First human Cerenkography

Antonello Enrico Spinelli
Marco Ferdeghini
Carlo Cavedon
Emanuele Zivelonghi
Riccardo Calandrino
Alberto Fenzi
Andrea Sbarbati
Federico Boschi
First human Cerenkography

Antonello Enrico Spinelli,* Marco Ferdeghini, Carlo Cavedon, Emanuele Zivelonghi, Alberto Fenzi, Andrea Sharbati, and Federico Boschi

San Raffaele Scientific Institute, Medical Physics Department and Centre for Experimental Imaging, Via Olgettina N. 60, Milan 20182, Italy
Verona University Hospital, Nuclear Medicine Department, Piazzale Ariste Stefanì 1, Verona 37126, Italy
Verona University Hospital, Medical Physics Unit, Piazzale Ariste Stefanì 1, Verona 37126, Italy
University of Verona, Department of Neurological, Neuropsychological, Morphological and Motor Sciences, Strada Le Grazie 8, Verona 37134, Italy

Abstract. Cerenkov luminescence imaging is an emerging optical preclinical modality based on the detection of Cerenkov radiation induced by beta particles when traveling through biological tissues with a velocity greater than the speed of light. We present the first human Cerenkography obtained by detecting Cerenkov radiation escaping the thyroid gland of a patient treated for hyperthyroidism. The Cerenkov light was detected using an electron multiplied charge coupled device and a conventional C-mount lens. The system set-up has been tested by using a slab of ex vivo tissue equal to a 1 cm slice of chicken breast in order to simulate optical photons attenuation. We then imaged for 2 min the head and neck region of a patient treated orally for hyperthyroidism before with 550 MBq of I-131. Co-registration between photographic and Cerenkov images showed a good localization of the Cerenkov light within the thyroid region. In conclusion, we showed that it is possible to obtain a planar image of Cerenkov photons escaping from a human tissue. Cerenkography is a potential novel medical tool to image superficial organs of patients treated with beta minus radiopharmaceuticals and can be extended to the imaging of beta plus emitters. © 2013 Society of Photo-Optical Instrumentation Engineers (SPIE) [DOI: 10.1117/1.JBO.18.2.020502]

Keywords: biomedical optics; emission; luminescence; medical imaging.

Paper 12772L received Dec. 4, 2012; revised manuscript received Dec. 29, 2012; accepted for publication Jan. 4, 2013; published online Jan. 18, 2013.

The emission of Cerenkov radiation (CR) is a well-known relativistic effect that takes place when a charged particle travels in a dielectric medium with a velocity greater than the speed of light in the medium. The in vivo detection of CR is an emerging optical preclinical modality called Cerenkov luminescence imaging (CLI); more precisely, CLI is based on the detection of CR using radiopharmaceuticals labeled with beta plus and beta minus emitters. An overview of the different applications of CLI can be found in Ref. 7.

The imaging of an animal thyroid using CLI was introduced in Ref. 8, injecting a mouse with 1–124 and I–131; in this case the authors showed a good correlation with PET and SPECT imaging.

In order to detect Cerenkov photons, we used a cooled (−80°C) electron multiplied charge coupled device (EMCCD) (Photometrics, Evolve 512) coupled with a conventional F = 1.4, 8 mm C-mount lens (Edmund Optics, LV 0814). The EMCCD has a back illuminated sensor with pixel size equal to 16 μm and a peak quantum efficiency of 90%. The images were acquired using binning=1, corresponding to a matrix size of 512 × 512 pixels and the gain of the EMCCD was set to 400. All the Cerenkov images were acquired using 2 min of exposure time.

Because of the direct interaction of gamma rays with the EMCCD sensor, a sparse spike noise pattern is generated during the Cerenkov image acquisition. The noise signal has been partially removed by applying a box median filter (size = 4 pixels) on the CLI data.

The EMCCD was placed inside a light tight dark room in order to avoid any visible light signal cross-contamination. More precisely, all the Cerenkov image acquisitions were performed in a hospital basement room with no windows. The neon lamps were removed and the entire door covered by a thick black curtain. The patient too has been surrounded by thick black curtains in order to stop any possible residue of external light.

In order to investigate the absence of any external light cross-contamination in the room, a set of 5 min images with objects (phantom or human person) without any Cerenkov sources were acquired. In this case, no light signal was detectable by the EMCCD; the mean value of the dark image was 1300 arbitrary units (AU) and the standard deviation was 350 AU. The dark image has been subtracted from the Cerenkov image.

The object to be imaged was placed at a distance equal to 50 cm from the EMCCD.

Before acquiring Cerenkov images of patients, the system set-up has been tested using a syringe filled with physiologic solution containing 2-[18F]fluoro-2-deoxy-D-glucose (FDG). The syringe was filled with 80 MBq of FDG in order to take into account the 30% to 40% thyroid uptake at 24 h and also that the emitted Cerenkov light of F-18 is 2.5 higher than I–131 as shown in Ref. 4. The dimensions of the syringe active FDG region were equal to: 1 × 1.3 cm. The EMCCD gain has been chosen by acquiring different images of the chicken phantom with different gain values. We found that a gain of 400 (range 0 to 1000) is a good compromise between signal and...
number of striking artifacts due to direct gamma interactions with the CCD.

The syringe has been covered with a 1 cm slice of chicken breast in order to simulate optical photons attenuation due to the tissues.

In order to provide a simple anatomical reference, a photographic image was always acquired before any Cerenkov image. In this case, the gain was set to 0, the exposure time was 0.5 s, the matrix dimensions were $512 \times 512$, and the object was illumined by a LED lamp. The matrices of both photographic and Cerenkov images were the same, and since nothing was moved between the two acquisitions, the two images are intrinsically co-registered.

Figure 1(a) shows a Cerenkov image of the FDG syringe without chicken phantom, and Fig. 1(b) shows the image obtained by placing a 1 cm slice of chicken breast on the syringe. As one can see, there is good localization between the CR emission escaping the chicken breast and the syringe containing FDG. This is an encouraging result considering the non-negligible thickness of the chicken breast traveled by the Cerenkov photons. The signal-to-noise ratio was, respectively equal to 13.1 and 3.3 with and without median filtering.

After having acquired the results presented in Fig. 1, we then imaged the thyroid region of a patient treated for hyperthyroidism with 550 MBq of I–131 after an uptake of 24 h. The patient was not fixed and, thus, any movement can introduce a kinematic blurring on the Cerenkov image. However, given that the acquisition time is comparable (or smaller) with respect to conventional gamma camera imaging performed without fixing the patient, we estimate a similar kinematic blur for the two methodologies.

Figure 2(a) shows the Cerenkov image of the patient and Fig. 2(b) shows the overlay between the photographic image and the Cerenkov image. As one can see there is a good localization of the visible Cerenkov light signal within the thyroid region. The localization has been performed by a nuclear medicine physician (MF) with a long experience in thyroid imaging.

To our knowledge, this is the first experimental evidence showing that it is possible to obtain a planar image of Cerenkov photons escaping from a human tissue.

We called this new imaging method Cerenkography to underline the peculiar source of visible light. The novel and interesting aspect of Cerenkography is that it provides a fast approach to image superficial organs (1 to 2 cm depth) of patients treated with beta minus radiopharmaceuticals. An interesting clinical application of Cerenkography is monitoring the iodine uptake in the thyroid in order to estimate the radiation dose given to the organ.

Generally speaking, Cerenkography can also be used to monitor the radiation dose of any other therapy for superficial organs using beta emitters like Y-90 and Lu-177.

Cerenkography can also be extended to the imaging of beta plus emitters.

As shown also by recent simulations, a CCD detector optimized for the near infrared wavelength range could significantly improve the detection of CR escaping biological tissues.

Considering the moderate costs of EMCCD’s with respect to the tomographs currently available in any nuclear medicine
department, we believe that Cerenkography could became an interesting cost-effective medical tool in future years.

Acknowledgments

The authors would like to acknowledge Prof. Carlo Tacchetti and Dr. Davide Mazza of the San Raffaele Centre for Experimental Imaging for providing the EMCCD used in our experiments. We also would like to acknowledge Prof. Alberto Del Guerra and Dr. Claudia Ceccarelli of Pisa University for the useful discussions.

References