

Cite this: *J. Mater. Chem. A*, 2025, **13**, 5252

MnTiO₃ as a carbon-free cathode for rechargeable Li–O₂ batteries†

Doaa Aasef Ahmed,^{id}*^a Mustafa Çelik,^{id}^{bc} Wernfried Mayr-Schmölzer,^a Abdulkadir Kızılaslan^{id}^{bc} and Gregor B. Vonbun-Feldbauer^{id}*^{ad}

Li–O₂ batteries (LOBs) are next-generation energy storage systems. However, their main challenges are the sluggish kinetics of oxygen reduction and evolution reactions (ORR/OER) and high charge overpotentials due to the formation of discharge product (Li₂O₂). To address this challenge, developing a catalyst with a unique structure and exceptional catalytic properties is crucial to enhancing the reversible cycling performance of LOBs, particularly under high current density conditions. Herein, the transition metal-based perovskite MnTiO₃ was examined as a carbon-free cathode catalyst using density functional theory (DFT) calculations and experimental techniques. The intrinsic advantages of MnTiO₃ stem from the coexistence of Mn and Ti energy levels near the Fermi level, as revealed by our density of states (DOS) analysis. This electronic structure facilitates ORR/OER, thus endowing MnTiO₃ with a bifunctional role in promoting battery performance. Our DFT-based investigation elucidates the surface stability and catalytic properties of MnTiO₃. Furthermore, Energy Dispersive Spectroscopy (EDS) and X-ray diffraction (XRD) confirm that the electrochemical reaction on MnTiO₃ follows a two-electron pathway. Our findings reveal that a LOB with MnTiO₃ exhibits a total overpotential of 1.18 V and 1.55 V using DFT and electrochemical measurements, respectively. High current densities up to 1 A g⁻¹ also highlight its potential as a cathode catalyst for LOBs.

Received 8th August 2024

Accepted 22nd November 2024

DOI: 10.1039/d4ta05571c

rsc.li/materials-a

1 Introduction

Over the last few years, extensive research and development efforts have resulted in deploying various renewable energy conversion and storage devices, some of which have reached commercialization.^{1,2} Among energy storage devices, rechargeable lithium–oxygen (Li–O₂) batteries (LOBs) are considered highly promising power sources due to their extraordinarily high theoretical energy density of 3500 W h kg⁻¹, low cost, and environmental friendliness.^{3,4}

A novel rechargeable LOB comprises a Li conductive electrolyte membrane sandwiched by a Li metal foil anode and an electroactive cathode. During discharge oxygen accessed from the environment is reduced on the cathode *via* the oxygen reduction reaction (ORR, 2Li⁺ + 2e⁻ + O₂ → Li₂O₂). The

charging process involves the decomposition of Li₂O₂ through the oxygen evolution reaction (OER, Li₂O₂ → 2Li⁺ + 2e⁻ + O₂).^{5,6}

Although LOBs exhibit significant theoretical energy potential, several challenges presently limit their effectiveness. Namely, the formation of solid and electronically insulating discharge products *i.e.* LiO₂ and Li₂O₂ during discharge significantly impedes fast reaction kinetics which overall leads to sluggish reactions.^{6,7}

To overcome this problem, carbonaceous materials including porous carbon,⁸ carbon nanotubes (CNTs), and graphene⁹ have been widely utilized to elevate the overall conductivity of electrodes. However, the inherent instability of carbon electrodes often leads to reactions with lithium peroxide (Li₂O₂), resulting in the formation of a non-conductive layer of lithium carbonate (Li₂CO₃) at the electrode-discharge product interface.¹⁰ To address these challenges and increase the overall catalytic performance, various carbon free cathode materials, such as noble metals and their oxides, transition metal carbides, and transition metal compounds, have been explored for LOBs.¹¹ Noble metal materials like Au,¹² Pt,¹³ IrO₂,¹⁴ and RuO₂ (ref. 15) exhibit remarkable electrocatalytic activity, effectively addressing the sluggish kinetics of the ORR or OER. However, their high cost, scarcity, and inferior cycling stability limit their practical application. In this context, it is of great importance to develop a material with a high catalytic effect that will strengthen OER and ORR reactions simultaneously. Recent

^aInstitute of Advanced Ceramics, Hamburg University of Technology, 21073 Hamburg, Germany. E-mail: doaa.ahmed@tuhh.de; gregor.feldbauer@tuhh.de

^bResearch, Development and Application Center (SARGEM), Sakarya University, Esentepe, Sakarya 54187, Turkey

^cDepartment of Metallurgical and Materials Engineering, Engineering Faculty, Sakarya University, Esentepe, Sakarya 54187, Turkey

^dInstitute of Soft Matter Modeling, Hamburg University of Technology, 21073 Hamburg, Germany

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4ta05571c>



endeavours in LOB catalysis have shifted focus towards developing mono-component bifunctional catalysts,^{16–18} aimed at facilitating both the ORR and OER during discharging and charging processes, respectively. Traditionally, these catalysts often involve metals or metal oxides coupled with carbonaceous materials like carbon, CNTs, or graphene.¹⁹ This coupling is intended to leverage synergistic effects, enhancing catalytic performance. However, such configurations not only promote the formation of lithium carbonate (LiCO₃) but also complicate electrode fabrication processes. Consequently, the exploration of mono-component bifunctional catalysts for LOBs stands as a critical pursuit to address these challenges and streamline electrode development, aiming for enhanced efficiency and simplified fabrication techniques.

Manganese dioxide (MnO₂) and titanium dioxide (TiO₂) are widely employed as cost-effective catalysts in LOBs. However, when used separately, manganese (Mn)²⁰ and titanium (Ti)²¹ facilitate redox reactions in battery processes in one way (either OER or ORR) because of their distinct oxidizing and reducing properties and the density of states concentrated at varying energy levels.²² Motivated by the specific electronic structure inherent in a MnTiO₃ perovskite, encompassing a broad spectrum of the density of states near the Fermi energy level owing to the simultaneous presence of both Mn and Ti constituents,²³ MnTiO₃ perovskite is introduced as an electrocatalyst in (LOBs) in this study. In light of its outstanding catalytic performance in diverse applications, including water purification,²⁴ oxidative degradation,²⁵ and photocatalytic activities,²⁶ this study constitutes the first systematic evaluation of the electrochemical performance of MnTiO₃ in LOBs. The analysis was conducted through a combination of experimental techniques, including Energy Dispersive Spectroscopy (EDS) and X-ray diffraction (XRD), alongside computational studies based on Density Functional Theory (DFT), providing a comprehensive understanding the electrochemical behaviour of MnTiO₃ as a carbon-free electrocatalyst in LOBs.

2 Computational and experimental methods

2.1 First principles calculations

Density functional theory (DFT) calculations were performed using the Vienna *Ab initio* Simulation Package (VASP, version 5.4.4).^{27–31} The calculations were carried out using the Perdew–Burke–Ernzerhof (PBE) parametrization³² to describe the exchange–correlation functional. The pseudopotentials are obtained by the projector augmented wave (PAW) method.^{33,34} The states in the electronic configurations 3p4s3d, 3p4s3d, 2s2p, and 1s2s for Ti, Mn, O, and Li, respectively, were treated as valence states.

Using generalized gradient approximation (GGA) functionals like PBE often underestimates band gaps for transition metal including compounds. For a better description of the band gap, an effective Hubbard $U_{\text{H,eff}}$ parameter following the Dudarev approach³⁵ was used to account for the localization of the Mn *d*-electrons. In this study, we examined the band gap of MnTiO₃

using several $U_{\text{H,eff}}$ values, namely 2, 2.5, 3, 3.5, and 4 eV. $U_{\text{H,eff}} = 3$ eV yields a band gap of 1.56 eV in excellent agreement with our experimentally determined band gap of 1.58 eV as illustrated in Fig. S1.†

An energy cutoff of 650 eV was used for the plane-wave basis. For slab calculations, a Gamma-centered $5 \times 4 \times 1$ *k*-point mesh was employed for sampling the Brillouin zone for the slab of the surface stability calculations. For ORR/OER calculations, the slab was doubled in the *x*-direction, and the *k*-points were tested to be converged for $3 \times 4 \times 1$. Ionic relaxations were performed until the convergence criterion of 0.01 eV Å⁻¹ in the forces was reached while for electronic self consistency cycles an energy convergence criterion of 10⁻⁶ eV was used. To avoid interactions between periodic images of the slab, an 18 Å thick vacuum region in the *c*-direction (perpendicular to the surface) was added to the system.

The adsorption energies ($E_{\text{adsorption}}$) between the reaction intermediates (adsorbate) and substrate were calculated as,³⁶

$$E_{\text{adsorption}} = E_{\text{sub/ads}} - E_{\text{sub}} - E_{\text{ads}}, \quad (1)$$

where $E_{\text{sub/ads}}$, E_{sub} , and E_{ads} are the total energies of the optimized substrate/adsorbate system, the clean substrate, and the adsorbate in the structure, respectively. The adsorbate atom or molecule will be referred to by (*).

In the calculations of free energy profiles for ORR/OER reaction, we adopted the Gibbs free energy difference (ΔG) between two consecutive steps for ORR as:

$$\Delta G = \Delta E_{\text{total}} + \Delta E_{\text{ZPE}} - T\Delta S - eU, \quad (2)$$

where ΔE_{total} is the DFT total energy difference between two systems, ΔE_{ZPE} is the zero-point energy differences, and $T\Delta S$ is entropy changes at the temperature $T = 298.15$ K and pressure $p = 1$ bar in each step. The term eU with the electron charge e has been added to show the effect of a potential U on electrons.^{37,38} The theoretical overpotential for the ORR η_{ORR} and the OER η_{OER} was obtained by

$$\eta_{\text{ORR}} = U_{\text{eq}} - U_{\text{DC}} \quad (3)$$

$$\eta_{\text{OER}} = U_{\text{eq}} - U_{\text{C}} \quad (4)$$

where U_{eq} refers to the equilibrium potential at which the change of free energy for the whole process is zero. U_{DC} represents the maximum voltage at which the free energy goes downhill in each step of the ORR process. U_{C} denotes the minimum voltage at which the free energy goes downhill in each step of the OER process.³⁹

2.2 Synthesis of MnTiO₃

MnTiO₃ powders were synthesized by the sol–gel method according to M. Enhessari *et al.*⁴⁰ Initially, 0.4 mol of stearic acid were liquefied at a temperature of 73 °C. Subsequently, 0.1 mol of manganese acetate were dissolved in the liquid stearic acid. The temperature was kept at 73 °C until a dark brown transparent solution was obtained. Following this, 0.1 mol of



stoichiometric tetrabutyl titanate was added to the solution and mixed at 73 °C until a uniform brown colloidal solution was formed. The solution was then left to naturally cool down to room temperature and subsequently dried in an oven for 12 hours. Finally, the gel underwent calcination at 1000 °C in four steps, to yield nanocrystallites of MnTiO₃. The first step involved heating the dried gel at a rate of 3 °C min⁻¹ up to 470 °C; this was followed by holding the temperature constant at 470 °C for 40 minutes in the second step. Subsequently, the temperature was increased up to 910 °C, and in the final step, the temperature was sustained at 910 °C for 2 h under ambient air conditions. Oxidation and crystallization temperatures were determined according to the Differential Thermal Analysis (DTA) measurement as given in ESI Fig. S2.†

2.3 Electrochemical characterization

The electrochemical assessment was conducted utilizing a Swagelok-type battery configuration, comprising a lithium foil anode with a carbon-free composite cathode. The electrolyte employed was 1.5 M LiTFSI (lithium trifluoromethanesulfonate) dissolved in TEGDME (tetraethylene glycol dimethyl ether), while a glassy fiber served as the separator. The fabrication process of the carbon-free cathode involved the preparation of a composite powder consisting of MnTiO₃, mixed with polyvinylidene fluoride (PVDF) in 1-methyl-2-pyrrolidone (NMP), without any conducting carbon. The composite was composed of 90 wt% MnTiO₃ and 10 wt% binder. Subsequently, the mixture was cast onto a nickel foam current collector and dried at 120 °C for 12 h. The mass loading of each cathode, following the drying process, was 0.5 mg cm⁻². The batteries were assembled inside an argon-filled glove box (H₂O content <0.1 ppm and O₂ content <0.1 ppm, manufactured by MBraun, Germany) to ensure an inert atmosphere. The charge–discharge tests were conducted at a current density of 0.1 mA cm⁻², facilitated by a continuous flow of pure O₂ gas at a rate of 8

ml min⁻¹, maintained at a pressure of 1.0 atm and a temperature of 25 °C in the voltage range of 2.0 V to 4.5 V. Additionally, cyclic voltammetry (CV) analysis was performed in the same voltage range, employing a scan rate of 0.1 mV s⁻¹ to elucidate the electrochemical reactions and their associated voltage profiles within the Li//MnTiO₃ cells. All electrochemical evaluations were executed utilizing a Gamry ref. 1000 potentiostat-galvanostat device under ambient temperature.

3 Results and discussion

3.1 Morphology, crystal, and electronic structure of MnTiO₃

A schematic route for the synthesis of MnTiO₃ powders is given in Fig. 1a. As shown in Fig. 1b, irregular shaped chunks with different sizes were observed at low magnifications. At higher magnifications, the powders were observed to be in nano-sized irregular shaped particles that were sintered to each others due to high temperature required for the synthesis.

Fig. 2a illustrates the optimized crystal structure of MnTiO₃. The structure consists of MnO₆ and TiO₆ octahedrons with corner-sharing oxygen atoms. Within this crystal lattice, Ti and Mn atoms occupy the 6a Wyckoff positions, while O atoms are located at the 18b Wyckoff positions. The DFT-optimized lattice parameters were obtained as $a = b = 5.23$ Å and $c = 13.98$ Å.

Synthesized powders were characterized *via* X-ray diffraction (XRD) analysis to determine their phase composition (Fig. 2b). Our results indicate that the synthesized powders can be indexed within the JCPDS 29-0902 (PDF#29-0902) reference card, exhibiting a trigonal crystal system with the *R3c* space group. Further analysis was performed by Raman technique to characterize the atomic local environment of the structure as given in Fig. S3.† The UV-vis absorption spectrum was recorded to obtain the band gap E_g , which can be obtained using the following equation;

$$(\alpha h\nu)^n = K(h\nu - E_g) \quad (5)$$



Fig. 1 (a) Schematic synthesis route for the synthesis of MnTiO₃ powders, (b) low magnification, and (c) high magnification FESEM images for the MnTiO₃ powders.



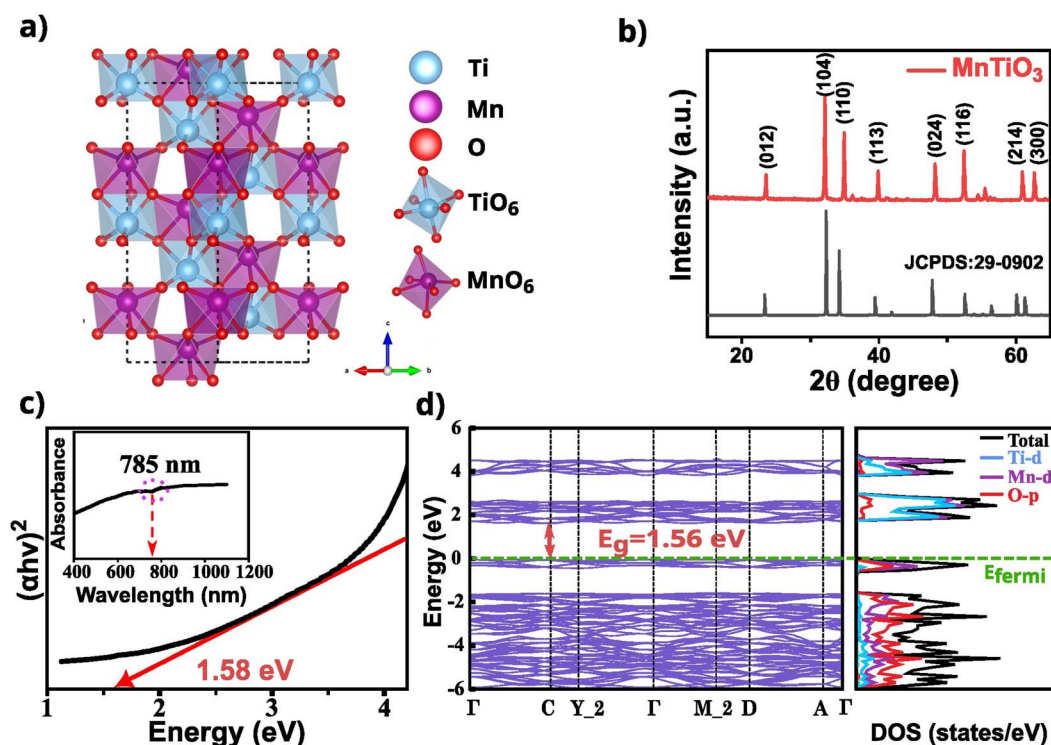


Fig. 2 (a) The crystal structure of MnTiO_3 with a trigonal unit cell. (b) The XRD patterns of MnTiO_3 , (c) $(\alpha h\nu)^2 - h\nu$ curves of the MnTiO_3 powders, and the corresponding absorption spectra in the inset obtained from the UV-visible measurements, (d) the calculated band structure, band gap (E_g), and the orbital-projected density of states (PDOS) from DFT.

where $h\nu$ is the photoenergy, α is the absorption coefficient, K is a constant related to the material, and n is either 2 for a direct transition or $1/2$ for an indirect transition. The experimental band gap was measured to be 1.58 eV as shown in Fig. 2c. A direct band gap of 1.56 eV was obtained from DFT calculations with a Hubbard parameter of $U_{\text{H,eff}} = 3$ eV which is in well alignment with the experimental measurement. The electronic structure of MnTiO_3 was further analyzed using the Projected Density of States (PDOS). The Fermi level was chosen as the reference energy and set to 0 eV. The conduction band (CB) primarily derives from the contribution of the Ti-d and Mn-d orbitals, whereas the valence band predominantly stems from the Mn-d and O-p orbitals with a smaller contribution of Ti-d orbitals (Fig. 2d). The existence of electronic states spanning a wide range in energy due to the coexistence of titanium and manganese is expected to promote both OER and ORR reactions simultaneously which take place at different energy levels. Taking into account the influence of orbital states in close proximity to the Fermi level on oxidation-reduction (ORR and OER) activities, it is anticipated that energy levels located on either side of the Fermi level will contribute to the bifunctional ORR and OER capabilities of the resulting material.

3.2 Surface reactivity

The obtained XRD results, depicted in Fig. 2b, reveal distinct peaks denoted by indices (104) and (110) at 38.7° and 39.8° , respectively. Notably, these peaks exhibit a pronounced degree

of crystallinity and are indexed to the crystal structure of MnTiO_3 . Ribeiro *et al.*⁴¹ used DFT calculations to investigate the stoichiometric polar and nonpolar surfaces of MnTiO_3 , specifically the surfaces indexed as (110), (012), (101), (100), (001), and (111). They determined that the (110) surface exhibited the highest degree of stability. Since our XRD analysis reveals the most pronounced reflections at the (104) plane, we conduct a comparative analysis between the (104) and (110) surfaces, considering their different terminations. For the MnTiO_3 -(104) surface and the MnTiO_3 -(110) surface, the atomic layers follow a stacking sequence of (...O2-O1-Mn-TiO...) and (...O2-O1-MnTi...), respectively, as illustrated in Fig. 3.

A comprehensive examination of surface stability for the (104) and (110) surfaces is presented in Section S3 of the ESI.† Our investigation (Fig. S4†) reveals that the (104) surface is significantly more stable than the (110) surface. Additionally, (Fig. S5†) elucidates that the (O1 and TiO) terminations exhibit the highest stability among the (104) terminations.

The cathode of a battery operates in an oxygen-rich environment. In order to examine the stability of the MnTiO_3 -(104) surface under these conditions, we have conducted an analysis of the capacity of the two MnTiO_3 -(104) terminations, namely O1 and TiO, to interact with oxygen atoms. This investigation was carried out by calculating the adsorption energy of oxygen atoms at multiple sites. The oxygen atoms are placed in high-symmetry points on the top of the two distinct MnTiO_3 (104) terminations, O1 and TiO, as shown in Fig. 4. Subsequently, the



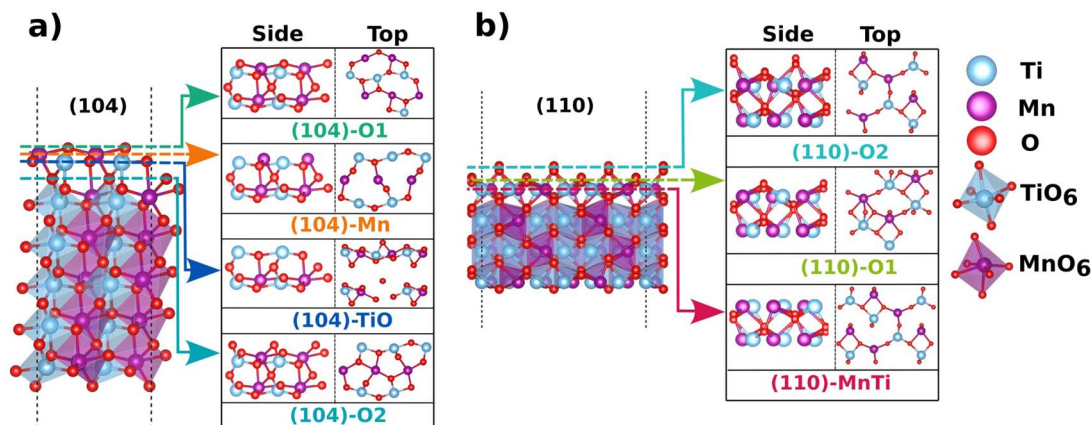


Fig. 3 Top and side views of MnTiO_3 slabs showing (a) a (104) surface with O1, O2, Mn and TiO terminations. (b) a (110) surface with O1, O2, and MnTi terminations.

structures were optimized using DFT calculations to minimize the forces acting on the atoms. Our results for the O1 termination show that regardless of the initial position after relaxation all oxygen atoms were adsorbed at a bridge position between Ti and Mn with an adsorption energy of -4.4 eV. This indicates the strong preference of this adsorption site. The adsorption results in the formation of bonds between the oxygen atom and the titanium and manganese atoms, with bond lengths of 1.72 Å and 2.06 Å, respectively. Similarly, for the TiO termination, all initial adsorption structures resulted in the same final geometry after relaxation. On the TiO termination, O atoms prefer the proximity of Ti leading to the formation of O-Ti bonds with a bond length of 1.63 Å and with an adsorption energy of -3.6 eV. Considering our results, it was found that oxygen adsorption energies are considerably more negative on the O1 termination compared to the TiO termination.

3.3 ORR/OER mechanism

The oxygen reduction reaction (ORR) and oxygen evolution reaction (OER) processes in a nonaqueous Li-O_2 battery can proceed in two reaction paths, namely the $2e^-$ and the $4e^-$ pathways.⁴² In the $2e^-$ pathway, oxygen is reduced to Li_2O_2 via the reaction $2(\text{Li}^+ + e^-) + \text{O}_2 \rightarrow \text{Li}_2\text{O}_2$ with $U_{\text{eq}} = 2.96$ V. In contrast in the $4e^-$ path, oxygen is fully reduced to Li_2O via $4(\text{Li}^+ + e^-) + \text{O}_2 \rightarrow 2\text{Li}_2\text{O}$ with $U_{\text{eq}} = 2.91$ V.

The adsorption of O_2 and Li species on the cathode surface can significantly alter the ORR pathway, $2e^-$ or $4e^-$, and the reaction kinetics.^{43,44} As illustrated in Fig. 5 and Table S1† the adsorption energy per adsorbate (calculated according to eqn (1)) for O_2 on the investigated cathode surface is -2.88 eV, which is significantly more negative than that for Li atoms, which is -1.50 eV. This higher energy gain from O_2 adsorption indicates that the initial lithiation process (involving the formation of Li_2O_2) follows the $2e^-$ pathway. In detail, the ORR process starts with the adsorption of an O_2 molecule at the clean surface, followed by the reaction of a Li-ion with O_2 to form LiO_2 . A subsequent reaction of another Li-ion with LiO_2 yields Li_2O_2 , that is the final product of the first lithiation process. Fig. 5 and Table S1† shows that all intermediates have negative adsorption energies, indicating thermodynamically favorable adsorption processes.

To elucidate the second lithiation process, which involves the formation of Li_4O_4 , the analysis was started with the independent adsorption of O_2 and Li on the Li_2O_2^* -covered surface obtained from the first lithiation. The adsorption energy of -2.83 eV for O_2 is considerably more negative than the -1.18 eV for Li atom. These adsorption energies were also calculated using eqn (1).

The significantly higher adsorption energy of O_2 suggests that the second lithiation also follows the $2e^-$ pathway. The reaction sequence for the second lithiation can be summarized as follows: $\text{Li}_2\text{O}_2^* + \text{O}_2 \rightarrow \text{Li}_2\text{O}_4^* + \text{Li} + e^- \rightarrow \text{Li}_3\text{O}_4^* + \text{Li} + e^- \rightarrow \text{Li}_4\text{O}_4^*$. The high adsorption energy of O_2 plays a crucial role in facilitating electron transfer during the electrochemical

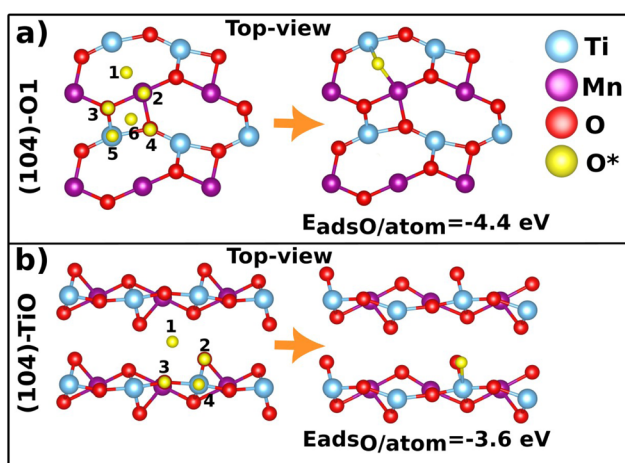


Fig. 4 Top views of (a) O1 and (b) TiO terminations of MnTiO_3 (104). In the left, the tested initial adsorption sites for oxygen atoms are shown, while in the right, the geometry after structural optimization is presented. The adsorption energies are given per adsorbed oxygen atom.



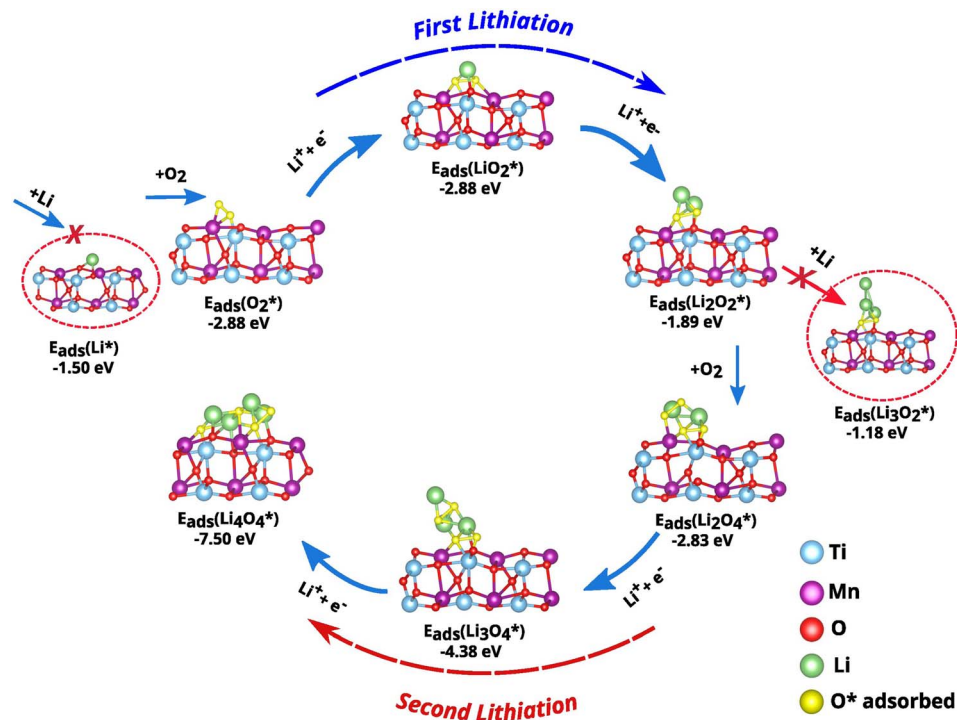


Fig. 5 The adsorption energies of ORR intermediates during the first and second lithiation.

reaction, thereby accelerating the reaction kinetics. Moreover, throughout the discharge process, the adsorption energy of intermediate LiO_2 increases (from -2.88 eV in the first lithiation to -4.38 eV in the second lithiation), promoting the growth of discharge products through a surface reaction pathway.³⁹

Fig. 6a and b display the simulated Gibbs free energy profiles for ORR/OER corresponding to the first and second lithiation. At zero potential ($U = 0$ V), all the reaction steps are downhill, and exhibit favourable thermodynamics, underscoring the spontaneity of Li_2O_2 and Li_4O_4 formation. Utilizing the data from the Gibbs free energy diagrams, we calculated the cathodic

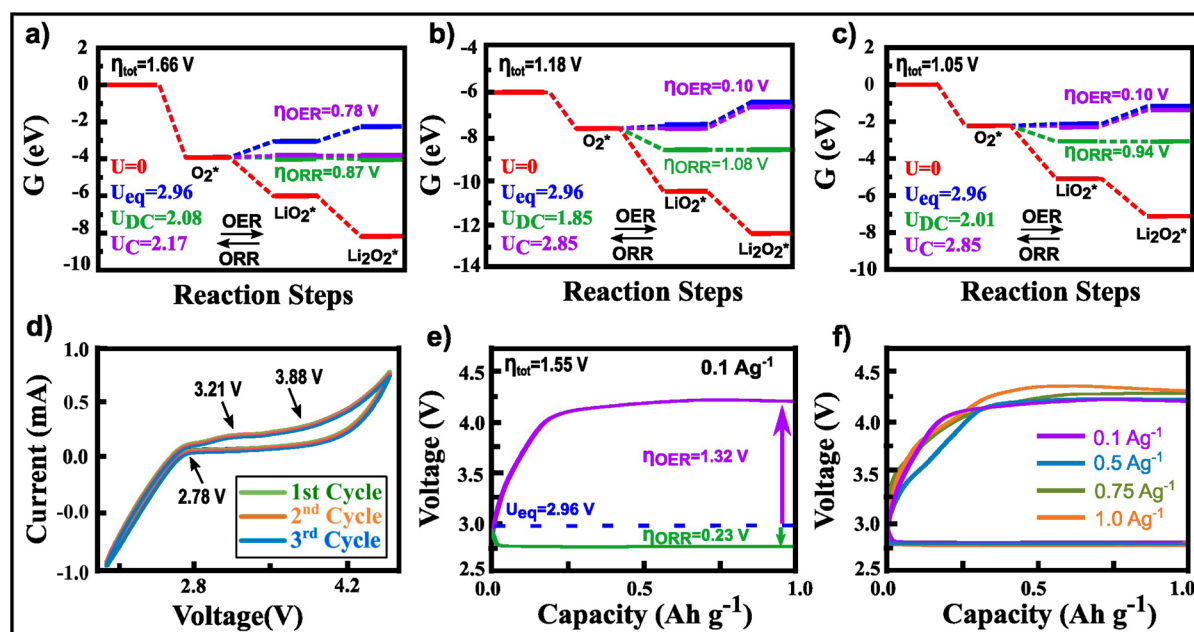


Fig. 6 Gibbs free energy diagrams for (a) the first lithiation and (b) the second lithiation for MnTiO_3 (104) with O1 termination. (c) Gibbs free energy diagrams after reducing the surface unit cell area of MnTiO_3 . (d) Cyclic voltammety curves illustrating electrochemical reactions. (e) Discharging and charging curves with a specific capacity limited to 1 Ah g^{-1} and a current density of 0.1 A g^{-1} . (f) Rate performance at various current densities ranging from 0.1 A g^{-1} to 1 A g^{-1} .



discharge (U_{DC}) and charge (U_C) potentials for the first and second lithiation as shown in Fig. 6(a) and (b). According to eqn (3) and (4), the total overpotentials η were calculated as 1.66 V and 1.18 V for the first and second lithiations, respectively.

Furthermore, we examined the first lithiation process on the $MnTiO_3$ surface by reducing the surface supercell along the X -direction until a single Li_2O_2 molecule fully covered the newly formed surface. The calculated free energy diagram is shown in Fig. 6c. The first lithiation process yields nearly identical values for U_{DC} , U_C , and the total overpotential as the second lithiation on the original surface. Additionally, the profiles for ORR/OER closely resemble that of the second lithiation in Fig. 6(b) and (c). Consequently, this observation suggests that, instead of employing a large surface cell area, reducing the surface area can yield equivalent results. Thereby optimizing the computational resources in the analysis of the lithiation process.

The cyclic voltammetry (CV) profile of $MnTiO_3$ shown in Fig. 6d exhibits a prominent cathode peak between 2.0 V and 2.8 V, indicating the formation of discharge products, specifically Li_2O_2 or LiO_2 . During the anodic scan, two oxidation peaks

are observed: one at 3.21 V and another starting at 3.88 V, where the peak at 3.21 V correspond to the oxidation of amorphous Li_2O_2 , particularly when the catalyst has a strong affinity for LiO_2 intermediates.^{45,46} The higher peak starting at 3.88 V is associated with the decomposition of bulk Li_2O_2 , which requires a higher dissociation potential.⁴⁷ The results highlight the importance of considering the role of the $MnTiO_3$ cathode in the formation and decomposition of Li_2O_2 , which can significantly affect charge–discharge characteristics of the battery.

To determine the voltages associated with the OER and ORR, the $MnTiO_3$ cathode underwent charging and discharging cycles with a 1 A h g^{-1} specific capacity limit, between the potentials from 2 V to 4.5 V, while operating at a current density of 0.1 A g^{-1} (Fig. 6e). The total overpotential was determined to be 1.55 V in close agreement with the predictions from DFT, which yielded values of 1.66 V and 1.18 V for the first and second lithiation, respectively.

Those results underscore the effectiveness of $MnTiO_3$ in facilitating the oxidation of Li_2O_2 during the charging phase.

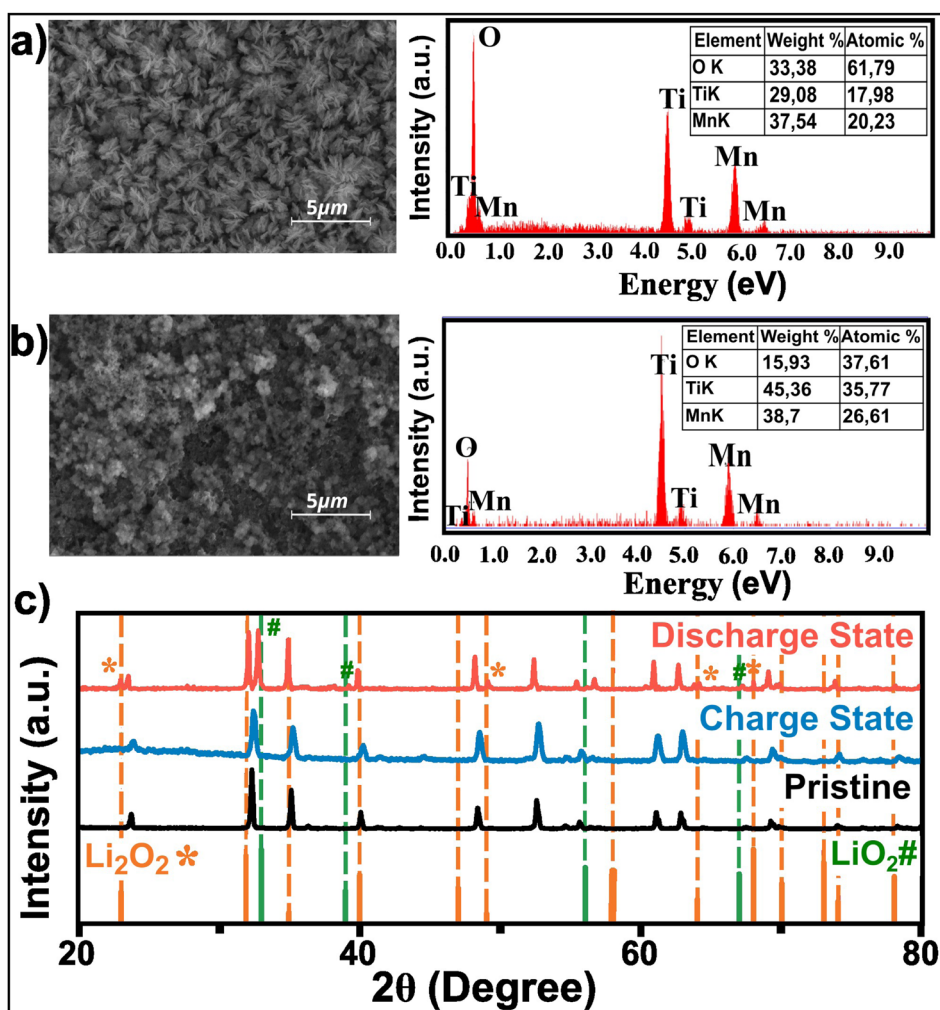


Fig. 7 FESEM images and EDS spectra of $MnTiO_3$ cathode after (a) first discharge and (b) first charge, and the ex situ. (c) XRD patterns after charge and discharge.



Moreover, the charge overpotentials of the MnTiO₃ cathode remained relatively constant for different current densities, ranging from 0.1 to 1 A g⁻¹ (Fig. 6f).

In order to gain deeper insights into the superior characteristics exhibited by Li–O₂ batteries incorporating the MnTiO₃ cathode, the configurations of discharge products were further analyzed through field-emission scanning electron microscopy (FESEM), energy-dispersive X-ray spectroscopy (EDS), and X-ray diffraction (XRD) analyses. Following the initial discharge, the FESEM images revealed the presence of discharge Li₂O₂ products (Fig. 7), confirming the adherence of the electrochemical reaction to the 2e⁻ pathway. Furthermore, these images demonstrate that the majority of discharge byproducts decomposed after the charging process (Fig. 7). The EDS spectrum of the electrode following discharge indicates a significant increase in oxygen content of 61.79%, resulting from the formation of Li₂O₂. Nevertheless, the oxygen ratio experienced a notable decline following the decomposition of Li₂O₂ during the charging phase (Fig. 7a and b). Subsequent XRD analysis corroborated the existence of LiO₂ and Li₂O₂ products, consistent with the reference patterns of JCPD-12-0254 and JCPD-09-0355, respectively (Fig. 7c). After the charging process, those characteristic peaks diminished as the discharge products decomposed, providing further evidence of the reversible nature of this process. Importantly, the XRD pattern did not reveal any additional reactions, emphasizing the effective role of the MnTiO₃ perovskite catalyst in preventing the formation of undesired byproducts.

4 Conclusions

In this work, a MnTiO₃ catalyst was used as a potential oxygen electrode for a Li–O₂ battery for the first time. Both *ab initio* calculations and experimental techniques were employed to investigate the electrochemical behavior. The electronic structure calculations demonstrate that the simultaneous presence of Mn and Ti orbitals near the Fermi level can facilitate the electron transport, thereby promoting ORR and OER reactions. Furthermore, our research elucidated the impact of the reactive surface area on the computational prediction of the electrochemical reactions. DFT adsorption energy and XRD analysis confirm that the electrochemical reaction proceeds following the 2-electron discharge pathway. As a result, the carbon-free MnTiO₃ cathode exhibits excellent electrochemical performance and a low overpotential of 1.55 V, with a small increase in overpotential at elevated current densities up to 1 A g⁻¹.

Data availability

The data supporting this article have been included as part of the ESI.†

Author contributions

Doaa Aasef Ahmed: conceptualization, formal analysis, investigation, methodology, project administration, visualization, writing – original draft, writing – review and editing, data

curation. Mustafa Çelik: visualization, investigation, methodology, formal analysis, writing – original draft, writing – review and editing. Wernfried Mayr-Schmölzer: methodology, supervision, writing – review and editing. Abdulkadir Kızılaslan: data curation, conceptualization, supervision, writing – original draft, writing – review and editing. Gregor B. Vonbunfeldbauer: conceptualization, funding acquisition, methodology, project administration, resources, supervision, validation, writing – review and editing.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Funded by the Deutsche Forschungsgemeinschaft (DFG, German Research Foundation) – Project number 192346071–SFB 986. Doaa Aasef Ahmed was co-funded by the European Union within the Erasmus+ programme. The numerical calculations were partially performed at TUBITAK ULAKBIM, High Performance and Grid Computing Center (TRUBA resources).

References

- H. Hou, Y. Cong, Q. Zhu, Z. Geng, X. Wang, Z. Shao, X. Wu, K. Huang and S. Feng, *Chem. Eng. J.*, 2022, **448**, 137684.
- X. Mu, C. Xia, B. Gao, S. Guo, X. Zhang, J. He, Y. Wang, H. Dong, P. He and H. Zhou, *Energy Storage Mater.*, 2021, **41**, 650–655.
- Q. Qiu, Z.-Z. Pan, P. Yao, J. Yuan, C. Xia, Y. Zhao and Y. Li, *Chem. Eng. J.*, 2023, **452**, 139608.
- J. Kang, J.-S. Yu and B. Han, *J. Phys. Chem. Lett.*, 2016, **7**, 2803–2808.
- M.-C. Sung, G.-H. Lee and D.-W. Kim, *InfoMat*, 2021, **3**, 1295–1310.
- J. G. Kim, Y. Kim, Y. Noh, S. Lee, Y. Kim and W. B. Kim, *ACS Appl. Mater. Interfaces*, 2018, **10**, 5429–5439.
- C. Xia, M. Waletzko, L. Chen, K. Pepller, P. J. Klar and J. Janek, *ACS Appl. Mater. Interfaces*, 2014, **6**, 12083–12092.
- D. Zhai, H.-H. Wang, J. Yang, K. C. Lau, K. Li, K. Amine and L. A. Curtiss, *J. Am. Chem. Soc.*, 2013, **135**, 15364–15372.
- Q. Li, P. Xu, W. Gao, S. Ma, G. Zhang, R. Cao, J. Cho, H.-L. Wang and G. Wu, *Adv. Mater.*, 2013, **26**, 1378–1386.
- B. D. McCloskey, A. Speidel, R. Scheffler, D. C. Miller, V. Viswanathan, J. S. Hummelshøj, J. K. Nørskov and A. C. Luntz, *J. Phys. Chem. Lett.*, 2012, **3**, 997–1001.
- D. Cao, S. Zhang, F. Yu, Y. Wu and Y. Chen, *Batteries Supercaps*, 2019, **2**, 428–439.
- C. Xu, B. M. Gallant, P. U. Wunderlich, T. Lohmann and J. R. Greer, *ACS Nano*, 2015, **9**, 5876–5883.
- Y.-C. Lu, Z. Xu, H. A. Gasteiger, S. Chen, K. Hamad-Schifferli and Y. Shao-Horn, *J. Am. Chem. Soc.*, 2010, **132**, 12170–12171.
- Y. Zhang, X. Li, M. Zhang, S. Liao, P. Dong, J. Xiao, Y. Zhang and X. Zeng, *Ceram. Int.*, 2017, **43**, 14082–14089.
- K. Li, H. Dong, Y. Wang, Y. Yin and S. Yang, *J. Colloid Interface Sci.*, 2020, **579**, 448–454.



- 16 H. Wu, W. Sun, J. Shen, Z. Mao, H. Wang, H. Cai, Z. Wang and K. Sun, *ACS Sustainable Chem. Eng.*, 2018, **6**, 15180–15190.
- 17 Y. J. Lee, D. H. Kim, T.-G. Kang, Y. Ko, K. Kang and Y. J. Lee, *Chem. Mater.*, 2017, **29**, 10542–10550.
- 18 Y. Xia, X. Yu, Y. Xu, X. Fan, B. Gao, C. Jiang, M. Zhang, X. Huang, H. Gong, J. He and T. Wang, *J. Mater. Chem. A*, 2023, **11**, 24918–24927.
- 19 S. Zhang, Y. Wang, D. Li, Z. Kang, F. Dong, H. Xie and J. Liu, *J. Alloys Compd.*, 2020, **825**, 154054.
- 20 S. H. Kim, Y. J. Lee, D. H. Kim and Y. J. Lee, *ACS Appl. Mater. Interfaces*, 2017, **10**, 660–667.
- 21 S. Xu, Q. Zhu, J. Long, H. Wang, X. Xie, K. Wang and J. Chen, *Adv. Funct. Mater.*, 2016, **26**, 1365–1374.
- 22 T. G. Yun, Y. Heo, H. Bin Bae and S.-Y. Chung, *Nat. Commun.*, 2021, **12**, year.
- 23 K. Yoshimatsu, H. Mashiko, N. Umezawa, K. Horiba, H. Kumigashira and A. Ohtomo, *J. Phys. Chem. C*, 2017, **121**, 18717–18724.
- 24 D. Wang, H. Xu, J. Ma, X. Lu, J. Qi and S. Song, *ACS Appl. Mater. Interfaces*, 2018, **10**, 31631–31640.
- 25 H. Wang, Q. Gao, H. Li, M. Gao, B. Han, K. Xia and C.-G. Zhou, *ACS Appl. Nano Mater.*, 2018, **1**, 2727–2738.
- 26 N. Kitchamsetti, P. N. Didwal, S. R. Mulani, M. S. Patil and R. S. Devan, *Heliyon*, 2021, **7**, e07297.
- 27 J. Hafner, *J. Comput. Chem.*, 2008, **29**, 2044–2078.
- 28 G. Kresse and J. Hafner, *Phys. Rev. B:Condens. Matter Mater. Phys.*, 1993, **47**, 558–561.
- 29 G. Kresse and J. Hafner, *Phys. Rev. B:Condens. Matter Mater. Phys.*, 1994, **49**, 14251–14269.
- 30 G. Kresse and J. Furthmüller, *Comput. Mater. Sci.*, 1996, **6**, 15–50.
- 31 G. Kresse and J. Furthmüller, *Phys. Rev. B:Condens. Matter Mater. Phys.*, 1996, **54**, 11169–11186.
- 32 J. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**, 3865–3868.
- 33 P. Blöchl, *Phys. Rev. B:Condens. Matter Mater. Phys.*, 1994, **50**, 17953–17979.
- 34 G. Kresse and D. Joubert, *Phys. Rev. B:Condens. Matter Mater. Phys.*, 1999, **59**, 1758–1775.
- 35 S. L. Dudarev, G. A. Botton, S. Y. Savrasov, C. J. Humphreys and A. P. Sutton, *Phys. Rev. B:Condens. Matter Mater. Phys.*, 1998, **57**, 1505–1509.
- 36 T. Zhu, C. Xia, B. Wu, J. Pan, H. Yang, W. Zhang and B. Y. Xia, *Appl. Catal., B*, 2024, **357**, 124315.
- 37 Y. Gong, W. Ding, Z. Li, R. Su, X. Zhang, J. Wang, J. Zhou, Z. Wang, Y. Gao, S. Li, P. Guan, Z. Wei and C. Sun, *ACS Catal.*, 2018, **8**, 4082–4090.
- 38 S. Ding, X. Yu, Z.-F. Ma and X. Yuan, *J. Mater. Chem. A*, 2021, **9**, 8160–8194.
- 39 Y. Cao, Z. Zhu, X. Li, J. Xi, D. J. Singh, L. Xi, J. Yang and W. Zhang, *J. Mater. Chem. A*, 2022, **10**, 11039–11045.
- 40 M. Enhessari, A. Parviz, E. Karamali and K. Ozaee, *J. Exp. Nanosci.*, 2012, **7**, 327–335.
- 41 R. A. Ribeiro, J. Andrés, E. Longo and S. R. Lazaro, *Appl. Surf. Sci.*, 2018, **452**, 463–472.
- 42 B. D. McCloskey, D. S. Bethune, R. M. Shelby, G. Girishkumar and A. C. Luntz, *J. Phys. Chem. Lett.*, 2011, **2**, 1161–1166.
- 43 F. Zheng, H. Dong, Y. Ji and Y. Li, *J. Power Sources*, 2019, **436**, 226845.
- 44 Y. Jing and Z. Zhou, *ACS Catal.*, 2015, **5**, 4309–4317.
- 45 K. R. Yoon, K. Shin, J. Park, S.-H. Cho, C. Kim, J.-W. Jung, J. Y. Cheong, H. R. Byon, H. M. Lee and I.-D. Kim, *ACS Nano*, 2017, **12**, 128–139.
- 46 Y. Dou, X.-G. Wang, D. Wang, Q. Zhang, C. Wang, G. Chen, Y. Wei and Z. Zhou, *Chem. Eng. J.*, 2021, **409**, 128145.
- 47 Y. Dou, R. Lian, Y. Zhang, Y. Zhao, G. Chen, Y. Wei and Z. Peng, *J. Mater. Chem. A*, 2018, **6**, 8595–8603.

