RNA interference for apoptotic signal-regulating kinase-1 (ASK-1) rescues photoreceptor death in the rd1 mouse

Daisuke Sekimukai, Shigeru Honda, Akira Negi

Department of Surgery, Division of Ophthalmology, Kobe University Graduate School of Medicine, Kobe, Japan

Purpose: To evaluate whether RNA interference against apoptotic signal-regulating kinase-1 (ASK-1), a gene involved in stress-induced apoptosis, inhibits photoreceptor death in retinal degeneration 1 (rd1) mice.

Methods: Retinal explants from rd1 mice were subjected to organ cultures on postnatal day 9 (P9). Short interfering RNA (siRNA) for ASK-1 was transfected into cultured retinas at the onset of experiments. Real-time PCR was performed to evaluate the natural expression of ASK-1 mRNA and its inhibition with siRNA. Retinal explants were fixed at P13 and P16, and consecutive cryosections were prepared. Histological and immunohistochemical examinations including TUNEL assays were performed.

Results: In preliminary experiments, the incorporation of fluorescent siRNA was found in cells in the outer nuclear and inner nuclear layers on the day following transfection. The expression of ASK-1 mRNA increased with time, which was suppressed more than 70% by siRNA. ASK-1 immunopositive cells were found mostly in the outer nuclear layers, and the number of immunopositive cells was remarkably reduced in retinas treated with siRNA for ASK-1 compared to untreated controls. The thickness of outer nuclear layers of control retinas decreased with time, while the thickness of siRNA transfected retinas was significantly preserved compared to control at P16 (p=0.0021). In TUNEL assays, siRNA for ASK-1 significantly decreased TUNEL-positive cells (49% and 42% of controls at P13 and 16, p=0.039 and 0.0028, respectively).

Conclusions: RNA interference against ASK-1 may provide a benefit by inhibiting photoreceptor apoptosis in rd1 mice.
ASK-1 works in retinal organ cultures and inhibits photoreceptor death in rd1 mice.

METHODS

Retinal organ cultures: The study adhered to the Association for Research in Vision and Ophthalmology (ARVO) Statement for the Use of Animals in Ophthalmic and Vision Research. Retinal explant cultures of C3H/HeN (rd1) mice and C57Bl/6 (Bl6) mice were prepared as described previously [24]. The mice were purchased from (CLEA Japan Inc., Tokyo, Japan) and kept in mouse cages with free access of solid food and water. Light-dark cycle was 12:12 h. Briefly, mice were euthanized (intraperitoneal injection of 15 mg of pentobarbital per mouse for euthanasia) and neural retinas were extracted on postnatal day (P) 8 or 9, depending on the experiment. Each retina was submerged in HBSS and extended to make whole flat-mounts on 30 mm diameter microporous membranes (Millicell-CM; Millipore, Bedford, MA) with ganglion cell layers facing up. Millicell-CM containing retinal explants were soaked in 1 ml of culture medium (Opti-MEM, Invitrogen, Carlsbad, CA) with or without 25% heat inactivated horse serum in six-well plates and incubated, allowing an air interface with the ganglion cell without 25% heat inactivated horse serum in six-well plates medium (Opti-MEM, Invitrogen, Carlsbad, CA) with or containing retinal explants were soaked in 1 ml of culture media was replaced with Opti-MEM culture media containing 25% serum. After 4 h at 34 °C, the transfaction media was replaced with Opti-MEM culture media containing 25% serum. After 24 h, organ culture cryosections were prepared with DAPI staining and observed with a Keyence Biozero fluorescence microscope (BZ-8000; Keyence, Osaka, Japan). For silencing ASK-1, Stealth RNAs, which included two different sequences of siRNA for ASK-1 (Table 1), were obtained from Invitrogen. Each siRNA was transfected into the retina with lipofectamine RNAiMAX using the same method as described above. Medium GC content scrambled control siRNA (Invitrogen; proprietary sequence) was also applied to examine off-target effects. Retinal explants were either transfected or untransfected with scrambled siRNA and cultured for 72 h before RNA extractions.

Real-time PCR: Total RNA was extracted using RNeasy Plus (Qiagen, Valencia, CA) from organ cultures for 0, 1, 3, 5, and 7 days according to the manufacturer’s recommendations. Total RNA was eluted from columns in 50 μl RNase-free water. The purity and concentration of RNA was determined by measuring the absorbance at 260 nm and 280 nm. RNA was reverse-transcribed into cDNA (High Capacity cDNA Reverse Transcription Kit, Applied Biosystems, Foster City, CA) in a total volume of 100 μl according to the manufacturer’s instructions. Real-time PCRs were performed in 96 well plates on an ABI Prism 7500 Sequence Detection System (Applied Biosystems StepOne™ and StepOnePlus™ Real Time PCR System and Taqman Fast Universal PCR Master Mix, Applied Biosystems) according to the manufacturer’s instructions. Final reaction volumes were 25 μl. Each sample was analyzed in triplicate. Thermal cycler conditions were as follows: 2 min at 50 °C, 10 min at 95 °C, followed by 40 cycles of 15 s at 95 °C and 1 min at 60 °C. Sequence Detector Software (Applied Biosystems) was used to extract PCR data, which were exported into Excel 2003 (Microsoft Corporation, Redmond, WA) for further analyses. The amount of targeted gene expressed was normalized to an endogenous reference and relative to a calibrator. \( \beta\)-Actin was used as an endogenous reference in these experiments. The formula \( \Delta \Delta^C T \) was used to calculate the amount of target gene expression normalized to the endogenous control and relative to the calibrator.

Histological procedures: After transfections and organ cultures for 0, 4, and 7 days (P9, 13 and 16, respectively), retinal explants were fixed with 4% paraformaldehyde. Cryosections were prepared and stained with DAPI so the thickness of the outer nuclear layer (ONL) could be measured. For immunohistochemistry, sections were blocked with 5% normal goat serum in 0.1 M phosphate buffered saline (PBS; 137 mM NaCl, 2.7 mM KCl, 10 mM Sodium Phosphate dibasic, 2 mM Potassium Phosphate monobasic and a pH of 7.4), and incubated in 1:200 diluted rabbit anti-human ASK-1 antibody (catalog# sc-7931; Santa Cruz Biotech, Santa Cruz, CA), with normal goat serum overnight at 4 °C. This rabbit ASK-1 antibody is compatible with mouse ASK-1. For secondary antibody, incubation with 1:1,000 diluted sheep anti-rabbit IgG (whole molecule), F(ab’); fragment–Cy3 antibody (Sigma-Aldrich, Tokyo, Japan) was applied for 30 min at room temperature. Nuclei were stained with 1:1,500 diluted TO-PRO-3 iodide (642/661; Invitrogen), and sections were examined using a LSM 5 Pascal confocal imaging system (Carl Zeiss Inc., Tokyo, Japan).

### Table 1. Sequences of Stealth siRNAs for ASK-1 Used in the Study.

| Title | Lot No. | Sequence |
|-------|---------|----------|
| siRNA-1 | MSS218535 | 5′-AAUUGCAGUGCUGCAGCCUCUUGG-3′ |
| siRNA-2 | MSS218536 | 5′-AAAUGCGUAAUGAACCUCAGUGG-3′ |
Terminal dUTP Nick End Labeling: Cryosections were incubated in 50 μl of reaction buffer containing terminal deoxynucleotidyl transferase (TdT; Promega, Southampton, UK) and fluorescein-12-dUTP (Roche, Basel, Switzerland) according to the manufacturer’s instructions. Sections were incubated at 37 °C for 1 h in a humidified chamber. After several washes in PBS and staining with DAPI, sections were observed with a Keyence Biozero fluorescence microscope. Under masked conditions, at least 500 cells in the ONL were counted using digital images at 200× magnification for each condition and the proportion of TUNEL-positive cells was determined.

Statistical analyses: Unpaired two-tailed Student’s t-tests were used for all statistical analyses in the present study. A p value less than 0.05 was considered statistically significant.

RESULTS

Incorporation of siRNA: Fluorescently labeled dsRNA oligonucleotides were found in most cells in outer nuclear layers (ONLs) and inner nuclear layers (INLs) 24 h after transfection (Figure 1). Labeled oligonucleotides were found only in cytoplasm and not in nuclei.

Expression of ASK-1 mRNA: The expression of ASK-1 mRNA in extracted retinas from rd1 mice increased in a time-dependent manner during experimental days 0 to 7.
Upregulation was significant on days 3 and 7 (Figure 2). The expression of ASK-1 mRNA in retinas from Bl6 mice was low until day 3 but significantly increased by day 7.

RNA interference for ASK-1: Since the effect of RNA interference is usually evaluated within 72 h [25-27], we determined the relative expression of ASK-1 mRNA in rd1 mice retinas at 24 h (P9) and 72 h (P11) after siRNA transfection. The expression of ASK-1 mRNA was suppressed more than 70% with each of two siRNA at both time points (Figure 3). Because of the equal inhibitory effect of both siRNAs, we used siRNA-2 for the subsequent experiments. In the control experiment, retinal explants treated with scrambled siRNA showed higher expression of ASK-1 (3.5 fold compared to starting point, n=3) than untreated subjects (threefold compared to starting point, n=3) at 72 h of incubation although the difference was not significant.

Immunoreactivity of ASK-1: Immunoreactivity of ASK-1 was found in the outer part of ONLs of control retinas (day 4 corresponding to P12; Figure 4). However, the immunoreactivity was very weak for the same time period with transfected siRNA-2.

Histological findings: ONL thickness decreased markedly during P9 to P16 in retinal explants from rd1 mice (Figure 5). The transfection of siRNA-2 markedly preserved the ONL thickness in rd1 mice. In semiquantitative analyses, ONL thickness in retinas transfected for siRNA-2 was preserved significantly compared with untreated control retinas from rd1 mice (p=0.012 at P13, and p=0.0021 at P16).

Apoptotic cells in retinas: Most TUNEL-positive cells were found in the ONL, while few were in the INL in untreated controls and siRNA-2 transfected retinas from rd1 mice. There were fewer TUNEL-positive cells in siRNA-2 transfected retinas than in untreated controls (Figure 6). The mean ratio of TUNEL-positive cells to total ONL cell counts in untreated control retinas of rd1 mice was 5.0% at P13 and 3.3% at P16. Ratios in retinas transfected with siRNA-2 were 2.4% at P13 and 1.4% at P16. The ratio of TUNEL positive cells to total ONL counts in siRNA-2-transfected retinas was about 48% of that seen in untreated retinas at P13, decreasing further to 42% at P16, which was statistically significant (p=0.039 at P13 and p=0.0028 at P16).

DISCUSSION

In this study, we confirmed that RNA interference is possible in a retinal explant. Further, we demonstrated that ASK-1 is mainly expressed in ONL cells in rd1 mice and plays an important role in inducing the apoptosis of photoreceptors.

Photoreceptor degeneration is known to occur due to various changes to molecules in signal transduction cascades, dysfunctions in energy metabolism, oxidative stress in outer retinas, or disturbances of phagocytic processes by RPE [28]. In the rd1 mouse, a mutation in the enzyme cGMP phosphodiesterase causes a remarkable elevation of cytoplasmic Ca2+ [6], which activates multiple pathways of apoptosis including the activation of Ca2+ dependent proteases, mitochondrial stress accompanied with the generation of ROS, and ER stress [7-11]. These events are followed by the activation of the ASK-1-JNK-c-Jun pathway (Figure 1) [12,16,18,19,29-31]. The continuous elevation of...
Ca\(^{2+}\) levels in retinas from rd1 mice from P9 to P13 was previously reported \[10\] to correspond with a gradual increase in the expression of \(\text{ASK-1}\) mRNA, as found in the present study. Moreover, the expression of \(\text{ASK-1}\) mRNA was increased in retinal explants of Bl6 mice after P11, three days from the start of organ culture. It is likely that the oxygen tension in normal atmosphere (about 21\%) is high enough to cause oxidative stress in cultured retinal explants and induce \(\text{ASK-1}\) \[32\].

RNA interference techniques are used in vitro to silence specific genes; Palffy et al. \[21\] applied RNA interference to knock down rds-peripherin in retinal organotypic culture. Zacks et al. \[22\] demonstrated that RNA interference for Fas can work in retinal organ cultures to rescue photoreceptors.

**Figure 3.** Suppression of \(\text{ASK-1}\) mRNA expression by siRNA. The expression of \(\text{ASK-1}\) mRNA in the retina of rd1 mice was inhibited at 24 and 72 h after the transfection of siRNA. The expression of \(\beta\)-actin was used as an internal control. Values are presented as the average±SEM of three independent experiments.

**Figure 4.** Immunohistochemistry of \(\text{ASK-1}\) on experimental day 4. A: Immunoreactivity of \(\text{ASK-1}\) was found in the outer part of ONL from control retina in rd1 mice (arrows). B: Immunoreactivity was weakly found (arrows) in retina transfected with \(\text{ASK-1}\) siRNA. Images are shown at 200× magnification.
from apoptosis. Lingor et al. [23] applied anti-Apaf-1 and anti-c-Jun siRNA into vitreous cavities of rats and successfully rescued retinal ganglion cells from axotomy-induced apoptosis. Since c-Jun is directly downstream of the ASK-1-JNK apoptosis pathway, the aforementioned findings are consistent with our results. In our study, siRNA was successfully incorporated in retinal explants, and siRNA for ASK-1 silenced ASK-1 mRNA expression. Considering that previous reports of RNA interference in vivo or ex vivo resulted in about 50%–80% suppression of target genes [30, 33-35], it was likely sufficient that we achieved more than 70% suppression of target genes in retinal explants. The immunoreactivity of ASK-1 was predominantly found in several ONL cells that represented nuclei of photoreceptor cells, which suggests several stresses are present in photoreceptors from rd1 mice to activate ASK-1. It is not clear why only the most distal cells in ONL expressed ASK-1, but it is possible that the increased oxygen tension in the ONL accelerates the expression of ASK-1 [36]. In contrast, ASK-1 mRNA was found predominantly in inner retinal cells in ischemic retinal injury models that cause strong oxidative stress in inner retinas [29]. ASK-1 immunoreactivity was undetectable in retinas treated with siRNA for ASK-1, which means that ASK-1 immunoreactivity in cultured retinas from rd1 mice was possibly regulated by RNA interference. Although the ONL thickness in untreated retinas from rd1 mice was markedly decreased at P16 compared with that at P9, the ONL thickness of retinas transfected with siRNA for ASK-1 was preserved significantly over the same course. However, all retinal explants tended to decrease in thickness including ONLs in the present study, probably due to the influence of organ culture conditions [37]. This finding was supported by TUNEL assays showing that TUNEL-positive cells were significantly fewer in retinas transfected with siRNA for ASK-1 at four days and seven days of incubation compared with untransfected control retinas. These findings suggest that ASK-1 is an important molecule for the apoptosis of photoreceptors in the rd1 mouse. However, the distribution of TUNEL-positive cells was not matched to that of ASK-1-positive cells, which suggested the existence of other apoptosis pathways independent of ASK-1.

Kim et al. [38] attempted to rescue photoreceptors from rd1 mice by inhibiting calcium ion dependent proteases or oxygen radicals, but this approach failed to halt apoptosis of...
Figure 6. TUNEL-positive cells. A: In untreated retinas from rd1 mice, almost all TUNEL-positive cells were found in ONLs while few were found in INLs. B: There were few TUNEL-positive cells in retinas transfected for ASK-1 siRNA-2. C: The transfection of ASK-1 siRNA-2 significantly decreased TUNEL-positive cells compared with untransfected retinas from rd1 mice on day 4 (P13) and day 7 (P16). Filled column denotes control rd1 mouse retina, and open column indicates rd1 mouse retina transfected for siRNA-2. Data are presented as the average±SD of three independent experiments.
photoreceptors. Other studies using p53 or p75NTR knockout rd1 mice did not stop photoreceptor apoptosis [39,40]. In contrast, increasing glutathione transferase levels, adding growth factors such asCNTF with BDNF, or inhibiting poly 
(ADP-ribose) polymerase promotes rescued rd1 photoreceptors [41-43]. These facts indicate that the mechanism of apoptosis in the rd1 mouse is quite complicated and multiple pathways may work simultaneously. Although apoptosis may occur by several signaling cascades, ASK-1 may be a good target, since it is involved in most stress-induced apoptosis pathways [29-31]. However, downstream signals of ASK-1 are unclear in photoreceptor apoptosis in the rd1 mouse. Harada et al. reported that p38 MAP kinase expression was suppressed in the retina of ASK-1 knockout mice compared with control mice after ischemic retinal injury, while JNK expression was unchanged [29,44]. Further investigations will be needed to determine the complete mechanism of photoreceptor apoptosis in the rd1 mouse.

Of course, possible side effects from inhibiting ASK-1 should be considered. A previous report demonstrated that retinal structures and some cell death during development were normal in ASK-1 knockout mice [29]. Interestingly, the knockouts were less susceptible to ischemic injury, and the number of surviving retinal neurons was significantly increased compared with that in wild-type mice. Another problem requiring a solution is the discovery of effective in vivo retinal drug delivery systems for siRNA. Since the in vivo duration of a single administration of siRNA is thought to be one to four weeks [45,46], repeated treatment is required to maintain healthy photoreceptors over time. Chemical modification of siRNA may improve the stability and lengthen in vivo effects [46].

In conclusion, RNA interference of ASK-1 may become a preventative strategy for retinal degeneration associated with mutations in the cGMP phosphodiesterase beta-subunit. However, further investigations are needed to define the complete effects of ASK-1 inhibition in vivo.

**ACKNOWLEDGMENTS**

This work was supported by a grant-in-aid from the Japanese Retinitis Pigmentosa Society (SH).

**REFERENCES**

1. Boughman JA, Conneally PM, Nance WE. Population genetic studies of retinitis pigmentosa. Am J Hum Genet 1980; 32:223-35. [PMID: 7386458]
2. Farrar GJ, Kenna P, Jordan SA, Kumar-Singh R, Humphries MM, Sharp EM, Sheils D, Humphries P. Autosomal dominant retinitis pigmentosa: a novel mutation at the peripherin/RDS locus in the original plinklinked pedigree. Genomics 1992; 14:805-7. [PMID: 1427912]
3. Bascom RA, Manara S, Collins L, Molday RS, Kalnins VI, McIlnnes RR. Cloning of the CDNA for a novel photoreceptor membrane protein (rom-1) identifies a disk rim protein family implicated in human retinopathies. Neuron 1992; 8:1171-84. [PMID: 1610568]
4. McLaughlin ME, Sandberg MA, Berson EL, Dryja TP. Recessive mutations in the gene encoding the beta-subunit of rod phosphodiesterase in patients with retinitis pigmentosa. Nat Genet 1993; 4:130-4. [PMID: 8394174]
5. Farrar GJ, McWilliam P, Bradley DG, Kenna P, Lawler M, Sharp EM, Humphries MM, Eiberg H, Conneally PM, Trofatter JA, Humphries P. Autosomal dominant retinitis pigmentosa: linkage to rhodopsin and evidence for genetic heterogeneity. Genomics 1990; 8:35-40. [PMID: 2081598]
6. Bowes C, Li T, Danciger M, Baxter LC, Applebury ML, Farber DB. Retinal degeneration in the rd mouse is caused by a defect in the beta subunit of rod cGMP-phosphodiesterase. Nature 1990; 347:677-80. [PMID: 1977087]
7. Lolley RN, Rong H, Craft CM. Linkage of photoreceptor degeneration by apoptosis with inherited defect in phototransduction. Invest Ophthalmol Vis Sci 1994; 35:358-62. [PMID: 8112981]
8. Portera-Cailliau C, Sung CH, Nathans J, Adler R. Apoptotic photoreceptor cell death in mouse models of retinitis pigmentosa. Proc Natl Acad Sci USA 1994; 91:974-8. [PMID: 8302876]
9. Rohrer B, Pinto FR, Hulse KE, Lohr HR, Zhang L, Almeida JS. Multidestructive pathways triggered in photoreceptor cell death of the rd mouse as determined through gene expression profiling. J Biol Chem 2004; 279:41903-10. [PMID: 15218024]
10. Doonan F, Donovan M, Cotter TG. Activation of multiple pathways during photoreceptor apoptosis in the rd mouse. Invest Ophthalmol Vis Sci 2005; 46:3530-8. [PMID: 16186330]
11. Doonan F, Donovan M, Cotter TG. Caspase-independent photoreceptor apoptosis in mouse models of retinal degeneration. J Neurosci 2003; 23:5723-31. [PMID: 12843276]
12. Takeda K, Matsuzawa A, Nishitoh H, Ichijo H. Roles of MAPK/ASK in stress-induced cell death. Cell Struct Funct 2003; 28:23-9. [PMID: 12655147]
13. Ichijo H, Nishida E, Irie K, ten Dijke P, Saitoh M, Moriguchi T, Takagi M, Matsumoto K, Miyazono K, Gotoh Y. Induction of apoptosis by ASK1, a mammalian MAPKKK that activates SAPK/JNK and p38 signaling pathways. Science 1997; 275:90-4. [PMID: 8974401]
14. Chang HY, Nishitoh H, Yang X, Ichijo H, Baltimore D. Activation of apoptosis signal-regulating kinase 1 (ASK1) by the adapter protein Daxx. Science 1998; 281:1860-3. [PMID: 9743501]
15. Saitoh M, Nishitoh H, Fujii M, Takeda K, Tobiume K, Sawada Y, Kawabata M, Miyazono K, Ichijo H. Mammalian thioredoxin is a direct inhibitor of apoptosis signal-regulating kinase (ASK) 1. EMBO J 1998; 17:2596-606. [PMID: 9564042]
16. Hatai T, Matsuzawa A, Inoshita S, Mochida Y, Kuroda T, Sakamaki K, Kuida K, Yonehara S, Ichijo H, Takeda K. Execution of ASK1-induced apoptosis by the mitochondria-dependent caspase activation. J Biol Chem 2000; 275:26576-81. [PMID: 10849426]
17. Tobiume K, Matsuzawa A, Takahashi T, Nishitoh H, Morita K, Takeda K, Minowa O, Miyazono K, Noda T, Ichijo H. ASK1 is required for sustained activations of JNK/p38 MAP kinases and apoptosis. EMBO Rep 2001; 2:222-8. [PMID: 11266364]
18. Nishitoh H, Matsuzawa A, Tobiume K, Saeugusa K, Takeda K, Inoue K, Horii S, Kakizuka A, Ichijo H. ASK1 is essential for endoplasmic reticulum stress-induced neuronal cell death triggered by expanded polyglutamine repeats. Genes Dev 2002; 16:1345-55. [PMID: 12050113]

19. Takeda K, Matsuzawa A, Nishitoh H, Tobiume K, Kishida S, Ninomiya-Tsuji J, Matsumoto K, Ichijo H. Involvement of ASK1 in Ca2+-induced p38 MAP kinase activation. EMBO Rep 2004; 5:161-6. [PMID: 14749717]

20. Corey DR. Chemical modification: the key to clinical application of RNA interference. J Clin Invest 2007; 117:3615-22. [PMID: 18060019]

21. Palfi A, Ader M, Kiang AS, Millington-Ward S, Clark G, Palfi A. Down-regulation of apoptosis mediators by RNAi inhibits axotomy-induced retinal ganglion cell death in vivo. Brain 2005; 128:550-8. [PMID: 1569426]

22. Hatakeyama J, Kageyama R. Retrovirus-mediated gene transfer to retinal explants. Methods 2002; 28:387-95. [PMID: 12507456]

23. Murphy N, Bonner HP, Ward MW, Murphy BM, Prehn JH, Henshall DC. Depletion of 14–3–3 zeta elicits endoplasmic reticulum stress and cell death, and increases vulnerability to kainate-induced injury in mouse hippocampal cultures. J Neurochem 2008; 106:978-88. [PMID: 18466333]

24. Comes N, Borrás T. Functional delivery of synthetic naked siRNA to the human trabecular meshwork in perfused organ cultures. Mol Vis 2007; 13:1363-74. [PMID: 17768383]

25. Davies JA, Lademery M, Hohenstein P, Michael L, Shafe A, Spraggon L, Hastie N. Development of an siRNA-based method for repressing specific genes in renal organ culture and its use to show that the Wt1 tumour suppressor is required for nephron differentiation. Hum Mol Genet 2004; 13:235-46. [PMID: 14645201]

26. Stone J, Maslim J, Ader M, Kiang AS, Millington-Ward S, Clark G, Barnett N, Provis J, Lewis G, Fisher SK, Bisti S, Gargini PF, Everitt B. Mechanisms of photoreceptor death and survival in mammalian retina. Prog Retin Eye Res 2007; 26:260-8. [PMID: 16419083]

27. Gorbatyuk M, Justilien V, Liu J, Hauswirth WW, Lewin AS. Suppression of mouse rhodopsin expression in vivo by AAV mediated siRNA delivery. Vision Res 2007; 47:1202-8. [PMID: 17292939]

28. Wang J, Jiang S, Kwong JM, Sanchez RN, Sadun AA, Lam TT. Nuclear factor-kappaB p65 and upregulation of interleukin-6 in retinal ischemia/reperfusion injury in rats. Brain Res 2006; 1081:211-8. [PMID: 16530172]

29. O'Reilly M, Palfi A, Chadderton N, Millington-Ward S, Ader M, Cronin T, Tuohy T, Auricchio A, Hildinger M, Tivnan A, McNally N, Humphries MM, Kiang AS, Humphries P, Kenna PF, Farrar GJ. RNA interference-mediated suppression and replacement of human rhodopsin in vivo. Am J Hum Genet 2007; 81:127-35. [PMID: 17564969]

30. Yu DY, Cringe SJ. Oxygen distribution in the mouse retina. Invest Ophthalmol Vis Sci 2006; 47:1109-12. [PMID: 16505048]

31. Johnson TV, Martin KR. Development and characterization of an adult retinal explant organotypic tissue culture system as an in vitro intraocular stem cell transplantation model. Invest Ophthalmol Vis Sci 2008; 49:3503-12. [PMID: 18408186]

32. Bernardini C, Fantinati P, Zannoni A, Forni M, Tamanini C, Bacci ML. Expression of HSP70/HSC70 in swine blastocysts: effects of oxidative and thermal stress. Mol Reprod Dev 2004; 69:303-7. [PMID: 15349842]

33. Azadi S, Johnson LE, Paquet-Durand F, Perez MT, Zhang Y, Ekström PA, van Veen T. Decreased glutathione transferase levels in rd1/rd1 mouse retina: replenishment protects photoreceptors in retinal explants. Neuroscience 2005; 131:935-43. [PMID: 15749346]

34. Nakamura K, Harada C, Okamura A, Namekata K, Mitamura Y, Yoshida K, Ohno S, Yoshida H, Harada T. Effect of p75NTR on the regulation of photoreceptor apoptosis in the rd mouse. Mol Vis 2005; 11:1229-35. [PMID: 16402023]

35. Ahuja P, Caffè AR, Ahuja S, Ekström P, van Veen T. Decreased expression of the p53 gene in rd mouse retina. J NeuroOphthalmol 2005; 25:208-15. [PMID: 15616881]

36. Nakamura K, Harada C, Okamura A, Namekata K, Mitamura Y, Yoshida K, Ohno S, Yoshida H, Harada T. Effect of p75NTR on the regulation of photoreceptor apoptosis in the rd mouse. J NeuroOphthalmol 2005; 25:208-15. [PMID: 15616881]

37. Johnson TV, Martin KR. Development and characterization of an adult retinal explant organotypic tissue culture system as an in vitro intraocular stem cell transplantation model. Invest Ophthalmol Vis Sci 2008; 49:3503-12. [PMID: 18408186]

38. Kim JH, Kim JH, Yu YS, Jeong SM, Kim KW. Delay of photoreceptor cell degeneration in rd mice by systemically administered phenyl-N-tert-butylnitrore. Korean J Ophthalmol 2005; 19:288-92. [PMID: 16491819]

39. Wu J, Trogadis J, Bremmer R. Rod and cone degeneration in the rd5 mouse is p53 independent. Mol Vis 2001; 7:101-6. [PMID: 11344337]

40. Azadi S, Johnson LE, Paquet-Durand F, Perez MT, Zhang Y, Ekström PA, van Veen T. Decreased expression of the p53 gene in rd mouse retina. J NeuroOphthalmol 2005; 25:208-15. [PMID: 15616881]

41. Nakamura K, Harada C, Okamura A, Namekata K, Mitamura Y, Yoshida K, Ohno S, Yoshida H, Harada T. Effect of p75NTR on the regulation of photoreceptor apoptosis in the rd mouse. Mol Vis 2005; 11:1229-35. [PMID: 16402023]

42. Ahuja P, Caffè AR, Ahuja S, Ekström P, van Veen T. Decreased glutathione transferase levels in rd1/rd1 mouse retina: replenishment protects photoreceptors in retinal explants. Neuroscience 2005; 131:935-43. [PMID: 15749346]

43. Azadi S, Johnson LE, Paquet-Durand F, Perez MT, Zhang Y, Ekström PA, van Veen T. CNTF+BDNF treatment and neuroprotective pathways in the rd1 mouse retina. Brain Res 2007; 1129:116-29. [PMID: 17156753]

44. Paquet-Durand F, Silva J, Talukdar T, Johnson LE, Azadi S, van Veen T, Ueffing M, Hauck SM, Ekström PA. Excessive activation of poly (ADP-ribose) polymerase contributes to inherited photoreceptor degeneration in the retinal degeneration 1 mouse. J Neurosci 2007; 27:10311-9. [PMID: 17881537]

45. Harada C. Role of apoptosis signal-regulating kinase 1 (ASK1)-mediated signaling pathway during ischemic retinal injury (Japanese). Nippon Ganka Gakkai Zasshi 2008; 112:965-74. [PMID: 19069379]

46. Takabatake Y, Isaka Y, Mizut M, Kawachi H, Takahara S, Imai E. Chemically modified siRNA prolonged RNA interference in renal disease. Biochem Biophys Res Commun 2007; 363:432-7. [PMID: 17880921]
silencing. Biotechnol Bioeng 2007; 97:909-21. [PMID: 17154307]