

A nanotube-based field emission x-ray source for microcomputed tomography

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Microcomputed tomography (micro-CT) is a noninvasive imaging tool commonly used to probe the internal structures of small animals for biomedical research and for the inspection of microelectronics. Here we report the development of a micro-CT scanner with a carbon nanotube-(CNT-) based microfocus x-ray source. The performance of the CNT x-ray source and the imaging capability of the micro-CT scanner were characterized. © 2005 American Institute of Physics. [DOI: [10.1063/1.2041589](https://doi.org/10.1063/1.2041589)]

The development of computed tomography (CT) technologies is one of the most important breakthroughs in the field of radiology.^{1,2} CT scanners are now widely used for diagnostic medical imaging and security screening. Microcomputed tomography (micro-CT), which is similar to CT but provides a better spatial resolution, has recently emerged as a powerful noninvasive imaging tool for biomedical research^{3,4} and for industrial inspection. It has been applied to the high-resolution imaging of bony structures and soft tissues of small animals, materials analysis, and inspection of microelectronics. A typical micro-CT scanner consists of a *microfocus* x-ray source, a sample stage, and an area x-ray detector. A three-dimensional (3D) image of the entire object is reconstructed using the cone-beam reconstruction algorithm⁵ from a set of two-dimensional (2D) images recorded over a wide range of viewing angles by either rotating the object or the x-ray source and the detector simultaneously. One of the most critical components of a micro-CT scanner is the microfocus x-ray source. The *spatial* resolution of the micro-CT scanner is largely determined by the focal spot size of the x-ray source. The *temporal* resolution depends on the switching time of the x-ray source, and is important for minimizing motion-induced blurring of moving objects and for gated imaging.

Current commercial microfocus x-ray sources use hot cathodes to generate electrons for x-ray production. The thermionic technology has several inherent limitations. The slow response time limits the *temporal* resolution of the x-ray

source. The high operating temperature results in a short lifetime and a large device size. It also requires complex electromagnetic optics to focus the spatially random thermal electrons to provide the small focal spot size required for high spatial resolution. The thermionic microfocus x-ray sources used in the commercial micro-CT scanners typically have an x-ray switching time of 10 ms, effective focal spot size of 5–100 μm at the beam current of 40–500 μA , and anode voltage of 30–160 kV.⁶

Field emission x-ray sources can, in principle, offer significantly improved temporal resolution because of the intrinsic instantaneous response time of the field emission process.⁷ The design of the microfocus x-ray source can potentially be simplified because of the small divergence of the field-emitted electrons. Field emission x-ray sources based on metal tips have been investigated and tested for clinical uses in the past, but have suffered from a high extraction field and a short lifetime.^{8,9} Carbon nanotubes (CNTs)¹⁰ have improved emission characteristics compared to the conventional field emitters.^{11–14} It has been shown that they can generate diagnostic quality x-ray radiation^{15–17} with temporal resolution up to nanoseconds,¹⁸ which is significantly better than that of the thermionic x-ray tubes.¹⁹ Here we report the development of a micro-CT scanner using a CNT-based microfocus x-ray source and discuss the preliminary results on the performances of the x-ray source and the imaging capability of the micro-CT scanner.

The micro-CT scanner is illustrated in Fig. 1. It com-

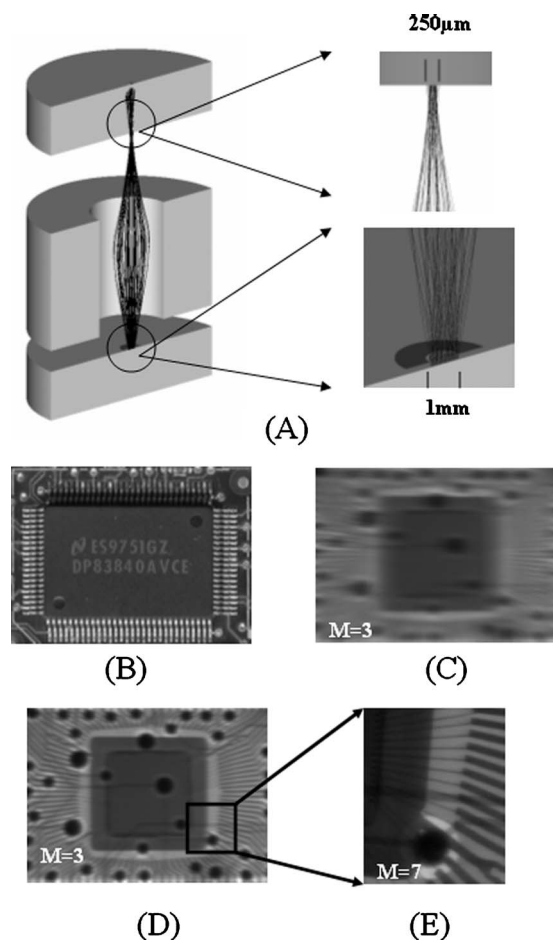


FIG. 3. (a) Computer-simulated electron trajectory using the actual dimension and operating parameters of the x-ray source. The emitting cathode (bottom part) diameter is 1 mm. With a gate at 800 V and a focusing electrode (middle section) at 600 V, the electron beam profile at the anode is 250 μm in size. An *actual* focal spot size of $<150 \mu\text{m}$ was obtained at $V_a=40 \text{ kV}$, $V_g=800 \text{ V}$, and $V_f=700 \text{ V}$. (b) Optical image of a computer chip. (c)–(e) Corresponding x-ray image of the same computer chip under different focusing voltages: $V_f=0 \text{ V}$, 500 V, and 700 V, respectively. M in the graph represents the magnification factor, defined as the ratio of the image and object dimension, of the x-ray image.

beam center needs to be determined to reduce artifacts in the reconstructed images. This was accomplished by analyzing the distortion of the x-ray projection images of a phantom that contained several equally spaced identical thin metal disks stacked inside a plastic cylinder.⁴ Tomographic images of a normal 8-week-old mouse carcass (25 g) were obtained using this micro-CT scanner. To increase the contrast for the soft tissues, an iodinated contrast agent was injected into the abdomen (0.2 ml) of the mouse. A set of 600 projection images was taken over 360° at 1 s exposure per image. The x-ray source was operated at 40 kVp, 100 μA , 150 \times 50 μm effective focal spot size, and cone beam geometry. As shown in Fig. 4(a), the projection images demonstrated good bone delineation and soft tissue contrast. A modified Feldkamp algorithm⁵ was used for 3D reconstruction. It took about 3 h to reconstruct a $325 \times 325 \times 500$ volume at 100 μm pixel size with a single processor on a PC running LINUX. The computing time was subsequently reduced by utilizing multiple processors with a MPI parallelized version of the code. Examples of the tomographic transverse and

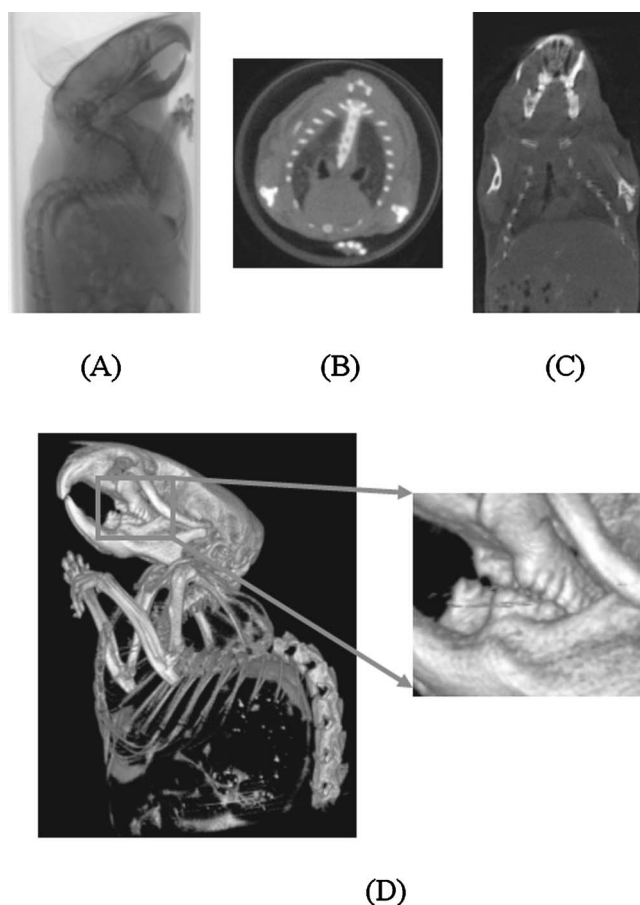


FIG. 4. CT image of a normal mouse carcass (25 g) obtained using the current imaging system. The imaging conditions are 40 kVp, 100 μA tube current, 1 s exposure time, and $150 \times 50 \mu\text{m}$ effective focal spot size without energy filter. (a) A sample projection image demonstrating good bone delineation and soft tissue contrast. (b), (c) Transverse and coronal micro-CT images of the same mouse clearly show the lungs' location and the separation between low-density fat and surrounding soft tissues. (d) 3D volume rendering of the whole-body skeletal dataset. The bony structures are well visualized, and even small subtle structures, such as the bony sutures in the skull and individual teeth, are readily apparent.

coronal cuts from the reconstructed volume are shown in Figs. 4(b) and 4(c), respectively. 3D skeletal renderings of the carcass are shown in Fig. 4(d). Due to a much higher x-ray attenuation coefficient in calcified tissue than in soft tissue, the bony structures are well visualized, and even small subtle structures, such as the bony sutures in the skull and individual teeth, are readily apparent. As shown in the figure inset, the image provides sufficient resolution to visualize the individual tooth of the mouse.

In summary, we demonstrate the generation of microfocus x-ray radiation using a CNT-based field emission x-ray source. An effective focal spot size of 50 μm was obtained using one active electrostatic focusing electrode, which can be further reduced with improved design. A micro-CT scanner with the field emission x-ray source was designed and its utility for small animal imaging was demonstrated. The temporal resolution of the system affords potentials for gated and dynamic CT imaging.

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