Photoacoustic microtomography using optical interferometric detection

Robert Nuster
Karl-Franzens-University of Graz
Department of Physics
Universitätsplatz 5
Graz, 8010 Austria

Markus Holotta
Christian Kremser
University Hospital of Innsbruck
Department of Radiodiagnostics
Innsbruck, 6020 Austria

Harald Grossauer
Peter Burgholzer
Günter Paltauf
Karl-Franzens-University of Graz
Department of Physics
Universitätsplaza 5
Graz, 8010 Austria

1 Introduction
In various areas of preclinical and biological research, but also in clinical practice, there exists a need to obtain three-dimensional (3-D) images of small samples (several millimeters to some centimeters) with micrometer resolution. Typical targets are whole small animals, isolated organs of small animals, or biopsy samples. An established device for this imaging task is the X-ray microtomograph, which was first introduced in the 1980s and has found applications not only in biomedical research but also in other areas such as materials research or electronics. While X-rays give excellent contrast in many kinds of biological samples, alternative contrast mechanisms are often desirable, especially for resolving minute changes of soft tissue composition. In recent years, photoacoustic (PA) (also called optoacoustic) imaging methods have been investigated intensively as a means to reveal structures with optical contrast and a resolution comparable with high-frequency ultrasound devices. This is achieved by using short light pulses in the nanosecond range to irradiate the investigated object. The diffusely propagating light is absorbed in the object, leading to a small temperature rise and associated thermoelectric pressure. The generated pressure is proportional to the locally deposited energy density, which is in turn given by the local optical absorption coefficient multiplied with the local radiant fluence. In this way, the acoustic wave field generated in the sample is strongly influenced by the distribution of tissue chromophores and thus contains important diagnostic information. Measurement of the acoustic field and applying some kind of reconstruction algorithm yields the distribution of initial pressure (right after the light pulse) or energy density. From this distribution also, the spatially dependent optical absorption coefficient can be reconstructed, or spectrally dependent values of optical absorption as they are required for quantitative measurements of local blood oxygenation values.

High-resolution PA imaging has been demonstrated using a microscopy approach with a fixed-focus acoustic lens transducer scanning across the surface of an object together with the pulsed illumination beam. Similar resolution for the imaging of superficial blood vessels has been demonstrated with a pointlike optical ultrasound sensor scanning over a surface close to an object. Both methods are optimized for the imaging of structures near the surface of an extended object, within a range of several millimeters. They are therefore particularly...
suited for visualizing blood vasculature in or directly below the skin.

For this kind of half space imaging, the detection aperture is always smaller than $2\pi$. This aperture is defined as the solid angle occupied by the detection surface, which is the area scanned or covered by detectors, as seen by a point in the imaged object. Exact 3-D imaging requires a detection aperture of at least $2\pi$ for each reconstructed point in the object. Obviously, this cannot be provided by a limited size, planar arrangement of ultrasound detectors. A curved array or a combination of several planar arrays can yield the desired $2\pi$ minimum aperture.

In tomographic PA imaging, where a single ultrasound detector scans a curve or surface that encloses the object, the resolution is always limited by the size of the detector. It was shown theoretically that for a circular or spherical scan around an object with a typical commercially available piezoelectric detector having a flat, finite-sized active area, the resolution is highest in the center of rotation and becomes worse at positions approaching the detection curve. Increasing the object-detector distance would provide approximately constant resolution throughout the object but at the same time the frequency-dependent damping would tend to decrease high-frequency components and therefore deteriorate resolution again. A solution is the use of pointlike detectors. In particular, optical detection can provide at the same time high resolution and sensitivity. Piezoelectric detectors, on the other hand, exhibit decreasing sensitivity with decreasing size.

The method proposed in this paper combines small-sized detectors and the ability to measure signals from a sufficiently large aperture around an object. It is based on the use of a focused laser beam propagating freely in a coupling liquid surrounding the object as an acoustic detector (Fig. 1). Minute changes of optical phase are integrated along the beam path when it is crossed by an acoustic wave and are detected interferometrically. A free laser beam as acoustic detector is completely transparent, both optically and acoustically. It can therefore disturb neither the incoming light pulse nor the outgoing acoustic wave. Furthermore, the resolution can be changed by varying the numerical aperture of the focusing lens.

In this paper we demonstrate the ability of the PA tomography device to create 3-D images with constant resolution throughout the volume enclosed by the scanning optical beam. Furthermore, two scanning modes of the beam around an object are compared, and finally the imaging of a complex biological sample (an isolated mouse heart) is demonstrated.

2 PA Tomography with Line Detectors

The principles of signal acquisition and image reconstruction have been described in detail elsewhere, therefore only a short overview is given here. For an exact reconstruction, it is necessary to scan the entire acoustic field around the object with the line detector. The line always moves in a way that it is perpendicular to one single axis, about which the object is rotated. For one rotational angle, the scanning path of the detector should preferentially be a closed curve in the plane perpendicular to the line direction, completely surrounding the object. From a set of such measurements it is possible to reconstruct a 2-D projection of the initial distribution of pressure in the object $p^{2D}(u,v,t=0)$ generated by a short laser pulse. Here, the variables $u$ and $v$ are coordinates perpendicular to the line direction in the frame rotated relative to the object. The rotation axis is parallel to the $v$ direction. Due to the integration of the acoustic field along the line, this reconstruction problem is strictly two-dimensional. When projections over a total angle of $\pi$ rad are acquired, the next step in 3-D image reconstruction is the application of the inverse Radon transform to all planes perpendicular to the rotation axis. Since the inverse Radon transform is a standard mathematical operation usually implemented in software such as MATLAB, our previous studies have focused mainly on the 2-D reconstruction problem. A single linear scan combined with a frequency domain reconstruction was proposed first. Later, we worked on refinements with other scan geometries such as a half circle, an L-shaped, or even arbitrarily shaped curve using time-domain back projection, frequency-domain, or time reversal reconstruction. We also addressed the problem of limited view data acquisition and found special weight functions that can be included in a back-projection algorithm to reduce the level of artifacts in the case where the complete object is in a region where the detection aperture is between $\pi$ and $2\pi$ rad, where $\pi$ is the minimum required aperture for exact reconstruction in two dimensions.

In this paper, two scan geometries are employed, which have turned out to be well suited for 3-D imaging. Both are open curves, providing access for holding and manipulating the sample. The first one uses a combination of three linear scans, forming an open box that encloses the object from three sides (Fig. 2). The advantage of using linear scans is the possibility to use a fast, frequency-domain (FD) reconstruction algorithm. It was first introduced for 3-D imaging, where a point detector scans a plane next to the object, and can easily be adapted to 2-D imaging, where the detector scans along a straight line. Because a finite scan length introduces strong artifacts due to missing data for points lying about $\frac{1}{4}$ scan length above the detection line, the idea is to acquire data along at least a second detection line perpendicular to the first one and also to add the resulting images, either in frequency or in real space. The third line scan is added to provide a detection aperture of at least $\pi$ rad for all points localized inside the open box. A problem here is the varying detection aperture within the imaging volume. We follow the
suggestion made by Xu et al. \textsuperscript{10} to weight the reconstructed image with $2\pi/\Psi(u, v)$, where $\Psi(u, v)$ is the spatially varying detection aperture angle.

The second imaging geometry is a half circular scan of the line detector around the sample (Fig. 2). Although for such a $\pi$-arc scanning curve also an FD reconstruction is possible using an algorithm for a closed circular scan \textsuperscript{14,22} with missing data, we prefer here the 2-D version of a filtered back projection algorithm. \textsuperscript{5,17} It was originally developed for 3-D imaging with point detectors where it was found to give accurate reconstructions for detection surfaces in the shape of a closed sphere, a closed cylinder, or an infinite plane. \textsuperscript{23} The advantage of this imaging procedure is that in the back projection for each reconstruction-detection point pair, an individual weight factor can be applied, which can be used to improve the image quality in limited view imaging, providing more accurate relative amplitude values and reducing the “duplicate direction” artifact. \textsuperscript{17} The applied weight factors reduce the contribution of the detectors, which lie on opposite points on the detection curve with respect to a reconstructed point in such a way that they complement each other. Detectors having no opposite point on the detection curve receive a weight equal to 1. Such individual weight factors are not possible for the FD reconstruction. However, to achieve a similar effect for points lying near the center of the box, the reconstructions from the two line scans on the sides of the box are weighted with 0.5, whereas the line scan at the bottom of the box receives a weight of 1.

The line detector used in this study is a light beam propagating in water. Its advantage over a detector formed by a strip of piezoelectric polymer film is its omnidirectional sensitivity in the plane perpendicular to the line. \textsuperscript{24} Imaging with such a line detector should give a 3-D image with constant speed of sound and excluding acoustic attenuation. The 2-D projection images were reconstructed on a grid of $200 \times 200$ thick solid line rectangular box. The number of signals along the arc was 200. For the box scan, the number of detection points on each side had to be 200, because the spatial increment of the measurement automatically becomes the increment of the reconstructed image. Finally, in both cases 200 projections were calculated while rotating the phantom 180 deg.

Figure 3\textsuperscript{a} shows a cross section of the original phantom. The corresponding sections of the reconstructed 3-D image are shown in Fig. 3\textsuperscript{b} for the $\pi$-arc and filtered back projec-
tion and in Fig. 3(c) for the box scan using FD reconstruction. Figures 3(d) and 3(e) compare horizontal and vertical profiles, respectively, through the center of the phantom. Both reconstructions show sharp edges of the spheres, even at the periphery of the reconstruction volume. The exactness of the back-projection reconstruction is generally slightly higher, as indicated by the more accurate amplitude values, the better uniformity of amplitudes over the imaging volume, and the absence of artifacts as they appear in the FD reconstruction mainly along the vertical direction. Also, a slight distortion of the spheres is seen in the FD reconstruction near the detection curve. The FD algorithm, on the other hand, is considerably faster than the filtered back projection with weighting. In the simulations, the reconstruction of the 200 × 200 points area of one projection took 6.7 s with the back-projection algorithm and 1.0 s with the FD algorithm. Taking into account that the latter used three times more detector points, the back projection turns out to be slower by a factor of 20.

4 Experiment

4.1 Imaging Device

As an approximate line sensor we used a light beam from a continuous laser, in a Mach-Zehnder interferometer configuration (Fig. 4). The temporal resolution of this detector is limited by the beam diameter of about 40 μm in the waist of the focused beam. The imaging range, which is the range along the line detector where this high resolution can be achieved, is given by the focal depth of 16 mm. These are the measured values for the used focal length and numerical aperture. Higher resolution would require tighter focusing at the expense of a smaller imaging range.

For scanning, the sample was moved relative to the laser beam using two computer-controlled linear stages. Additionally, the sample was mounted on a rotation stage. At each position of the line detector relative to the object, only a single signal was acquired, without averaging. Samples were illuminated by pulses from an optical parametric oscillator (OPO) with a pulse duration between 5 and 8 ns.

4.2 Samples

A phantom consisting of black human hairs [photograph in the inset of Fig. 5(a)] was used to demonstrate the constant resolution. It was imaged using the π-arc scan with 91 detector positions along an arc with a radius of 10 mm and 200 angular positions of the arc relative to the sample, spanning a range of 180 deg. The acoustic waves were excited by illuminating the hair phantom with laser pulses at a wavelength of 502 nm from two directions, as shown in Fig. 4. Limited by the 10-Hz pulse repetition rate of the laser, the data acquisition for the 3-D image took about 30 min.

The second object was an isolated mouse heart prepared as follows. Immediately after excision the still beating heart of an anesthetized mouse was placed in a 4% formaldehyde solution for fixation. Since the heart continues beating for a short time after removal, the remaining blood is flushed out and replaced by the formaldehyde solution. For PA imaging, the heart was finally embedded into agarose to improve acoustic coupling. Illuminated with pulses in the near-IR range at a wavelength of 750 nm inner structures of the mouse heart were also visualized because of the increased penetration of the light at this wavelength. Pulses with a fluence of 12 mJ/cm² illuminated one side of the object from a direction parallel to the detection laser beam. The mouse heart was rotated to 179 angular positions over a range of 360 deg.

Fig. 4 Mach-Zehnder interferometer as an acoustic line detector: BS, beamsplitter; L, lens; M, mirror; M+PT, piezoactuator with mounted mirror; BPD, balanced photodetector (bandwidth 80 MHz); BPF, optical bandpass filter; and HPF, electronic high-pass filter.

Fig. 5 Maximum amplitude projections in three orthogonal directions, (a) to (c), of the reconstructed image of the human hair phantom. Inset of (a) shows a photograph of the phantom. (d) The width of the hair that was located in the center of the phantom as a function of position along the hair in the vertical (thick solid line) and in the horizontal direction (dashed line).
At each rotational position, the detector scanned a box formed by three linear scans of 18 mm length and an increment of 72 μm, giving 750 detector positions in total. Using those scanning parameters, the data acquisition took about 4 h.

5 Results

5.1 Hair Phantom

Reconstructed images of the hair phantom are shown in Fig. 5 as maximum amplitude projections in three orthogonal directions. The inset in Fig. 5(a) shows a photograph of the phantom. The width of the hair that was located in the center of the phantom is drawn in Fig. 5(d) as a function of position along the hair. This diagram shows the dependence of width in the vertical and in the horizontal directions. Considering the thickness of the hair (60 μm), the measured width is consistent with the expected resolution of about 40 μm, given by the laser beam diameter in the focal region.

5.2 Mouse Heart

Figure 6 shows four sections through the 3-D image of the mouse heart. Sections in Figs. 6(b)–6(d) are perpendicular to the section in Fig. 6(a), where the positions of the three other sections are indicated as black arrows. To see the relatively faint structures formed by the heart muscle tissue together with the more strongly absorbing structures probably caused by absorption in blood remnants, we chose to display the square root of the reconstructed relative energy density. The inner structures of the heart, such as the left and right ventricles, the left atrium, and the papillary muscles can be clearly distinguished. Also, the very fine structure of a mitral valve can be seen in two sections.

6 Discussion

PA microtomography for biological or preclinical research is mainly designed for imaging of excised biological samples or of anesthetized small animals. Other than in PA in vivo imaging, speed has not such a high priority as accuracy and resolution. It is therefore feasible to use a single scanning detector for acquisition of PA signals. The proposed focused laser beam detector requires the sample to be submerged in a coupling liquid, but there is no contact of a physical transducer with the sample. In this kind of "noncontact" detection mode, the position of the sensor relative to the sample can be controlled with great accuracy, as is required to resolve structures with micrometer precision. A drawback is that the high resolution, below 100 μm, can be achieved only in the focal range of the sensor beam. The sample must therefore be carefully positioned next to the focus. For larger samples, where this is not possible, an alternative is the use of a guided laser beam, either in an optical fiber or in a planar waveguide.27–29

Comparing the image quality, the back-projection reconstruction of π-arc data turned out to give better results, partly due to the employed weight factors that give a more accurate reconstruction in a case of limited view imaging. The FD reconstruction using box scan data was considerably faster and therefore provides easier handling of larger amounts of data. This is an advantage in cases with more complex samples, where small scan increments and therefore a large number of detection points are essential. The box scan was therefore chosen for imaging of the mouse heart.

The data set of the mouse heart shown in Fig. 6 was recorded for a currently ongoing study of a cardiac infarct model. In this study, several artificially infarcted mouse hearts are investigated using different imaging modalities [PA, micro computed tomography (CT), histology, magnetic resonance imaging (MRI)]. This study is being performed to compare the different image representations of infarcted tissue and to determine whether the different imaging modalities can provide complementary information. Furthermore, PA measurements of the same heart are performed at different wavelengths to evaluate the possibility of an accurate discrimination of different tissues by their absorption spectra. The results of this study will be presented in forthcoming publications.

Currently, using a single line detector the imaging time is in the range of hours. It is mainly determined by the repetition rate of the laser and by the amount of detection points along the scanning curve. An obvious improvement, providing a reduction of data acquisition time by a factor of about 100, is therefore the use of a pulsed laser with a repetition rate in the kilohertz range. Another approach, which we are currently pursuing, is to speed up the data acquisition by using some kind of parallel detection to avoid the time-consuming scanning around the object. In conclusion, PA tomography with a scanning free laser beam and interferometric detection can provide 3-D images with a spatially constant resolution in the range of tens of micrometers and was shown to provide im-
ages revealing fine anatomic structures in an excised biological sample.

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References