

1 Selective-Area Deposition of Indium and Its Plasmonic Properties

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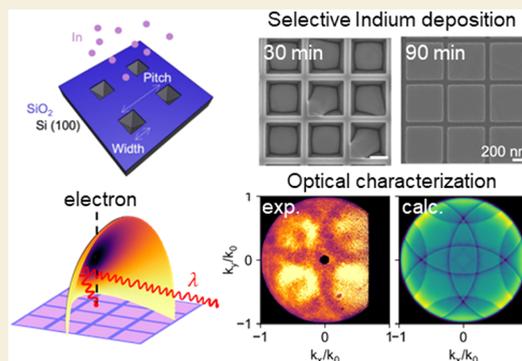
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Supporting Information

5 **ABSTRACT:** We present an effective process sequence for the deposition of
6 indium nanostructures using molecular beam epitaxy (MBE) on a silicon
7 substrate. Using a template structure composed of inverted pyramids and V-
8 grooves, we deposit indium nanostructures with various dimensions. Spatially
9 resolved cathodoluminescence spectroscopy (CL) using an electron-beam
10 energy of 30 keV electrons shows a localized surface plasmon (LSPs)
11 resonance in spherical particles with a peak wavelength at 300 nm and a full
12 width at half-maximum of 70 nm for the smallest particles (diameter of 85
13 nm), showing high optical quality of the grown indium. V-groove template
14 structures create indium nanowires for which CL spectroscopy reveals efficient
15 propagation of surface plasmon polaritons (SPPs), and angle-resolved CL on
16 the periodic inverted pyramids reveals optical lattice resonances arising from
17 the array's periodicity. The high optical quality of these nanostructures enables
18 further applications of plasmonic nanostructures in the ultraviolet (UV) spectral range.

19 **KEYWORDS:** indium nanoparticles, selective area, molecular beam epitaxy, plasmonics, cathodoluminescence



20 ■ INTRODUCTION

21 Noble metal nanostructures show strong interaction with light
22 due to collective oscillations of the free electrons in their
23 conduction band. When excited, they can support surface
24 plasmon polaritons (SPPs) and localized surface plasmons
25 (LSPs).^{1–3} Metal nanogrids, for instance, can function as
26 transparent conductors for photovoltaics.⁴ Furthermore, SPPs
27 can serve as information carriers in integrated optical elements
28 with length scales well below the optical diffraction limit, while
29 LSPs exhibit strong absorption and scattering resonances in the
30 ultraviolet (UV)–visible–near-infrared spectral range. These
31 resonances enable a variety of applications in, for example,
32 sensing, photo, or thermally enhanced catalysis driven by
33 strong optical near-fields.⁵ The LSP resonance wavelengths are
34 tunable with particle size, morphology, metal composition, and
35 the surrounding dielectric environment. This tunability offers
36 significant design flexibility, offering a versatile platform for
37 exploiting plasmonic properties and enriching applications
38 involving metal nanostructures.⁶

39 Assembling multiple metal nanoparticles into ordered arrays
40 further enables advanced functionalities. When metal particles
41 are positioned at controlled distances, the coherent interaction
42 between excited plasmons can create strong optical near-fields,
43 and as a result, controlled coupling to the far-field.^{1,7–9} Various
44 methods enable the formation of plasmonic lattices. One
45 favorable bottom-up technique is a sequence of substrate
46 patterning, metal deposition at exposed areas by evaporation or

sputtering, and lifting off the residual resist.¹⁰ This relatively
47 simple approach enables the formation of regular arrays of
48 metal nanostructures with defined distances, sizes, thicknesses,
49 and morphologies. Colloidal self-assembly is another method
50 to form nanoparticle arrays.^{11,12}

51
52 Noble metal nanostructures composed of silver, gold, and
53 copper are among the most extensively studied and exhibit
54 optical resonances in the visible and near-infrared spectral
55 range.^{13–15} Aluminum, a non-noble metal, displays a resonance
56 in the visible to deep ultraviolet (UV) spectral range, providing
57 compatibility with CMOS process technology.^{7,16–18} Bismuth
58 nanodisk arrays deposited by pulsed laser deposition have also
59 emerged as a UV plasmonic material.⁶ An unconventional
60 plasmonic metal that has not been explored widely is indium,¹⁹
61 which has a plasmon resonance in the near-UV visible spectral
62 range.¹⁶ Previously, indium nanoparticle arrays were made by
63 evaporation on a silicon substrate, followed by dewetting,
64 resulting in small grains that do not coalesce.^{20,21} The optical
65 properties of these geometries have been characterized by
66 transmittance and reflectance spectroscopy.^{20,22,23} Colloidal

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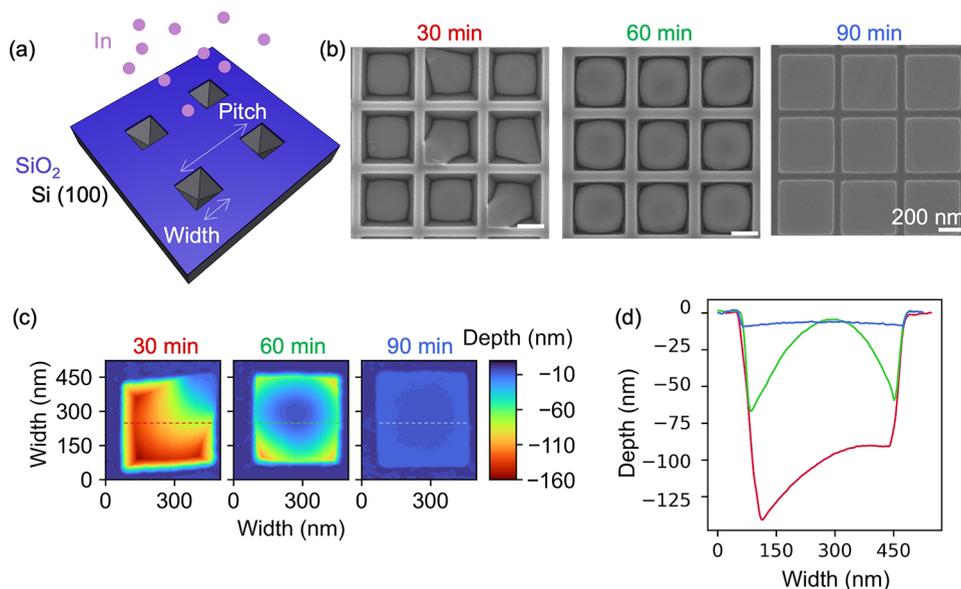


Figure 1. (a) Schematic showing the etched substrate covered with oxide and indium deposition within inverted pyramids; (b) SEM images of temporal evolution of indium deposition for 30, 60, and 90 min (scale bars are 200 nm); (c) corresponding AFM images of single structures shown in (b); and (d) line scans taken at the dashed lines shown in (c). The structures have a nominal width of 140 nm and a pitch of 500 nm.

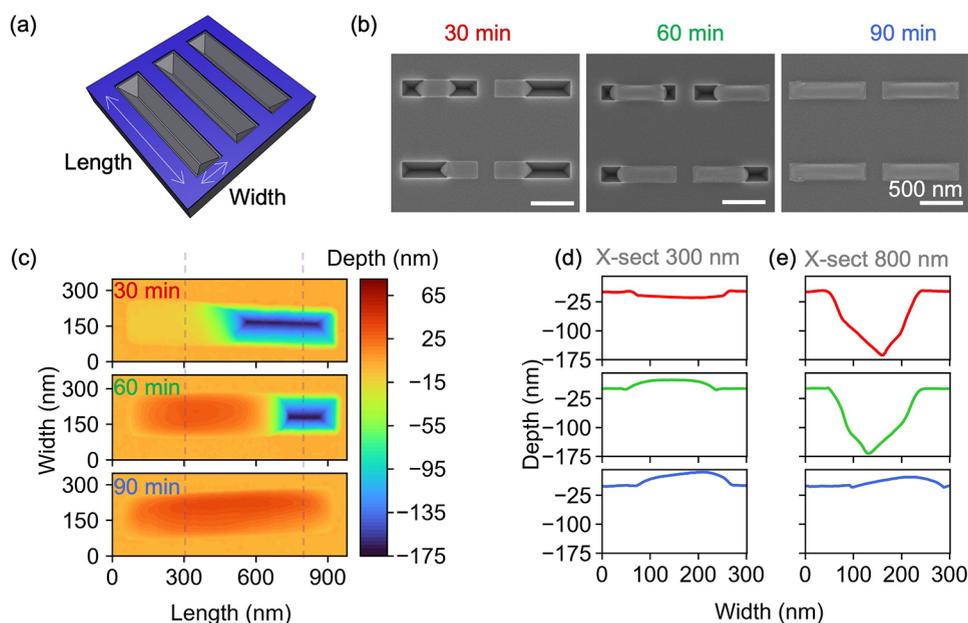


Figure 2. (a) Schematic of the substrate showing the width and length of a groove; (b) SEM images of temporal evolution of indium deposition for 30, 60, and 90 min (the scale bars are 500 nm); (c) AFM contour plots of one of the representative grooves shown in (b); and (d, e) line scans taken from the dashed lines shown in (c) at lengths of 300 and 800 nm, respectively.

indium nanostructures with a sub-10 nm diameter have also been synthesized. These particles showed unusual redshift in absorption with decreasing particle size, attributed to ligand and confinement effects.²⁴ Furthermore, due to oxidation, a 3 nm oxide shell forms around the particles,²⁵ for decreasing particle sizes (<30 nm), this has been shown to quench the LSP resonance.²⁶ Finally, indium arrays have been made by combining substrate conformal imprint lithography (SCIL) with electrodeposition, offering a means of producing scalable plasmonic lattices with a strong plasmonic LSP resonance due to enhanced in-plane light scattering.²⁷

In this paper, we present a method to selectively deposit indium with a high optical quality. Similar to selective area epitaxy used for the deposition of III–V nanostructures,²⁸ the process includes the patterning of silicon (Si) substrates with predefined widths, pitches, and lengths. Both inverted pyramids and V-groove geometries were made by chemical etching in silicon. Subsequently, indium is deposited by using molecular beam epitaxy (MBE), ensuring high purity. Atomic-force microscopy (AFM) and scanning electron microscopy (SEM) measurements are performed to characterize the morphology of the indium droplets after deposition. Under our deposition conditions, indium fully wets the V-grooves.

90 This wetting behavior results from confinement provided by
91 the patterned substrate combined with the optimization of
92 surface energy, resulting in nanostructure arrays with smooth
93 and flat surfaces. We characterize the optical properties of the
94 indium arrays with cathodoluminescence (CL) spectroscopy,
95 which provides a high spatial imaging resolution. Individual
96 indium structures show a clear LSP resonance that shifts with
97 an increasing particle size. Indium nanowires, grown inside V-
98 grooves, we find clear CL interference characteristics for the
99 generation of SPPs, and the angular distribution of the CL
100 emission clearly shows the interference due to the square
101 periodicity of the structure. Overall, our findings suggest that
102 indium deposition by MBE enables the fabrication of well-
103 defined, uniform, and reproducible metal nanostructure arrays
104 that can find use in optical applications in the ultraviolet
105 spectral range.

106 ■ RESULTS AND DISCUSSION

107 Indium Deposition

108 A Si (100) substrate is covered with an 18 nm thick SiO₂ mask
109 featuring diverse patterns of varying widths, pitches, and
110 lengths. Our focus lies on two different shapes: inverted
111 pyramids and V-grooves aligned along <110> directions
112 (illustrated in Figures 1a and 2a, respectively). Along these
113 directions, the trenches have a 111 family of side facets
114 resulting from directional KOH etching. These facets are not
115 fully formed for inverted pyramids with larger widths for which
116 a flat (100) bottom is present. The presence or absence of flat
117 (100) surfaces at the bottom within the studied width range
118 did not yield significant differences in the deposition of the
119 observed structures.

120 We varied the deposition temperature between 500, 550,
121 and 600 °C to find an optimized selective deposition
122 temperature (see SI, Figure S1). At 500 °C, the growth was
123 not selective, and indium droplets covered the substrate. Above
124 550 °C, we started to observe selectivity, and for the rest of the
125 experiments, we conducted our depositions at 600 °C. We did
126 not investigate the upper temperature limit of indium
127 deposition, as this would require a systematic study quantifying
128 the deposited volume within the structures. Furthermore,
129 having an oxide-free Si surface and an ultrahigh vacuum
130 environment of MBE enhances the wetting of indium metal.

131 Figure 1 shows the selective indium metal deposition
132 process within inverted pyramids. In Figure 1b, SEM images
133 depict the temporal evolution of the incorporation of indium
134 within inverted pyramids subjected to deposition times of 30,
135 60, and 90 min. We clearly see that the deposition is selective,
136 with no crystallites on the mask region. Longer deposition
137 times result in a higher liquid indium volume within the holes.
138 To study the temporal depth evolution, Figure 1c presents the
139 3D AFM contour plots of the individual structures for
140 deposition times of 30, 60, and 90 min. Indium predominantly
141 wets the bottom; however, in some grooves, metal pinning by
142 the mask occurs at the edge of the SiO₂. By 90 min, the liquid
143 reaches almost the surface level for this width. Horizontal line
144 scans along the width direction, as indicated by dashed lines in
145 Figure 1c, are shown in Figure 1d. Until the pyramid volume is
146 filled, the contact angle of the liquid decreases, resulting in a
147 flat surface morphology that thoroughly wets the surface.
148 Within inverted pyramids, initial droplet formation occurs
149 either at the bottom of the structures or at the interface
150 between oxide and Si surfaces. There is no notable difference

in filling amount among the various pitches analyzed for
inverted pyramids; however, these structures exhibit similar
heights, indicating no pitch dependency (see Supporting
Information, Figure S2a).

Next, we studied the indium deposition in the V-grooves.
Figure 2a shows the schematic of the deposition process, and
Figure 2b exemplifies the temporal evolution of the
incorporation of indium within the V-grooves. Height
information is depicted in the 3D contour plots captured by
AFM in Figure 2c, where the top-to-bottom progression
corresponds to 30, 60, and 90 min deposition intervals. Figure
2d,e shows the cross sections taken from the dashed lines at
300 and 800 nm in Figure 2c. As the deposition time increases,
metal fills the entire groove, nearly filling it up. Within V-
grooves, the initial droplet formation occurs at one of the two
side corners, with it occasionally commencing in the middle of
the groove. As the V-groove length increases, the probability of
indium droplets forming at multiple locations increases.
Furthermore, at a given time, increasing lengths result in
height variations for a constant width (Figure S2b). Under
optimized deposition conditions, once droplets form, instead
of swelling out, they form a horizontal metal nanowire with a
smooth top surface. This indicates that as the deposition time
increases, incoming indium adatoms preferably incorporate an
inclined growth front rather than at the top facet. After
reaching a certain volume with an increasing deposition time,
the droplet contact angle increases, extending beyond the
mask. After certain conformal filling, an increase in volume
results in a perturbation in droplet stability, which causes
droplet swelling, collecting more indium adatoms, and acting
as a sink (see Figure S2c). This behavior is easily observed in
smaller-width trenches at 90 min of deposition due to the
relatively low volume of the trench (see Figure S3).

The equilibrium wetting behavior of indium droplets inside
the trenches was modeled using Surface Evolver, assuming a
uniform contact angle of 72° on all surfaces (see Supporting
Information, Section S5). Two initial configurations were
simulated: a “one-sided” droplet at a terminating face and a
“two-sided” droplet positioned away from both faces. As the
droplet volume increased, it advanced along the trench while
maintaining nearly constant mean curvature, with distinct
transitions depending on trench length—either continuous
advancement or division and later coalescence. After the triple
line became fully pinned at the trench’s top edges, the droplet
evolved by forming a bulge with an increasing apparent contact
angle until depinning occurred. In V-groove trenches, a similar
evolution was observed, but the droplet advanced more rapidly
due to the geometry of the sidewalls.

Optical Properties

To study the optical properties of the grown indium
nanostructures, we used cathodoluminescence (CL) spectroscopy
with a 30 keV electron-beam energy. By scanning the
electron beam over the sample with a pixel size of 10 nm, we
built up a CL excitation map for the indium geometries.

Figure 3a shows the CL spectra for spherical indium
nanoparticles with diameters of 85 (blue), 100 (orange), 110
(green), and 150 nm (red), excited by the electron beam at the
edge of the particle. The inset displays the SEM image of the
smallest particle. Figure 3b shows the scattering efficiency for
the indium particles calculated using Mie theory and the
dielectric constants from the reference Palik.²⁹ The scattering
spectrum exhibits long-wavelength resonant electric dipole

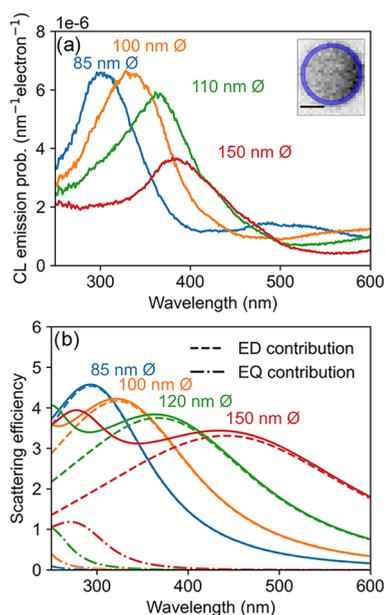


Figure 3. (a) Measured CL spectra for spherical indium nanoparticles on a Si substrate covered with an 18 nm SiO₂ layer, with diameters of 85 nm (blue), 100 nm (orange), 110 nm (green), and 150 nm (red). The inset shows an SEM image of the 85 nm-diameter particle. The scale bar is 50 nm. The nanospheres are excited by 30 keV electrons, and the CL is integrated over the pixels at the edge of the particle, indicated by the blue circle in the inset. (b) The scattering cross-section was calculated with Mie theory and normalized to the geometrical cross-section for spherical indium nanoparticles in vacuum with the same diameter as in (a). The contributions of electric dipole (ED) and quadrupole (EQ) resonant modes are indicated by dashed lines.

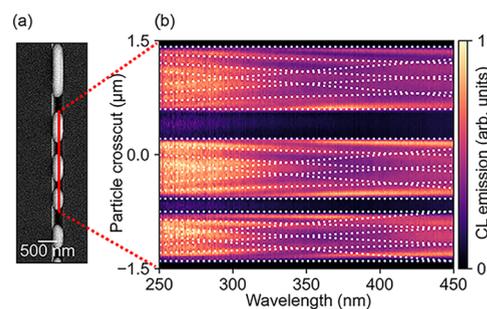


Figure 4. (a) SEM map showing the indium structure within V-grooves and (b) corresponding CL spectra along the line indicated in (a). The dashed white lines show the expected maxima of standing SPP waves with a period of half the wavelength of the SPP mode.

the electron beam from the top to the bottom with a step size 242 of 10 nm, and the CL spectrum was collected at every pixel. 243 This results in the CL map of Figure 4b that shows a spectrum 244 for every pixel on the *y*-axis corresponding to the red line in 245 Figure 4a. In the data, we clearly see minima and maxima due to 246 the destructive and constructive interference of SPP modes 247 launched by the electron beam and reflected on the edge of the 248 particle. The period of the standing waves is expected to match 249 half the SPP wavelength given by³¹ 250

$$\lambda_{\text{SPP}} = \lambda_0 / n_{\text{eff}} \quad (1) \quad 251$$

where the effective refractive index (n_{eff}) is given by³¹ 252

$$n_{\text{eff}} = \sqrt{\epsilon_d \epsilon_m / (\epsilon_d + \epsilon_m)} \quad (2) \quad 253$$

Overlaying the expected spatial period with the measurements 254 (dashed white line), we find good agreement. At a wavelength 255 of 350 nm, the calculated SPP wavelength is 330 nm. In the 256 experimental data, the average distance between the minima 257 was found to be 160 nm at a wavelength of 350 nm (see 258 Supporting Information, Figure S7 for a linescan at 350 nm 259 wavelength), corresponding to half the SPP wavelength. The 260 propagation length of the SPP is given by³² 261

$$\delta_{\text{SPP}} = \frac{\lambda_0}{2\pi} \left(\frac{\epsilon_m''}{\epsilon_m'} \right) \left(\frac{\epsilon_m' + \epsilon_d}{\epsilon_m' \epsilon_d} \right)^{3/2} \quad (3) \quad 262$$

At a wavelength of 350 nm, the calculated SPP propagation 263 length at a planar indium-air interface is 1500 nm. In the 264 experimental data, we see two periods in the CL map at 350 265 nm wavelength, and the visibility decreases toward the center 266 of the wire. We estimate the propagation length to be two 267 periods, corresponding to 640 nm, which is twice the SPP 268 wavelength. This indicates that the grown indium has 269 somewhat higher losses than expected based on the literature, 270 in agreement with the larger line widths observed in the 271 experiments of Figure 3a. Coupling of SPPs across the indium 272 nanowire surface can also reduce the SPP propagation length. 273

Finally, we investigate the angular distribution of the CL 274 emission of square indium arrays embedded inside inverted 275 pyramids. Angle-resolved CL measurements are performed 276 using a 70 nm bandwidth filter centered at a wavelength of 400 277 nm. The angular emission is represented in reciprocal-space 278 coordinates, where k_x and k_y denote the in-plane components 279 along the *x*- and *y*-axis, normalized to the free-space 280 wavevector ($k_0 = 2\pi/\lambda$). Figure 5a–c shows the result for an 281 indium array with a pitch of 2000 nm. Figure 5a,b shows an 282 SEM image of the structure and the corresponding angle- 283

213 (ED) and shorter-wavelength quadrupole (EQ) modes. Note 214 that for very small particles, Ohmic dissipation increases due to 215 surface scattering effects; yet the present work focuses on 216 much larger particles.

217 Comparing the measurements and the calculations in Figure 218 3, we see similar trends. The peak wavelength (300 nm) and 219 line width (70 nm FWHM) for the 85 nm-diameter particle 220 agree well between measurement and simulation. The EQ 221 mode seen in the simulations is not observed in the 222 experiments. Our earlier work has shown that the modal 223 excitation efficiency of plasmonic nanoparticles depends 224 strongly on the electron position on the particle, as that 225 determines the phase-matching condition at which efficient CL 226 excitation occurs.³⁰ At the particle edge, the electron beam 227 couples mostly to the ED mode, and hence, the EQ mode is 228 not visible in the CL spectrum. Furthermore, we find that both 229 experiments and simulations show a redshift with increasing 230 particle size, as expected due to weaker restoring forces on the 231 plasmonic charge displacement at larger sizes. Discrepancies 232 between the two may be due to differences in the optical 233 constants of indium for the experimental particles compared 234 with the literature values used. Additional data for excitation at 235 the particle center and corresponding boundary element 236 method calculations to calculate CL emission are shown in 237 the Supporting Information (Figure S6).

238 Next, we studied the SPP modes of the indium structure 239 within the V-grooves. Figure 4a shows the SEM image of 240 indium deposited inside a V-groove. Note that the indium 241 deposition is noncontinuous within the V-groove. We scanned

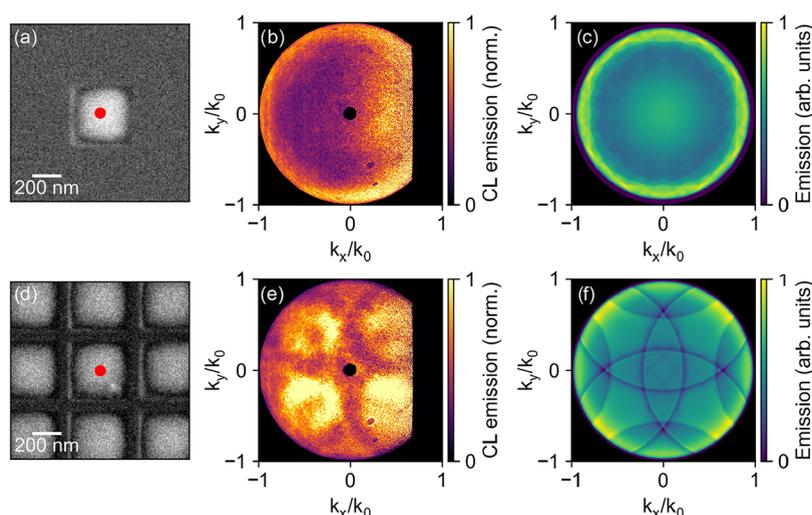


Figure 5. (a) SEM image of the indium-filled array of inverted pyramids with a width of 370 nm and a pitch of 2000 nm, showing the electron excitation position (red dot). (b) Experimentally measured angle-resolved CL emission intensity at a wavelength of 400 nm, and (c) simulated emission profile obtained from RCWA. (d–f) Corresponding results for a similar array with a pitch of 500 nm. The emission arrays are plotted as a function of the normalized in-plane wavevector along the x - (k_x/k_0) and y -direction (k_y/k_0), where $k_0 = 2\pi/\lambda$.

284 resolved CL intensity, measured with electron beam excitation
 285 at the center of the structure (indicated by the red dot in
 286 Figure 5a). Figure 5c shows the calculated emission profile for
 287 this geometry using rigorous coupled-wave analysis (RCWA),
 288 which models the electromagnetic wave diffraction through a
 289 periodic structure.³³ The simulation calculates the absorption
 290 of the array when it is illuminated with a plane wave at varying
 291 angles. Due to reciprocity, the absorption profile is analogous
 292 to the emission pattern. While we do not expect a perfect
 293 correspondence between the RCWA model and the CL
 294 emission—since CL involves localized, point-source excita-
 295 tion—the diffraction features (i.e., angular grating conditions)
 296 are expected to be similar. The primary difference lies in the
 297 amplitude, which reflects the coupling efficiency to the optical
 298 modes. Unlike a plane wave, which contains both s - and p -
 299 polarized components, an electron beam predominantly excites
 300 modes, emitting p -polarized light. Comparing Figure 5b,c, we
 301 observe the same trend of strong emission at the high angles.
 302 This is due to the z -oriented dipole excited at the center of the
 303 structure. Due to the large pitch, the diffraction bands are only
 304 visible at very high emission angles in the simulations. In the
 305 measurements, the angular resolution is insufficient to resolve
 306 them.

307 For comparison, we also study an array with a 500 nm pitch,
 308 shown in Figure 5d–f. Figure 5d shows the SEM image of the
 309 structure, while Figure 5e,f represents the angle-resolved CL
 310 measurement (again with center excitation, marked by the red
 311 dot) and the corresponding RCWA simulation, respectively.
 312 The angle-resolved CL emission distribution clearly shows a
 313 far-field interference pattern characteristic of square symmetry
 314 of the lattice. Comparing the measurements to the simulations,
 315 we clearly see similar dark diffraction bands, where the light
 316 cannot couple to the far-field. Furthermore, the intensity is
 317 homogeneously distributed across the upper hemisphere, in
 318 contrast to the large-pitch structure, which radiated more
 319 strongly at high emission angles. Additional data for arrays with
 320 intermediate pitches (750 and 1000 nm) are provided in the
 321 Supporting Information (Figure S8) and show how the
 322 diffraction bands move depending on the pitch of the lattice.

CONCLUSION

323

In conclusion, we have presented a process sequence for the
 324 selective deposition of indium on patterned Si(100) substrates.
 325 Both inverted pyramid structures and V-grooves form effective
 326 templates for the formation of indium nanostructures. The
 327 indium filling and wetting behavior depend on the pattern
 328 length-to-width ratio. Cathodoluminescence spectroscopy on
 329 the indium nanoparticles shows a strong electric dipole
 330 plasmon resonance, indicating a good optical quality of the
 331 grown indium material. Indium nanowire structures show SPPs
 332 with standing waves due to interference, showing sizable SPP
 333 propagation lengths along the wires. Overall, our findings
 334 highlight the efficacy of MBE in creating controlled indium
 335 nanostructures and arrays with high optical quality in the
 336 ultraviolet spectral range.
 337

EXPERIMENTAL METHODS

338

Substrate Fabrication

339

Intrinsic Si (100) wafers served as substrates for indium deposition. A
 340 25 nm thick thermal SiO₂ mask layer was grown on the as-received
 341 wafers to achieve selective deposition. The desired pattern was
 342 created employing electron-beam lithography (EBL) on a ZEP
 343 positive resist and cold development with n -amyl acetate. Following
 344 the transfer of the desired pattern to the mask via reactive ion etching
 345 (RIE) with CHF₃/SF₆, the polymeric resist was removed using a high-
 346 power oxygen plasma. Subsequently, the native oxide was removed
 347 with a 1% HF solution for 20 s, and the wafer was immersed for 8 min
 348 in a 40% KOH solution to form the V-grooves within the mask
 349 openings. The resulting wafer was cleaved into 1 cm² chips. Note that
 350 there is a difference between nominal and actual width values due to
 351 the fabrication process, including EBL patterning and subsequent
 352 etching. Unless stated otherwise, the values given will be the nominal
 353 ones.
 354

MBE Deposition

355

After a final immersion in a 1% HF solution for 1 min to remove any
 356 remaining native oxide, the chips were immediately introduced into
 357 the ultrahigh vacuum environment of a Veeco GENxplor MBE
 358 system. After the predegassing step at 400 °C for 2 h, samples are
 359 transferred into the growth chamber. Annealing is performed at a
 360 temperature of 800 °C for 30 min. Then, the temperature is decreased
 361 to the indium deposition temperature. The indium beam equivalent
 362

363 pressure (BEP) is 3.6×10^{-7} Torr, which is enough to achieve good
364 selectivity, and the deposition time is 60 min. After that, the substrate
365 is cooled to 100 °C. Note that the temperatures reported are
366 manipulator-set temperatures.

367 Cathodoluminescence Measurements

368 CL measurements were performed using a FEI Quanta FEB 650 SEM
369 (Thermo Fisher Scientific Inc., MA) and a SPARC Spectral CL
370 collection system (DELMIC BV, The Netherlands).³⁴ The measure-
371 ments used 30 keV electrons at a beam current of 2.3 nA. For the
372 spectral measurements, an exposure time of 1 s was used at two center
373 wavelength settings of the grating (350 and 500 nm). For every
374 spherical particle, a map was made of the entire particle and the signal
375 was integrated over the pixels at the edge of the particle. To correct
376 for the background, a spectrum taken with the electron beam off was
377 subtracted, and the total signal was normalized by using the system
378 response function.

379 For the angle-resolved CL measurements, we directly imaged the
380 light emanating from the sample through the parabolic collection
381 mirror onto an imaging camera and correlated each pixel on the
382 camera to an azimuthal and radial emission angle from the sample.
383 The angular emission for the inverted pyramids was measured by
384 using a 400 nm filter with a bandwidth of 70 nm and a 5 s exposure
385 time, while the electron beam was placed at the center of one of the
386 structures. The measured size of the inverted pyramids was 370 nm,
387 and the pitches were 500, 750, 1000, and 2000 nm.

388 RCWA Simulations

389 The angle-resolved emission intensities were computed using rigorous
390 coupled-wave analysis (RCWA) simulations from an open-source
391 software package S⁴: the Stanford Stratified Structure Solver by Liu
392 and Fan.³³ RCWA simulations compute the absorption in a periodic
393 system for incoming plane waves under different angles, ϕ and θ .
394 According to the reciprocity principle, the computed angle-resolved
395 absorption profiles equal the emission profile at the same wavelength
396 from the absorbing material, in this case, the indium within inverted
397 pyramids. We perform the simulation at a wavelength of 400 nm for
398 an indium array with a width of 370 nm (nominally 240 nm),
399 embedded in a Si substrate with 18 nm of SiO₂ on top. For all three
400 materials, we use the dielectric constants from Palik.²⁹

401 ■ ASSOCIATED CONTENT

402 ■ Supporting Information

403 The Supporting Information is available free of charge at
404 <https://pubs.acs.org/doi/10.1021/acsaoam.5c00373>.

405 Dependence of the indium deposition on temperature
406 (Figure S1); pitch, length, and width (Figures S2 and
407 S3); AFM and SEM images of unfilled structure (Figure
408 S4); modeling the indium deposition (Figure S5);
409 additionally, to further elaborate on the optical
410 characterization, there are additional figures: CL spectra
411 for spherical particles for both edge and center excitation
412 (Figure S6); observation of SPP waves within V-grooves
413 (Figure S7); and the angular distribution of CL emission
414 for a square lattice of indium structures depending on
415 the pitch of the lattice (Figure S8) (PDF)

416 Wetting behavior (Videos S1 and S2) (ZIP)

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426 Author Contributions

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428 Notes

The authors declare the following competing financial interest(s): Albert Polman is a cofounder and co-owner of Delmic BV, a company that produces commercial cathodoluminescence systems like the one used in this work.

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