Renal progenitors derived from human iPSCs engraft and restore function in a mouse model of acute kidney injury

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Acute kidney injury (AKI) is one of the most relevant health issues, leading to millions of deaths. The magnitude of the phenomenon remarks the urgent need for innovative and effective therapeutic approaches. Cell-based therapy with renal progenitor cells (RPCs) has been proposed as a possible strategy. Studies have shown the feasibility of directing embryonic stem cells or induced Pluripotent Stem Cells (iPSCs) towards nephrogenic intermediate mesoderm and metanephric mesenchyme (MM). However, the functional activity of iPSC-derived RPCs has not been tested in animal models of kidney disease. Here, through an efficient inductive protocol, we directed human iPSCs towards RPCs that robustly engrafted into damaged tubuli and restored renal function and structure in cisplatin-mice with AKI. These results demonstrate that iPSCs are a valuable source of engraftable cells with regenerative activity for kidney disease and create the basis for future applications in stem cell-based therapy.

A c u t e  k i d n e y  i n j u r y  ( A K I )  i s  a n  i m p o r t a n t  c a u s e  o f  m o r b i d i t y  a n d  m o r t a l i t y  w o r l d w i d e  a f f e c t i n g  m o r e  t h a n  1 3  m i l l i o n  p e o p l e ,  e s p e c i a l l y  i n  p o o r  r e s o u r c e  c o u n t r i e s 1,2.  H e r e ,  A K I  i s  g e n e r a l l y  d i s e a s e  o f  y o u n g  a d u l t s  a n d  c h i l d r e n  w h o  c o n t i n u e  t o  d i e  a s  a  c o n s e q u e n c e  o f  t h i s  d i s o r d e r  b e c a u s e  r e n a l  r e p l a c e m e n t  t h e r a p y  i s  n o t  a v a i l a b l e .  T h e r e f o r e  t h e  d e v e l o p m e n t  o f  n e w  s t r a t e g i e s  d i r e c t e d  t o  s u p p o r t  a n d  i m p r o v e  r e n a l  f u n c t i o n  i s  g r e a t l y  d e m a n d e d .  O n e  p r o m i s i n g  a p p r o a c h  a i m e d  a t  k i d n e y  r e g e n e r a t i o n  i s  c e l l - b a s e d  t h e r a p y  t h a t  r e q u i r e s  t h e  a v a i l a b i l i t y  o f  c e l l s  a b l e  t o  e f f i c i e n t l y  r e p l a c e  c e l l u l a r  c o m p o n e n t s  o f  t h e  d i v e r s e  r e n a l  c o m p a r t m e n t s .  S t u d i e s  h a v e  s u g g e s t e d  r e n a l  p r o g e n i t o r s ,  i s o l a t e d  f r o m  a d u l t  t i s s u e 3–6,  a s  a  p o s s i b l e  o p t i o n ,  a l t h o u g h  t h e i r  c l i n i c a l  t r a n s l a t i o n  i s  l i m i t e d  b y  t h e  i n a c c e s s i b i l i t y  o f  t h e  c e l l - s o u r c e  i n  p a t i e n t s ,  p o o r  e x p a n s i o n  c a p a b i l i t y  a n d  r e s t r i c t e d  d i f f e r e n t i a t i o n  p o t e n t i a l .  I P S C s ,  p l u r i p o t e n t  c e l l s  t h a t  p o s s e s s  e x t e n s i v e  s e l f - r e n e w a l  c a p a c i t y ,  o f f e r  t h e  p o s s i b i l i t y  o f  p a t i e n t - s p e c i f i c  t r e a t m e n t s  a n d  m a y  r e p r e s e n t  a  c e l l  t y p e  w o r t h  i n v e s t i g a t i o n  f o r  t h e r a p y .  U n d i f f e r e n t i a t e d  m u r i n e  i P S C s  h a v e  b e e n  d e s c r i b e d  t o  a t t e n u a t e  k i d n e y  i n j u r y  i n  i s c h e m i a  r e p e r f u s i o n - i n d u c e d  A K I  o n l y  w h e n  a d m i n i s t e r e d  v i a  i n t r a - a r t e r i a l  r o u t e ,  b y  p a r a c i n e  a c t i o n 7.  H o w e v e r ,  t h e  p l u r i p o t e n t  n a t u r e  o f  i P S C s  r a i s e  c o n c e r n s  o n  h i g h  r i s k  o f  t u m o u r  d e v e l o p m e n t  w h e n  t h e s e  c e l l s  a r e  a d m i n i s t e r e d  w i t h o u t  p r e - d i f f e r e n t i a t i o n .  A n  a t t r a c t i v e  a l t e r n a t i v e  c o n s i s t s  i n  t h e  d e r i v a t i o n  o f  r e n a l  p r o g e n i t o r s  f r o m  i P S C s .  I n i t i a l  a p p r o a c h e s  w i t h  m u r i n e  e m b r y o n i c  s t e m  c e l l s  ( E S C s )  a n d  i P S C s  h a v e  s h o w n  t h e  f e a s i b i l i t y  t o  g e n e r a t e  c e l l s  e x p r e s s i n g  i n t e r m e d i a t e  m e s e d e r m  ( I M )  a n d  M M  m a r k e r s 8–22  b y  e x p o s u r e  t o  s p e c i f i c  f a c t o r s  i n c l u d i n g  A c t i v i n  A  a n d  r e t i n o i c  a c i d  ( R A )  i n  c o m b i n a t i o n  w i t h  b o n e  m o r p h o g e n e t i c  p r o t e i n s  ( B M P s )  o r  G D N F ,  i m p o r t a n t  f a c t o r s  i n  k i d n e y  d e v e l o p m e n t 23.  R e c e n t l y ,  h u m a n  E S C s  a n d  i P S C s  h a v e  b e e n  d e s c r i b e d  t o  d i f f e r e n t i a t e  i n t o  n e p h r o g e n i c  c e l l s ,  r e n a l  p r o x i m a l  t u b u l a r  a n d  p o d o c y t e  l i n e a g e s 16–22,24.  H o w e v e r ,  r o b u s t  e n g r a f t m e n t  o f  h u m a n  i P S C - d e r i v e d  R P C s  i n  a n  a n i m a l  m o d e l  o f  A K I  a s s o c i a t e d  w i t h  r e s t o r a t i o n  o f  k i d n e y  f u n c t i o n  a n d  s t r u c t u r e  h a s  n e v e r  b e e n  s h o w n .

Results

Here, we report an efficient and reproducible two-stage protocol to obtain renal progenitors from human iPSCs (Fig. 1a). Initially, we performed differentiation studies on commercially available iPSCs obtained from human foreskin fibroblasts transfected with retrovirus encoding for Oct4, Sox2, Klf4, and c-Myc. In stage 1, to induce differentiation toward mesoderm (ME) and IM-like cells, we exposed undifferentiated iPSCs for 6 days to RA and...
to the small molecules RhoA inhibitor (CCG1423) and PI3K inhibitor (LY294002). The employment of small molecules has been suggested as a simple and direct method to address stem cell differentiation22,25. Activin A, known to stimulate the expression of IM markers in ESCs8,14, was added for two days starting on day 2. In stage 2, by day 6 up to 19, iPSCs were exposed to a cocktail of three nephrogenic factors FGF2, BMP7 and GDNF to commit cells towards MM. Undifferentiated iPSCs expressed pluripotency markers SSEA4, TRA-1-81, and Nanog (Fig. 1b) that progressively decreased after exposure to stage 1-medium. SSEA4 and TRA-1-81 were almost completely absent on day 6 whereas only few cells expressed a weak Nanog staining (Fig. 1b). By cytofluorimetric analysis, when we analyzed undifferentiated iPSCs, we found that the mean percentage of cells expressing TRA-1-81 was 88.5% ± 0.9 whereas it decreased at 24.7% ± 1.39 and 22.2% ± 0.93 respectively on day 12 and 19 of differentiation, exhibiting a very low fluorescence intensity per cell. As early as 3 days, we detected the expression of T (Brachyury), a transient marker of ME, that decreased on day 6 to a faint signal (Fig. 1b). Osr1, one of the earliest transcription factors specifying a multipotent population of IM cells giving rise to MM cells26, increased after 3 days and progressively decreased at 6 days (Fig. 1b). Other IM-specific markers including Wt1, Pax8 and Pax2 were observed by day 6 (Fig. 1c). At the same time, we found initial expression of Six2 and Sall1 (Fig. 1b) indicating the establishment of precursor MM fate. As a result of the exposure to stage 2-medium, on day 12 we found stable expression of Wt1, Pax8, Pax2, Six2 and Sall1 that decreased on day 19 except for Sall1 that maintained high expression. The developmental progression towards renal progenitor phenotype was confirmed by the presence of specific markers such as NCAM, CD133 and CD24 that during differentiation acquired cellular-specific localization. NCAM, normally expressed in the nephrogenic zone of developing kidney27 was induced by day 12 showing a clear pattern of expression at the cell periphery (Fig. 1d). CD133 and CD24, markers found in MM cells and glomerular progenitor cells28,29, were expressed on almost all the cells at any differentiation time considered (Fig. 1d) however, their distribution changed from a punctuated staining in undifferentiated iPSCs to a characteristic well-organized peripheral pattern by day 12 (Fig. 1d). Finally, we characterized the expression of Claudin1, a marker of glomerular epithelial cells and Aquaporin 1 (AQP1) and gamma-glutamyl transferase 1 (GGT1) markers of proximal tubular cells. By day 12, we observed a weak induction of Claudin1 and an
intense AQP1 staining (Fig. 1d) whereas GGT1 was weakly expressed on day 19 (Fig. 1d). Differentiated iPSCs did not express alpha-feto-protein (AFP), Pax6 and Nkx2.5, markers of endoderm, neuroectoderm and cardiac mesoderm, respectively (Fig. 1e).

Next, we tested the robustness of the two-step differentiation protocol on iPSCs generated in our laboratory by reprogramming human fibroblasts using the STEMCCA Cre-excisable polycistronic lentivirus with excisable reprogramming cassette. Specifically, the generated iPSC clones showed all the properties of the pluripotent stem cells in terms of expression of pluripotency markers, karyotype and ability to differentiate both in vitro and in vivo into derivatives of the three germ layers (Supplementary Fig. 1a-f). The fully characterized clone IV was used to test the differentiation protocol. Immunofluorescence analysis showed a pattern of protein expression similar to that observed with commercial iPSCs. Indeed, down-regulation of the pluripotency marker Nanog was observed going from undifferentiated cells to day 6 of differentiation. Conversely, upregulation of markers specific to the early ME, which included T, for the IM such as Lhx1, Osr1 (Supplementary Fig. 2a) and later Wt1, Pax8 and Pax2, and for the MM such as Six2 and Sall1 was detected (Supplementary Fig. 2b). To better define the phenotype of iPSC clone IV-derived RPCs, we evaluated the co-expression of several markers which defines renal commitment. Our data show the specification into a homogeneous IM cell population from day 3 to 6 as demonstrated by Lhx1/Osr1 co-expression (Fig. 2a). Six2 co-staining with Pax2, Pax8 and Wt1 at day 6 and 12 confirms differentiation toward MM cell phenotype (Fig. 2b). Protein expression of CD24 - already present in undifferentiated cells - Claudin1 and GGT1 was evident throughout the differentiation protocol (Fig. 2c). We next performed gene expression analysis by real-time PCR and found that the expression of gene and corresponding protein paralleled over time (Fig. 2d). Collectively, our results demonstrate that the protocol we established supports the efficient commitment of the two different human iPSC cell lines towards RPCs and provides the basis for functional studies.

The derivation of RPCs from iPSCs offers new platforms for drug screening and for in vitro tissue engineering however, the major challenge resides on their employment in regenerative medicine to in vivo generate new functional cells. Therefore, we next assessed the renoprotective potential of human iPSC-derived RPCs, obtained on day 12, in the murine model of cisplatin-induced AKI which is characterized by severely compromised renal function and tubular damage 4 days after drug injection. Intravenously infused iPSC-derived RPCs were found in the kidney as early as 24 h. At 4 days, a sustained number of iPSC-derived RPCs was observed in the cortical region of the kidney of cisplatin mice as proved by the presence of human mitochondria (h-Mito), human nuclear antigen (HNA) or PKH26 cell tracker (Fig. 3a and b). Human cells were found integrated into murine tubuli identified by wheat germ agglutinin (WGA)-lectin staining and cortical areas showed extensive engraftment accounting for 42% ± 12 HNA positive cells per high power field (HPF). By immunofluorescence staining with AQP1, we provided evidence that human PKH26 positive iPSC-derived RPCs mainly engrafted in proximal tubuli and expressed a marker of proximal tubular epithelial cells.
imal tubular cells (Fig. 3b, left). This result was confirmed by immunoperoxidase technique showing the co-localization of the signal for human mitochondria and AQP1 in proximal tubular cells (Fig. 3b, central and right panels). No human iPSC-derived RPCs were found in peanut agglutinin (PNA)-lectin positive distal tubuli while scattered human cells were observed in AQP3-positive collecting ducts. Engrafted human iPSC-derived RPCs, co-stained for the proliferation marker Ki-67 and HNA, showed 30% of human cells proliferating in proximal tubuli (Supplementary Fig. 3). Tropism of transplanted human cells for the kidney was supported by the observation that a paltry number of engrafted cells was found in other organs including liver (0.26 ± 0.06 cells/HPF), lung (0.26 ± 0.06 cells/HPF), heart (0.22 ± 0.08 cells/HPF) and spleen (0.16 ± 0.06 cells/HPF) 24 h after cell infusion (Supplementary Fig. 4) while no human cells were found at 4 days in all the organs studied. Mice that experienced renal engraftment exhibited a significant improvement of renal function to the extent of a 55% reduction of blood urea nitrogen (BUN) levels as compared with cisplatin-treated mice given saline (Fig. 3c). Renal histology of mice receiving human iPSC-derived RPCs was significantly improved, as shown by the remarkable reduction of renal tubular damage in terms of cell swelling, cast deposition, loss of brush border and cell necrosis compared to mice receiving saline (Fig. 3d). Proliferation of tubular cells, assessed by Ki-67 expression, was increased primarily in the proximity of engrafted human cells compared to cisplatin mice given saline (Ki-67 positive cells/HPF: iPSC-derived RPCs, 14 ± 0.86 vs saline, 8 ± 0.06; p < 0.01).

Injection of human undifferentiated iPSCs, here used as control, failed to exert any protective effect on renal function (Fig. 3c) and renal histology (Fig. 3d). Undifferentiated iPSCs were not present in the kidney and in the other organs at both 24 h and 4 days after cisplatin injection. No signs of inappropriate differentiation and tumor formation by iPSC-derived RPCs were noted in the renal tissues either at 4 days or 8 weeks after cisplatin injection.

Herein, we defined a successful differentiation protocol to efficiently derive renal progenitor cells from human iPSCs that consisted of the exposure of the cells to specific small molecules and growth factors resulting in the acquisition of markers sequentially expressed in different phases of kidney development. During the first stage of the differentiation process, we obtained a cell population with wide-spread expression of IM markers, whereas later on cells acquired MM phenotype consistent with the presence of early renal progenitor cells.
fibroblasts were plated on a plastic 35-mm culture plate. The next day cells were supplemented with 10% FBS (Lonza), 2 mM Glutamine (Invitrogen), 1 mM Sodium (NHDF- Neo) were purchased from Lonza and were maintained in MEM medium in the presence of Rock inhibitor (Sigma). Human dermal neonatal fibroblast cells derived from human dermal neonatal fibroblasts (MEFs), used as feeder cells for iPSCs, were isolated from embryos of 13.5 days (E13.5) from CD1 mice and cultured in DMEM medium supplemented with 0.1% penicillin and streptomycin. For passaging, human iPSCs were incubated with 1% trypsin-EDTA (Gibco) until the first colonies started to appear and then every day. iPSC colonies were manually picked about 30 days post-infection, transferred into 0.5 ml of human iPSC-medium. The medium was changed every other day.

**Methods**

Cell culture. Commercially available human iPSCs (SC101A-1) were purchased from Stem Biosciences International and maintained in DMEM/F12 medium containing 20% Knockout Serum Replacement (Invitrogen), 0.1 mM nonessential amino acids (Invitrogen), 0.1 mM 2-mercaptoethanol (Sigma), 10 ng/ml human basic fibroblast growth factor (FGF2, PeproTech) and 1% penicillin and streptomycin. Mouse embryonic fibroblasts (MEFs), used as feeder cells for iPSCs, were isolated from embryos of 13.5 days (E13.5) from CD1 mice and cultured in DMEM medium containing 10% FBS (Invitrogen), 0.1 mM nonessential amino acids (Invitrogen) and 1% penicillin and streptomycin. For passaging, human iPSCs were incubated with accutase (Millipore). Cell suspension was seeded on mitomycin C-inactivated MEFs in the presence of Rock inhibitor (Sigma). Human dermal neonatal fibroblast cells (NHDF- Neo) were purchased from Lonza and were maintained in MEM medium supplemented with 10% FBS (Lonza), 2 mM Glutamine (Invitrogen), 1 mM Sodium Pyruvate (Invitrogen) and 1% penicillin and streptomycin.

**Derivation and characterization of human iPSC clone IV.**

**Generation of human iPSC cell line.** Human iPSCs were derived from human dermal neonatal fibroblasts by StemCCA Cre-Excisable Constitutive Polycistronic (OKMS) lentivirus (Millipore) following manufacturer’s instructions. Briefly, neonatal fibroblasts were plated on a plastic 35-mm culture plate. The next day cells were infected with 200 MOI of virus in human fibroblast medium in the presence of polybrene (5 μg/ml). The medium was replaced 24 hours after transduction with fresh virus-containing medium for a second cycle of infection. Six days after the first infection, the plaques were trypsinized and plated at different densities on MEF-feeder coated dishes (eg. 20,000 or 50,000 per 10-cm culture dish). The next day the medium was changed to human iPSC-medium. The medium was changed every other day until the first colonies started to appear and then every day. iPSC colonies were manually picked about 30 days post-infection, transferred into 0.5 ml of human iPSC-medium and seeded to small clumps by pipetting and seeded onto a new MEFs feeder in a well of six well-plate. The colonies were mechanically split for 3 passages and then cultured as described above. Clone IV has been chosen for characterization and differentiation studies.

**Karyotyping.** Metaphase spreads were prepared after treatment with 10 μg/ml Colcemid and processed for karyotype analysis. At least 20 metaphases of each sample were counted. Karyotype analysis was performed in collaboration with the Genetic Medicine Laboratory of Azienda Ospedaliera Papa Giovanni XXIII, Bergamo (Italy).

**Embryoid body (EB) formation.** For EB formation, human iPSCs clone IV were harvested by treating with DMEM/F12 containing 1 mg/ml collagenase IV at 37 °C. When the perimeter of the colonies appeared highlighted and the cells started to peal up collagenase was removed and the colonies were scraped in human iPSCs medium and centrifuged. The clumps of the cells were suspended and cultured in DMEM/F12 medium containing 20% Knockout serum replacement, 0.1 mM nonessential amino acids, 0.1 mM 2-mercaptoethanol and 1% penicillin and streptomycin and then transferred to low attachment dish. The medium was replaced every other day.

**After 8 days as floating culture, EBs were transferred onto gelatin-coated plate and cultured in the same medium for another 8 days. At day sixteen the plates were fixed with 4% PFA solution and processed by immunostaining.**

**Teratoma formation and analysis.** Undifferentiated human ipSC clone IV (10/site) were injected subcutaneously into dorsal flank of NOD-SCID mice (n = 3). Eight weeks after injection, teratomas were dissected, fixed in formalin, embedded in paraffin and stained for hematoylin and eosin.

**Human iPSC differentiation into renal progenitor cells.** To induce differentiation toward mesoderm (ME) and intermediate mesoderm (IM)-like cells human iPSC-growth medium was replaced with basal DMEM/F12 medium containing 5% FBS (HyClone), 0.1 mM nonessential amino acids, 0.1 mM 2-mercaptoethanol supplemented with 0.1 μM All-Trans Retinoic Acid (Sigma), 1 μM CCG4123 (RhoA Inhibitor, Vinci Biochem) and 5 μM LY294002 (PI3K inhibitor) for 6 days. Ten ng/ml Activin A (Prepotech) was added for two days starting on day 2. To induce metanephrine mesenchyme (MM) specification and renal progenitor phenotype by day 6 up to 19 the medium was replaced with fresh basal medium containing niphrogenic factors: 50 ng/ml BMP7 (Prepotech), 10 mg/ml FGFB and 15 ng/ml GDNF (Abcam). The medium was changed every 2 days.

**RNA isolation and real-time PCR assays.** Total RNA was isolated by Rneasy plus Mini Kit (Qagen). First-strand cDNA was produced from 2 μg total RNA using Transcript II First-Strand Synthesis System Kit (Invitrogen) using a mixture of oligo-dT and random hexamers oligonucleotides following manufacturer’s instructions. The purity of human iPSCs clone IV was evaluated by TaqMan gene expression assay (Life Technologies) using predesigned probes for pluripotent and undifferentiated gene markers (Supplementary Table 1) according to the supplier’s instructions. RNA from commercially available human iPSCs was used as reference sample and the gene expression level was normalized to RnAF1 housekeeping gene.

**For gene expression analysis in renal induction experiments TaqMan Array Human Endogenous Control Plate (Life Technologies) was used in order to evaluate the best housekeeping gene and ELF1 gene was chosen to normalize the expression level. Real-time PCR assays were performed using Power Syber Green PCR master mix (Life Technologies). The primers employed are listed in Supplementary Table 2. The AAC1 technique was used to calculate cDNA content in each sample using the cDNA expression of the pluripotent state (d0) as a calibrator. The experiments were performed in triplicate and the results were expressed as mean ± SD.**

**Cell immunostaining.** Cells were fixed in 4% PFA in PBS for 30 min at room temperature. When appropriate, cells were permeabilized in 0.5% Triton X-100 for 30 min at room temperature followed by incubation in blocking solution (5% BSA). The primary antibodies were diluted according to manufacturer’s recommendations and incubated overnight at 4 °C, followed by incubation with the appropriate secondary antibody for 1 h at room temperature. Primary antibodies included anti-OCT4 (Santa Cruz); anti-NANOG (Santa Cruz); anti-TRA-1-60 (Millipore); anti-TRA-1-81 (Millipore); anti-SSEA3 (Santa Cruz); anti-SSEA4 (Santa Cruz); anti-AFP (Santa Cruz); anti-s-SMA Cy3- conjugated (Sigma); anti-β III Tubulin Alexa Fluor 488- conjugated (Millipore), anti-T (Abcam), anti-Osr1 (Santa Cruz), anti-Lhx1 (Abcam), anti-W1 (R&D System), anti-Pax2 (Novus biologicals), anti-Six2 (Proteintech, Abnova), anti-Pax2 (Invitrogen), anti-Sall1 (R&D System), anti-CD24 (Abcam), anti-NCAM (Millipore), anti-AQPI (Abcam), anti-GF1 (Santa Cruz), anti-Claudini (Thermo Scientific), anti-Pax6 (Novus biologicals) and anti-Nkx2.5 (Thermo Scientific). For CD131 staining, cells, after fixing, were blocked with 1.5% normal goat serum for 1 h at RT and incubated with anti-CD131 antibody (Miltenyi Biotech, AC133/1) for 1 h at 37 °C. Subsequently the cells were washed twice with PBS. Alexa Fluor 488 secondary antibodies (Molecular Probes) were used followed by nuclear counterstaining with 4',6-diamidino-2-phenylindole (DAPI) for 10 minutes at room temperature. Images (representative of n = 3 experiments) were taken by Apotome Axio Imager Z1 (Zeiss).

**Flow cytometric analysis.** TRA-1-81 expression was performed by flow cytometry FACs Canto (BD Bioscience). Cells were detached and incubated for 30 min with 1% BSA. Cells were then stained for 1 h with TRA-1-81 (Abcam), primary antibody diluted according to manufacturer’s recommendations followed by the appropriate secondary antibody.

**In vivo experiments.** Animal care and treatment were in accordance with institutional guidelines in compliance with national (D.L. n.116, G.U., suppl 40, 18 February 1992, Circolare No. 8, G.U. 14 July 1994) and international laws and policies (ECC Council Directive 86/609, OIL 358, Dec 1987; NIH Guide for the Care and Use of Laboratory Animals, U.S. National Research Council, 1996). Animal studies were submitted to and approved by the Institutional Animal Care and Use Committee of “Mario Negri” Institute (Milan, Italy). Animals were housed in a constant temperature room with a 12:12 h-dark-light cycle and fed a standard diet.

**Cisplatin-induced AKI mouse model and iPSC injection.** Two-month-old female NOD-SCID mice (Charles River Italia S.p.a.) received a subcutaneous injection of 13.9 mg/kg cisplatin (Ebewe Italia Srl) and after 24 h were divided into three groups receiving an intravenous injection, through the tail vein, of saline, undifferentiated iPSCs or iPSC-derived RPCs (5 × 10⁶ cells/mouse). In selected experiments, iPSC-derived RPCs have been pre-labeled with cell tracker PKH26 (Sigma Aldrich) following manufacturer’s instructions. Mice were sacrificed 4 days after cisplatin and kidneys were used for histology and immunohistochemistry. Human cell engraftment in kidney, liver, lung, heart and spleen was evaluated in cisplatin-treated
mice 24 hours after infusion of iPSCs or iPSC-derived RPCs. Renal function was assessed as blood urea nitrogen (BUN) by the Reflotron test (Boehringer Diagnostics Corporation). BUN levels exceeding 30 mg/dl were considered abnormal. To evaluate any possible cell maldifferentiation or tumor formation, an additional group of cisplatin-mice injected with iPSC-derived RPCs (n = 3 animals) were sacrificed 8 weeks after cisplatin injection and kidney, heart, liver and lung were examined. Normal, untreated mice served as controls.

Immunohistochemical analysis of tissues from kidney and other organs. Kidney samples were fixed in Duboscq-Brazil and paraffin sections were stained with hematoxylin and eosin, or periodic acid-Schiff’s reagent (PAS). Luminal hyaline casts and tubular necrosis (denudation of tubular basement membrane) were assessed in non-overlapping fields (up to 28 for each section) (40X, high power field, HPF).

To identify human iPSCs in murine renal tissues, acetone-fixed cryosections were incubated with anti-human nuclei antigen (HNA) Alexa Fluor 488 (Millipore) or anti-human mitochondria (Millipore) primary antibody diluted according to manufacturer’s recommendations followed by Cy3-conjugated donkey anti-mouse antibody (Jackson Lab, Bar Harbor). Sections were then stained with Rhodamine or FITC-labeled wheat germ agglutinin and DAPI. Fluorescence was examined by ApoTome Axio Image Z2 (Zeiss). The percentage of HNA positive cells per field was calculated by counting HNA- positive cells on total DAPI positive nuclei in 20 non-overlapping random areas per section. Human iPSC-derived RPCs, pre-labeled with PKH26, have been studied in PLP fixed renal sections co-stained with anti-AQP1 antibody (Abcam) followed by the appropriate Cy5-conjugated secondary antibody, FITC-WGA lectin and DAPI staining. In additional experiments, to identify human cells in distal tubuli and collecting ducts, sections have been stained with anti-PNA lectin (Vector Laboratories) or anti-AQP5 antibody (AbCam), respectively.

Duboscq-Brazil-fixed paraffin-embedded kidney sections (3 μm) were used to study immunophenotype of human cells in the kidinse of cisplatin-injected mice. Briefly, sections were deparaffinized, rehydrated, and then incubated for 30 minutes with 0.3% H2O2 in methanol to quench endogenous peroxidase. After antigen retrieval and blocking with 1% bovine serum albumin (BSA), serial sections were incubated with the appropriate anti-OCT3/4, OCT4A, anti-EP300, anti-Ki67 (Abcam), followed by the appropriate Cy5-conjugated secondary antibody, rhodamine-labeled WGA lectin and DAPI. Cell proliferation has been quantified as the number of Ki-67-positive cells/HPF (25–30 fields/section, n = 4 animals/group).

Human iPSCs and iPSC-derived RPC engrafment was analyzed in formalin-fixed, paraffin-embedded lung, kidney, heart and spleen. Three sections for each animal (n = 3/group) have been stained with anti-human mitochondria (Abcam) as previously described. RPC engrafment was quantified as the number of human mitochondria-positive cells/HPF (20–35 fields/section).

Statistical analysis. Results are expressed as mean ± SEM. Data were analyzed by ANOVA followed by Tukey-Citchetti test for multiple comparisons or t test for unpaired data, as appropriate. P < 0.05 was considered to represent statistically significant differences.

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analysis, data analysis and interpretation; R.N., P.R.: immunohistochemical analysis; A.B.: conception and study design, data analysis and interpretation; G.R.: data analysis and interpretation, writing the manuscript; M.M.: conception and study design, data analysis and interpretation and writing the manuscript.

**Additional information**

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Competing financial interests: CX is a co-founder and managing director of BIORENOVO; however, he received no compensation for his role.

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