Temperature dependence of afterglow in zirconia and its optically-stimulated luminescence by bone-through irradiation for biological temperature probe

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Development of minimally invasive and site-selective biological temperature sensing is quite important in medical field. This study presents a novel temperature sensing technique based on afterglow and optically-stimulated luminescence (OSL). The dependence of afterglow photoluminescent intensity on the environmental temperature of zirconia (ZrO2) phosphor is examined to validate its use as a sensing probe. In addition, assuming the measurement in deep-part of human body, we have applied the information gathered from our validation to observe OSL from the ZrO2 by irradiation with near-infrared laser through a bone sample. This study demonstrates an alternative medical application of phosphor, and introduces an elemental-technology for the temperature sensing.

Body temperature is main vital sign, which is closely related to immune system and metabolism. A technique for biological temperature monitoring is a powerful tool for medical treatment and monitoring human health. In addition, targeted temperature management has the potential to reduce the risk of brain dysfunction after a head injury and cardiopulmonary arrest. Although clinical thermometers and thermography are non-invasive, the methods measure the temperature on the surface of body. The gold standard for measuring temperature in critically ill patients is the Swan-Ganz catheter (Suppl. Fig. S1). The catheter must be inserted into the body in order to monitor the temperature at the desired location. Consequently, there is a need for accessible and minimally invasive temperature sensing methods which can monitor local internal temperature.

Optical/spectroscopic measurement is recognized to be a minimally invasive and contactless method, and particularly, fluorescent and afterglow phenomena have been attempted to utilize for sensing-/imaging-applications so far9,10. Quantum-dots and fluorescent polymers have been proposed as temperature sensors for cell-imaging due to the relationship between temperature and fluorescence intensity/lifetime11,12. The observation of these effects is only possible with the magnification of a microscope. Chermont et al. developed a bio-imaging technique using afterglow phosphor of Eu2+, Dy3+, Mn2+-tri-doped Ca0.2Zn0.9Mg0.9Si2O66. Their study results inspired us to use the afterglow phenomenon to measure the temperature within the human body. In this manuscript, the fundamental measurements necessary to validate the use of the decay curve in the afterglow phosphor to measure temperature are presented. In addition, optically-stimulated luminescence (OSL) phenomena was also considered as irradiation of near-infrared (NIR) laser light would enable site-selective photoluminescence (PL), making it possible to obtain temperature information at internal sites in the human body.

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Figure 1. Temperature sensing concept for measuring local internal temperature using afterglow phosphor.

Conception and Fundamental Principle of Temperature Measurement

We propose a new concept for minimally invasive biological temperature sensing method using afterglow phosphor\(^7\). In Fig. 1, we display a rough sketch of our concept: Afterglow originates in the recombination between holes and electrons that are trapped at a metastable site and subsequently released thermally. Since the release of electrons depends on thermal excitation, the environmental temperature determines the afterglow lifetime (or slope of the afterglow curve). If only one kind of the electron-trapping site is presented in afterglow phosphor, its temperature dependence on the intensity of afterglow PL follows to simple exponential relationship\(^8\):

\[
I(t)/I(0) \propto \exp(-at),
\]

where \(I(t)\) is the afterglow PL intensity, \(I(0)\) is the initial intensity, \(t\) is the time, \(a\) is the probability of thermal activation of a trapped electron to the conduction band, \(E\) is the thermal activation energy of an electron in the trap, \(s\) is the frequency factor, \(k_B\) is the Boltzmann constant, and \(T\) is the temperature. Since the \(a\) corresponds to an inverse of lifetime for the PL, observing the temperature dependence on the \(a\) (=1/\(\tau\)), we obtain the activation energy. As a result, we are able to estimate the environmental temperature of the phosphor from the afterglow measurement. In order to estimate the temperature, a suspension consisting of afterglow nanophosphor is prepared and is excited prior to injection into the region of interest. After injection, afterglow PL is detected from the outside of body to obtain information about the internal temperature. For the afterglow, the PL wavelength should be in the NIR region, which is called to “biological window,” (~650–1100 nm)\(^9\). Alternatively, the trapped-electrons can be released optically, for instance using irradiation of a NIR-laser, and then the recombination provides the luminescence, i.e., OSL. Therefore, we believe that OSL can also be used to obtain temperature information. Since a laser is a coherent light-source, site-selective temperature sensing at any location in the body should be possible using external irradiation with a NIR-laser.

Temperature measurement based on afterglow zirconia phosphor. In this study, we have focused on afterglow zirconia (\(\text{ZrO}_2\)) as a sensing probe. \(\text{ZrO}_2\) is an accessible material and possesses a high chemical-stability and bio-compatibility\(^10\). In addition, the size, morphology, physical and optical properties can be controlled by choice of appropriate synthetic conditions and dopants\(^11,12\). Pure \(\text{ZrO}_2\) has different polymorphs, and the monoclinic phase exhibits a remarkable PL\(^13\). Furthermore, monoclinic \(\text{ZrO}_2\) acquires a prominent afterglow property through the introduction of oxygen-defects, which are activated by the annealing\(^14,15\). Iwasaki et al. have reported that thermally-treated \(\text{ZrO}_2\) powder shows pronounced afterglow property\(^16\). Therefore, we selected the thermally-treated \(\text{ZrO}_2\) powder as our sensing probe (see Methods).

In Fig. 2, we show the afterglow PL and thermoluminescent (TL) properties of the \(\text{ZrO}_2\) sample. Excitation and detected-afterglow wavelength were \(\lambda_{\text{exc}} = 280\) nm and \(\lambda_{\text{emi}} = 480\) nm, respectively. When the decay curve is plotted on a semi-log-y graph, a linear relationship between time and intensity should be obtained. The slope should be steeper for higher environmental temperatures, show faster decay, according to the Eqs. (1)–(2). However, the linearity was not observed at any temperature [Fig. 2(a)]. The decay curves showed the complex behavior at times less than 200 s when reversal of the intensity and intersection of the decay curves is seen. The TL spectrum revealed six distinguishable peaks (indicated by A to F) in the temperature range tested [Fig. 2(b)]. The decay curve and TL spectrum indicates the presence of various kind of electron-trapping sites in the \(\text{ZrO}_2\) sample, suggesting that the plural trapping sites cause the non-linearity and complicated decay curves. In order to release the electrons form the trapping sites corresponding to the TL peaks A–D appearing below room temperature (RT), so-called de-trapping, a pre-heating process at 273 K for 2.5 h before UV-excitation for 1 h was conducted prior to TL measurement. As the result, the peaks A–D vanished and indicating the release of electrons at the traps which are active below RT. The pre-heating process successfully produced a de-trapped \(\text{ZrO}_2\) sample [red curve, Fig. 2(b)]. The de-trapped sample exhibited the expected response (Eqs. (1)–(2)) of linear decay with slopes that increased with increasing temperature [Fig. 2(c)]. The response at 323 K (30 °C) deviated from the theoretical
This departure was probably due to a partial electron-release from a deeper-trapping site corresponding to the peak F with peak at ~353 K. The pre-heating process is an effective technique to minimize the non-linearity in the decay curve. Using the Arrheniusian approach [Fig. 2(d)] the activation energy for afterglow PL related to the peak E for the pre-heated samples was found to be $E \approx 0.48$ eV. Thus, we demonstrated the fundamental temperature sensing principle based on the afterglow measurement in the ZrO$_2$ sample.

Optically-stimulated luminescence by bone-through light.

Since NIR can minimize its absorption loss for the tissue texture (i.e., biological window), laser irradiation with a NIR wavelength enables us to stimulate the PL and to obtain temperature information based on the OSL. Until now, OSL phosphor has been extensively studied for the application in detection of charge particle, and particularly the ZrO$_2$ is a candidate for the OSL phosphor toward scintillator/dosimeter material because of being chemically and thermally stable. Assuming the observation of a body part covered by bone (e.g., brain), it is necessary to stimulate the luminescence via bone by external laser irradiation. This is a significant issue that should be clarified for the OSL application in human body. In order to confirm that stimulation via bone was possible, we prepared a simulated bone sample and examined the OSL from the ZrO$_2$ sample.

In Fig. 3, we show the OSL properties of the ZrO$_2$ sample. By irradiation of the NIR-laser light ($\lambda_{st}=980$ nm), we visually confirmed the bluish PL spot in the irradiated area [Fig. 3(a)]. The excited ZrO$_2$ sample exhibited a decrease in PL with time (i.e., afterglow), and a sudden increase in the PL intensity and its subsequent decay were confirmed as the ZrO$_2$ sample was irradiated with NIR-laser (808 nm) [Fig. 3(b)]. Both the OSL and PL spectrum shown in Fig. 3(c) have a broad band with peak at ~480 nm, which is similar to the usual PL spectrum (see Suppl. Fig. S2) demonstrating obvious OSL in the ZrO$_2$ sample.

In Fig. 4, we show the results of OSL by irradiation of NIR through the simulated bone sample. We have cut and polished a rectangular section (dimension: $20 \times 15 \times 0.5$ mm$^3$) of bone from a cattle femur [Fig. 4(a,b)]. The spectral features of the optical extinction spectrum for the cattle bone was quite similar what has been reported for human bone: the broad absorption bands are attributed to water and lipid molecules [Fig. 4(b)]. An optically-transmittable region in ~650–1000 nm was found, leading us to speculate that OSL by NIR light through bone is possible. The Raman spectra in the range of 300–3000 cm$^{-1}$ (Suppl. Fig. S3) also resembled the spectral patterns reported for human bone. The Raman bands could be assigned to vibration modes related to phosphate, amide, and C–H bonds. These similarities show that the cattle femur bone is an appropriate animal model for human bone.
Observation of the OSL by NIR-laser irradiation through the simulated bone sample was performed and we could visually confirm bluish PL from the ZrO$_2$ sample [Fig. 4(c,d)]. In addition, the sudden increase and subsequent decay of the PL intensity was also confirmed [Fig. 4(e)]. OSL by the bone-through irradiation. The intensity of OSL presumably exhibits the similar temperature dependence to that of afterglow phenomenon because the OSL also originates from the release of electrons in trapping sites. Therefore, estimation of the environmental
temperature is could be possible. The combination of OSL and afterglow phosphor by means of external stimulation with NIR-laser is a promising technique for site-selective temperature measurement. Indeed, the investigation of temperature dependence of OSL in the ZrO₂ sample is now in progress.

Discussion
We have proposed a new concept for internal temperature measurement using a harmless and accessible inorganic crystal, afterglow ZrO₂, as a sensing probe. In this study, the fundamental principle of temperature sensing based on afterglow/OSL and its feasibility were demonstrated. Again, we want to stress that the OSL by means of NIR-laser enables us to measure the localized temperature at internal locations (e.g., the brain) non-invasively and site-selectively. The temperature dependence of OSL intensity and the red-shift of afterglow PL and OSL in ZrO₂ phosphor must be investigated. The authors’ group has already succeeded in fabricating ZrO₂ phosphor showing afterglow PL in red region. While this fundamental study leaves some important issues unresolved, this work is a novel and promising application of phosphor in the medical field.

Methods
Commercial reagent-grade ZrO₂ powder (purity: 99.9%, Soekawa Chem. Co., Ltd., white powder with size of ~3 μm) was thermally-treated at 1400°C. The heating rate was 10 K/min and the target temperature was maintained for 6 h in an electric furnace under atmospheric conditions. After the treatment, the ZrO₂ powder was cooled in the furnace, producing defect-activated ZrO₂. The ZrO₂ sample could be identified to be monoclinic by means of a powder X-ray diffraction (XRD; Cu-Kα) analysis. The ZrO₂ sample exhibited the broad PL and PLE bands with peaks at ~480 nm and ~280 nm, respectively (Suppl. Fig. S2). In addition, the visible PL and bluish afterglow was also confirmed. The PL and afterglow features were identical to the features previously reported indicating that the ZrO₂ prepared in the earlier study was properly reproduced. A cattle femur bone that is commercially available is used as the simulated human bone sample in this study. Length and width of the cattle bone are approximately 31 cm and 13 cm, respectively [Fig. 4(a)].

The PL and PL excitation (PLE) spectra, and afterglow PL intensity as a function of time (decay curve) was obtained using a spectrophotometer with a xenon lamp as the excitation source. The environmental temperature of the ZrO₂ sample during decay curve measurement was controlled by a cryostat (Temperature range: 293–323 K; 20–50°C). Afterglow PL intensity as a function of temperature (TL spectrum) was also measured by means of the spectrophotometer conjugated with the cryostat (Temperature range: 77–450 K, heating rate: 1 K/min). Prior to the TL measurement, the ZrO₂ sample was subjected to UV-excitation (280 nm) for 1 h. The cattle-bone sample’s optical properties were characterized using a spectrophotometer with an integrating sphere and Raman scattering spectroscopy (Excitation wavelength: 532 nm).

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Author contributions
M.O. fabricated and characterized the materials. Y.T., N.T. and T.F. designed the study; N.O. and T.S. contributed the conception of this study, and Y.T. and T.F. wrote the manuscript. All authors discussed the results and commented on the manuscript.

Competing interests
The authors declare no competing interests.

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