

Supporting Information

Mo₂CT_x MXene–Based Non-Enzymatic Electrochemical Sensor for Selective Detection of Hydrogen Peroxide in Colon Cancer Cells

Shruthi Venkataraman^{1,#}, Vasanth Magesh¹, Chandramohan Govindasamy², Raji Atchudan^{3,#},
Sandeep Arya⁴, and Ashok K. Sundramoorthy^{*,1}

¹Centre for Nano-Biosensors, Department of Prosthodontics and Materials Science, Saveetha
Dental College and Hospitals, Saveetha Institute of Medical and Technical Sciences, India.

²Department of Community Health Sciences, College of Applied Medical Sciences, King
Saud University, P.O. Box 10219, Riyadh 11433, Saudi Arabia.

³School of Chemical Engineering, Yeungnam University, Gyeongsan 38541, Republic of
Korea.

⁴Department of Physics, University of Jammu, Jammu 180006, Jammu and Kashmir, India.

* Corresponding author.

E-mail addresses: ashok.sundramoorthy@gmail.com (A.K. Sundramoorthy).

#These authors equally contributed.

List of Supplementary Figures

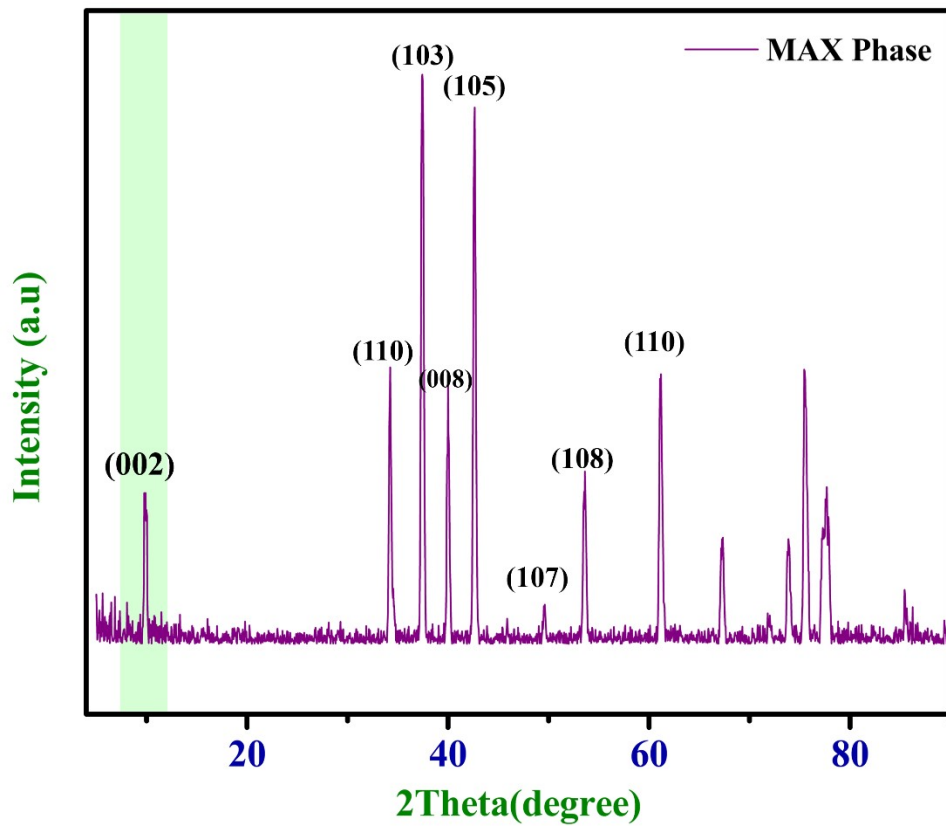


Figure S1. XRD data of Mo₂Ga₂C MAX phase.

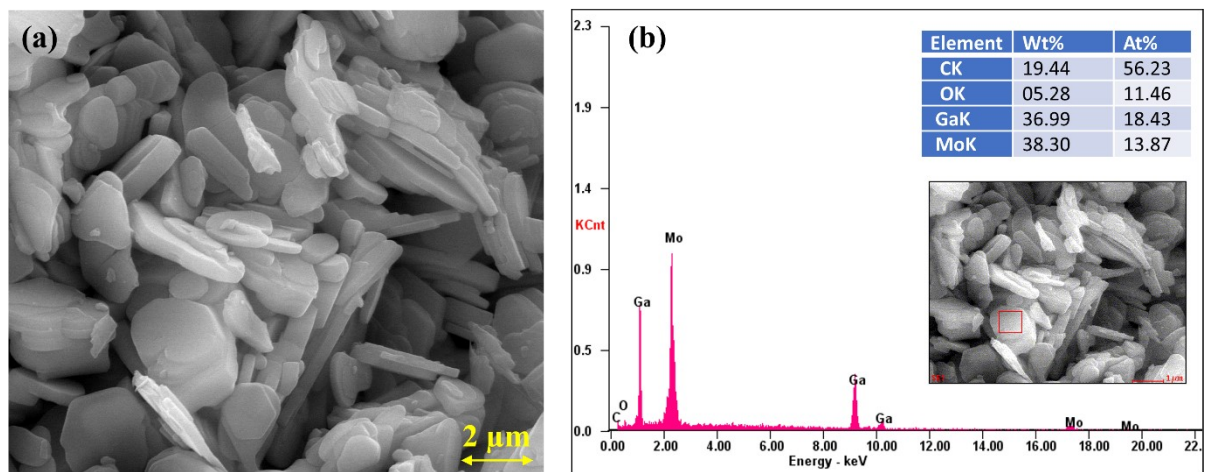


Figure S2. (a) SEM micrographs of Mo₂Ga₂C MAX phase precursor (Scale bar - 2 μm), (b) EDAX spectrum of Mo₂Ga₂C MAX phase. Inset: Table shows the list of elements and their wt% and at% along with SEM micrograph of Mo₂Ga₂C.

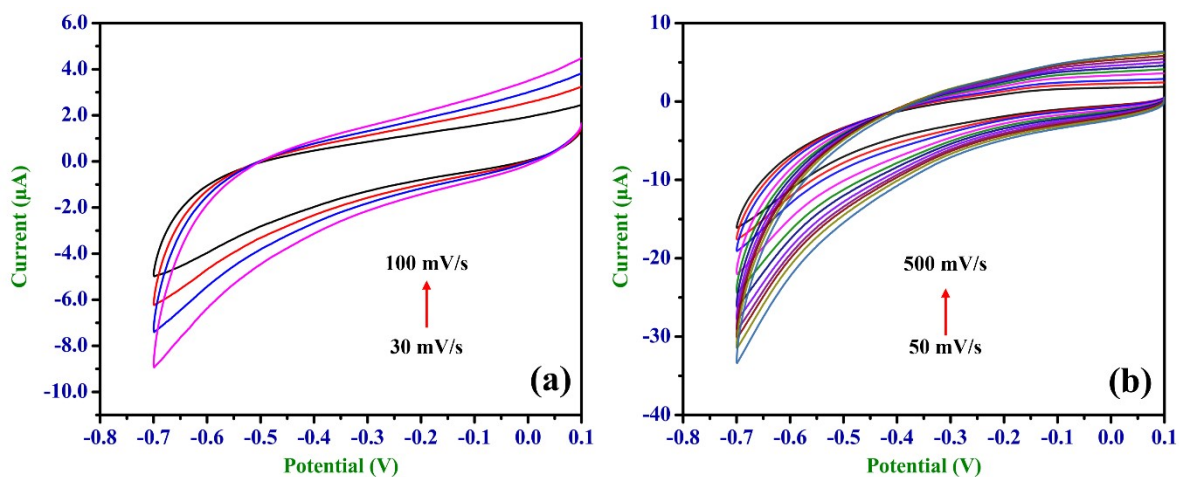


Figure S3. (a) CVs were recorded at different scan rates using an MX-GCE (without H₂O₂) in 0.1 M PBS (pH=7.4) from 30 mV/s to 100 mV/s. (b) CVs were recorded at different scan rates from 50 mV/s to 500 mV/s using an MX-GCE in 0.1 M PBS (pH=7.4) containing 284 mM H₂O₂.

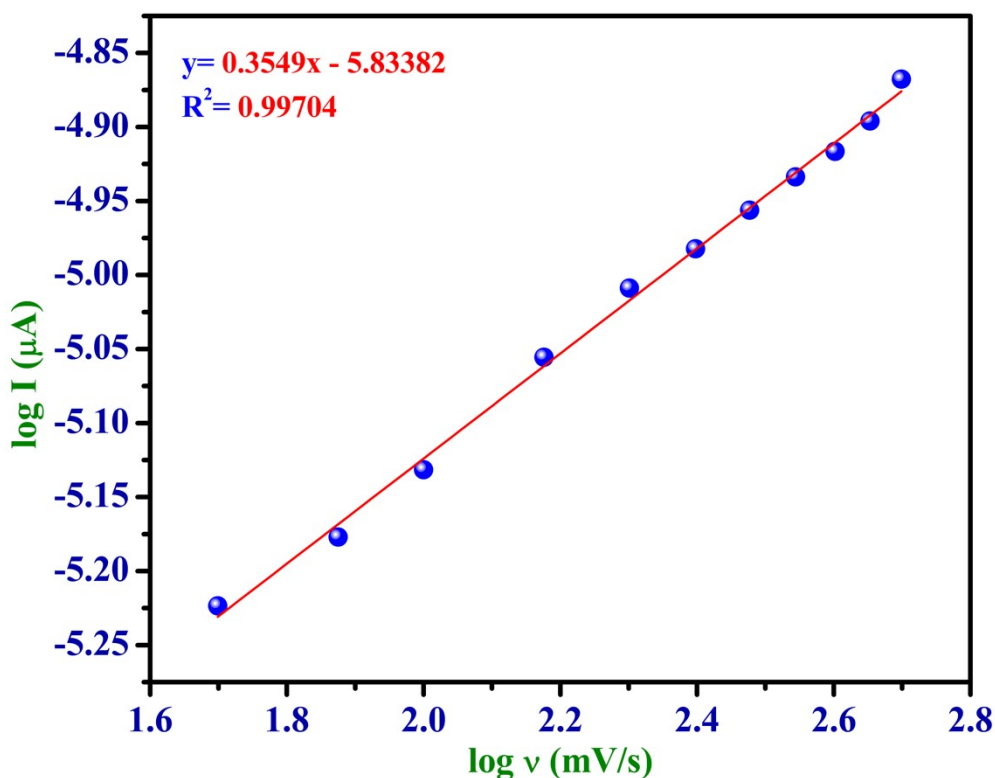


Figure S4. Linear plot of log I (µA) vs log v (mV/s) with different scan rates at a fixed potential of -0.45 V.

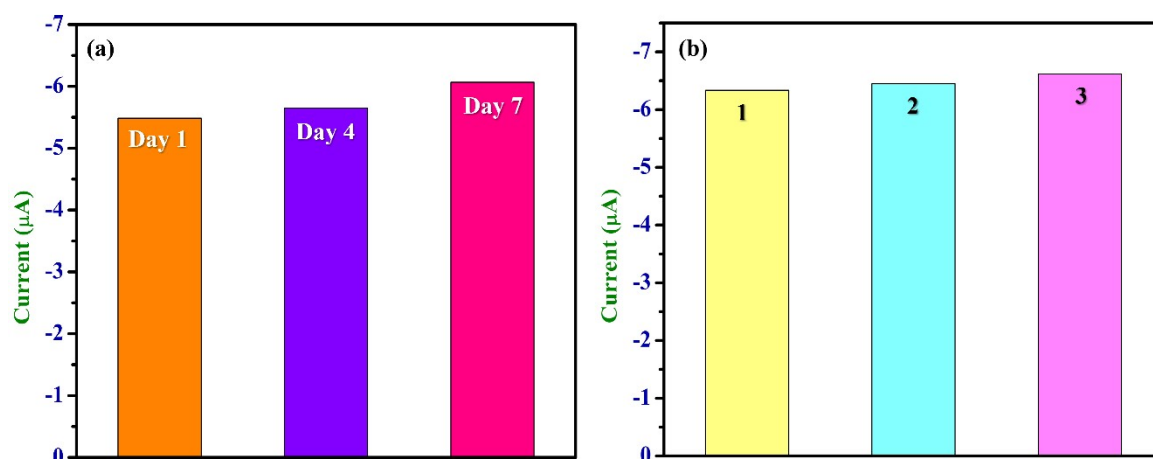


Figure S5. (a) CVs were recorded using MX-GCE in 0.1 M PBS (pH=7.4) at a scan rate of 50 mV/s on day 1, day 4, and day 7. (b) CVs were recorded using MX-GCE in 0.1 M PBS at a scan rate of 50 mV/s ($n=3$) for intra-day stability measurements.

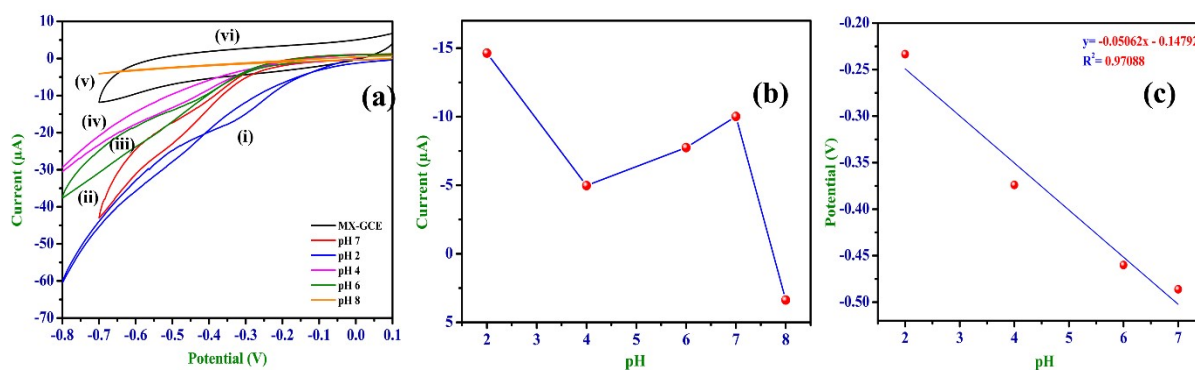


Figure S6. (a) CVs were recorded using a MX-GCE in the presence of 284 mM H_2O_2 with various buffer solutions at different: pH = (i) 2, (ii) 7.4, (iii) 6, (iv) 4, (v) 8, in comparison with CVs recorded with MX-GCE in the absence of H_2O_2 in 0.1 M PBS (pH = 7.4) at a scan rate of 50 mV/s. (b) Cathodic current response vs different pH in the presence of 284 mM H_2O_2 . (c) A linear plot shows the corresponding H_2O_2 reduction potential vs different pH in the presence of 284 mM H_2O_2 .

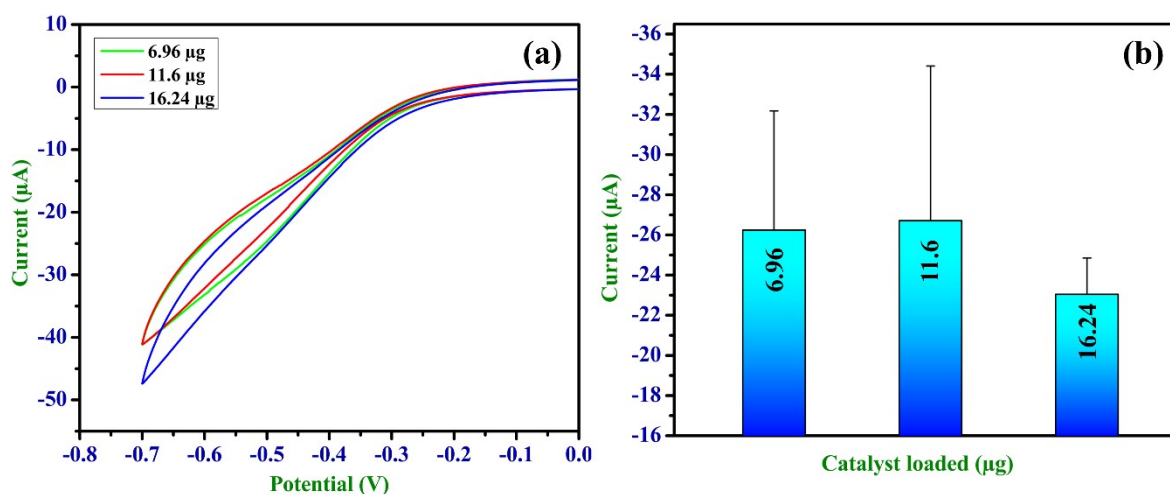


Figure S7. (a) CVs were recorded using an MX-GCE modified with different amount of catalyst (6.96, 11.6, 16.24 µg) in 0.1 M PBS (pH 7.4) in the presence of 284 mM H₂O₂ at a scan rate of 50 mV/s. (b) Corresponding bar plot of cathodic current vs different catalyst concentrations in 0.1 M PBS (pH = 7.4) in the presence of 284 mM H₂O₂ with standard deviation (SD) of 5.94×10^{-6} , 7.70×10^{-6} , 1.81×10^{-6} for 6.6 µg, 11.6 µg, and 16.2 µg, respectively.

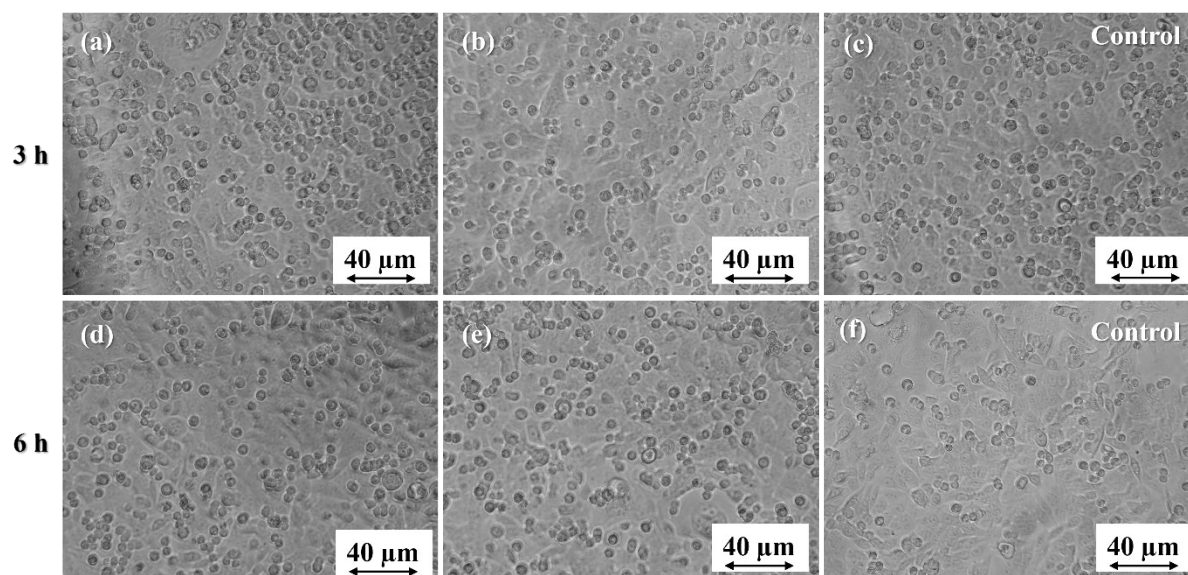


Figure S8. Phase contrast optical microscopy images of SW480 colorectal cancer cells following treatment with 0.5 mM ascorbic acid for 3 h at 37 °C, (a,b) different fields of view

from AA-treated SW480 cells for 3 h, (c) untreated controls (SW480 cells) without 0.5 mM AA for 3 h at 37 °C, (d, e) different fields of view from AA-treated SW480 cells for 6 h, and (e) untreated controls without 0.5 mM AA for 6 h at 37 °C.