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## Fabrication and characterization of a focused ion beam milled lanthanum hexaboride based cold field electron emitter source

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We report on a method of fabricating lanthanum hexaboride (LaB<sub>6</sub>) cold field emission tips with sub-100-nm apices by using a combination of electrochemical etching and focused ion beam milling. The primary advantage of combining the two methods is rapid fabrication while maintaining reproducibility. The LaB<sub>6</sub> tips have low work functions and high mechanical stabilities and are chemically inert to residual gases. Field emission characterization was performed on three tips, with apex sizes of 15, 85, and 80 nm yielding 10 nA cold field emission currents at 0.76, 3.9, and 3.6 kV extraction potentials, respectively. All three tips showed excellent emission current stability for periods exceeding 30 min in a  $5 \times 10^{-9}$  mbar vacuum. *Published by AIP Publishing.*

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A field emission electron source is an essential component in a modern transmission electron microscope (TEM), scanning TEM (STEM), and scanning electron microscope (SEM), as it provides an electron spot size down to the sub-Angstrom range in a STEM or SEM and excellent phase contrast in a high-resolution TEM.<sup>1,2</sup> Whereas a Schottky thermally assisted field emission (STFE) source has a high emission current and good long-term stability, a cold field emission source (CFES) has a higher source brightness and a lower energy spread.<sup>3</sup> CFES technology based on crystalline tungsten (W) has long been implemented<sup>4</sup> but exhibits poorer emission current stability and higher vacuum sensitivity than STFE devices. These issues are related to the work function and robustness of the tips, as well as chemical inertness to residual gases.<sup>5</sup> The advent of new material fabrication techniques, such as chemical vapor deposition, has allowed the fabrication of carbon-based devices, including carbon nanotubes,<sup>6</sup> carbon cones,<sup>2</sup> and doped silicon carbide<sup>7</sup> emitters, which have excellent brightness, peak emission current, and stability. However, the disadvantages of these sources include difficulties associated with their axial alignment in W wire mounts,<sup>8</sup> as well as Coulomb effects following emission from such small tips.<sup>9</sup> An alternative strategy is to employ compound materials such as lanthanum hexaboride (LaB<sub>6</sub>),<sup>10–12</sup> whose low work function of 2.7 eV, low vapor pressure,<sup>13</sup> chemical inertness,<sup>14</sup> and high mechanical robustness potentially make it a superior CFES compared to W.<sup>15</sup> Although functional LaB<sub>6</sub> nano-tip

emitters have been prepared using various methods,<sup>12,14,16</sup> these generally involve many steps, from fabrication to mounting of the tips, making them time-consuming in production and difficult to use as practical CFES devices. Here, we present a simple, fast, versatile, and robust method for the fabrication of LaB<sub>6</sub> nano-emitter devices. We employ a two-step process, involving electrochemical etching and focused ion beam (FIB) milling of single crystal LaB<sub>6</sub> rods that are permanently fitted to thermally robust, electrically conducting fixtures, resulting in ready-mounted monolithic LaB<sub>6</sub> structures. This method in comparison to others<sup>2,3,6,24</sup> allows easy and superior control over axial alignment as well as over the tip apex size and shape, thereby ensuring a high degree of performance and reliability. Fabricated LaB<sub>6</sub> tips have shown excellent stability in comparison to conventional W field emitters<sup>24</sup> and earlier LaB<sub>6</sub> monolithic field emitters.<sup>12</sup> Nondecaying emission current is also observed for 24 h operation (see [supplementary material](#)).

Single crystalline LaB<sub>6</sub> rods with diameters of 0.60 mm, lengths of 5 mm, and  $\langle 100 \rangle$  orientation, so chosen because of their low work function and vapor pressure,<sup>10</sup> were purchased from APTech. Each LaB<sub>6</sub> rod was fitted tightly into a short tube made from tantalum (Ta), protruding by about 2 mm from one end. The Ta tube was wrapped with 0.25 mm diameter W wire, and W wire leads were laser-welded to the pins of a ceramic mount, as shown schematically in Fig. 1(a). This fixture formed the basis of all subsequent processing and characterization steps. Electrochemical etching of the LaB<sub>6</sub> rod was then performed using a method similar to that described by Wang *et al.*,<sup>16</sup> yielding a tip with an apex of sharpness 1–10  $\mu\text{m}$ , as shown in Fig. 1(b).

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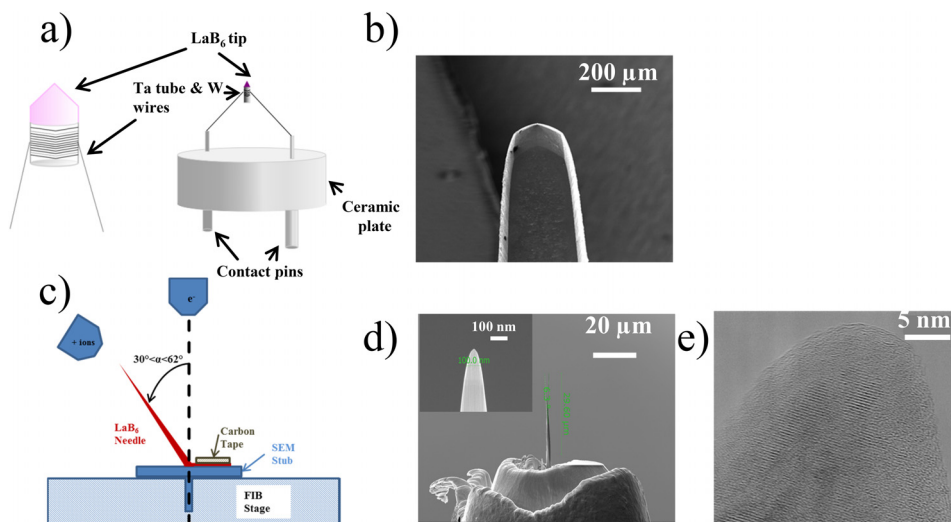


FIG. 1. (a) Schematic diagram of the mounted LaB<sub>6</sub> tip protruding outside a Ta tube surrounded by W wires that are laser welded to contact pins on a ceramic mount; (b) SEM image of an electrochemically etched LaB<sub>6</sub> tip of size 1–10 μm; (c) schematic diagram of FIB milling of a LaB<sub>6</sub> tip; (d) SEM image of a LaB<sub>6</sub> tip after FIB milling. Inset: magnified view of the tip apex; (e) HRTEM image of the tip apex, showing 5 nm irregularities on the surface.

Further apex size reduction and shaping were carried out using FIB milling, as shown schematically in Fig. 1(c). The rod was aligned on-axis with the ion beam and milled centre-symmetrically from the top. This procedure is similar to the annular milling method used for the preparation of atom probe tomography specimens.<sup>17</sup> Rough milling (with a beam current of 21–60 nA at 30 kV) was used to sharpen the tip by iteratively removing ring-shaped areas, thereby decreasing the inner diameter. This step was followed by finer milling, involving gradual reduction of the ion current (typically from 2.5 nA to 80 pA), as well as the outer and inner diameters of the milling pattern. Finally, a polishing step using an ion beam of 5 kV and a current 40 pA for approximately 20 s with an inner diameter mask setting of zero was applied to reduce the level of gallium implantation.<sup>17</sup>

Figure 1 shows the LaB<sub>6</sub> tip morphology subject to the chosen processing steps. An SEM image of the tip after electrochemical etching is shown in Fig. 1(b), revealing an apex size of approximately 10 μm. Figure 1(d) shows one of the etched tips after FIB milling. This process resulted in conical structures with lengths of approximately 20–30 μm and base diameters of 2–5 μm. The apex diameters were 85, 15, and 80 nm for tips 1, 2, and 3, respectively. The combined LaB<sub>6</sub> fabrication procedure (electrochemical etching and FIB milling) yielded field emission structures with large bases (tens of μm) and good mechanical stabilities and electrical conductance. The largely non-selective rate of FIB milling preserves the material stoichiometry and eliminates potential surface contamination and chemical changes caused by electrochemical etching. A high degree of reproducibility is also ensured by this process.<sup>18</sup> A possible concern is destruction of the LaB<sub>6</sub> crystallinity due to ion bombardment during FIB milling. HRTEM images of the tip apex recorded in a TEM fitted with a customized sample transfer arm confirmed the presence of damage to the tip in the form of irregularities. A more detailed study of the tip surface is deferred to a future report.

The field emission behaviour of the emitters was then studied. The entire assembly [Fig. 1(a)] was transferred to a field emission characterization chamber with a base pressure of approximately  $5 \times 10^{-9}$  mbar, as shown schematically in Fig. 2. A precision high voltage power supply

(iseq SHQ 226L) was used to set up the field emission voltage, and the emission current was detected using an ammeter (Keithley 6487) connected to a grounded retractable stainless steel plate positioned at a distance of 30 mm from the tip. A single microchannel plate (MCP) and phosphor assembly that was lens-coupled to a CCD camera was placed 20 mm behind the steel plate to image the electron emission pattern. In order to optimize the shape of the tip apex and to achieve stable emission, thermal conditioning was applied.<sup>19</sup> At temperatures above 800 °C, surface tension forces drive surface migration, rounding, and smoothing of the LaB<sub>6</sub> emitter tip.<sup>12</sup> Finally, in a process known as dulling, the apex radius increases through transport of material from the apex towards the shank.<sup>19</sup> The temperature was typically fixed in the 900 °C–1500 °C range for approximately 2 min, well below the point at which changes in the surface composition may occur.<sup>12</sup> Field emission is described by the well-known Fowler-Nordheim (F-N) equation,<sup>20</sup> which can be written in the simplified form<sup>21,22</sup>

$$I = AV^2 \exp\left(-\frac{B}{V}\right), \quad (1)$$

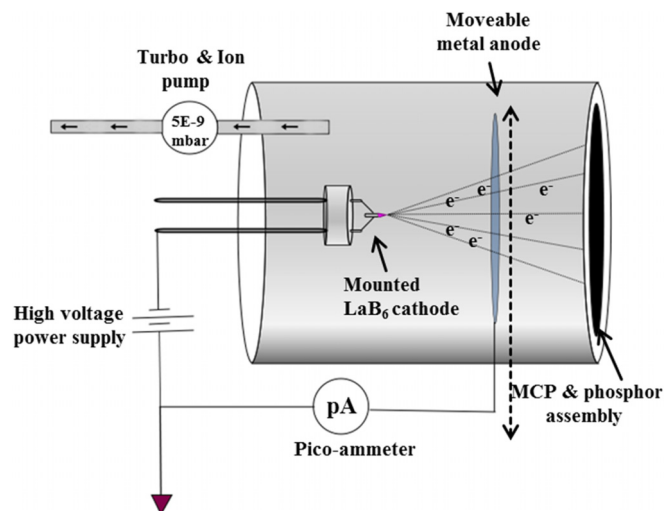


FIG. 2. Schematic diagram of the present field emission characterization setup.

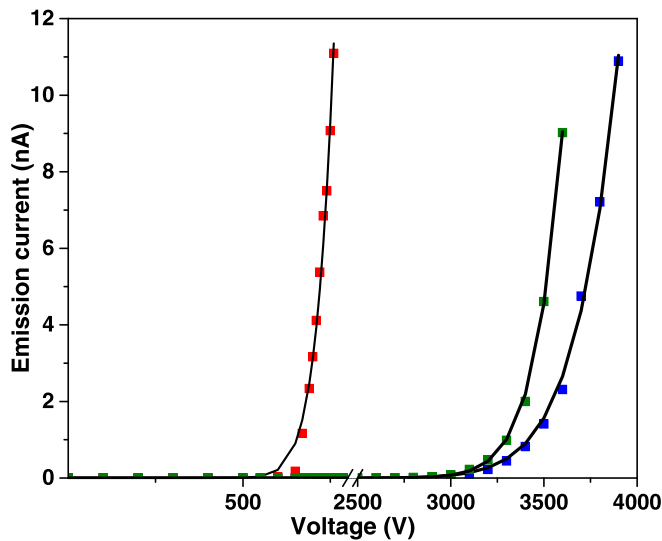


FIG. 3. Current vs voltage ( $I$ - $V$ ) curves recorded from the LaB<sub>6</sub> tips, showing the onset of field emission for tip 1 (blue), 2 (red), and 3 (green) at 3.0 kV, 550 V, and 2.8 kV, respectively, confirming the sharpness of tip 2 in comparison with tip 1 and tip 3.

$$A = 1.5 \times 10^{-6} \frac{S\beta^2}{\phi} \exp\left(10.4/\sqrt{\phi}\right),$$

$$B = 6.44 \times 10^9 \phi^{3/2}/\beta,$$

where  $V$  is the applied voltage,  $I$  the emission current,  $S$  the emitting area, and  $\beta$  the field enhancement factor. All of the quantities are given in S.I. units, while the work function  $\phi$  is in eV. From Eq. (1), it is clear that a high emission current can be achieved by choosing a material that has a low work function, as well as by fabricating a sharper tip, as the field enhancement factor is typically inversely proportional to the tip apex radius, and hence,  $\beta \propto S^{-1/2}$ .  $I$ - $V$  curves recorded after conditioning are shown in Fig. 3 for all three tips. Least-square fits of Eq. (1) are in excellent agreement with the data for  $B_{\text{FN}}^{(1)} \approx 5.9(4) \times 10^4$  V,  $\beta^{(1)} \approx 4.8 \times 10^5$  m<sup>-1</sup>, for tip 1,  $B_{\text{FN}}^{(2)} \approx 9.9(5) \times 10^3$  V,  $\beta^{(2)} \approx 2.9 \times 10^6$  m<sup>-1</sup> for tip 2, and  $B_{\text{FN}}^{(3)} \approx 5.6(3) \times 10^4$  V,  $\beta^{(3)} \approx 5.1 \times 10^5$  m<sup>-1</sup> for tip 3. The apex radii can be estimated from the empirical relation  $\beta \approx 1/5r$  to be  $r^{(1)} \approx 410$  nm,  $r^{(2)} \approx 70$  nm, and  $r^{(3)} \approx 395$  nm. By using the fit result, the electric field  $E$  at the apex of each tip can be estimated to be 1.9 GV/m at 3.9 kV for tip 1, 2.2 GV/m at 760 V for tip 2, and 1.8 GV/m at 3.6 kV for tip 3. Here, we assumed that the bulk work function of LaB<sub>6</sub> is 2.7 eV. The effective barrier height subject to this electric field, also known as the Schottky reduced tunneling barrier,<sup>23</sup> is then estimated to be 1.0 eV for tip 1, 0.9 eV for tip 2, and

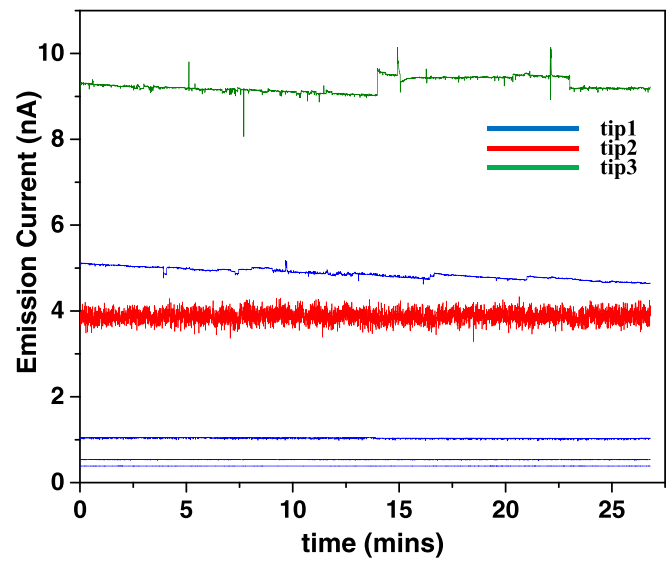


FIG. 5. Temporal stability of the emission current for LaB<sub>6</sub> tips. Note the slight jumps for tip 1 and tip 3.

1.1 eV for tip 3. The imaging results for the emission pattern are shown in Fig. 4. Gaussian beam spots are suggestive of the Gaussian virtual source. We attribute the distorted appearance of the emission pattern from tip 1 to the imperfect alignment of the tip with respect to the optical axis.

Long-term emission stability is a key factor in the practical utilization of electron emitter devices. In this respect, LaB<sub>6</sub> nanowires have been shown to significantly outperform metallic tips.<sup>24</sup> Figure 5 shows temporal stability profiles of our LaB<sub>6</sub> emitters. We observed an almost negligible decay of emission current compared to metallic emitters (typically  $\sim 35\%$  decay over 1 min) over  $\sim 30$  min and relative RMS noise values of 0.2%, 0.6%, 1.1%, and 2.5% (drift-subtracted) for tip 1 at 300 pA, 500 pA, 1 nA, and 5 nA, respectively, 3.5% for tip 2 at 4 nA, and 5% for tip 3 at 9 nA. All measurements were performed at a frequency of 3 Hz. Different traces of emission current for tip 2 in comparison to tip 1 and tip 3 are due to the smaller emission area of tip 2 in comparison to tip 1 and tip 3, which results in high fluctuations.<sup>26</sup> When performing longer stability measurements at currents exceeding approximately 4 nA, jumps in current with intermittent plateaus were occasionally observed. These plateaus are attributed to bombardment of the tip with ions desorbed from the anode.<sup>25</sup>

We have presented a method of fabricating LaB<sub>6</sub> field emission tips, based on a combination of electrochemical etching, FIB milling, and thermal conditioning. In addition to its simplicity and rapidity of execution, the technique

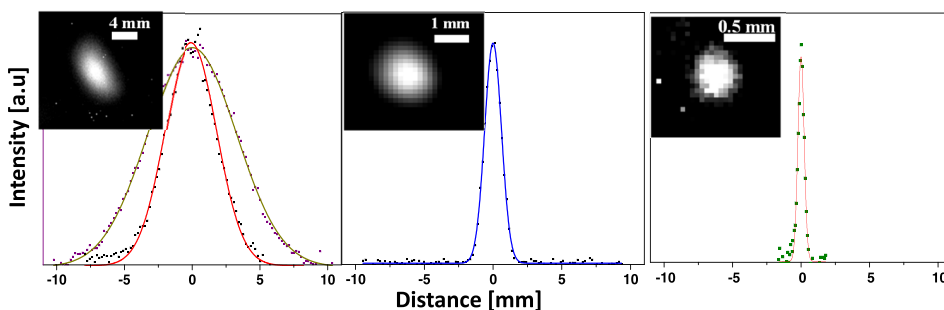


FIG. 4. Field electron micrograph (FEM) images of tip 1 (left), tip 2 (middle), and tip 3 (right) corresponding to emission currents of approximately 10 nA at 3.9 kV, 760 V, and 3.6 kV, respectively. Gaussian fits (solid line) to the profiles give FWHM values of approximately 7.7 and 4.5 mm along the long and short axes, respectively, for tip 1, 1.4 mm for tip 2, and 0.4 mm for tip 3.

allows for high reproducibility and customizability in terms of emitter geometry. We characterized the field emission behavior of three tips with different parameters, yielding 10 nA of emission current at 3.9 kV, 760 V, and 3.6 kV. Emission current stability measurements over 30 min confirmed excellent stability in terms of current drift and noise at a modest vacuum level of approximately  $5 \times 10^{-9}$  mbar, suggesting that the chemical inertness of LaB<sub>6</sub> is a key advantage over metals for field emission. This property, in combination with the low energy spread expected for a low work function material, makes our emitter a promising candidate for use in electron beam instruments. Further investigations, comprising field ion microscopy, virtual source size measurement, and pulsed emission, will be presented in upcoming articles.

See [supplementary material](#) for 24 h continuous emitter operation (Fig. S1) and the noise spectrum (Fig. S2).

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