Josephson junctions via anodization of epitaxial Al on an InAs heterostructure
Josephson junctions via anodization of epitaxial Al on an InAs heterostructure

Cite as: Appl. Phys. Lett. 119, 172601 (2021); doi: 10.1063/5.0060757
Submitted: 22 June 2021 · Accepted: 13 October 2021 · Published Online: 25 October 2021

A. Jouan,1,a) J. D. S. Witt,1 G. C. Gardner,2 C. Thomas,2,3 T. Lindemann,2,3 S. Gronin,2 M. J. Manfra,2,3 and D. J. Reilly1,4,a)

AFFILIATIONS
1ARC Centre of Excellence for Engineered Quantum Systems, School of Physics, The University of Sydney, Sydney, NSW 2006, Australia
2Microsoft Quantum Purdue, Purdue University, West Lafayette, Indiana 47907, USA
3Department of Physics and Astronomy, Purdue University, West Lafayette, Indiana 47907, USA
4Microsoft Quantum Sydney, The University of Sydney, Sydney, NSW 2006, Australia

a)Authors to whom correspondence should be addressed: alexis.jouan@ens-lyon.fr and david.reilly@sydney.edu.au

ABSTRACT

We combine electron beam lithography and masked anodization of epitaxial aluminum to define tunnel junctions via selective oxidation, alleviating the need for wet-etch processing or direct deposition of dielectric materials. Applying this technique to define Josephson junctions in proximity-induced superconducting Al-InAs heterostructures, we observe multiple Andreev reflections in transport experiments, indicative of a high quality junction. We further compare the mobility and density of Hall-bars defined via wet etching and anodization. These results may find utility in uncovering better fabrication approaches to junction-based qubit platforms.

Published under an exclusive license by AIP Publishing. https://doi.org/10.1063/5.0060757

Zero energy modes in a one-dimensional (1D) topological superconductor are candidate building blocks for high-fidelity quantum computation.1,2 One proposed approach to realizing a topological superconductor is via a two-dimensional electron gas (2DEG) with large spin–orbit interaction, proximity coupled to a conventional superconductor and confined to 1D using electrostatic gates and tunnel barriers.3 Such a configuration is experimentally challenging to realize however, since many of the needed parameters also compete with each other. Inducing superconductivity, for instance, requires that the 2DEG be close to the superconductor, where disorder from surface states can also greatly degrade the mobility.

Josephson junctions or tunnel barriers separating topological segments from normal regions are particularly sensitive to disorder. Typically, crystalline superconductors, grown using molecular beam epitaxy (MBE), are etched away using invasive wet chemical processing or damaging dry etchant techniques.9–11 This approach is often used to create, for example, the weak link of a Josephson junction via wet etching of Al (usually with transene).12 Although there is opportunity to further optimize the etching process,13 circumventing the need for it entirely would likely lead to junctions with less disorder. An associated challenge is the patterning of features on the length-scale of a few tens of nanometers in device structures, for example, to be able to reach the Josephson short-junction limit.14 At these length-scales, wet-etching can be difficult to control.

Here, we report the use of anodization to define a Josephson junction via the selective oxidation of epitaxial aluminum on an InAs heterostructure. In addition to being largely noninvasive, the technique offers a means of defining structures with a resolution limited, in principle, only by the nanometer precision of the electron beam lithography. In a straight up comparison, we also show that Hall bars defined via anodization have a moderately higher mobility than devices that are shaped by wet-etch processing.

Anodization is a standard industrial method used extensively to protect or modify the properties of surfaces.15 It has a wide range of applications, most abundantly for protecting metallic objects from uncontrolled oxidation. In the field of device physics, it has been used to define Nb Josephson junctions.8

The anodization circuit is realized such that the target metal constitutes the anode. By driving a current through the target metal submerged in a suitable electrolyte, an oxydo-reduction reaction can be induced. In the case of Al, alumina (Al2O3) is formed,

\[ 2\text{Al}(s) + 3\text{H}_2\text{O}(l) = \text{Al}_2\text{O}_3(s) + 6\text{H}^+ + 6\text{e}^-. \]  (1)
This process allows for the controlled oxidation of Al to proceed to a specific depth beyond the 3 nm native oxide, at a rate fixed by the applied current.17

Turning to quantum devices, we make use of this reaction to define tunnel barriers in materials comprising superconducting heterostructures. Unlike conventional fabrication approaches that remove the Al superconductor and back-fill with dielectric, here we rather use a second liftoff patterned layer of evaporated Al (Al-mask) on top of the epi-Al and employ the anodization process. In this way, we are able to retain the high-resolution pattern of the evaporated Al-mask and also the high-quality interface between the heterostructure and the newly formed dielectric near the shallow quantum well.

Starting from an InAlAs/InAs/InGaAs heterostructure (quantum well 10 nm below the surface and 7 nm of in situ grown epi-Al), a poly(methyl methacrylate) resist is spin-coated and patterned using electron beam lithography, as shown in Fig. 1(a). The Al-mask layer—the same thickness as the epi-Al layer—is then evaporated and the pattern formed using the liftoff technique. Immediately prior to the evaporation of the Al-mask layer, in the same vacuum cycle, an Ar ion mill step is performed to remove the native oxide from the epi-Al, ensuring good electrical contact between the two Al layers. The sample is then placed in a setup as shown in Fig. 1(b), submerged in an electrolyte—a mixture of ammonium pentaborate tetra-hydrate and ethylene glycol— and a current of 50 mA flows between the needle and surface of the sample. (c) Optical microscope image of the 2μm wide planar Josephson junction after the ALD-grown Al2O3 layer was deposited. The junction was aligned along the 110 orientation of the InAs heterostructure beneath. It is 2μm wide, surrounded by normal 2DEG regions, and has a gap of 150 nm.

FIG. 1. (a) In0.9Ga0.1As/InAs/In0.9Al0.1As quantum well covered with an in situ grown 7 nm epi-Al layer. An Al-mask (circled in black) was patterned using electron beam lithography. Finally, the sample was anodized and covered with ALD-grown Al2O3 and Au top gate. (Inset) A schematic representation of the quantum well (in red) and the wave function of the first filled level. (b) Schematic of the jig used to anodize the Al. A mixture of ammonium pentaborate tetra-hydrate with ethylene glycol, following Ref. 5, is poured into the jig, and a current of 50 μA flows between the needle and surface of the sample. (c) Optical microscope image of the 2 μm wide planar Josephson junction after the ALD-grown Al2O3 layer was deposited. The junction was aligned along the 110 orientation of the InAs heterostructure beneath. It is 2 μm wide, surrounded by normal 2DEG regions, and has a gap of 150 nm.
as shown in Fig. 2(a). By monitoring \( V_{\text{anod}} \) as a function of time, the effective depth of the anodized Al can be determined.

The anodization is stopped at a known voltage, \( V_{\text{anod}} \) which corresponds to the thickness of the Al-mask. In this way, the Al-mask has effectively been anodized instead of the epi-Al in the specifically patterned regions. The regions that were not covered by the Al-mask are now fully insulating, while the parts of the epi-Al, which were covered with the Al-mask are now covered with a layer of alumina [Fig. 1(c)]. An anodization rate of 100 mV/s for both the Al-mask and epi-Al has been measured. It is worth emphasizing that for this method to work, the rates for both must be equal. This ensures that the Al-mask and the uncovered epi-Al layer will be fully anodized at the same point in time.

Following the anodization, a mesa structure is etched in order to avoid top-gate leakage. The alumina around the mesa is removed using a developing solution, AZ726, followed by a dilute phosphoric acid etch. Finally, a 5 nm thick alumina layer is grown by atomic layer deposition (ALD) and the Ti/Au top-gates are deposited, as shown in Fig. 1(a).

The anodization ratio of oxide thickness, \( d \), to measured voltage, \( V_{\text{anod}} \) was determined by performing ellipsometry on test samples anodized to different depths. A ratio of 1.37 nm V\(^{-1}\) was found, which is consistent with values in the literature.\(^{15,16}\) A further corroboration of the extrapolated rate is the clear kink in \( V_{\text{anod}} \) vs \( t \) of the epi-Al anodization, which can be seen in Fig. 2(a). This voltage translates to an epi-Al thickness of 7 nm, which matches that measured by x-ray diffraction. The change in rate of anodization corresponds to a higher critical electric field, \( E_c \), needed to force ions through the InAlAs layer. Consistent with this picture, we estimate the electric field of the oxide by measuring the capacitance, \( C = \varepsilon_0 \varepsilon_r S/d \), between two patterned pads of surface, \( S \), with different anodized alumina of known thickness, \( d \), see Fig. 2(b). The relative permeability was found to be, \( \varepsilon_r = 8.9 \pm 0.4 \).

Next, we compare the mobility and electron density of Hall bars defined using standard wet etching to those defined by anodization, as shown in Figs. 2(c) and 2(d). Here, data are taken at an electron temperature of \( T_e = 55 \) mK. For the Hall-bar defined using anodization, the mobility at \( (V_g = 0 \text{ V}) \) was found to be \( \mu \sim 12,000 \text{ cm}^2/(\text{V s}) \), with electron density \( (n = 1.2 \times 10^{12} \text{ electrons/cm}^2) \), comparable to values given in the literature.\(^17\) Application of a negative gate voltage leads to a peak mobility of \( \mu \sim 15,000 \text{ cm}^2/(\text{V s}) \). The variation in mobility as a function of applied top-gate voltage is consistent with screening of charge impurities by electrons in the 2DEG.\(^17\)

For comparison, a sample from the same wafer was patterned into a Hall bar using an optimized standard transene type-D etch\(^3\) (otherwise processed identically) and was found to have a peak mobility of \( \mu \sim 7000 \text{ cm}^2/(\text{V s}) \). We underscore the improvement in mobility of the device fabricated using anodization, indicating the sensitivity of these device structures to disorder induced by chemical processing. A more detailed statistical study would have to be done to quantify this improvement. Also of note, for the anodized Hall bar, the electron density at \( V_g = 0 \text{ V} \) is around half that of the chemically etched device. A similar anodization method has also recently yielded an enhancement of the mobility.\(^20\)

Finally, to demonstrate the utility of this technique for nanoscale device fabrication, a Josephson junction with a barrier region of length \( \sim 150 \text{ nm} \) was fabricated using anodization, as shown in Fig. 1(c). The device was characterized using a standard dc transport making use of a current-bias with four-point measurement in a dilution refrigerator with an electron temperature of \( T_e = 55 \) mK (determined using an normal insulator superconductor (NIS) junction\(^1\)). The epi-Al layer has a critical temperature above \( T \sim 1.5 \text{ K} \) and an in-plane critical magnetic field, \( B_c \sim 2.5 \text{ T} \).\(^2\) The mobility obtained from the Hall measurements corresponds to an elastic scattering length, \( l_s \sim 200 \text{ nm} \), which suggests that the junction is in the quasi-ballistic limit.

The current–voltage characteristics of the device as a function of applied top-gate voltage are shown in Fig. 3. Despite the relatively low critical current compared to similar devices in the literature,\(^*\) multiple Andreev reflections (MARs) are clearly resolved, which indicates that the electrons remain phase coherent over multiple times the length of the weak link.

With the increasing negative applied top-gate voltage, the 2DEG is depleted and the critical current decreases, as shown in the lower panel of Fig. 3. Below \( V_g = -1.5 \text{ V} \), the junction loses phase coherence and a finite voltage appears across the weak link. The characteristic voltage or \( I_l R_l \) product is constant between 0 and \(-1.5 \text{ V} \) (see the supplementary material) at around 3 \( \mu\text{V} \), which is a few percent of the Al gap, \( \Delta_{\text{Al}} = 1.76k_BT_e \simeq 230 \mu\text{V} \). As an indication, the excess current at \(-1 \text{ V} \) is measurable and is \( I_{\text{ac}} \approx 0.95 \text{ nA} \), which is approximately 15% of the critical current. Although lower than the value expected for a fully transparent interface,\(^20\) further investigations into the exact nature of the induced superconducting gap, \( \Delta' \), in this particular type of heterostructure need to be performed in order to fully understand this discrepancy. The concentration of Al in the InAlAs layer is especially important in this regard.
Finally, we apply a perpendicular magnetic field and investigate the differential resistance of the junction, as shown in Fig. 4. Fraunhofer patterns are evident—with some asymmetry between positive and negative applied bias. The critical current goes to zero twice in the range shown (three lobes) at the field values where an integer multiple of the flux quantum threads through the weak link. The measured field at which the critical current is cancelled by currents of opposite direction due to the flux through the junction is ~1 mT, which is lower than the expected value given the dimensions of the junction. This discrepancy can be explained either by magnetic flux focusing due to the Meissner effect in the superconducting leads or junction. This discrepancy can be explained either by magnetic flux focusing due to the Meissner effect in the superconducting leads or junction. This discrepancy can be explained either by magnetic flux focusing due to the Meissner effect in the superconducting leads or junction.

The authors thank S. Pauka and M. Cassidy for useful conversations and technical help. This research was supported by the Microsoft Corporation and the Australian Research Council Centre of Excellence for Engineered Quantum Systems (EQUSS, No. CE170100009). The authors acknowledge the facilities as well as the scientific and technical assistance of the Research & Prototype Foundry Core Research Facility at the University of Sydney, part of the Australian National Fabrication Facility.

AUTHOR DECLARATIONS
Conflict of Interest
The authors declare no conflict of interest.

DATA AVAILABILITY
The data that support the findings of this study are available from the corresponding authors upon reasonable request.

REFERENCES

1. A. Y. Kitaev, Phys.-Usp. 44, 131 (2001).
2. C. Nayak, S. H. Simon, A. Stern, M. Freedman, and S. D. Sarma, Rev. Mod. Phys. 80, 1083 (2008).
3. R. M. Lutchyn, J. D. Sau, and S. D. Sarma, Phys. Rev. Lett. 105, 077001 (2010).
4. Y. Oreg, G. Refael, and F. von Oppen, Phys. Rev. Lett. 105, 177002 (2010).
5. V. Mourik, K. Zuo, S. M. Frolov, S. R. Plissard, E. P. A. M. Bakkers, and L. P. Kouwenhoven, Science 336, 1003 (2012).
6. M. Kjaergaard, F. Nichele, H. J. Suominen, M. P. Nowak, M. Wimmer, A. A. Akhmerov, J. A. Folk, K. Flensberg, J. Shabani, C. J. Palmstrøm, and C. M. Marcus, Nat. Commun. 7, 12841 (2016).
7. A. Hatke, T. Wang, C. Thomas, G. Gardner, and M. Manfra, Appl. Phys. Lett. 111, 142106 (2017).
8. H. Kroger, L. Smith, and D. Jille, Appl. Phys. Lett. 39, 280 (1981).
9. A. Richter, P. Baars, and U. Merkt, Physica E 12, 911 (2002).
10. Heida, B. Van Wees, T. Klappwig, and G. Borghs, Phys. Rev. B 57, R5618 (1998).
11. L. Mur, C. Harmans, J. Moosj, J. Carlin, A. Rudra, and M. Ilegems, Phys. Rev. B 54, R2327 (1996).
12. A. Fornieri, A. M. Whiticar, F. Setiawan, E. Portolés, A. C. C. Drachmann, A. Keselman, S. Gronin, C. Thomas, T. Wang, R. Kallalher, G. C. Gardner, E. Berg, M. J. Manfra, A. Stern, C. M. Marcus, and F. Nichele, Nature 569, 89 (2019).
13. S. Pauka, J. Witt, C. Allen, B. Harlech-Jones, A. Jouan, G. Gardner, S. Gronin, T. Wang, C. Thomas, M. Manfra et al., J. Appl. Phys. 128, 114301 (2020).
14. Y. Peng, F. Pienitka, E. Berg, Y. Oreg, and F. von Oppen, Phys. Rev. B 94, 085409 (2016).
15. J. W. Diggle, T. C. Downie, and C. W. Goulding, Chem. Rev. 69, 365 (1969).
16. N. Mott, Trans. Faraday Soc. 43, 429 (1947).
17. M. S. Hunter and P. Fowle, J. Phys. E 101, 481 (1954).
18. T. Tschirky, S. Mueller, C. A. Lechner, S. Fält, T. Ihn, K. Ensslin, and W. Wegscheider, Phys. Rev. B 95, 115304 (2017).
19. V. Umansky, R. de Picciotto, and M. Heiblum, Appl. Phys. Lett. 71, 683 (1997).
20. A. C. C. Drachmann, R. E. Diaz, C. Thomas, H. J. Suominen, A. M. Whiticar, A. Fornieri, S. Gronin, T. Wang, G. C. Gardner, A. R. Hamilton et al., Phys. Rev. Mater. 5, 013805 (2021).
21. A. Fedchenko, L. Casparis, I. Khaymovich, D. Maradan, O.-P. Saira, M. Palma, M. Meschke, J. Pokola, and D. Zumbühl, Phys. Rev. Appl. 4, 034001 (2015).
22. H. Suominen, J. Danon, M. Kjaergaard, K. Flensberg, J. Shabani, C. Palmstrøm, F. Nichele, and C. Marcus, Phys. Rev. B 95, 035307 (2017).
23. W. Mayer, J. Yuan, K. S. Wickramasinghe, T. Nguyen, M. C. Dartiailh, and J. Shabani, Appl. Phys. Lett. 114, 103504 (2019).
24. M. Kjaergaard, H. J. Suominen, M. Nowak, A. Akhmerov, J. Shabani, C. Palmstrøm, F. Nichele, and C. M. Marcus, Phys. Rev. Appl. 7, 034029 (2017).
25. G. E. Blonder, M. Tinkham, and T. M. Klapwijk, Phys. Rev. B 25, 4515 (1982).
26. M. Tinkham, Introduction to Superconductivity (Courier Corporation, 2004).