

Support Information: Supported two- and three-dimensional vanadium oxide species on the surface of β -SiC

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S.1) Set of data to accurately measure:

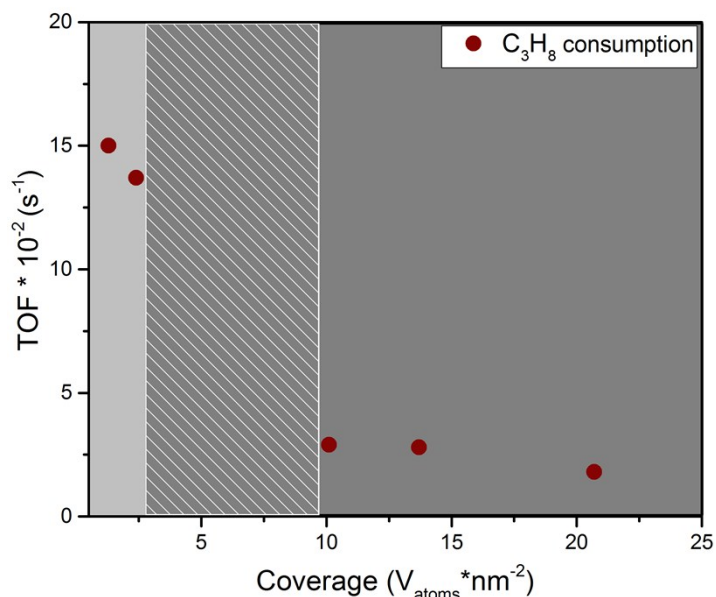
BET surface area (m²/g)

Sample	BET-I	BET-II	BET-III	Average	STDEV
β -SiC _p	32.1	28.3	29.4	29.9	1.6
2V/ β -SiC _p	25.6	28.5	25.8	26.6	1.3
β -SiC _o	29.3	29.5	29	29.3	0.2
1V/ β -SiC _o	27.1	26.6	26.2	26.6	0.4
2V/ β -SiC _o	24.6	26.3	26.8	25.9	0.9
4V/ β -SiC _o	14.9	12.4	15.8	14.4	1.4
6V/ β -SiC _o	14.2	14.7	13.7	14.2	0.4
8V/ β -SiC _o	13.5	11.8	12.9	12.7	0.7
10V/ β -SiC _o	12.1	10.4	11.9	11.5	0.8

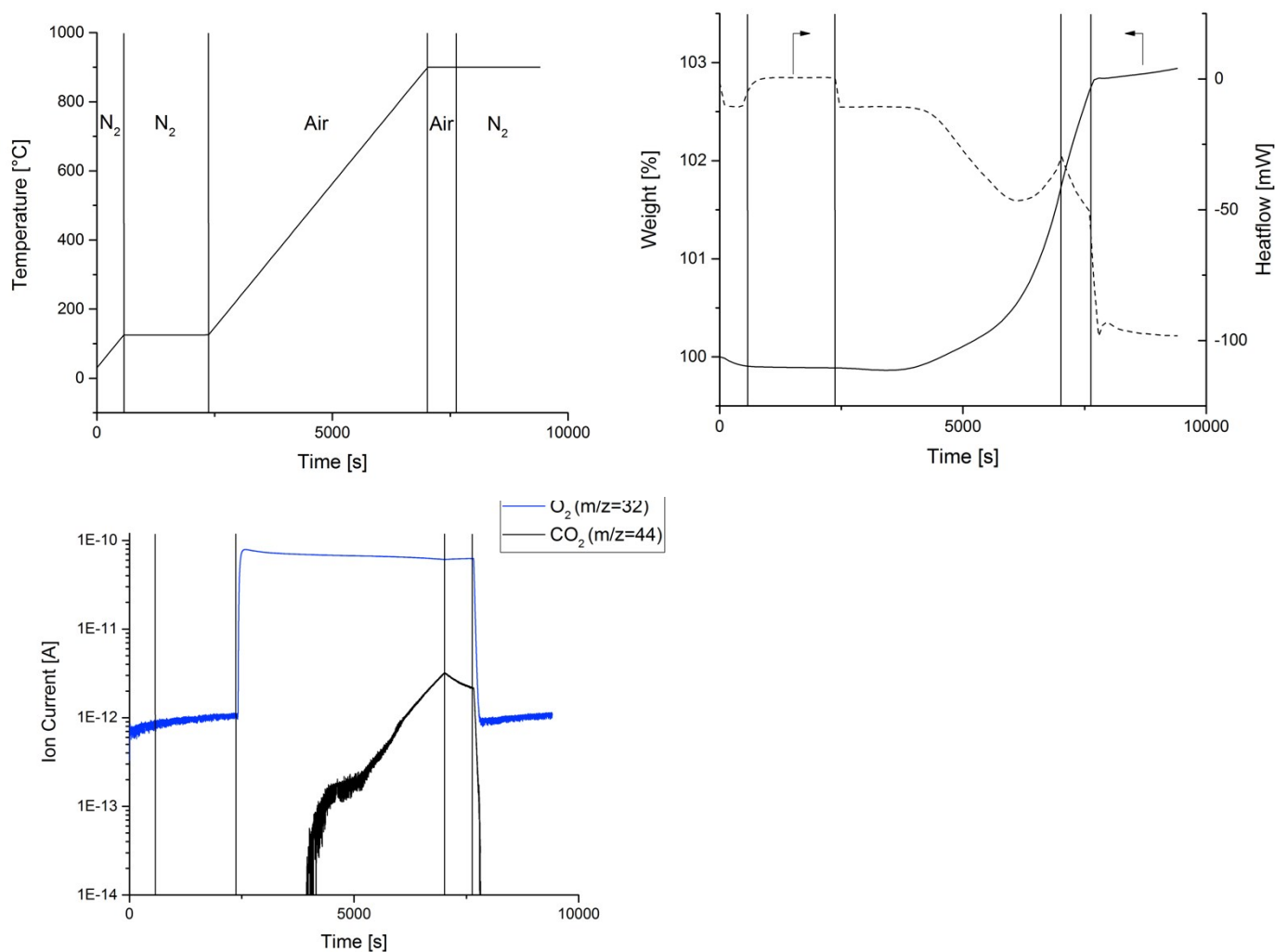
V loading (w.t. %)

Sample	ICP-I	ICP-II	ICP-III	Average	STDEV
β -SiC _p	-				
2V/ β -SiC _p	1.5	2.1	1.9	1.8	0.2
β -SiC _o	-				
1V/ β -SiC _o	1.4	0.9	0.7	1.0	0.3
2V/ β -SiC _o	1.3	2.1	2.5	2.0	0.5
4V/ β -SiC _o	4.1	4.3	4.5	4.3	0.2
6V/ β -SiC _o	6.1	6.3	5.4	5.9	0.4
8V/ β -SiC _o	7.9	7.8	7.9	7.9	0.1
10V/ β -SiC _o	9.6	9.8	10.1	9.8	0.2

S.2) Turn over frequency (TOF) for propane consumption as a function of V coverage for the studied V/ β -SiC₀ catalysts at 490 C. C₃H₈/O₂/N₂ = 6:3:11 composition. The inverse weight hourly space velocity (WHSV⁻¹) was varied between 40-250 kg_{cat}·s/m³.

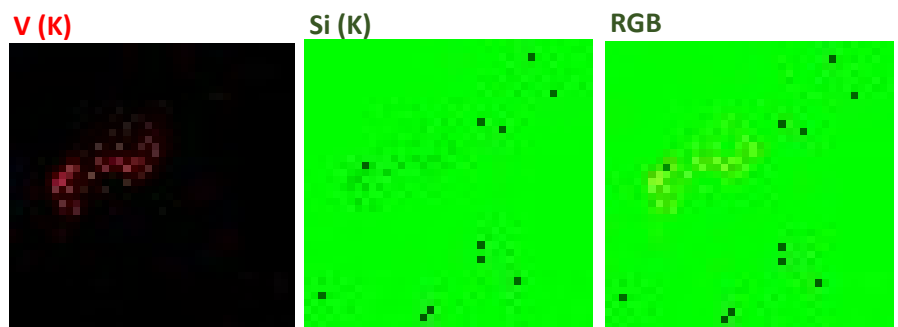
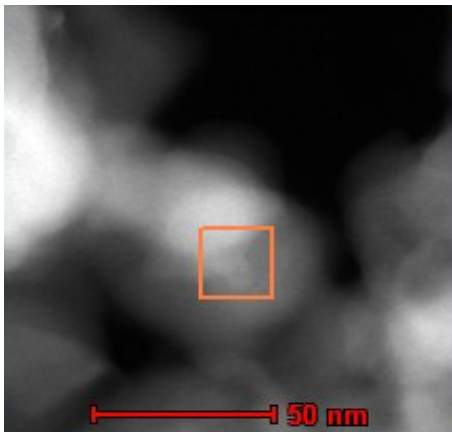
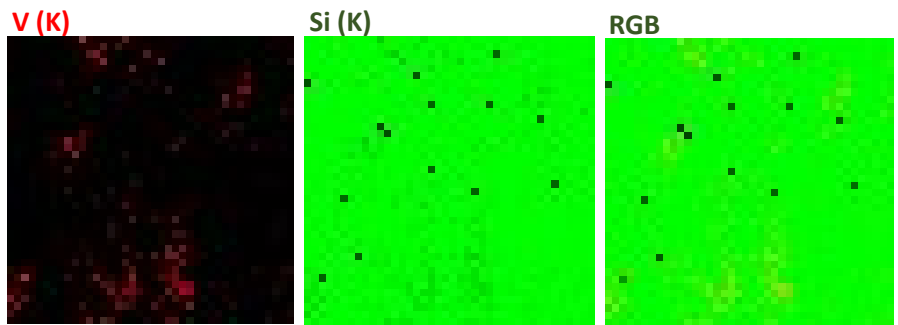
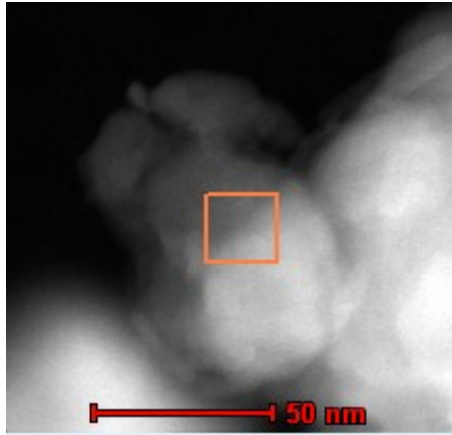


S.3) The used temperature program (top left), TGA-DSC data (top right), and the MS data for O₂ and CO₂ (bottom left). At around 400 °C under air, the formation of CO₂ begins and steadily increases. Once air is switched to N₂, CO₂ production stops. The increase in mass, despite of the formation of CO₂, occurs basically because SiC + 2O₂ -> SiO₂ + CO₂. As the molar mass of SiC is lower than SiO₂, this results in an increase in total mass.

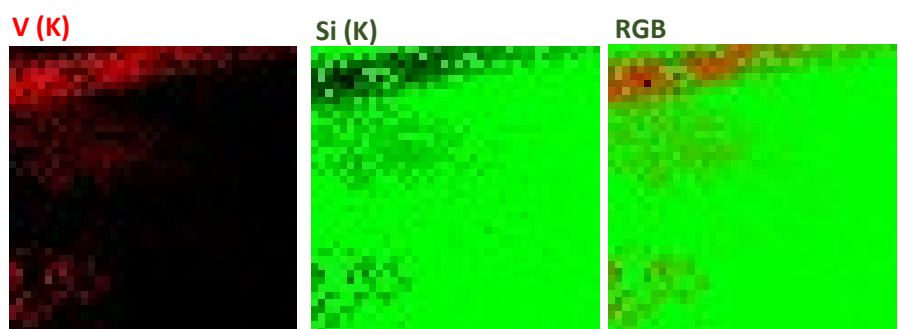
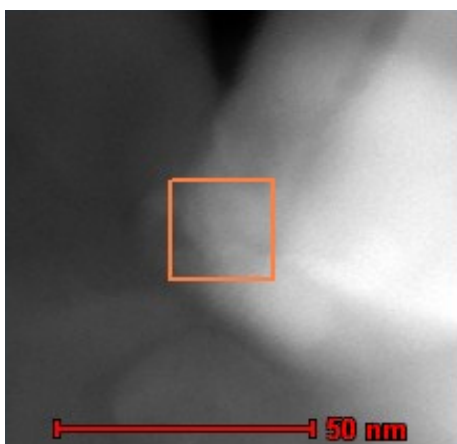


S.4) EDS Analysis for V/ β -SiC and V/SiO₂ samples. Whereas the formation of “V-islands” can be observed both at low ($2.4 V_{\text{atoms}} \cdot \text{nm}^{-2}$) and high ($13.7 V_{\text{atoms}} \cdot \text{nm}^{-2}$) V coverage on β -SiC, a more uniform V distribution is observed on V/SiO₂ samples with similar V coverage.

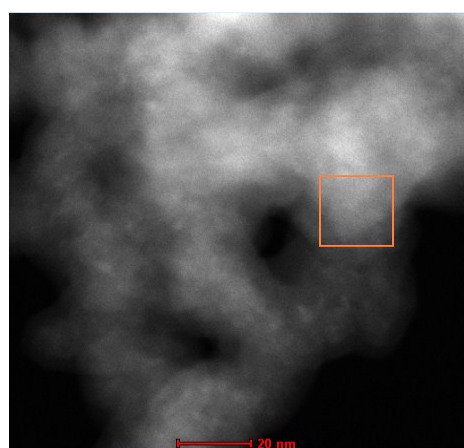
2V/ β -SiC ($\theta = 2.4 V_{\text{atoms}} \cdot \text{nm}^{-2}$)



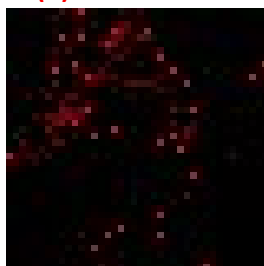
6V/ β -SiC ($\theta = 13.7 V_{\text{atoms}} \cdot \text{nm}^{-2}$)



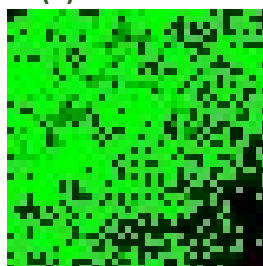
2V/SiO₂ ($\theta = 1.3 \text{ V}_{\text{atoms}} \cdot \text{nm}^{-2}$)



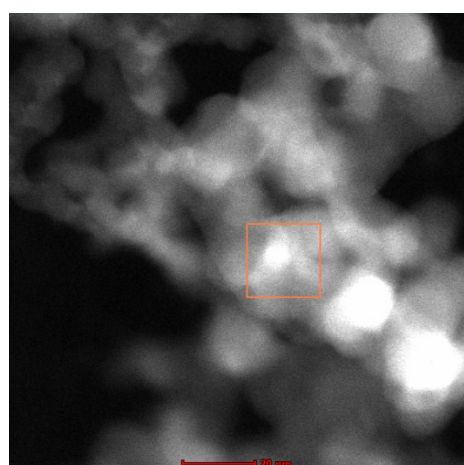
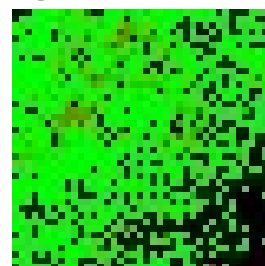
V (K)



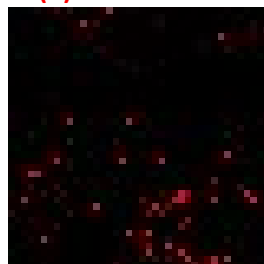
Si (K)



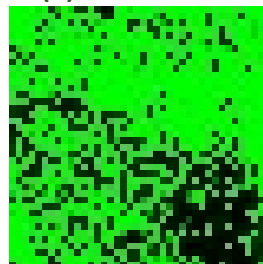
RGB



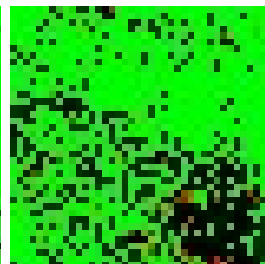
V (K)



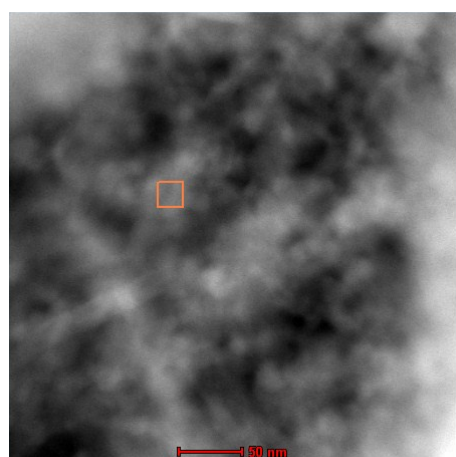
Si (K)



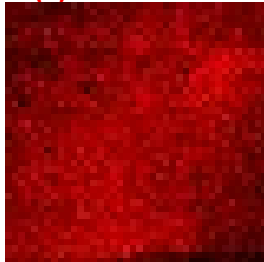
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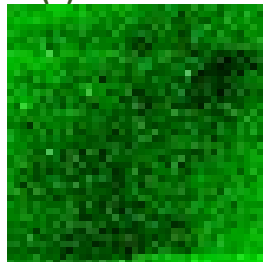
7V/SiO₂ ($\theta = 13.4 \text{ V}_{\text{atoms}} \cdot \text{nm}^{-2}$)



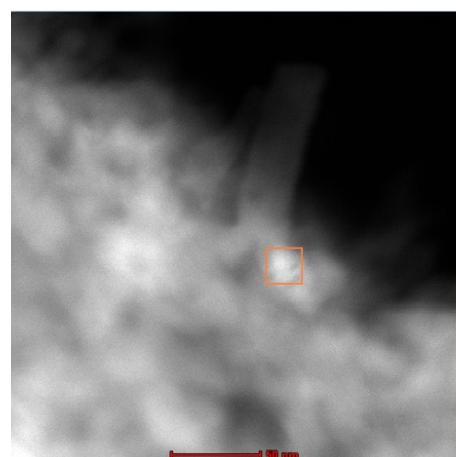
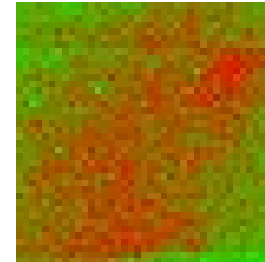
V (K)



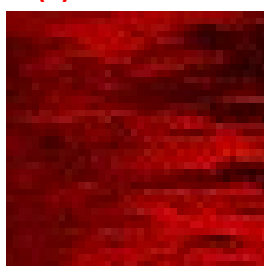
Si (K)



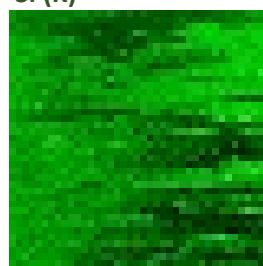
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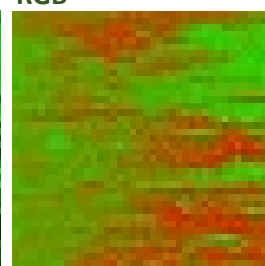
V (K)



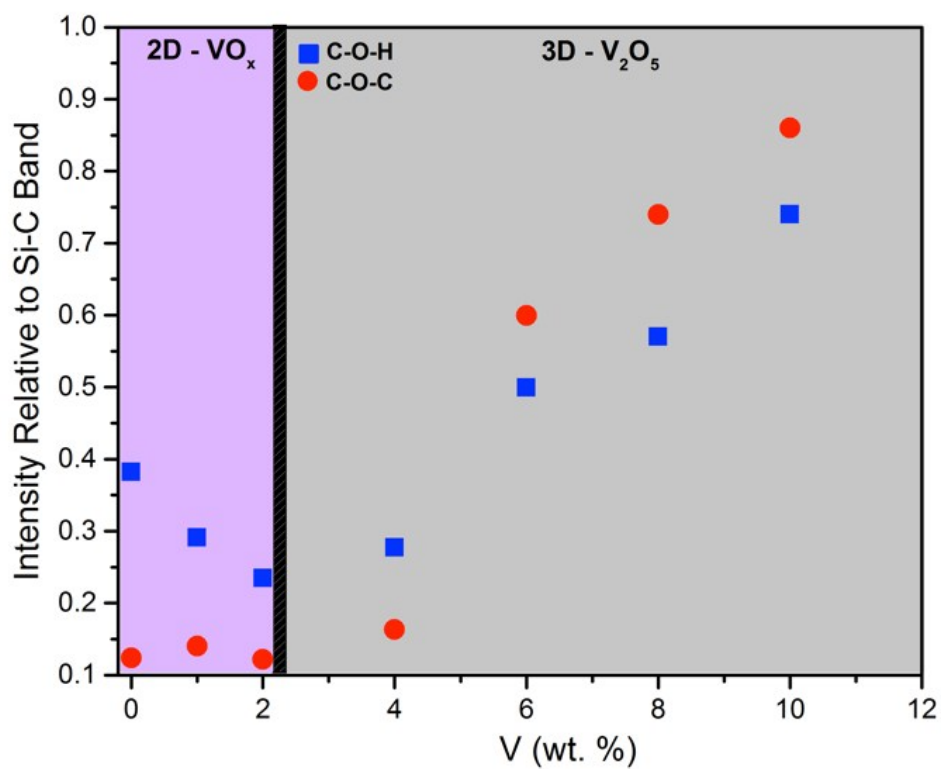
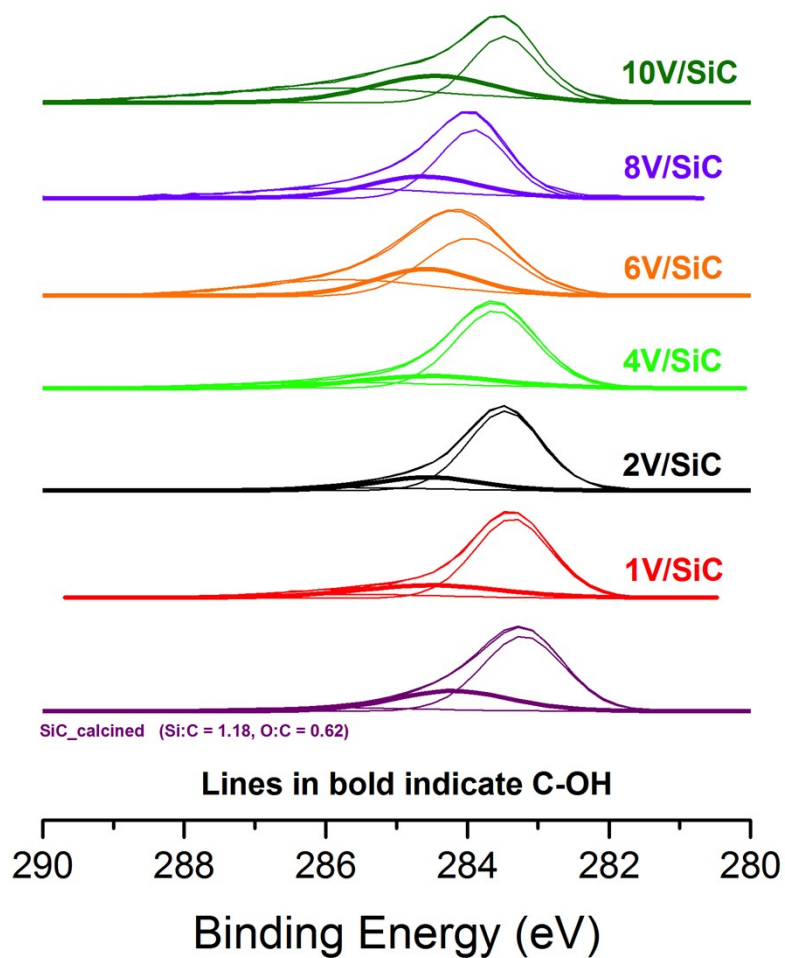
Si (K)



RGB



S.5) XPS C1s spectra of β -SiC with various vanadium oxide loading.



S.6) ^{51}V MAS NMR spectra of dehydrated $\text{V}/\beta\text{-SiC}_o$ materials containing 1, 2, 4, and 6 wt.% V, with signal intensity scaled to an equivalent number of scans. The 4 and 6 wt.% $\text{V}/\beta\text{-SiC}_o$ exhibit a single isotropic shift (denoted by an asterisk, *) at -616 ppm, whereas no signal is observed in the spectra for 1 and 2 wt.% V.

