Facet modulation selective epitaxy—a technique for quantum-well wire doublet fabrication

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The technique of facet modulation selective epitaxy and its application to quantum-well wire doublet fabrication are described. Successful fabrication of wire doublets in the $Al_xGa_{1-x}As$ material system is achieved. The smallest wire fabricated has a crescent cross section less than 140 Å thick and less than 1400 Å wide. Backscattered electron images, transmission electron micrographs, cathodoluminescence spectra, and spectrally resolved cathodoluminescence images of the wire doublets are presented.

Semiconductor structures exhibiting quantum confinement in two or three dimensions have attracted considerable attention for their potential in improving optoelectronic devices^{1,2} and in revealing new phenomena in solidstate physics, such as polarization anisotropy in quantum wires.³ Many approaches to fabricate these quantum-confined structures have been studied. Already grown quantum-well material has been physically patterned by a combination of lithography and etching⁴⁻⁶ or has been selectively disordered by ion implantation⁷ to achieve lateral confinement. In situ formation of nanostructures during epitaxial growths has also been studied. Migration-enhanced epitaxy on tilted substrates has exhibited quantumconfinement effects,8 and stimulated emission from quantum wires⁹ has been demonstrated by performing metalorganic vapor-phase epitaxy (MOVPE) growths on etched substrates. Recently, wire and dot structures have been selectively grown on substrates covered with patterned dielectric masks.¹⁰⁻¹² The formation of crystal facets in single precursor chemistry (trimethyl) selective growth has also been studied as a potential method for quantumwell wire fabrication.¹³

In this letter, we describe a new technique, facet modulation selective epitaxy, and present its application to quantum-well wire doublet fabrication in the $Al_xGa_{1-x}As$ material system. Facet modulation selective epitaxy, as it is defined here, is the application of different precursor chemistries to layers within a single growth to alter sequentially the appearance of facets on a growing structure and to thereby form heterostructures of novel geometry. Two precursor chemistries are used here: a combination of diethylgallium chloride (DEGaCl) and arsine (AsH₃) for GaAs growth and a combination of trimethylaluminum (TMAl), trimethylgallium (TMGa), and arsine (AsH_3) for $Al_xGa_{1-x}As$ growth. The morphology of GaAs selective growth using DEGaCl and AsH₃ is dominated by the appearance of the {111} and {110} families of slow growth planes, with the appearance of a particular plane dependent on the growth temperature and the mask opening orientation.^{10,11,14} The morphology of $Al_xGa_{1-x}As$ selective growth using TMAl, TMGa, and AsH₃ is similar, but the bounding planes include higher-index-number planes (one or more indices greater than one) in addition to the {111} and {110} families of slow growth planes.

One application of facet modulation selective epitaxy is the fabrication of quantum-well wire doublets in one single growth as illustrated in Fig. 1. Using DEGaCl and AsH₃, a GaAs buffer bounded by the low-index-number facets is grown on a substrate covered with a stripe patterned mask, then an $Al_xGa_{1-x}As$ layer is grown using TMAl, TMGa, and AsH₃. Using DEGaCl and AsH₃ again, a GaAs wire doublet is formed as the higher-index-number $Al_xGa_{1-x}As$ facets are partially filled in by the GaAs growth attempting to form low-index-number GaAs facets. The wire doublet is buried in situ by another layer of $Al_rGa_{1-r}As$ growth to eliminate any free surface that would produce nonradiative recombination centers. The width of the starting $Al_xGa_{1-x}As$ facets and the amount of GaAs deposited determine the size of the wires and the spacing between wires in a wire doublet. In the work presented here, successful fabrication of wire doublets by facet modulation selective epitaxy is achieved. Analysis of these

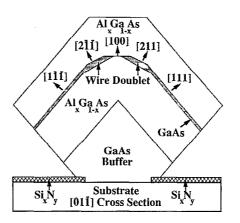


FIG. 1. Schematic illustration of quantum-well wire doublet fabrication by facet modulation selective epitaxy.

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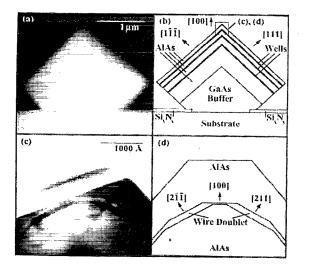


FIG. 2. (a) $[01\overline{1}]$ cross-sectional backscattered electron image of a complete structure grown by facet modulation selective epitaxy. (b) Schematic illustration of the complete structure. (c) $[01\overline{1}]$ cross-sectional transmission electron micrograph of the wire doublet. (d) Schematic illustration of the wire doublet.

wire doublet structures by backscattered electron microscopy, transmission electron microscopy, and cathodoluminescence scanning electron microscopy is presented.

The substrates used in this study contained a $2-\mu m$ Si-doped Al_{0.4}Ga_{0.6}As layer, an undoped 100-nm Al_{0.2}Ga_{0.8}As layer, and a 10-nm GaAs cap layer all deposited by MOVPE on Si-doped (100) GaAs substrates. The dielectric mask was 175 Å of silicon nitride deposited by plasma-enhanced chemical vapor deposition. Arrays of 5mm-long stripe openings, with stripe widths varying from 1 to 8 μm and center-to-center spacing between stripes of 250 μm , were patterned into the silicon nitride mask by photolithography and reactive-ion etching in CF₄ plasma. The stripes were oriented along the [011] direction.

Sample growth was performed in an atmospheric-pressure MOVPE reactor with graphite susceptor temperature set to 730 °C. Precursors for the buffer and the wells were DEGaCl and AsH₃. Precursors for the barriers were TMAl and AsH₃. AlAs barriers were chosen here to test the extreme case. Growth interruptions were placed between layers while AsH₃ flow was maintained. No dopant was intentionally introduced.

Figures 2(a) and 2(b) show a complete structure grown in a stripe opening 1.2 μ m wide. The structure exhibits a faceted profile bounded mainly by the {111} family of planes. Measured along the [100] crystal direction from the substrate to the crystal vertex, the growth consists of 1.3 μ m of GaAs, 0.6 μ m of AlAs, first well, 0.4 μ m of AlAs, second well, 0.2 μ m of AlAs, third well, and finally 0.13 μ m of AlAs. Measured along the [111] crystal direction on the (111) facet, the first, second, and third wells are approximately 10, 20, and 5 nm thick, respectively.

The wire doublet is shown clearly in Figs. 2(c) and 2(d). The transmission electron micrograph in Fig. 2(c) reveals the higher-index-number $\{211\}$ -type AlAs facets near the crystal vertex. The micrograph also shows the

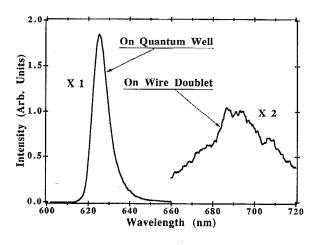


FIG. 3. Cathodoluminescence spectra of the region near the wire doublet. The sample is at 14 K.

third-well growth partially filling in the two $\{211\}$ -type facets, thereby forming a wire doublet as predicted earlier. The two wires in the wire doublet are almost identical in shape and size. Both wires are approximately 140 Å thick at the center and 1400 Å wide, although the effective width of these wires is probably smaller than 1400 Å, because the wires have a tapered cross section.

The luminescence properties of these wire doublet structures were investigated by applying low-temperature cathodoluminescence (CL) scanning electron microscopy.¹⁵ Figure 3 contains the CL spectra of the sample region near the wire doublet. Two distinct peaks appear in the CL spectra: one from the wire doublet at 690 nm and another from the side-wall quantum wells at 625 nm. The red shifting of the wire doublet peak with respect to the side-wall quantum-well peak is as expected from the thickness modulation illustrated in Fig. 2(c). However, it is clear from the CL spectra and the thicknesses measured from Fig. 2(c) that the wire doublet and the side-wall quantum wells are most likely Al_{0.2}Ga_{0.8}As, not GaAs, because a 140-Å GaAs quantum well at 14 K luminesces at about 800 nm, not at 690 nm as in the CL spectra, and a 50-Å well luminesces at about 710 nm, not at 625 nm. Here the wire doublet is approximated as a quantum well because the widths of the wires are much larger than their thicknesses. The presence of aluminum is due to interlayer mixing, presumably caused by incomplete purging of TMAl before the DEGaCl growth had begun. The wire doublet luminescence peak at 690 nm also appears broadened. Possible reasons for this broadening may include the fluctuations in the width of the {211}-type AlAs facets, the surface roughness on the {211}-type AlAs facets, or some aluminum segregation from mixing the two precursor chemistries.

Figures 4(a)-4(c) are spectrally resolved CL images of the wire doublet structure in cross section. Figure 4(a) is imaged at 625 nm where the side-wall quantum wells and the substrate $Al_{0.4}Ga_{0.6}As$ luminesce. Figure 4(b) is imaged at 693 nm, the luminescence peak of the wire doublet. Figure 4(c) is imaged at 820 nm, the luminescence

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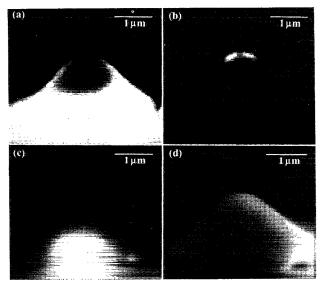


FIG. 4. Spectrally resolved cathodoluminescence images of a wire doublet structure in cross section. The sample is at 12 K, and the wavelengths are set at (a) 625 nm, (b) 693 nm, and (c) 820 nm. (d) Scanning electron micrograph of the same structure in cross section.

wavelength of the buffer and substrate GaAs. The scanning electron micrograph in Fig. 4(d) shows the cross section imaged in Figs. 4(a)-4(c). Figures 5(a)-5(d) are spectrally resolved CL images of structures seeded by different stripe opening widths on the same sample. These images

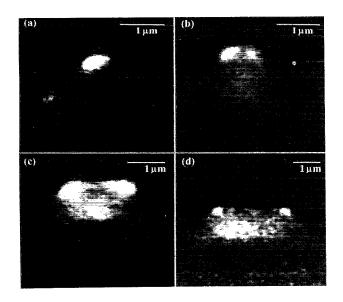


FIG. 5. Spectrally resolved cathodoluminescence images of different wire doublet structures on the same sample. The sample is at 77 K, and the wavelengths are set at (a)-(c) 690 nm and (d) 680 nm.

demonstrate the various wire doublet configurations.

In conclusion, the technique of facet modulation selective epitaxy and its application to quantum-well wire doublet fabrication have been described. Successful fabrication of wire doublets in the $Al_xGa_{1-x}As$ material system has been achieved. The smallest wire fabricated has a crescent cross section less than 140 Å thick and less than 1400 Å wide. Further reduction in wire size will be pursued by reducing the width of the starting $Al_xGa_{1-x}As$ facets and the amount of GaAs deposited. By doping the layers above and below the wire doublet p and n type, respectively, and by embedding the wire doublet in an $Al_xGa_{1-x}As$ optical waveguide, an injection laser may be fabricated. Other applications will also be explored, including the fabrication of quantum dots and the extensions of this technique to other systems besides $Al_xGa_{1-x}As$, such as material $In_xGa_{1-x}As$, by using analogous precursor chemistries.¹¹

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- ¹Y. Arakawa, K. Vahala, and A. Yariv, Appl. Phys. Lett. 45, 950 (1984).
- ²M. Asada, Y. Miyamoto, and Y. Suematsu, IEEE J. Quantum Electron. **QE-22**, 1915 (1986).
- ³P. C. Sercel and K. J. Vahala, Appl. Phys. Lett. 57, 545 (1990).
- ⁴K. Kash, A. Scherer, J. M. Worlock, H. G. Craighead, and M. C. Tamargo, Appl. Phys. Lett. 49, 1043 (1986).
- ⁵H. Temkin, G. J. Dolan, M. B. Panish, and S. N. G. Chu, Appl. Phys. Lett. 50, 413 (1987).
- ⁶B. I. Miller, A. Shahar, U. Koren, and P. J. Corvini, Appl. Phys. Lett. 54, 188 (1989).
- ⁷J. Cibert, P. M. Petroff, G. J. Dolan, S. J. Pearton, A. C. Gössard, and J. H. English, Appl. Phys. Lett. **49**, 1275 (1986).
- ⁸M. Tsuchiya, J. M. Gains, R. H. Yan, R. J. Simes, P. O. Holtz, L. A. Coldren, and P. M. Petroff, Phys. Rev. Lett. **62**, 466 (1989).
- ⁹E. Kapon, D. M. Hwang, and R. Bhat, Phys. Rev. Lett. 63, 430 (1989).
- ¹⁰J. A. Lebens, C. S. Tsai, K. J. Vahala, and T. F. Kuech, Appl. Phys. Lett. 56, 2642 (1990).
- ¹¹ T. F. Kuech, M. S. Goorsky, M. A. Tischler, A. Palevski, P. Solomon, R. Potemski, C. S. Tsai, J. A. Lebens, and K. J. Vahala, J. Cryst. Growth 107, 116 (1991).
- ¹² Y. D. Galeuchet, H. Rothuizen, and P. Roentgen, Appl. Phys. Lett. 58, 2423 (1991).
- ¹³S. Ando and T. Fukui, J. Cryst. Growth 98, 646 (1989).
- ¹⁴T. F. Kuech, M. A. Tischler, and R. Potemski, Appl. Phys. Lett. 54, 910 (1989).
- ¹⁵M. E. Hoenk and K. J. Vahala, Rev. Sci. Instrum. 60, 226 (1989).