Fiberoptic surgery by ultrabright lamp light

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Abstract. We report the first realization of interstitial surgery by ultrabright lamp light, on the kidneys and livers of live animals. A high-flux optic concentrates lamp emissions into an optical fiber for power delivery inside the body. The trials reveal surgical efficacy comparable to corresponding laser fiber optic treatments, as well as pronounced delayed tissue death. © 2006 Society of Photo-Optical Instrumentation Engineers. [DOI: 10.1117/1.2363354]

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Laser fiberoptic surgery is a minimally invasive technique for the rapid and localized necrosis of growths in internal organs.^{1–4} The considerable cost of laser surgical systems prompts the quest for less expensive noncoherent photonic alternatives, especially since the key essential laser characteristic often is immense power density rather than monochromaticity or coherence.

Recently, we demonstrated that highly concentrated solar radiation can generate the same degree, rate, and type of photothermal tissue damage as lasers.⁵ However, its value is restricted by the ephemeral nature of sunlight. Here, we demonstrate the first exploitation of artificial noncoherent light for photothermal surgery on live rats, with an inexpensive device as efficacious as many surgical lasers.

A compact tabletop optical system was developed⁶ [Fig. 1(a)] that reconstitutes the power density from the radiant plasma of a commercial ultrabright lamp inside an optical fiber that delivers light inside the body [Figs. 1(b) and 1(c)]. The light source was a 150-W xenon short-arc discharge lamp,⁷ with a 2.0-mm electrode gap, which emits an approximately 6000-K graybody spectrum enhanced by strong xenon emission peaks at wavelengths from 800 to 1000 nm. This source is suitable for photothermal surgery because 1. it subsumes visible and near-infrared wavelengths where tissue optical penetration is greatest,^{1,2} and 2. its brightness is high enough to provide the required power density, a feat unattain-

able with other light sources such as light-emitting diodes or incandescent lamps.

A dual-mirror compact aplanatic optic [Fig. 1(a)] concentrates lamp emissions into an optical fiber. The (silvered specular) reflector contours enable high-flux transfer by eliminating spherical aberration and coma for a relatively small light source [Fig. 1(a)], and are essentially achromatic. The analytic equations that describe the reflectors were reported in Ref. 8. A hemispherical mirror recycles lamp emissions and heightens delivered power by about 50%.

The fiber had a quartz core diameter of 1.0 mm, a teflon cladding, and a numerical aperture (NA) of 0.66. The fact that high power density mandates high NA, i.e., large delivery angle, follows from the principle of phase space conservation,⁹ and limits effective lamp surgery to the fiber being inside the organ or maintained close to its surface. The largest NA value in commercially available fibers that are also transmissive over the lamp's emission spectrum (350 to 2500 nm) is 0.66. A pragmatic benefit of large-angle fiber emission is that lamp surgery does not engender the risk of eye injury to the surgical team.

The fiber was fit snugly into a protective Pyrex sleeve, via which light was injected to the organs. A sleeve thickness of about 1 mm separated the fiber tip from the irradiated tissue, such that the spot size on the tissue was around 2 mm.

Recent *ex-vivo* lamp surgery experiments⁶ guided our choice of irradiation times and flux values for generating immediate necrosis of the order of hundreds of mm³: a compromise between the large lesions often needed in human surgery and the small size of rat kidney and liver. Lamp surgery with irradiation times varying from 60 to 240 s and radiative input from 1.7 to 2.1 W was conducted on the livers and kidneys of 13 anesthetized healthy adult rats divided into groups for histopathology examination at 0 (immediate group), 24, and 48 (delayed group) hours after surgery. Input power was purposely limited toward minimizing the risks of carbonization while producing sizable necrosis with relatively short exposures. Optical losses from the light source to the net delivered radiative power were assessed by ray-trace simulation in Ref. 8. Given the additional inefficiency in converting electricity to utilizable visible and near-IR light of sufficient brightness,⁷ lamp nominal power ratings of around 100 W and higher are needed for effective photothermal surgery.

Healthy adult rats of body mass 300 to 400 g were used in all experiments. The targeted organs were exposed to view with a longitudinal midline scalpel abdominal incision. Experiments were approved by the Institutional Animal Care and Use Committee of the Soroka Medical Center, Ben-Gurion University, Beersheva, Israel, and the surgeons underwent the health protection program for handling research animals. All organs were irradiated in regions that would not interfere with large veins, thus avoiding major vascular thrombosis that could otherwise contribute to lesion size.

Both the thick and thin lobes of the liver were irradiated in all procedures. Only one kidney was treated in each of the rats in the delayed group to avoid complete renal failure. For rats in the immediate group, a single exposure was applied to both kidneys. Those animals intended for delayed pathology were revived, and functioned without complication.

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Fig. 1 Optics, surgical procedure, and pathology on living rats. (a) Drawing of the compact high-flux optic that concentrates ultrabright lamp emissions into an optical fiber. Intense noncoherent (white) light being delivered to the center of (b) the kidney and (c) liver of a live anesthetized rat, with NADH-diaphorase stain pathology for sections from rats in the immediate and 24-h groups. Irradiation of 1.9 W was applied for 60 and 90 s to these kidneys and livers, respectively. Viable tissue stains blue (dark). Nonviable (necrosed) tissue appears transparent (color online only).

Liver and kidney pathology for all procedures was investigated with NADH-diaphorase stain. Pathology for the 24 and 48 h groups was also performed with haematoxylin-and-eosin (H and E) stain for independent confirmation. The H and E technique was constrained to the delayed group, because it was previously shown to lead to dramatic underestimates for immediate pathology, the shortfall being mitigated when histological examination is postponed for 24 h or longer.¹⁰

Pathology revealed the coagulative and ablative tissue transformations characteristic of laser surgery^{1–4} [Figs. 1(b) and 1(c)]. Lesion volumes were calculated from the lateral and longitudinal cross sectional diameters observed in the pathology slides, and varied from 100 to 800 mm³ with shapes that were roughly hemiellipsoidal for liver and fully ellipsoidal for kidney. In samples where the necrosis traversed the entire organ, lesion volume was estimated from the linear dimensions of the actual asymmetric coagulated region. The extent of necrosis was distinctly larger after 24 h [Figs. 1(b) and 1(c)] but did not increase further at 48 h.

For a given time after surgery, the volume of dead tissue was approximately proportional to both exposure time and input power. Surgical efficacy E (lesion volume per delivered light energy) is then a suitable figure of merit for assessing and comparing results. Since no significant differences were

discerned between animals in the 24 and 48 h groups, those results were lumped into a single delayed category.

We found $E_{\text{kidney}} = 1.4 \pm 0.4$ and $E_{\text{liver}} = 1.0 \pm 0.2$ mm³/J immediately after irradiation—values commensurate with corresponding results for laser fiberoptic^{1–4} and solar⁵ surgery. For the delayed groups, $E_{\text{kidney}} = 2.3 \pm 0.4$ and $E_{\text{liver}} = 2.7 \pm 1.2$ mm³/J. Some lesions traversed the entire liver lobe [Fig. 1(c)]. They represent a lower bound for surgical efficacy, and account for the relatively large variance in E_{liver} .

When induced photothermally, delayed tissue death is an athermal process of sizable degree.^{11–14} The phenomenon is only sparsely documented for livers,^{10–13} and the understanding of the underlying mechanisms is incomplete.¹² There are no analogous results of which we are aware for kidneys.

The comparable efficacy of lamp and laser procedures stems from their sharing similar optical and biophysical properties.^{1–3,5,6} As such, the virtues of lamp surgery relative to hot-wire techniques, radiofrequency ablation, and cryoablation should be the same as those for laser surgery compared to the same methods.¹⁴

The confluence of maximum-flux optics and the suitability of ultrabright lamp light for interstitial photothermal surgery bodes well for the feasibility of a photonic tool that is potentially far less expensive than corresponding laser surgical systems and offers comparable effectiveness.

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