Application of femtosecond ultrashort pulse laser to photodynamic therapy mediated by indocyanine green

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EXTENDED REPORT

Backgrounds/aims: To evaluate treatment with high peak power pulse energy by femtosecond ultrashort pulse laser (titanium sapphire laser) delivered at an 800 nm wavelength for corneal neovascularisation using photodynamic therapy (PDT) mediated by indocyanine green (ICG).

Methods: Using a gelatin solid as an in vitro corneal model, the safety of laser power was studied to determine if it degenerated gelatin with or without ICG. The authors then induced corneal neovascularisation in rabbit eyes by an intracorneal suturing technique. Fluorescein angiography was used to evaluate occlusion before PDT and 0, 1, 3, and 10 days after PDT. The authors performed light microscopy with haematoxylin eosin staining and transmission electron microscopy to determine thrombosis formation in the neovascular regions.

Results: The threshold of peak laser power density ranged from 39 to 53 W/cm². Laser irradiation was started 30 seconds after a 10 mg/kg ICG injection, and all irradiated segments were occluded at 0, 1, 3, and 10 days at 3.8 J/cm². Light and electron microscopy documented thrombosis formation in the neovascular region.

Conclusion: Femtosecond pulse laser enhanced by ICG can be used for PDT. Because of effective closure of corneal neovascularisation at a low energy level, the high peak power pulse energy of the femtosecond pulse laser might be more efficacious than continuous wave laser for use with PDT.
was measured by spectrophotometry (Diaphot 300, Nikon, Tokyo, Japan). Absorbance was measured at the 805 nm wavelength with each concentration of the ICG solution.

**Threshold of laser power with gelatin with and without ICG**

We investigated the irradiation effects of the femtosecond laser on gelatin containing ICG. The laser was focused on the inside of the gelatin target using a 250 mm focal length refracting mirror to a focal spot of about 0.2 μm in full width at half maximum (focal spot size area, 0.047 cm²), providing a maximum laser intensity of about 10¹⁴ W/cm² with 150 femtosecond pulses (maximum laser fluency of 1 J/cm²). The laser intensity was varied by inserting neutral density filters into the beam. The laser beam focused in the gelatin and formed the denatured area.

Gelatin (Sigma Chemical, St Louis, MO, USA) (20 mg) was dissolved in 180 ml distilled water. Gelatin was coagulated in a 1 cm cell. The mean power density was measured to degenerate the gelatin. Using gelatin mixed with 1.5 mg/dl of ICG, the mean power density was also measured.

**Corneal neovascularisation**

This study was conducted in accordance with the Association for Research in Vision and Ophthalmology Statement for the Use of Animals in Research.

To evaluate the effect of PDT using a femtosecond pulse laser mediated by ICG, a corneal neovascularisation model in rabbit eyes was used, which is an established model using the technique of intracorneal sutures. Thirty one albino rabbits weighing 2–2.5 kg were used. Corneal neovascularisation was induced in one eye of each animal using an intracorneal suturing technique as previously described. Briefly, the rabbits were anaesthetised with intramuscular injections of ketamine hydrochloride (35 mg/kg) and xylazine hydrochloride (5 mg/kg). Silk sutures (size 7–0) were placed to knots was buried. Topical ofloxacin ointment was applied at the positions on the cornea. The sutures started from the limbus and advanced toward the centre of the cornea. None of the vessels had closed in 10 rabbits. Five rabbits at day 0, two at day 1, and three at day 3 were sacrificed by intravenous injection of an overdose of pentobarbital sodium. The eyes were enucleated and fixed in paraformaldehyde for electron microscopy and 4% glutaraldehyde for electron microscopy for 30 minutes. The corneas and iris in the irradiated areas were then dissected from the fixed tissue. For light microscopy, the specimens were immersed again in paraformaldehyde and fixed overnight at room temperature. They then were dehydrated in ethanol, stained with uranyl acetate and lead citrate were observed by electron microscopy (H-7000, Hitachi, Tokyo, Japan).

**Photothermolysis**

Indocyanine green was reconstituted with sterile distilled water and administered at a dose of 10 mg/kg as a photosensitizer. General anaesthesia was induced in the animals as previously described. Local anaesthesia was also induced with oxybuprocaine before each experiment.

ICG was injected through the marginal ear vein. Laser irradiation was started at 30 seconds in 23 areas and 5 minutes in 22 areas after ICG injection. Because the laser instrument was not attached to the ocular probe, the laser light was guided with refracting mirrors. The irradiation parameters were as follows: peak power density, 80 W/cm² (mean power density, 38 mW/cm²); focal spot size area, 0.047 cm²; repetition rate, 1 kHz; pulse width, 150 femtoseconds; total irradiation time, 100 seconds; and total irradiation energy, 3.8 J/cm². Before each period of irradiation, the mean power density was calibrated to 38 mW/cm².

Photodynamic therapy was usually applied to two corneal neovascularisation lesions and one corneal neovascularisation lesion was left untreated as a control (fig 1). To evaluate the femtosecond laser effect alone, four eyes were irradiated at a peak power density of 80 W/cm² without ICG injection. Four eyes were left untreated to observe the natural course.

**Histologic examination**

We performed a histological evaluation to determine that the vessels had closed in 10 rabbits. Five rabbits at day 0, two at day 1, and three at day 3 were sacrificed by intravenous injection of an overdose of pentobarbital sodium. The eyes were enucleated and fixed in paraformaldehyde for light microscopy and 4% glutaraldehyde for electron microscopy for 30 minutes. The corneas and iris in the irradiated areas were then dissected from the fixed tissue. For light microscopy, the specimens were immersed again in paraformaldehyde and fixed overnight at room temperature. They then were dehydrated in ethanol, stained with haematoxylin and eosin. For electron microscopy, the samples were immersed again and fixed overnight at 4°C. They then were dehydrated in ethanol, fixed in osmium tetroxide, and embedded in epoxy resin. Thin sections stained with uranyl acetate and lead citrate were observed by electron microscopy (H-7000, Hitachi, Tokyo, Japan).

**RESULTS**

**Absorbance of ICG**

The optical spectrum using any ICG concentration consisted of two main peaks, mainly at 800–805 nm (fig 2). The absorbance value was significantly correlated with the ICG concentration at 805 nm (by Pearson product moment correlation, correlation coefficient = 0.979, p = 0.000678) (fig 3).

**Threshold of laser power using gelatin with and without ICG**

Although gelatin without ICG did not degenerate at a peak power density of 42 W/cm² (fig 4B), it started to do so at a peak power density of 53 W/cm² (fig 4C). Gelatin mixed with 1.5 mg/dl of ICG started to degenerate at a peak power density of 38 mW/cm² (fig 3).
density of 39 W/cm² and degenerated at 42 and 53 W/cm² (fig 4D–F).

From both these examinations in an in vitro model, the threshold power levels at which the gelatin did not degenerate without ICG and at which it did degenerate with ICG, ranged from 39 W/cm² to 53 W/cm².

ICG angiography of corneal neovascularisation
Thirty seconds after ICG injection, ICG angiography clearly defined the network of corneal neovascularisation and iris vessels. One minute after ICG injection, the hyperfluorescence from the iris vessels decreased, and the hyperfluorescence from the corneal neovascularisation increased with enlargement of the area of leakage. Five minutes after ICG injection, fluorescence from the corneal neovascularisation weakened, and disappeared from the iris.

Fluorescein angiography in PDT using ICG
The irradiation time was 30 seconds and 5 minutes after ICG injection. Twenty three lesions were irradiated after 30 seconds and 22 lesions after 5 minutes.

Fluorescein angiography showed that 3 hours after PDT all 23 lesions, irradiated 30 seconds after ICG injection, were closed (fig 5B). The remaining 18 of 18 lesions were still closed at day 1 in all remaining 18 eyes (fig 5C). Irradiated lesions continued to be closed until day 10 in all remaining 12 eyes.

Of the 22 lesions, irradiated 5 minutes after ICG injection, 21 lesions were closed and only one lesion was open 3 hours after PDT. Fourteen of 16 lesions closed on day 3 and they closed in seven of 13 eyes on day 10. Four lesions treated using the titanium sapphire laser alone did not close in any eyes (table 1).

Corneal thinning or bubble formation was not detected by slit lamp biomicroscopy in any eyes that received laser irradiation. No photosensitivity developed in any treated animals.

Histology
Light microscopy revealed that the corneal neovascularisation was packed with erythrocytes with no invasion of inflammatory cells 3 hours after PDT (fig 6, top left). Thrombi were observed at day 3 and neovascularisation decreased (fig 6, top right). Thermal damage around the neovascularisation was not apparent. Light microscopy showed patent blood vessels without alterations in the iris 3 hours after PDT (fig 6C) and on day 3 (fig 6D).

Electron microscopy on day 3 after PDT showed cell remnants and some amorphous materials suggestive of erythrocytes in the vascular lumen (fig 7). Swelling of the endothelial cells and nuclei were detected. Nuclear alterations of the endothelial cell suggesting cell death were found. Vacuolisation of the endothelial cell cytoplasm was also observed. Neither ablation nor disorganisation of collagen fibrils was detected.
DISCUSSION

Our results show the effects of femtosecond pulse laser with PDT enhanced by ICG using a corneal neovascularisation model. We succeeded in closing the corneal neovascularisation at 3.8 J/cm², which is very low energy. In previous reports of in vitro experiments with PDT mediated by ICG using a continuous wave laser, the authors reported using a 830 nm wavelength diode laser at a radiance of 30–100 J/cm² (100–200 mW/cm²) on melanoma cells¹¹ and a 805 nm wavelength continuous wave diode laser at a radiance of 30 J/cm² (40 mW/cm²) on colon cancer cells.¹²

In in vivo experiments, the choriocapillaris of normal rabbits closed successfully using PDT mediated by ICG at doses of 10 and 20 mg/kg with a diode laser tuned to 810 nm at a radiance of 6.3 J/cm².¹⁴ In a clinical trial, occult choroidal neovascularisation in age related macular degeneration was successfully closed using 1.5 mg/kg of ICG at 3000 mW/cm² for 95 seconds (calculated as 285 J/cm²).¹³ In Kaposi’s sarcoma, 2.0 and 4.0 mg/kg of ICG were used to successfully close the lesions with a diode laser (805 nm wavelength and total energy of 100 J/cm²).¹⁷ Compared with previous reports, the energy level used in the present study was exceptionally low, which is the advantage of the high peak power pulse energy of the femtosecond pulse laser.

Femtosecond pulse laser has been used during refractive surgery using a photodisruption phenomenon and subthreshold laser to the retina in a few femtoseconds of irradiation. Photodisruption using the femtosecond laser is more localised than that caused by picosecond or nanosecond pulses.¹⁸ Photodisruption occurs when laser light by femtosecond pulses is focused to power intensities ranging from 10¹¹ to 10¹² W/cm².³ Optical breakdown is the process by which transparent material is ionised by multiphoton absorption.¹⁹ A shock wave is created by the plasma ignition and explosive expansion. When this process occurs in a fluid, bubbles develop in a cavity. From the power range causing photodisruption, the power of this study was very low and did not generate bubble formation or ablate the tissue, making the femtosecond pulse laser much safer than other lasers.

In an in vivo study in which venous varicosities were treated by 808 nm continuous diode laser enhanced by ICG,
the histological results of light microscopy showed an area of focal thermal damage surrounding the vessels. Temperature increases ranging from 5–25°C within tumours were reported after PDT using haematoporphyrin derivative, and a thermal effect on normal tissue was suggested. The ideal effect of PDT is no or minimal damage to the surrounding normal tissue from the secondary thermal effect. Our histological results did not show thermal damage in surrounding tissue.

From a report that analysed time resolved fluorescence spectroscopy using ICG, the response time from irradiation to the maximum signal fluorescence was approximately 1 nanosecond, and no fluorescence signal was detected at approximately 4 nanoseconds. The interval between each laser peak in the present study was 1 millisecond calculated from the repetition rate of 1 kHz, which might be sufficient to cool down surrounding normal tissue to prevent thermal spread. Continuous wave laser has no cooling effect. Considering this, femtosecond pulse laser might be the ideal laser for PDT. Interruptions in the femtosecond pulse laser really help to reduce the thermal effect without decreasing the photodynamic effect, compared with the continuous wave laser.

The disappearance rates of ICG in healthy subjects ranged from 18% to 24% per minute and the normal biological half life is approximately 3 minutes. The ICG concentration at 5 minutes was estimated to be approximately 30% of the initial ICG concentration. At day 10, occlusion of new vessels was seen in 100% (12/12) when tissue was exposed to the laser for 30 seconds after ICG injection, and the closure rate was 54% (7/13) when the laser was applied 5 minutes after ICG injection. It is reasonable that laser exposure is more effective at 30 seconds than 5 minutes after ICG injection when one considers ICG metabolism and the correlation between ICG absorption and concentration (fig. 3).

To obtain the optimal goal of closing only the neovascular lesions and not injuring the surrounding normal tissue, various parameters must be examined—for example, the photosensitiser dose, the timing of the start of irradiation, laser power, and laser duration. In preliminary experiments, we performed PDT using ICG at a dose of 1.0 mg/kg and at a peak power density of 53 W/cm². With these parameters, no occlusion was obtained in any eyes (unpublished data). We used ICG at a dose of 10 mg/kg, the concentration reported by Costa et al. An intravenous LD50 of ICG in a rat was reported to range from 50–80 mg/kg, concentrations that are much higher than that used in our study. Determining the optimal concentration of ICG should be addressed before a clinical trial is performed, although less ICG might induce less thermic reaction following less of a photodynamic reaction.

We chose the time to start the first irradiation at 30 seconds following the results of ICG angiography of corneal neovascularisation. We set the second irradiation at 5 minutes because of the total irradiation time of 100 seconds. For example, the irradiation time starting at 1 minute overlaps 70 seconds with the time starting at 30 seconds. In this study, we used a laser power in the in vivo experiments that was determined from the results of threshold experiments performed in vitro using gelatin mixed with 1.5 mg/dl of ICG. Injected ICG was diluted within the rabbit body and the highest concentration soon after the injection was estimated to be approximately 1 mg/dl of ICG. Based on the correlation between absorption and ICG concentration (fig 3), the peak power was set at 80 W/cm², which is 53 W/cm² multiplied by 1.5, from the difference of the ICG concentrations between the in vitro and the in vivo experiments.

Further investigations are required to confirm that PDT using the femtosecond pulse laser and ICG can be used clinically to treat neovascular tissues. Further, this study did not prove long term efficacy, as the follow up time was only 10 days. Longer follow up is needed to show advanced efficacy.

The femtosecond pulse laser might be a potential treatment modality used for PDT with ICG and it may herald an era of new treatments.

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