Supporting Information

Effective surface diffusion of nickel on single crystal β -Ga₂O₃ for Schottky barrier modulation and high thermal stability

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Deposition methods	Doping concentration (N_d - N_a , cm ⁻³)		
	As-dep.	400 °C	
E-beam	2.66 x 10 ¹⁶	2.47 x 10 ¹⁶	
CMFS	4.52 x 10 ¹⁶	3.88 x 10 ¹⁶	

Fig. S1. Capacitance-voltage (*C-V*) measurement on the CMFS deposited and E-beam evaporated Ni/ β -Ga₂O₃ SBDs with post-annealing. The doping concentration (N_d - N_a) of drift layer was calculated about 3 x 10¹⁶ cm⁻³ shown in the table.



Fig. S2. (a) The variations of the onset voltage (V_{ON}) and (b) reverse breakdown voltage (V_{BR}) on CMFS deposited and E-beam evaporated Ni/ β -Ga₂O₃ SBDs. The V_{BR} of CMFS-Ni SBDs was varied -443 to -496 V, while which of E-beam evaporated Ni-based SBDs was varied -577 to -585 V with post-annealing temperature. In general, the reverse leakage current in the SBDs is related to the SBH, because the thermionic emission current across the SBH is a dominant component of the leakage current. In addition, some phenomena, such as field-emission, image force lowering effect, and barrier inhomogeneity can lower the SBH value. The 400 °C annealed CMFS-Ni devices have relatively higher and homogeneous SBH rather than the E-beam devices, which can lead to the low leakage current. However, it seems that there are some carrier trap regions in CMFS-Ni/ β -Ga₂O₃ interface, which can be accelerated to the impact ionization, lowering the V_{BR} value.



Ni (111)	2θ (deg)	FWHM (deg)	Lattice constant (nm)	Average grain size (nm)
CMFS as-dep.	44.65	0.73	0.410	11.51
CMFS 400 °C	44.69	0.73	0.388	11.59
E-beam as-dep.	44.60	0.64	0.439	13.03
E-beam 400 °C	44.63	0.69	0.423	12.14

Fig. S3. The XRD analysis of the Ni with CMFS and E-beam evaporation methods. The diffraction peaks at $2\theta = 44.6^{\circ}$ and 51.9° were assigned to the (111) and (200) crystal planes of the Ni. (JCPDS #04-0850) The table shows the parameters of Ni (111) peaks, which are the 2 θ center of peak, full width at half maximum (FWHM), lattice constant, and average grain size calculated using the Scherrer equation. All Ni peaks were slightly shifted to the increasing 2 θ angle after post-annealing due to the decrease in the lattice constant with relaxation of crystal.



Fig. S4. The XPS depth-profiling analysis with the E-beam evaporated Ni/ β -Ga₂O₃ interface. (a,c,e) As-deposited and (b,d,f) post-annealed at 400 °C with Ni2p, Ga2p, and O1s spectrums, respectively. (g, h) The shift of binding energies of Ni2p and Ga2p peaks as increasing interfacial depth. No additional Ni diffusion was identified with post-treatment.