cGMP-grade human iPSC-derived retinal photoreceptor precursor cells rescue cone photoreceptor damage in non-human primates

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Abstract

Background: Retinal regenerative therapies hold great promise for the treatment of inherited retinal degenerations (IRDs). Studies in preclinical lower mammal models of IRDs have suggested visual improvement following retinal photoreceptor precursors transplantation, but there is limited evidence on the ability of these transplants to rescue retinal damage in higher mammals. The purpose of this study was to evaluate the therapeutic potential of photoreceptor precursors derived from clinically compliant induced pluripotent stem cells (iPSCs).

Methods: Photoreceptor precursors were sub-retinally transplanted into non-human primates (Macaca fascicularis). The cells were transplanted both in naïve and cobalt chloride-induced retinal degeneration models who had been receiving systemic immunosuppression for one week prior to the procedure. Optical coherence tomography, fundus autofluorescence imaging, electroretinography, ex vivo histology and immunofluorescence staining were used to evaluate retinal structure, function and survival of transplanted cells.

Results: There were no adverse effects of iPSC-derived photoreceptor precursors on retinal structure or function in naïve NHP models, indicating good biocompatibility. In addition, photoreceptor precursors injected into cobalt chloride-induced retinal degeneration NHP models demonstrated an ability both to survive and to mature into cone photoreceptors at 3 months post-transplant. Optical coherence tomography showed restoration of retinal ellipsoid zone post-transplantation.

Conclusions: These findings demonstrate the safety and therapeutic potential of clinically compliant iPSC-derived photoreceptor precursors as a cell replacement source for future clinical trials.

Keywords: Cell therapy, Induced pluripotent stem cells, Non-human primates, Photoreceptor precursors

Background

The global incidence of inherited retinal diseases (IRDs) is approximately 1 in 2000 [1]. These disorders are amongst some of the leading causes of blindness worldwide [2, 3]. IRDs are a group of genetically and clinically heterogeneous diseases with progressive retinal damage leading to vision loss [4]. Retinitis pigmentosa (RP,
rod-cone dystrophy) is the most common with approximately 1.5 million people affected worldwide [5].

The introduction of gene therapy, such as Voretigene Neparvovec-ryzl-Luxturna™ for the treatment of biallelic Retinal Pigment Epithelium 65 (RPE65) associated Leber Congenital Amaurosis (LCA), has been a significant development [6–9]. However, the sheer extent of genetic heterogeneity, with more than 260 genes implicated in IRDs [3], limits the widespread application of gene therapy for all IRDs. Moreover, gene therapy will have limited efficacy in clinical cases of advanced retinal degeneration in which significant photoreceptor cell death has already occurred [10].

With the advent of induced pluripotent stem cell (iPSC) and embryonic stem cell (ESC) technology, regenerative stem cell therapy has the potential to be an alternative treatment for end-stage retinal degeneration, independent of the underlying genetic defect. ESC- and iPSC-derived retinal progenitor cells (RPCs) or photoreceptor precursors can differentiate into photoreceptors [11], making them an attractive cell therapy resource [12–14]. A number of studies in rodent models of photoreceptor degeneration have demonstrated that photoreceptors precursors injected sub-retinially are capable of differentiating into photoreceptors, albeit with limited survival and integration with host tissue [10, 15–17]. More recent efforts, focused on transplanting retinal sheets consisting of both neural retina and retinal pigment epithelium (RPE) derived from iPSC/ESC into both rodent and non-human primate (NHP) models [18–21], have suboptimal outcomes due to tissue disorganization (inappropriate photoreceptor orientation) and poor integration with host bipolar cells [19, 21].

The use of foetal and embryonic stem cells is restricted by limited access to new donor tissue and potential ethical concerns. Thus, allogenic iPSC RPC presents an attractive alternative cell resource. However, there is a potential risk of genetic instability introduced during reprogramming [10] and it is imperative to develop Current Good Manufacturing Practice (cGMP) standards to ensure the generation of clinically compliant and safe iPSC.

This study evaluates the safety and efficacy of cGMP-iPSC-derived photoreceptor precursors injected sub-retinally into naive NHP models, and their ability to promote endogenous photoreceptor structural recovery in cobalt chloride (CoCl₂)-induced NHP models of retinal degeneration [21]. The photoreceptor precursors were injected as a cell suspension as they were produced from a two-dimensional (2D) cell culture system. The methodology used, which incorporates in vivo optical coherence tomography (OCT), examines biocompatibility with the naïve NHP eye; ability to promote ellipsoid zone recovery following CoCl₂ damage; and ability of photoreceptor precursors to survive in the outer nuclear layer (ONL) and differentiate into cone photoreceptors.

Methods
RPC maintenance and maturation into photoreceptor precursors
The cGMP-grade iPSC line used in this study was obtained from RxCell Science as described previously [22, 23]. This is the first USFDA-approved iPSC line developed under cGMP-compliant conditions including tissue sourcing, manufacturing, testing and storage. The line was differentiated into RPC for 8 weeks at RxCell under cGMP conditions, cryopreserved and shipped to Singapore. Subsequent maturation for another 4 weeks (12 weeks old in total) before transplantation was performed in Singapore under non-cGMP conditions. These RPCs were plated at a cell density of 1 to 2 million in 6-well plates pre-coated with CTG-Retinal Coating Substrate (XR-CTG-101-SUBS, RxCell Science, US) in cGMP-grade Retinal Differentiation Medium (RD-001-M100, RxCell Science, US) for 12 weeks to obtain photoreceptor precursors for transplantation.

Characterization of RPCs and photoreceptor precursors by immunocytochemistry
Cells were cultured in 12-well microscopy chamber slides (#81201, Ibidi, Munich, Germany) and stained for stem cell, retinal progenitor and cone/rod photoreceptor markers (Additional file 1: Table S1). The cells were fixed with 4% paraformaldehyde (PFA) for 10 min at room temperature, blocked in 1% bovine serum albumin (BSA) and incubated with primary antibodies (diluted in 1% BSA) overnight at 4 °C. The cells were counterstained with Alexa Fluor 488, Alexa Fluor 568, Alexa Fluor 647 and Hoechst 33342 at room temperature for 30 min with Alexa Fluor 488, Alexa Fluor 568, Alexa Fluor 647 and Hoechst 33342 at room temperature for 30 min and mounted in ProLong™ Gold Anti-fade Mountant (Thermo Fisher Scientific, P36930, Waltham, Massachusetts, USA). Fluorescence micrographs were collected using a Zeiss LSM 800 confocal microscope (Carl Zeiss GmBH, Jena, Germany). The expression of each marker was quantified from at least five micrographs using Fiji version 1.46r [29].

Lentivirus packaging
Lentivirus pCDH-GFP was packaged in H293T cells using TransITLT1 transfection reagent (Mirus Bio, MIR 2305, Madison, Wisconsin, USA). Briefly, pCDH-GFP plasmid was co-transfected into H293T cells together with tat, rev, gag/pol and vsv-g packaging plasmids. Virus particles were harvested at 48 h and 72 h post-transfection, filtered with a 0.45 mm filter (17,574-K, Sartorius, Göttingen, Germany) and concentrated by...
ultracentrifugation at 23,000 rpm for 90 min. The virus was stored in 100 µL aliquots at −80 °C.

GFP labelling of photoreceptor precursors and dissociation for transplantation

Lentivirus infection was performed one week prior to transplantation. 5 × 10^7/mL (Multiplicity of infection = 5) of pCDH-GFP virus was mixed with 4 µg of polybrene (Sigma-Aldrich, TR-1003-G, St. Louis, Missouri, USA) in 1 mL of culture medium and added to the cells for 6 h, after which it was replaced by fresh culture medium. The photoreceptor precursors were dissociated into single cells on the day of transplantation using TrypLE™ Express Enzyme (Thermo Fisher Scientific, 12604013, Waltham, Massachusetts, USA) as previously described [15]. Infection efficiency was analysed by flow cytometry (BD FACS LSR II, BD Bioscience, San Jose, California, USA), and culture plates were also imaged using a Zeiss Axio Vert A1 inverted microscope (Carl Zeiss GmBH, Jena, Germany) to detect GFP expression.

Animal studies

A total of six Macaca fascicularis, aged 3 to 5 years old and weighing 3 to 4.5 kg were sourced from SingHealth Experimental Medicine Centre, Singapore (IACUC numbers: 2015/SHS/1092, 2015/SHS/1044). All animal experiments were approved by the Institutional Animal Care and Use Committee (IACUC) of SingHealth (Singapore) (IRB number: 2019-009), performed in an American Association for Accreditation of Laboratory Animal Care (AAALAC) approved facility, and according to the Statement from the Association for Research in Vision and Ophthalmology (ARVO) for the Use of Animals in Ophthalmic and Vision Research.

Cobalt chloride-induced NHP retinal degeneration model

Cobalt chloride hexahydrate (CoCl_2·6H_2O, molecular weight of 237.93, Nacalai tesque, Kyoto, Japan) was dissolved in 0.9% saline solution to obtain 0.3 mg/mL solution for induction of retinal injury [21]. Animals were sedated with ketamine (10 mg/kg, body weight (BW)) and atropine (0.05 mg/kg, BW) followed by induction of general anaesthesia (GA) with 2% isoflurane and maintenance with 0.5–2% isoflurane [24, 25]. A 25-gauge (G) 3 port-vitreectomy (infusion, chandelier endo-illumination, and working port) was performed with a Bausch & Lomb Stellaris® PC machine (Bausch & Lomb, Rochester, New York, USA). Local retinal detachment (1- to 2-disc diameter size) within macular region was created by manual sub-retinal injection of 40 to 50 µL of 0.3 mg/mL CoCl_2 solution using an extendible 38 G sub-retinal injection cannula [26] (MedOne Surgical Inc., Sarasota, Florida, USA). Retinal and RPE damage was observed by OCT and Fundus Autofluorescence (FAF) imaging.

Immunosuppression

All NHPs received systemic immunosuppression 7 days prior to photoreceptor precursor transplantation, and this was continued throughout the study period [25]. Sirolimus was administered orally with a loading dose of 2 mg, followed by a 1 mg daily dose. Doxycycline and minocycline (7.5 mg/kg, BW) were delivered orally, twice daily [24, 27].

Transplantation of photoreceptor precursors into NHPs

In both naïve and CoCl_2-induced diseased NHPs, a suspension of photoreceptor precursors in media was manually injected sub-retinally using an extendible 38 G injection cannula (MedOne Surgical Inc., Sarasota, Florida, USA)(Additional file 2). Seven eyes from 6 NHPs received photoreceptor precursors at two different dosages (40,000 to 60,000 cells, n = 3, for initial safety test in naïve NHPs; 100,000 to 300,000 cells, n = 4 for rescue in diseases NHPs). The retinal bleb was created by injection of 40 to 50 µL cell suspension medium with about 1- to 2-disc diameter size. Details are provided in Table 1.

In vivo follow up by ophthalmic imaging

Cross-sectional images of the central 30 degrees of the retina were acquired non-invasively using spectral domain-optical coherence tomography (SD-OCT, Spectralis® Heidelberg Engineering, Inc., Heidelberg, Germany). Real-time eye-tracking system and image registration capabilities on the Spectralis® were used to mitigate motion artefacts. Repeat OCT scans were performed at the same location over consecutive evaluations. Fundus auto-fluorescence (FAF) and infrared (IR) images were obtained using the same device.

Table 1 Summary of sub-retinal injection of cGMP-iPSC-derived photoreceptor precursors in both naïve and disease NHP models

| Period of CoCl_2 damage | No. of cells | GFP labelled | Period of rescue | Eyes |
|-------------------------|--------------|--------------|------------------|------|
| Naïve                   | 40,000–60,000| Non-GFP labelled | 3 months | n = 3 |
| Disease/rescue          |              |              |                  |      |
| 1 month                 | 100,000–300,000 | GFP labelled | 1 month | n = 1 |
| 1 month                 | 100,000–300,000 | GFP labelled | 3 months | n = 1 |
| 4 months                | 100,000–300,000 | GFP labelled | 3 months | n = 2 |
Qualitative SD-OCT analysis
Precise mapping of the area and extent of sub-retinal CoCl2 and photoreceptor precursor injection was achieved by extrapolating the information from surgical videos super-imposed onto the OCT images, using vessel markings as landmarks. Treated areas of the retina were analysed for structural changes. The ellipsoid zone (EZ), identified as the second hyper-reflective band on the OCT image, was used as a surrogate imaging biomarker of photoreceptor integrity. Disruption of the EZ was the indicator of the presence of photoreceptor damage, and restoration of the EZ is an indicator of photoreceptor recovery.

Electroretinography
Global retinal function was assessed by full-field electroretinography (ERG) using an Espion system (Diagnosys LLC, Lowell, Massachusetts, USA) and Jet recording electrodes with protocols based upon those recommended for humans by the International Society for Clinical Electrophysiology of Vision (ISCEV) [28] but using a flash strength of 5 cd s m\(^{-2}\) for the photopic single flash responses.

Histopathological examination
Animals were sacrificed under deep anaesthesia and perfused with 10% formalin at 3 months after photoreceptor precursor transplantation. The eyes were enucleated and fixed overnight. Full-thickness foveal samples (retina → sclera) were collected and embedded in paraffin. 10 µm sections were cut with a microtome (Leica RM2255, Wetzlar, Germany) and stained with hematoxylin and eosin (H&E).

Immunofluorescence staining of NHP retinal sections
Paraffin-embedded sections were deparaffinized in xylene and rehydrated in ethanol. Following antigen retrieval at 95 °C for 20 min in citrate buffer pH 7.0 or tris-EDTA buffer pH 9.0 (depending on the antibody), the samples were blocked for 1 h in 10% donkey serum. Samples were incubated with primary antibodies overnight at 4 °C (Additional file 1: Table S1). The samples were counterstained with Alexa 488, Alexa 568 and Alexa 647, mounted in ProLong™ Gold Antifade Mountant (Thermo Fisher Scientific, P36930, Waltham, Massachusetts, USA) and imaged using a Zeiss LSM800 upright confocal microscope (Carl Zeiss GmBH, Jena, Germany). Z-stack images were collected every 0.56 µm, and the micrographs were analysed using Fiji version 1.46r [29].

Results
Characterization of photoreceptor precursors in vitro prior to transplantation
Cells were thawed and cultured for an additional 4 weeks prior to transplantation. The cells, thus 12 weeks old, showed the absence of pluripotency marker OCT4 (Fig. 1A) and only a minority of cells express the retinal progenitor marker LHX2 (Fig. 1B and B’). The majority of the cells express pan-photoreceptor markers CRX (37 ±7% of total Hoechst 33342 stained cells) (Fig. 1C and C’), Recoverin (41 ±3% of total Hoechst 33342 stained cells) (Fig. 1E and E’) and OTX2 (46.3 ±13% of total Hoechst 33342 stained cells) (Fig. 1D and D’). These markers were previously used to characterize photoreceptor precursors derived from ESC and retinal organoids [30–32]. The expression levels of the pan-photoreceptor markers were comparable to previously published data for photoreceptor precursors derived from the same cGMP-grade iPSCs [15]. In addition, the presence of early rod-specific transcription factor NRL (45 ±16% of total Hoechst 33342 stained cells) (Fig. 1F and F’) and the cone-specific markers cone-arrestin (43 ±2.6% of total Hoechst 33342 stained cells) (Fig. 1G and G’) and Opsin M/L (46.65 ±1.67% of total Hoechst 33342 stained cells) (Fig. 1I and I’) was observed, indicating that these cells at 12 weeks old were photoreceptor precursors. Neither the bipolar cell marker PKCa (Fig. 1K) nor the RPE marker RPE65 (Fig. 1J) was observed, confirming the commitment to the photoreceptor lineage. Of note, these photoreceptor precursors did not express late rod specific marker rhodopsin, suggesting that they had not achieved the level of maturity needed for rhodopsin expression (Fig. 1H).

Tolerance of transplantation of photoreceptor precursors into naïve NHP eyes
Three naïve NHPs eyes received sub-retinal injections of 40,000 to 60,000 photoreceptor precursors for initial safety and biocompatibility evaluation (n=3). Colour fundus and FAF images revealed a clear vitreous cavity with no overt signs of inflammation (Fig. 2A1 and A2). The initial disruption of the retinal layers at the injection site (Fig. 2A3) showed limited recovery at 3 months with residual scarring only at the original site of retinotomy, with normal surrounding retina. The structure of the macula was well maintained over the 3 months as shown by OCT imaging (Fig. 2A4) and histology (Fig. 2B2). Full-field ERG at 3 months post-transplantation showed no significant abnormality of retinal function in either cone or rod derived responses (Fig. 2C). All showed slightly lower DA 0.01 dim flash dark-adapted responses, but no definite clinically significant differences compared to the pre-treatment data.
The effects of CoCl₂ on the photoreceptors and retinal pigment epithelium in NHP eyes

In the NHP model of retinal degeneration established by sub-retinal injection of CoCl₂ [33], mild to moderate ONL loss was observed on OCT imaging. High-resolution OCT scans revealed persistent outer retinal alterations and disruption of EZ for up to 4 months post-CoCl₂ injection (Fig. 3E1 to E3). This was consistent with the histology data, which showed both ONL and inner nuclear layer (INL) damage compared to non-operated controls (Fig. 3F4 and G4). Immunofluorescence staining with the pan photoreceptor marker recoverin (E, E') and the rod photoreceptor marker rhodopsin also showed disorganization of the outer nuclear later and photoreceptor outer segments 1 month after CoCl₂ damage (Fig. 3G1-G3) compared with non-operated controls (Fig. 3F1-3). Corresponding FAF images showed persistent hyper-autofluorescence at the site of CoCl₂ injection up to 4 months (Fig. 3B and C), in keeping with structural damage caused by CoCl₂ treatment to both the photoreceptors and RPE.

Effect of transplanted photoreceptor precursors on outer retinal layers in CoCl₂ disease models

Photoreceptor precursors were injected sub-retinally into the damaged retinas of four eyes (100,000 to 300,000) to assess the ability of the transplanted cells to rescue retinal damage. In eyes rescued at 1 month post-CoCl₂ damage, high-resolution, tracked SD-OCT line scans showed a reappearance of EZ signals 1 month post-injection, with continued improvement and near-complete restoration by 3 months post-injection (Fig. 4F1-3). In addition, at 3 months post-photoreceptor precursor injection, the hyper-auto fluorescence seen at 1 month post-CoCl₂ injection (Fig. 4B) was reduced in intensity (Fig. 4C). This was consistent with OCT images showing continuity of the RPE layer in the regions rescued with photoreceptor precursor injection at 3 months (Fig. 4F).

Maturation of transplanted photoreceptor precursors within the ONL of NHP retina

Lentiviral transduction of the photoreceptor precursors was used to generate stable GFP expression to
Fig. 2 Safety of sub-retinal transplant photoreceptor precursors into naïve NHP eyes. **A** Photoreceptor precursors were transplanted subretinally in yellow circle zone (colour fundus, Column A1, and autofluorescence images, Column A2) in naïve NHP eyes and followed at 2 days, 1, 2 and 3 months post-transplantation. Limited changes were observed at transplantation zone on autofluorescence images and showed time-dependent recovery. OCT line scan images on transplantation zone (Column A3, indicated as yellow arrows in Column A2) showed interrupted ONL at retinal incision site (red arrows) at day 2 and recovered during follow up. Normal macula structures were shown on OCT scans (Column A4) as indicated the scanned position at white arrows in Column A2. **B** Histology images (H&E) on transplantation zone and macula both showed normal retinal structure. Scale bar, 200 μm in Column A3 and A4, B1 and B2. **C** Full-field electroretinography (ERG) was normal at 3-month follow-up in the eyes with transplanted photoreceptor precursors. C1, C2 and C3 each represent ERG recordings from 3 different NHPs. The terms DA and LA refer to dark-adapted and light-adapted. The numbers refer to flash strength in cd s m⁻².
track transplanted cell survival in vivo and ex vivo. This was performed 1 week prior to transplantation with more than 90% efficiency as assessed by flow cytometry (Fig. 5A3, Additional file 1: figure S1). GFP-labelled photoreceptor precursors were transplanted sub-retinally into four eyes in CoCl₂-induced diseased NHPs. Post-injection, the GFP signal was tracked by FAF imaging. GFP-positive cells were observed within the injection zone at 2 days post-surgery (Fig. 5B2), with gradual reduction by 1 to 2 weeks (Fig. 5B3 and B4) and were virtually undetectable by 4 weeks (Fig. 5B5). Live cell detection was likely limited by the relative low sensitivity of this method for low cell numbers. Long-term survival of GFP-labelled photoreceptor precursors was confirmed by immunofluorescence staining (Fig. 5C2) of retina sections at 3 months post-transplantation. GFP-positive cells were identified in the ONL and co-stained with a human cytoplasmic marker (SC121), confirming that these cells were of human origin (Fig. 5C).

At 1 month post-transplantation, there was evidence of the precursors maturing into cone photoreceptors. Human-specific makers SC121 and anti-human mitochondria (AMA) were used to identify transplanted cells. SC121-positive (Fig. 6A) and AMA-positive (Fig. 6B) cells were identified in the ONL, and they co-localized with medium/long (M/L) opsin-positive staining, indicating maturation into cone photoreceptors (Fig. 6A1, B1 and C1). Human-derived cells were surrounded by SC121 and AMA negative cone photoreceptors, likely native NHP cones (Fig. 6A-A2 and B-B2). Transplanted photoreceptor precursor cells positive for AMA did not appear to co-localize with rhodopsin (Fig. 6C2), suggesting a propensity to mature into cone photoreceptors. Retinal tissue sections collected from non-operated NHP controls were examined to confirm that the SC121, AMA and GFP staining was specific to the transplanted photoreceptor precursors; specific staining for these markers was not detected in the non-operated controls (Additional file 1: figure S2 E-H), or in operated eyes where primary antibodies were omitted (Additional file 1: figure S2 A-D).

**Discussion**

This study demonstrates the safety and efficacy, in a naïve immunosuppressed non-human primate model, of sub-retinal transplantation of a cGMP-iPSC line previously demonstrated in immunodeficient murine host retina to generate a high percentage of photoreceptor precursors [15]. Upon sub-retinal transplantation of such cells, they displayed photoreceptor morphology and expressed specific markers of mature photoreceptors. Sub-retinal injection of these cells is well-tolerated, with no adverse effects on structure (OCT) or function (ERG) of the retina for 3 months following transplantation. Importantly, there was no evidence of an undesirable immune response, as shown by the lack of retinal oedema, inflammatory infiltrates or retinal/ choroidal lesions when examined by OCT and fundus imaging, the standard tools used to observe ocular inflammation in rodents [34], primates [35] and humans [36].

In addition, sub-retinal injection of photoreceptor precursors into NHP models with CoCl₂-induced retinal degeneration was able to induce structural recovery and restored the EZ line on the OCT at 1 and 3 months post-transplantation. The EZ line corresponds to the mitochondria found within the ellipsoid layer of the outer portion of the photoreceptor inner segments [37]. Recovery of the EZ (Fig. 4) suggests that sub-retinally injected retinal photoreceptor precursors successfully mediated structural recovery of the CoCl₂ damaged photoreceptor layer. Although, a contribution from altered refractive and reflectance characteristics [38] cannot be excluded with certainty, restoration of the EZ was not present in controls up to 4 months following the procedure. This almost certainly suggests that transplanted photoreceptor precursors were able to mediate photoreceptor damage rescue. Unfortunately, correlation with histological results was technically difficult to achieve, as the region of interest was too small (1 mm² in area) to be identified accurately on histology specimens.

Possible mechanisms by which photoreceptor recovery occurs include: (1) neuroprotective effects from secreted immune-modulatory factors, (2) maturation...
Fig. 3  (See legend on previous page.)
and integration of stem cell-derived photoreceptors to replace their damaged counterparts, and/ or (3) intracellular material exchange between transplanted cells and host photoreceptors [10, 39–41]. In this study there is evidence of photoreceptor structural recovery, possibly mediated by survival of photoreceptor cells, and the transplanted photoreceptors were able to survive in the NHP retina for up to 3 months in the presence of immunosuppression. This was demonstrated by the expression of GFP and two other human-specific cytoplasmic markers, SC121 and anti-mitochondrial antibody (AMA). Use of FAF enabled tracking of these GFP labelled photoreceptor precursors in the NHP eye and demonstrated that GFP fluorescent cells were detectable for up to 4-week post-transplantation by FAF and up to 3 months by IHC. In addition, these surviving...
photoreceptor precursors showed evidence of maturation into cone photoreceptors, with suggestion of integration into host ONL (as shown by the Hoechst 33342 negative gaps in the ONL that stained for human SC121/AMA cytoplasmic staining). Confocal microscopy identified the presence of photoreceptors expressing both the human-specific markers and the cone-specific marker opsin M/L, but did not identify cells co-expressing AMA and the rod specific marker rhodopsin. This is in contrast to previous work where cGMP-iPSC-derived photoreceptor precursors transplanted into immune-deficient murine models predominantly differentiated into rod photoreceptors [15]. In other NHP studies whereby retinal cell sheets and photoreceptor precursors were transplanted, the transplanted cells differentiated into cone photoreceptors first, with only one study reporting differentiation into rod photoreceptors only after an additional 28 days after the detection of cone photoreceptors [21,50]. Thus, it is possible that with longer follow-up times post-transplantation, we might be able to observe transplanted cells differentiating into rod photoreceptors. It is probable also that this difference observed between NHP and murine transplants is due to the fundamental differences in retinal anatomy of the host retina. This
difference in photoreceptor precursor differentiation will be investigated in future studies by staining with other rod markers, at longer post-transplantation time points.

Studies in NHP are of fundamental importance because assessment of biocompatibility in higher order animals that closely resemble human retinal physiology and structure is crucial for pre-clinical evaluation. Traditionally, rodent models have predominantly been used for pre-clinical evaluation of photoreceptor precursor transplants [42, 43]. However, rodents lack the cone-rich macula [44] found in humans, and as patients with macular degeneration will be one eventual target population for potential therapeutic intervention, the presence of a macula is desirable. Other large mammals such as cats [45] and pigs [46, 47] have been used for pre-clinical evaluation of retinal sheets, RPE monolayer and photoreceptor precursors but are also not ideal. Limitations of the cat model include the presence of the reflective tapetum lucidum, [48], while porcine eyes lack a fovea and are not optimal for pre-clinical studies due to inconsistencies in disease development patterns [48, 49]. In contrast, the ocular anatomy of NHPs resemble humans the most, making them most suitable for the evaluation of retinal cell-based therapies [44, 50–53]. To date there has been only one other study of photoreceptor precursor transplants in NHPs. CRX+/tdTomato ESC-derived photoreceptor precursors [50] were transplanted sub-retinally into NHPs with laser ablated ONL. Similarly, they demonstrated cell survival in the sub-retinal space, with limited evidence of synaptic formation between photoreceptor precursors and host bipolar cells. Our present study is unique in that the photoreceptor precursors are derived from cGMP-grade iPSCs, which are cryo-preserved and shipped from USA. The cells are subsequently thawed in Singapore for further in vitro culture prior to transplantation. This proof-of-concept study therefore also
demonstrates the feasibility of shipping cell products manufactured in one location to other countries, once approved, for clinical application.

One limitation of this study is the inability to co-stain the transplanted cells with human-specific nuclear markers such as human-specific nuclear antigen (HuNu) or human specific lamin B2 (hLMNB2) nuclear membrane protein to confirm transplanted cell survival and integration [15]. This difficulty arose from cross reactivity with the host NHP tissue (Additional file 1: figure S3). Thus, the possibility of retinal structural recovery due to indirect neurotrophic effects from the transplanted photoreceptor precursors or cytoplasmic material transfer or cell fusion cannot definitely be excluded [39–41]. To overcome this, alternative cell tracking methods such as CRISPR/Cas9 based genome detection will be explored [54, 55]. Future work also includes optimizing other human specific nuclear antibodies to differentiate host versus transplanted photoreceptor precursors, and the use of presynaptic markers to demonstrate better integration with the host retina [15, 56–58]. In addition, functional rescue should be examined by other modalities including multi-focal electro-retinography and visual evoked potentials. Another limitation is the inability to assess any immune privilege in the subretinal space, due to the need for immunosuppression for the xenograft transplantation. Future studies might include an allogenic cell resource with and without immunosuppression.

Conclusions
This study firstly demonstrates the safety of subretinally transplantation of photoreceptor precursors derived from cGMP-grade iPSCs into healthy NHPs. When transplanted into a CoCl2-induced retinal degeneration model, the photoreceptor precursors could survive for up to 3 months post-transplant, mediating the structural recovery of the chemically damaged retina. Importantly, the transplanted photoreceptor precursors were able to mature and differentiate into cone photoreceptors, essential for the technique to have therapeutic potential in patients. This study provides proof of concept demonstrating the potential applicability of subretinal photoreceptor precursors suspension transplantation as a therapeutic strategy for IRDs.

Abbreviations
IRDs: Inherited retinal diseases; RP: Retinitis pigmentosa; RPE: Retinal pigment epithelium; iPSC: Induced pluripotent stem cell; ESC: Embryonic stem cell; RPC: Retinal progenitor cell; cGMP: Current good manufacturing practice; OCT: Optical coherence tomography; NHP: Non-human primate; ONL: Outer nuclear layer; INL: Inner nuclear layer; PFA: Paraformaldehyde; BSA: Bovine serum albumin; FAF: Fundus autofluorescence; IR: Infrared; EZ: Ellipsoid zone; ERG: Electroretinography; AMA: Anti-mitochondrial antibody; DA: Dark-adapted; LA: Light-adapted.

Supplementary Information
The online version contains supplementary material available at https://doi.org/10.1186/s13287-021-02539-8.

Additional file 1. Table S1. List of primary antibodies used in the study. figure S1. Estimation of GFP expression by FACS. After lentivirus infection, 95% of the RPC cells expressed GFP (left panel). There was no GFP in the non-infection control (right panel). figure S2. Controls for GFP, SC121 and AMA staining. (A-D) Secondary antibody only controls for Alexa 488 (B), Alexa 568 (C) and Alexa 647 (D). The background for all secondary antibodies appears to be high. (E-H) GFP, SC121 and anti-human mitochondrial antibody staining in non-operated NHP retina. None of the antibodies specifically stained anything in the non-operated control retina. The brightness/contrast was artificially increased to identify any weakly stained cells. Hoechst 33342 was used to stain the INL and ONL layers (A, E). Scale bar 10 µm. figure S3. Cross reactivity of human lamin B2 (hLMNB2) antibody with the NHP retina. Tissue sections collected from NHP eyes post-photoreceptor precursor transplantation were stained with the human marker SC121 (B) and the human specific nuclear marker lamin B2 (C). The human nuclear marker antibody showed high cross-reactivity with the NHP tissue (C, D). Nuclei were stained with Hoechst 33,342 (H33342). Scale bar: 10 µm.

Additional file 2. Video: Surgical video of photoreceptor precursors subretinal injection under the NHP retina.

Acknowledgements
We would like to acknowledge the veterinary team at the Translational Pre-Clinical Model Platform (Singapore Eye Research Institute, Singapore) for providing support in animal surgery preparation and follow-up.

Authors’ contributions
SL and BHP performed collection and/or assembly of data, data analysis and interpretation, manuscript writing and final approval of manuscript. ZL, BY and GL contributed to conception and design, collection and/or assembly of data, data analysis and interpretation, manuscript writing and final approval of manuscript. JYO and DSLW contributed to collection and/or assembly of data, data analysis and interpretation and final approval of manuscript. DG performed data interpretation and final approval of manuscript. WW helped with data analysis and interpretation, manuscript writing and final approval of manuscript. QSWT and GSWT were involved in collection and/or assembly of data, data analysis and interpretation, manuscript writing and final approval of manuscript. VAB contributed to conception and design, manuscript writing and final approval of manuscript. WH contributed to conception and design, manuscript writing and final approval of manuscript. RK performed collection and/or assembly of data and final approval of manuscript. VAB was involved in administrative support, manuscript writing and final approval of manuscript. GEH contributed to conception and design, manuscript writing and final approval of manuscript. XZ and XS contributed to conception and design, data analysis and interpretation, manuscript writing and final approval of manuscript. ZL, BY and GL contributed to conception and design, data analysis and interpretation, manuscript writing and final approval of manuscript. All authors read and approved the final manuscript.

Funding
This study was supported by IAF-PP (HMBS Domain) (OrBiD): Ocular Biomaterials and Device, A*STAR, Singapore (H17/01/a0/013), and NUHS Clinician Scientist Program Grant.

Availability of data and materials
All data generated or analysed during this study are included in this published article and its supplementary information files.

Declarations
Ethics approval and consent to participate
All animal experiments were approved by the Institutional Animal Care and Use Committee (IAUC) of SingHealth (Singapore) (IRB number: 2019-009), performed in an American Association for Accreditation of Laboratory Animal Care (AAALAC) approved facility, and according to the Association
for Research in Vision and Ophthalmology’s (ARVO) Statement for the Use of Animals in Ophthalmic and Vision Research.

Consent for publication
Not applicable.

Competing interests
SL, ZL, BY, JYO, WW, DSLW, QSWT, BHP, DG, GSWT, GEH, RK, VAB, WH, GL and XS have no competing interest. XZ was the founder and CEO of RxCell.

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Received: 15 June 2021   Accepted: 31 July 2021

Published online: 19 August 2021

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