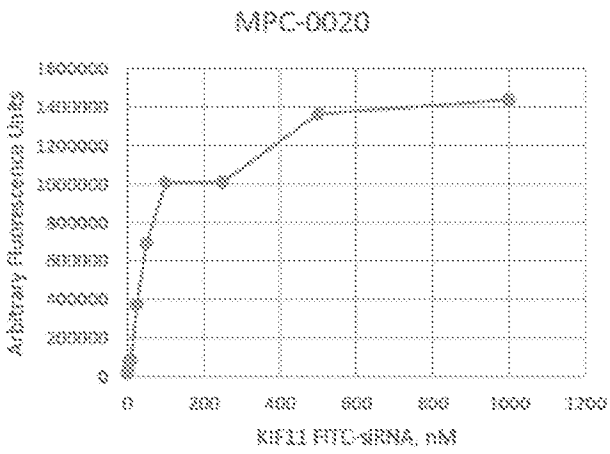


FIGURE 28

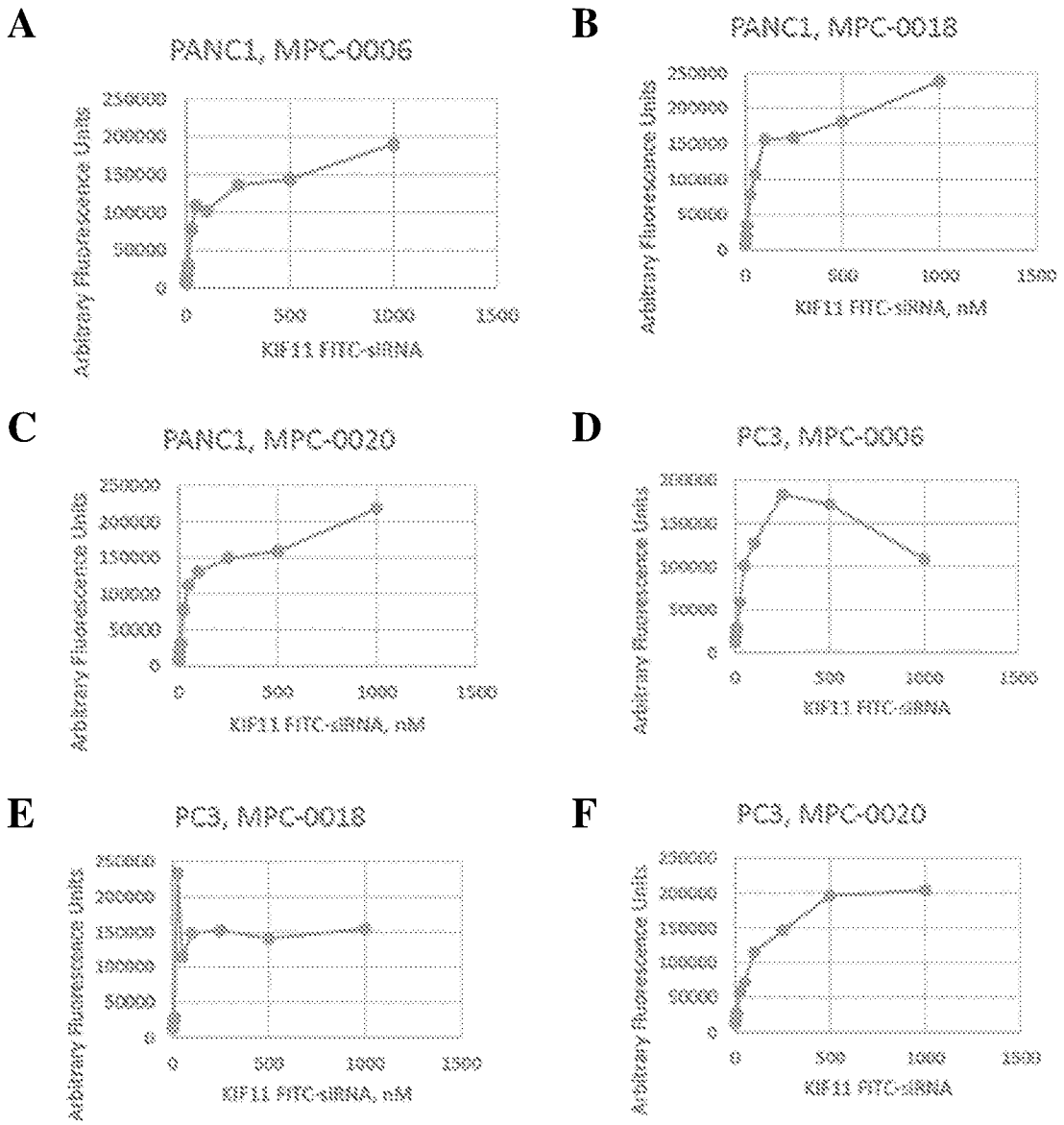

A
PANC1, MPC-0006
Arbitrary fluorescence Units
250000
200000
150000
100000
50000
0
0
500
1000
1500
KIF11 FITC-siRNA
B
PANC1, MPC-0018
Arbitrary fluorescence Units
250000
200000
150000
100000
50000
0
0
500
1000
1500
KIF11 FITC-siRNA, nM
C
PANC1, MPC-0020
Arbitrary fluorescence Units
250000
200000
150000
100000
50000
0
0
500
1000
1500
KIF11 FITC-siRNA, nM
D
PC3, MPC-0006
Arbitrary fluorescence Units
0
500
1000
1500
KIF11 FITC-siRNA
E
PC3, MPC-0018
Arbitrary fluorescence Units
250000
200000
150000
100000
50000
0
0
500
1000
1500
KIF11 FITC-siRNA, nM
F
PC3, MPC-0020
Arbitrary fluorescence Units
0
500
1000
1500
KIF11 FITC-siRNA

FIGURE 29

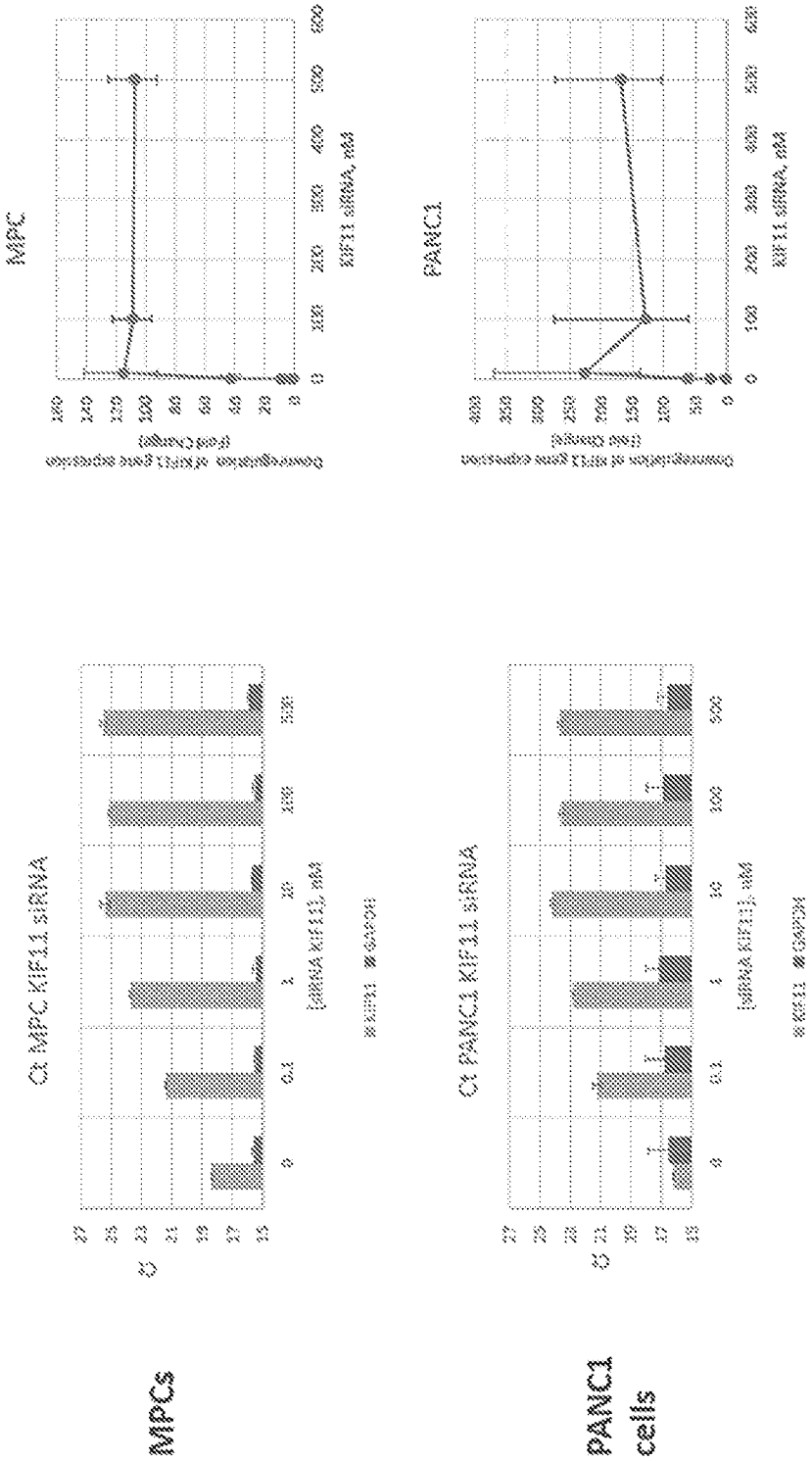


FIGURE 30

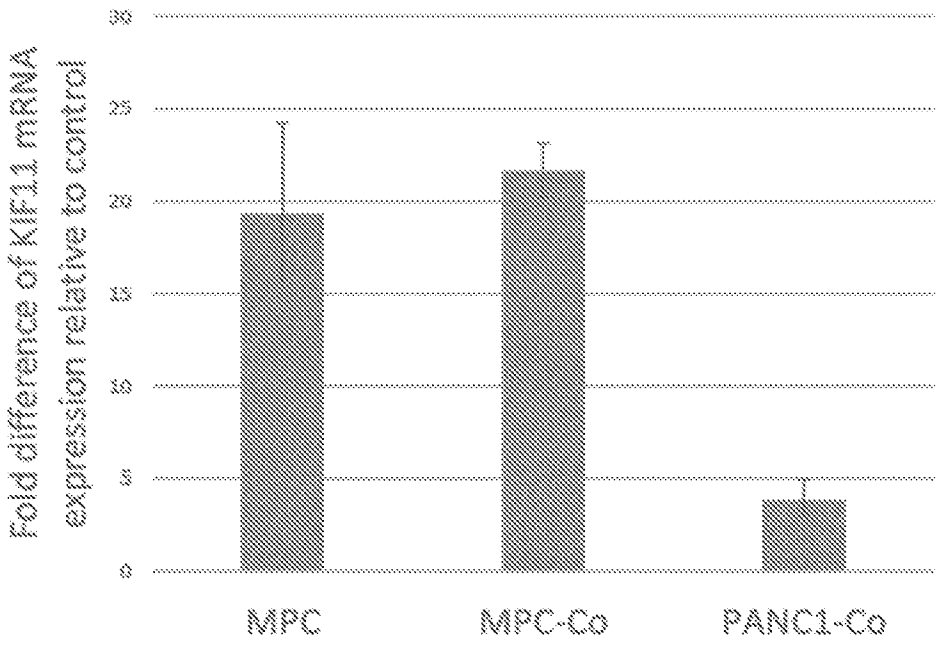

Fold difference of KIF11 mRNA expression relative to control
0
5
10
15
20
MPC
MPC-Co
PANC1-Co

FIGURE 31

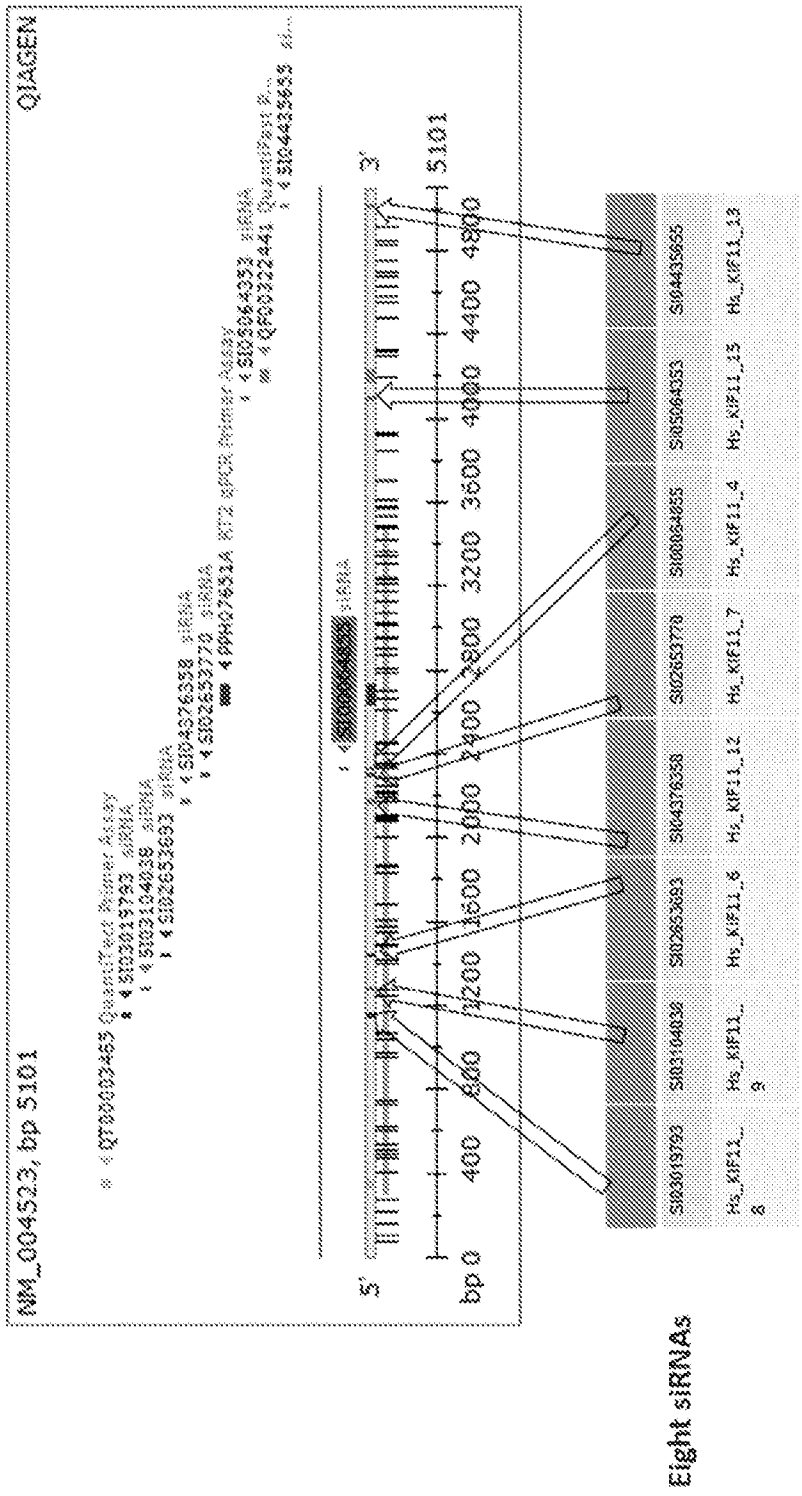

QIAGEN
NM_004523, bp 5101
5'
3'
bp 0
400
800
1200
1600
2000
2400
2800
3200
3600
4000
4400
4800
5101
Eight siRNAs

FIGURE 32

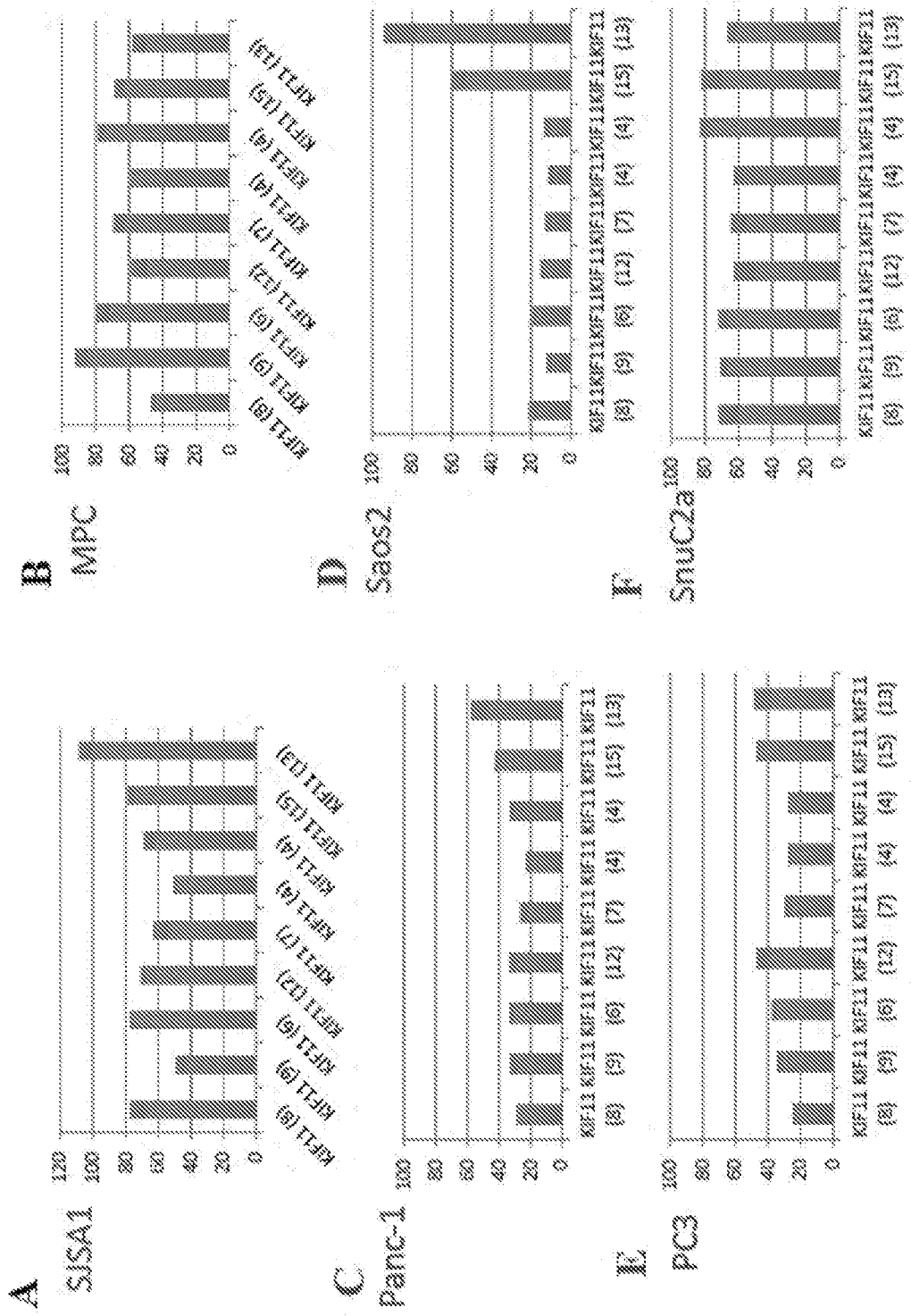


FIGURE 33

Top 10 down regulated genes in MPCs

ID	Trans Avg (log2)	Ctrl Avg (log2)	Trans Expressed	Ctrl Expressed	Fold Change	P-val	FDR P-val	Gene Symbol	Description
TCONS_00004471_hg1	5.92	10.13	1	1	-16.47	3.77E-06	4.09E-05	KIF11	kinesin family member 11
TCONS_00012245_hg1	13.24	14.93	1	1	-3.22	8.42E-06	0.0015	FSTL3, MIR199	follistatin like 3, microRNA 199
TCONS_00015269_hg1	8.98	10.6	1	1	-3.07	0.0002	0.1211	ADAM33	adhesion metalloprotease 33 [Source:HGNC Symbol;Acc:HGNC:24333]
TCONS_00008535_hg1	9.82	11.28	1	1	-3.02	2.56E-06	0.0211	EIF3	EIF3 CTD-domain protein [GO:0071986 eukaryotic translation initiation factor 3 complex]
TCONS_00013169_hg1	8.48	9.97	1	1	-2.98	6.55E-07	0.0047	SYP12	synaptophysin-like 1
TCONS_00007036_hg1	5.93	7.2	1	1	-2.59	0.0047	0.376	GINS1	GINS complex subunit 1 (Psf1 homolog)
TCONS_00007697_hg1	5.11	6.49	1	1	-2.58	0.0058	0.4947	CHP2	dysferlin related protein 2
TCONS_00010244_hg1	6.02	8.55	1	1	-2.54	2.17E-05	0.0083	CARKMT1	carnosine N-methyltransferase 1
TCONS_00010269_hg1	6.78	7.82	1	1	-2.22	0.0007	0.1858	MYO5C	myosin heavy chain superfamily protein containing a
TCONS_00013432_hg1	8.4	10.51	1	1	-2.13	4.37E-06	0.0188	MMGT2	high mobility group box 2

Top 10 down regulated genes in MSCs

ID	Trans Avg (log2)	Ctrl Avg (log2)	Trans Expressed	Ctrl Expressed	Fold Change	P-val	FDR P-val	Gene Symbol	Description
TCONS_00004471_hg1	5.19	9.51	1	1	-20.07	5.24E-05	0.0011	KIF11	kinesin family member 11
TCONS_00012245_hg1	11.34	15.22	1	1	-8.71	7.11E-05	0.1286	FSTL3, MIR199	follistatin like 3, microRNA 199
TCONS_00008181_hg1	5.52	7.28	1	1	-3.39	1.06E-06	0.0147	PLPPR4	phospholipid phosphatase related 4
TCONS_00011800_hg1	8.13	9.96	1	1	-3.13	6.47E-05	0.1336	SYP12	synaptophysin-like 1
TCONS_00008535_hg1	9.01	10.84	1	1	-3.11	6.46E-06	0.0946	EIF3	EIF3 CTD-domain protein [GO:0071986 eukaryotic translation initiation factor 3 complex]
TCONS_00011317_hg1	9.2	10.93	1	1	-2.93	0.0002	0.1359	TAPT1	transmembrane anterior posterior transformation 1
TCONS_00013169_hg1	8.59	10.36	1	1	-2.55	1.42E-06	0.0147	ADAM33	ADAM metallopeptidase domain 33 [Source:HGNC Symbol;Acc:HGNC:24333]
TCONS_00008166_hg1	5.46	6.73	1	1	-2.43	0.0028	0.3538	KCTD17	potassium channel tetramerization domain containing 17
TCONS_00011111_hg1	11.28	12.5	1	1	-2.34	0.0001	0.1336	TOR1	torsin family 1 member A (torsin 1)
TCONS_00016693_hg1	10.88	11.89	1	1	-2.31	0.0002	0.0823	GST1	glutathione S-transferase theta 1

FIGURE 34

INTERNATIONAL SEARCH REPORT

International application No
PCT/IB2018/001439

A. CLASSIFICATION OF SUBJECT MATTER
INV. C12N5/0775 C12N15/113 C12N15/87 A61P35/00
ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
C12N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
EPO-Internal, BIOSIS, WPI Data, COMPENDEX, EMBASE, FSTA

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2017/165641 A1 (UNIV COLUMBIA [US]; UNIV NEW YORK STATE RES FOUND [US]) 28 September 2017 (2017-09-28) pages 33-34, paragraph 69-70; claims 1-5	1-8, 12-24, 27-31
X	WO 2016/086020 A1 (CYTOSTORMRX LLC [US]) 2 June 2016 (2016-06-02) paragraph [0011] - paragraph [0013]; claim 10	1-8, 12-24, 27-31
X	WO 2010/111522 A2 (UNIV CALIFORNIA [US]; NOLTA JAN A [US]; OLSON SCOTT [US]; WIRTHLIN LOU) 30 September 2010 (2010-09-30) claims 1-67	1-8, 12-24, 27-31

-/--

[X] Further documents are listed in the continuation of Box C.

[X] See patent family annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
4 March 2019	06/05/2019

Name and mailing address of the ISA/	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Piret, Bernard

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Form PCT/ISA/210 (second sheet) (April 2005)

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2008/141177 A1 (UNIV COLUMBIA [US]; LEE FRANCIS Y [US]) 20 November 2008 (2008-11-20) claims 1-12	1-8, 12-24, 27-31
X	WO 2008/011133 A2 (UNIV COLUMBIA [US]; GAUDETTE GLENN [US]; COHEN IRA S [US]; POTAPOVA IR) 24 January 2008 (2008-01-24) claims 1-28	1-8, 12-24, 27-31
X	WO 2005/059111 A2 (UNIV COLUMBIA [US]; UNIV NEW YORK STATE RES FOUND [US]; BRINK PETER R) 30 June 2005 (2005-06-30) claims 1-23	1-8, 12-24, 27-31
A	BRINK PETER R ET AL: "In vivo cellular delivery of siRNA", I DRUGS: THE INVESTIGATIONAL DRUGS JOURNAL, CURRENT DRUGS LTD, GB, vol. 13, no. 6, 31 May 2010 (2010-05-31), pages 383-387, XP009511415, ISSN: 2040-3410	1-8, 12-24, 27-31
A	PETER R. BRINK ET AL: "Can gap junctions deliver?", BIOCHIMICA ET BIOPHYSICA ACTA (BBA) - BIOMEMBRANES, vol. 1818, no. 8, 1 August 2012 (2012-08-01), pages 2076-2081, XP055559150, AMSTERDAM, NL ISSN: 0005-2736, DOI: 10.1016/j.bbamem.2011.09.025	1-8, 12-24, 27-31
A	L. PEREBOEVA ET AL: "Approaches to Utilize Mesenchymal Progenitor Cells as Cellular Vehicles", STEM CELLS, vol. 21, no. 4, 1 July 2003 (2003-07-01), pages 389-404, XP055559170, ISSN: 1066-6099, DOI: 10.1634/stemcells.21-4-389	1-8, 12-24, 27-31
A	LINA RIMKUTE ET AL: "The role of neural connexins in HeLa cell mobility and intercellular communication through tunneling tubes", BMC CELL BIOLOGY, vol. 17, no. 1, 13 January 2016 (2016-01-13), XP055559171, DOI: 10.1186/s12860-016-0080-1	1-8, 12-24, 27-31

-/--

2

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	SHUO LIU ET AL: "Connexin43 Mediated Delivery of ADAMTS5 Targeting siRNAs from Mesenchymal Stem Cells to Synovial Fibroblasts", PLOS ONE, vol. 10, no. 6, 15 June 2015 (2015-06-15), page e0129999, XP055559173, DOI: 10.1371/journal.pone.0129999 -----	1-8, 12-24, 27-31
A	MIROSLAVA MATUSKOVA ET AL: "Intrinsic properties of tumour cells have a key impact on the bystander effect mediated by genetically engineered mesenchymal stromal cells : MSC-mediated gene therapy depends on tumour cells properties", JOURNAL OF GENE MEDICINE, vol. 14, no. 12, 1 December 2012 (2012-12-01), pages 776-787, XP055559184, US ISSN: 1099-498X, DOI: 10.1002/jgm.2684 -----	1-8, 12-24, 27-31

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No
PCT/IB2018/001439

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 2017165641 A1	28-09-2017	AU 2017238505 A1	06-09-2018
		BR 112018069090 A2	29-01-2019
		CA 3018150 A1	28-09-2017
		CN 108882705 A	23-11-2018
		EP 3432714 A1	30-01-2019
		JP 2019509297 A	04-04-2019
		KR 20180123052 A	14-11-2018
		SG 11201807223U A	27-09-2018
		WO 2017165641 A1	28-09-2017
WO 2016086020 A1	02-06-2016	EP 3223830 A1	04-10-2017
		JP 2017536127 A	07-12-2017
		US 2016317583 A1	03-11-2016
		WO 2016086020 A1	02-06-2016
WO 2010111522 A2	30-09-2010	AU 2010229872 A1	03-11-2011
		CA 2756670 A1	30-09-2010
		CN 102439136 A	02-05-2012
		EP 2411505 A2	01-02-2012
		US 2012114618 A1	10-05-2012
		US 2016263160 A1	15-09-2016
		WO 2010111522 A2	30-09-2010
WO 2008141177 A1	20-11-2008	NONE	
WO 2008011133 A2	24-01-2008	US 2010047216 A1	25-02-2010
		WO 2008011133 A2	24-01-2008
WO 2005059111 A2	30-06-2005	EP 1709194 A2	11-10-2006
		JP 2007514445 A	07-06-2007
		US 2007224176 A1	27-09-2007
		US 2011076773 A1	31-03-2011
		WO 2005059111 A2	30-06-2005

INTERNATIONAL SEARCH REPORT

International application No.
PCT/IB2018/001439

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.

2. ☐ As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of additional fees.

3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

4(completely); 1-3, 5-8, 12-24, 27-31(partially)

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.

☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.

☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. claims: 4(completely); 1-3, 5-8, 12-24, 27-31(partially)

   A method (or composition) for delivering an oligonucleotide into a target cell, the method comprising contacting a target cell with a mesenchymal lineage precursor or stem cell, wherein said cell has been modified by the introduction of the oligonucleotide or a vector expressing the oligonucleotide, wherein the modified mesenchymal lineage precursor cell or stem cells comprise sufficient levels of inhibitory oligonucleotide to effect transfer of between 0.5 to 5 nM of the inhibitory oligonucleotide to a target cell.

   ---

2. claims: 11(completely); 1-3, 5-10, 12-24, 27-31(partially)

   A method (or composition) for delivering an oligonucleotide into a target cell, the method comprising contacting a target cell with a mesenchymal lineage precursor or stem cell, wherein said cell has been modified by the introduction of the oligonucleotide or a vector expressing the oligonucleotide, wherein the oligonucleotide is a KIF11 siRNA.

   ---

3. claims: 1-3, 5-10, 12-24, 27-31(all partially)

   A method (or composition) for delivering an oligonucleotide into a target cell, the method comprising contacting a target cell with a mesenchymal lineage precursor or stem cell, wherein said cell has been modified by the introduction of the oligonucleotide or a vector expressing the oligonucleotide, wherein the oligonucleotide is a PLK1 siRNA.

   ---

4. claims: 25, 26(completely); 1-3, 5-8, 12-24, 27-31(partially)

   A method (or composition) for delivering an oligonucleotide into a target cell, the method comprising contacting a target cell with a mesenchymal lineage precursor or stem cell, wherein said cell has been modified by the introduction of the oligonucleotide or a vector expressing the oligonucleotide, wherein the target cell is a syncitial cell.

   ---

5. claims: 32-35(completely); 1-3, 5-8, 12-24, 27-31(partially)

   A method (or composition) for delivering an oligonucleotide into a target cell, the method comprising contacting a target cell with a mesenchymal lineage precursor or stem

cell, wherein said cell has been modified by the introduction of the oligonucleotide or a vector expressing the oligonucleotide, wherein the mesenchymal lineage precursor or stem cell has been modified using a viral vector.

---

6. claims: 36(completely); 1-3, 5-8, 12-24, 27-31(partially)

A method (or composition) for delivering an oligonucleotide into a target cell, the method comprising contacting a target cell with a mesenchymal lineage precursor or stem cell, wherein said cell has been modified by the introduction of the oligonucleotide or a vector expressing the oligonucleotide, wherein the mesenchymal lineage precursor or stem cell expresses CD46.

---

7. claims: 37-42(completely); 1-3, 5-8, 12-24, 27-31(partially)

A method (or composition) for delivering an oligonucleotide into a target cell, the method comprising contacting a target cell with a mesenchymal lineage precursor or stem cell, wherein said cell has been modified by the introduction of the oligonucleotide or a vector expressing the oligonucleotide, wherein the mesenchymal lineage precursor or stem cell has been treated to modify the cell surface glycans.
