

Biological Activity and Bio-Sorption Properties of the Ti₂C Studied by Means of Zeta Potential and SEM

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The expanded Ti₂C MXene phase was synthesized from the commercial Ti₂AlC MAX phase using classical acidic aluminium extraction method. The individual expanded Ti₂C sheets were characterized by the diameters of between 26 and 61 nm and formed the specific network of slit-shaped pores in the size range of 3–5 nm. The BET specific surface area of expanded Ti₂C was ca. 42% higher in comparison with layered Ti₂AlC. Our investigations results revealed that both Ti₂AlC and Ti₂C phases did not influence negatively the gram-positive bacteria commonly occurring in the natural environment. Moreover, the slightly intensified growth of *Bacillus sp.* strain was observed in the vicinity of tested Ti₂AlC and Ti₂C. The SEM investigations of the preferential sites for bacteria adsorption indicated the presence of minor apoptosis for only *Bacillus sp.* strain, especially when the cells were located between individual sheets of the expanded Ti₂C. The zeta potential curves obtained for Ti₂AlC and Ti₂C in distilled water differed significantly in shape. The intensive peak towards zero zeta potential was observed for Ti₂C. Adsorption of bacteria cells on Ti₂C resulted in changing of its zeta potential to that of bacteria cells.

Keywords: Ti₂C, MXene, zeta potential, bio-activity, biocidal

1. INTRODUCTION

Recently, the layered early transition metal carbides and nitrides (called MAX phases) have been first introduced by Barsoum et al. [1]. Since that time, MAX phases have attracted the interest of academia and industry due to the fact that they allow to achieve a dramatic improvement in mechanical

properties of materials [2, 3]. Chemically expanded structures of early transition metal carbides, known as MXenes, are novel and poorly studied family of materials characterized by specific properties which can be described as characteristic for both metals and ceramics [4]. These unique properties made the applicability of MXenes very attractive in the high-tech area of science. It should be however noted, that the potential for the biological activity of MXenes have not yet been fully discovered, highly limiting the possibility of their application in many fields of bioscience and biotechnology. So far, only Wang *et al.* [5] discovered that MXenes exhibited effective immobilization of enzymes and biocompatibility for red-ox proteins. This indicates the promise for applications in electrochemical biosensors. However still, there are no reports on antimicrobial and bio-sorption properties of MXenes. The present study, for the first time, investigates the biological activity of Ti_2C , which is the expanded form of layered titanium carbide (Ti_2AlC) against gram-positive bacteria such as *Bacillus sp.*, *S. aureus* and *Sarcina*. The aim of this study was also to describe the preferential sites for adsorption of the bacteria cells using scanning electron microscope examinations as well as analysis of changes of zeta potential of expanded Ti_2C MXene as a result of bacteria cells adsorption.

2. EXPERIMENTAL

2.1. Synthesis procedure of the expanded Ti_2C (MXene)

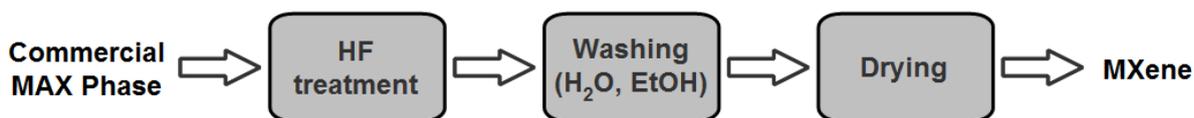


Figure 1. Flowchart of the technological stages of the synthesis process of Ti_2C MXene phase from Ti_2AlC commercial MAX phase.

The layered Ti_2AlC was purchased from Kanthal (the Sandvik AB group), Sweden. The composition of Ti_2AlC as declared by a manufacturer was as following: 5-10 wt% of carbon, 20-25 wt% of aluminium and balanced wt% of titanium.

The technological stages of the synthesis process of Ti_2C MXene phase from Ti_2AlC commercial MAX phase were presented in Fig. 1. The Ti_2AlC commercial MAX phase powder was immersed in 48% hydrofluoric acid (Sigma-Aldrich) for 24 h at room temperature. About 10 ml of hydrofluoric acid was used for 1 g of starting material. The resulting sediment was washed four times using deionized water and four times with technical grade ethanol. The solid product of the layered Ti_2C was dried overnight in room temperature.

2.2. Characterization of the morphology and structure

The morphologies of the surface of layered Ti_2AlC (MAX phase) and its expanded form - Ti_2C (MXene phase) were analyzed using scanning electron microscope (SEM) of LEO 1530, Zeiss, USA.

The microscope operated at 2.0kV. The tested powders were directly deposited on the surface of a carbon tape and subsequently coated with a thin carbon layer (BAL-TEC SCD 005 duster with a CEA 035 unit). Then, the obtained samples were subjected to the analysis of morphology. The elemental composition in layered Ti_2AlC and expanded Ti_2C was analyzed using energy dispersive X-ray spectroscopy (EDS, coupled with SEM) allowing to obtain the information on the elements content in a given region. The X-ray diffraction (XRD) was used for phase analysis at room temperature (Bruker D8 Advance) with Cu $K\alpha$ radiation ($\lambda=0.154056$ nm) and voltage of 40 kV (40 mA). The angle 2θ range was chosen from 15° to 120° . The step $\Delta 2\theta$ was 0.025 together with counting time of 3 s. Subsequently, the obtained XRD patterns were analysed using the software of Bruker EVA.

2.3. Analysis of physical properties and porous structure

The analysis of physical properties and porous structure of layered Ti_2AlC and expanded Ti_2C were examined using the physical nitrogen sorption isotherms $V=f(p/p^o)$ which relates to the volume of the adsorbed gaseous N_2 as a function of N_2 relative pressure. The isotherms were measured experimentally using a Quadrasorb-SI device from Quantachrome Instruments, USA. Before the measurements, the samples were degassed in vacuum at a temperature of $300^\circ C$ for 24 hours. The isotherms of adsorption and desorption of gaseous N_2 on the surface of tested samples were recorded for the entire relative pressure range (p/p^o from 0 to 1) at a temperature of liquid nitrogen bath i.e. $-195.8^\circ C$. The specific surface area, S_{BET} was obtained using the method of Brunauer, Emmett, and Teller (BET) within the relative pressure range of 0,05-0,35. The shape of the obtained isotherms was also analysed. It represents the shapes of pores that are present in the tested material. The total volume of the pores, V_{pores} and the surface area occupied by the pores S_{pores} was determined using the method of Barret, Joyner, and Halenda (BJH). The assumption related to the BJH method is that, at the relative pressure $< 0,4$, during the capillary condensation in pores the only effect that is observed together with a further increase of the pressure is the thickening of the N_2 mono-layer. This mono-layer is formed on the pore walls surface. The average pore size, D_{BJH} was estimated from the distributions of the pore sizes.

2.4. Analysis of potential antimicrobial properties

The bioactivities of layered Ti_2AlC and expanded Ti_2C were tested qualitatively (agar diffusion method) using the selected gram-positive bacteria strains of *Bacillus sp.*, *Staphylococcus aureus* and *Sarcina* which were taken from the private collection of microorganisms of the Biology Department, Faculty of Environmental Engineering, Warsaw University of Technology. On the surface of Petri dishes with solid nutrient agar (Merck) there were inoculated lines of microorganisms. The tested powders were then located in the area of the spread lines and the samples were then incubated for 48h at different temperatures i.e. $26^\circ C$ or $37^\circ C$, which were dependent on the bacterial species. After incubation, the Petri plates were photographed which allowed for observations of the growth inhibition zones.

2.5. Analysis of preferential sites for bacteria adsorption

The qualitative analysis of preferential sites for adsorption of bacteria cells on the surface of expanded Ti_2C was investigated using scanning electron microscope examination with the same bacteria strains used as in agar diffusion method i.e.: *Staphylococcus aureus*, *Bacillus sp.*, and *Sarcina*. Before analysis, microorganisms were spread on the nutrient agar (Merck) and cultivated for 24 h at $37^\circ C$. Drinking water was filtered before use with $0,2 \mu m$ sterile syringe filter (Whatman). Then, the bacterial cells were harvested from Petri plates and transferred to the drinking water due to obtain cells density of about 10^5-10^6 cells/ml. Subsequently, the such prepared bacteria stock suspension was incubated at $37^\circ C$ for 5 minutes. Subsequently, $200 \mu L$ of each suspensions of bacteria were transferred to 5×10^{-5} g/ $200 \mu L$ suspensions of Ti_2C powder in drinking water. The such prepared mixtures of Ti_2C powder and bacteria cells were incubated in $37^\circ C$ for 30 minutes. After that time, $100 \mu L$ of 20 wt% water solution of glutaraldehyde was added to each suspension which was then left for spontaneous sedimentation. After 1 hour, the clarified solution from above each of the the sedimented powder was removed and changed for the $500 \mu L$ of the first of series of graded concentrations of ethanol (20 wt%). Subsequently, 40, 60, 80 and 98 wt% ethanol was used for rinsing the powder samples. The samples were then dried and deposited on a carbon tape. Next, the samples were coated with a thin carbon layer and were subjected to SEM analysis (LEO 1530, Zeiss, USA). The SEM examinations were conducted under 3.0kV.

2.6. Zeta potential analysis

The zeta potential (ζ) of the layered Ti_2AlC and expanded Ti_2C powder in the distilled and drinking water environment was analyzed using Zetasizer Nano ZS (Malvern Instruments), equipped with an MPT-2 automatic titrator and a titration media degasser. Drinking water was purified as described in 2.5. Then, the zeta potential was determined using Smoluchowski's rule based on the electrophoretic mobility (μ).

The ζ analysis was first performed both in distilled water as well as drinking water environment in the pH range from 4 to 10 which enabled the determination of IEP value (isoelectric point) for each powder in the specific environment. The test samples were $0,5 \text{ mg}/10 \text{ mL}$ suspensions of Ti_2C and Ti_2AlC . The Ti_2AlC was used as a non-expanded reference material. Before the measurement, the samples were mixed for 5 seconds and then transferred to a polystyrene U-shaped cells (Malvern Instruments). In the next step, the cells were stabilized at $25^\circ C$ for 5 seconds. Finally, every measurement was conducted at $25^\circ C$, and repeated 10 times. The sodium hydroxide (0.1 M NaOH) and hydrochloric acid (0.1 M HCl) were used as the titration media.

Due to allow the observation of the changes of electrical charge which is formed on the surface of the analyzed powders in presence of bacterial cells, the measurements of the zeta potential were conducted in drinking water environment, separately for: (i) $0,5 \text{ mg}/10 \text{ mL}$ suspensions of Ti_2AlC and Ti_2C powders; (ii) $100 \mu L/10 \text{ mL}$ suspensions of pure bacteria cells; (iii) mixed suspensions of powders and bacteria cells. The samples of pure bacteria cells were stabilized before measurement at $25^\circ C$ for 1 minute. The mixed suspensions of powders and bacteria were incubated at $37^\circ C$ for 5

minutes with 100 rpm stirring rate. Then, the samples were stabilized at a temperature of 25 °C for 1 minute and the electrophoretic mobility was measured. It should be noted, that each of the zeta potential measurements was an average of 100 repetitions.

3. RESULTS AND DISCUSSION

The morphologies of layered Ti_2AlC and expanded Ti_2C were examined using a scanning electron microscope (SEM) and were presented in Fig. 2.

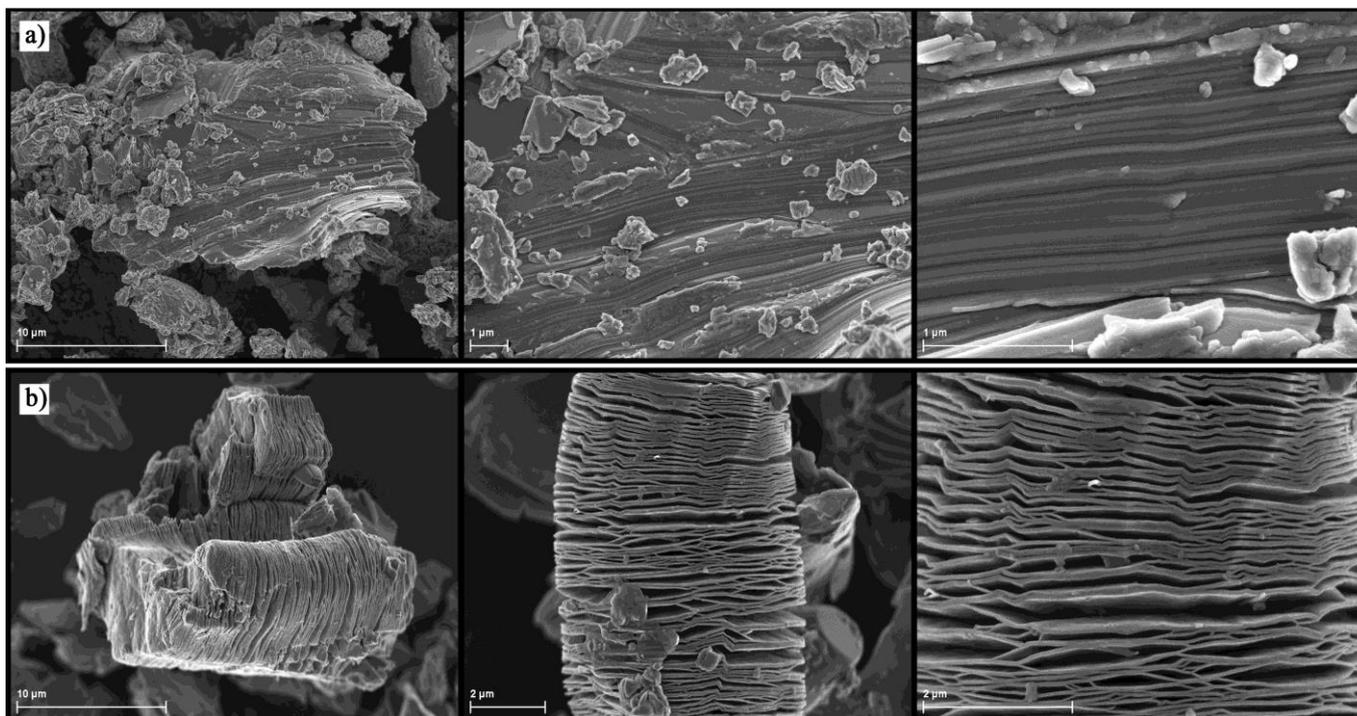


Figure 2. The set of SEM images obtained for the Ti_2AlC a) and expanded Ti_2C b).

One can notice that the Ti_2AlC grains possessed non-regular shapes (Fig. 2a). The high magnification images confirmed the layered structure of Ti_2AlC used as a substrate for Ti_2C synthesis. The SEM observations also showed that the sizes of the large Ti_2AlC grains were generally below 20 μm. It can be also noticed the large amount of fraction of small particles (below 2 μm) which were not characterized by the layered structure. It should be noted that they were almost completely removed during the rinsing procedure due to obtain the pure Ti_2C material which morphology was presented in Fig. 2b. As can be seen, the layers of the dense Ti_2AlC grains were successfully split into individual sheets of the expanded Ti_2C . The SEM results also indicate that the widths of the grains located perpendicular to the surface of the sheets were significantly extended in comparison to Ti_2AlC grains.

The high magnification SEM images presenting in detail the morphologies of the Ti_2C sheets were shown in Fig. 3. The qualitative analysis of sheets diameters indicated that the measured thickness varied between 26 and 61 nm. The sheets possessed the relatively sharp edges and were

connected to each other by specific bridges, forming the stable stack of nano-sized sheets with the slit-shaped pores.

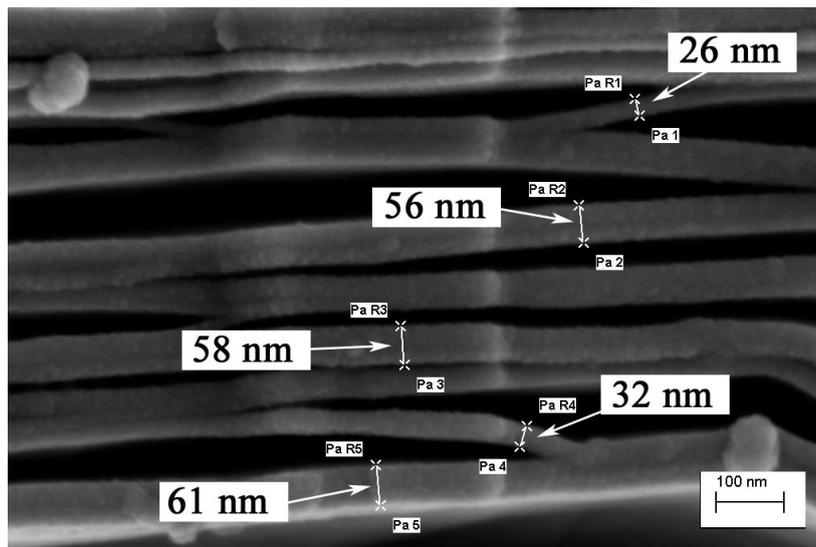


Figure 3. High magnification SEM image obtained for the expanded Ti_2C , added with the analysis of sheets diameters.

The elemental composition of the layered Ti_2AlC and expanded Ti_2C was also confirmed using the EDS analysis (Energy Dispersive X-ray Spectroscopy). The corresponding electron diffractions were presented in Fig. 4. The EDS results of the Ti_2AlC showed the presence of titanium, aluminium and carbon (Fig. 4a) whereas for Ti_2C almost no aluminium was detected (Fig. 4b).

Table 1. Summary of EDS results obtained from the surface of Ti_2AlC particle.

element	Series	C [at%]	C[wt%]	C error [wt%]
titanium	K-series	68,03	42,24	6,08
aluminium	K-series	15,55	17,14	2,28
carbon	K-series	16,41	40,63	7,39

The EDS results for each of the samples were also summarized in Tab. 1 and 2. As can be seen, the amount of aluminium was reduced from c.a. 17 wt% in Ti_2AlC (Tab. 1) to almost 0 wt% in Ti_2C (Tab. 2) which confirmed the efficiency of the chemical process of acidic aluminium extraction from Ti_2AlC . The EDS results obtained for Ti_2C have also shown the presence of significant amounts of elements such as: oxygen (19,6 wt %) and fluorine (12,2 wt %) which were previously described by other researchers [4, 6]. Other elements such as: calcium, sodium, sulfur, phosphorus and magnesium were present in Ti_2C in amounts around 1 wt% which were within the standard deviation of measurement (Tab. 2). Their presence may be considered as negligible impurities introduced during chemical process of aluminium extraction or rinsing.

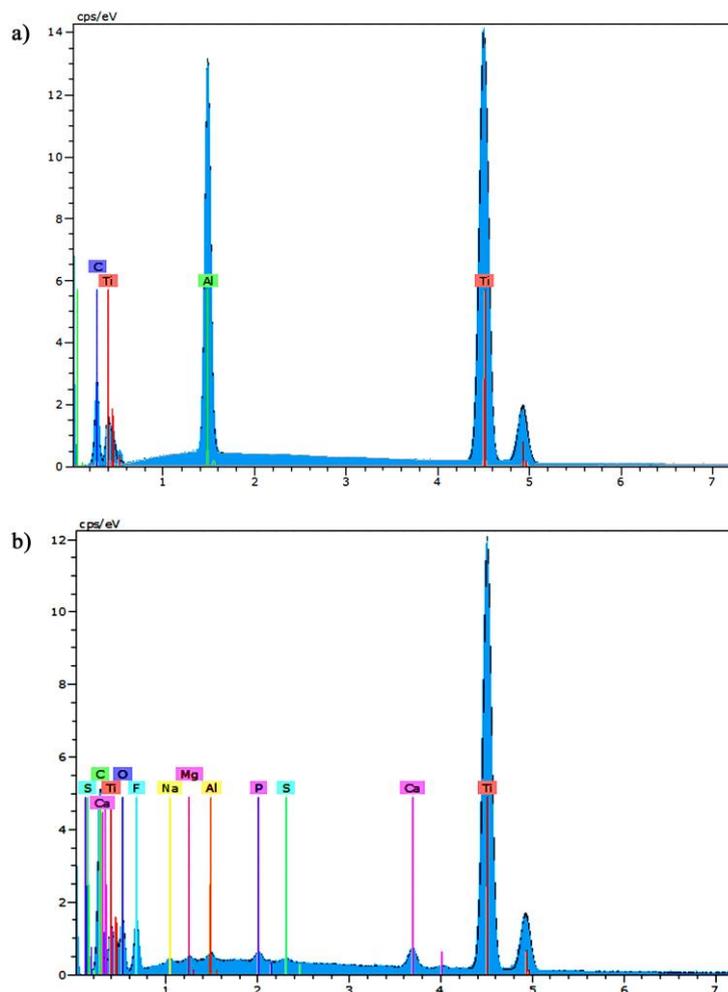


Figure 4. Results of EDS analysis obtained from the surface of Ti_2AlC grain a) and expanded Ti_2C sheets b).

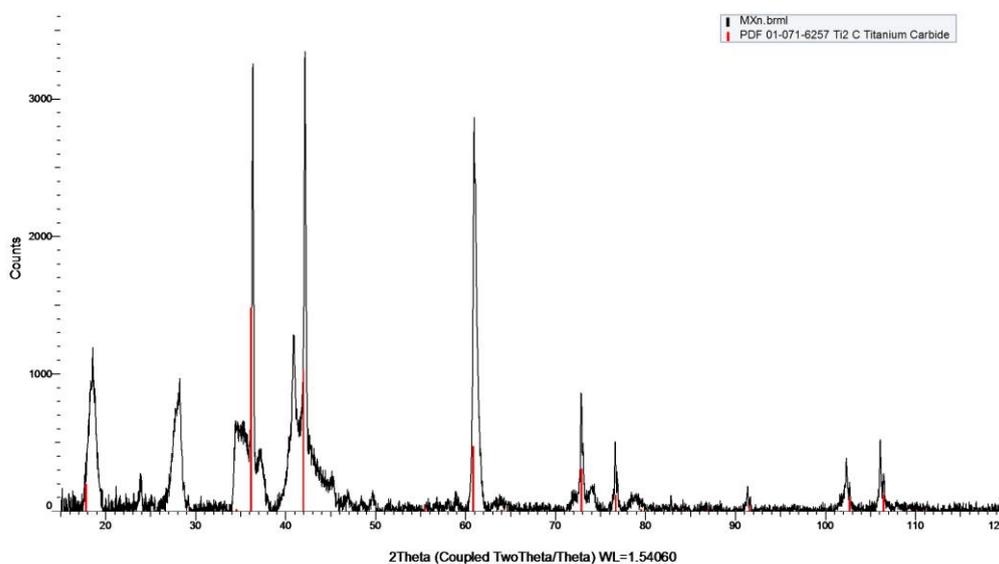


Figure 5. The XRD pattern obtained for Ti_2C sample.

It is also very highly possible, that those impurities were actually present from the beginning in the commercial MAX phase, but could not be detected by EDS due to the high amount of aluminium. After its removal, they become detectable by EDS detector. It should be noted, that we are the first who presented the EDS results showing that, in spite of using the pure HF for aluminium extraction, some minor impurities can be detected by EDS analysis in the final product. The XRD pattern obtained for the analysed sample was presented in Fig. 5. The obtained diffractogram confirmed the presence of the Ti_2C phase.

Table 2. Summary of EDS results obtained from the surface of Ti_2C sheets.

element	Series	C [at%]	C [wt%]	C error [wt%]
titanium	K-series	55,76	28,01	5,53
aluminium	K-series	0,24	0,21	0,13
carbon	K-series	19,23	38,50	8,55
oxygen	K-series	13,05	19,62	6,91
fluorine	K-series	9,64	12,20	5,03
calcium	K-series	1,33	0,80	0,23
sodium	K-series	0,17	0,17	0,13
sulfur	K-series	0,15	0,11	0,10
phosphorus	K-series	0,31	0,24	0,13
magnesium	K-series	0,14	0,14	0,11

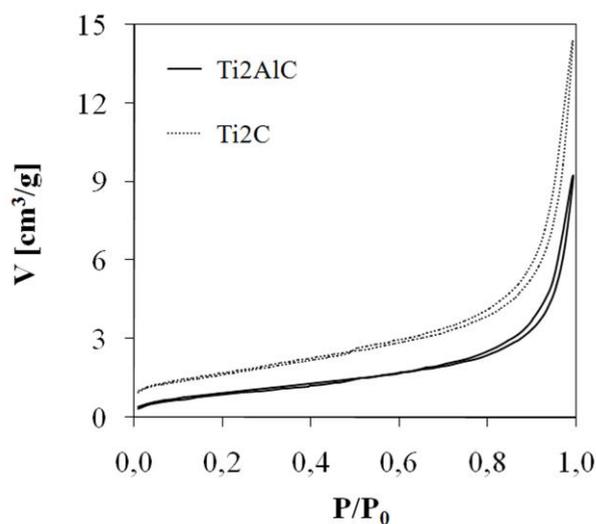


Figure 6. Nitrogen sorption isotherms obtained for the Ti_2AlC a) and expanded Ti_2C b).

The analysis of physical properties and porous structure of layered Ti_2AlC and expanded Ti_2C were examined based on the isotherms of the physical nitrogen sorption. The obtained isotherms are given in Fig. 6. The obtained results indicate, that the expanded Ti_2C adsorbed significantly more nitrogen in comparison go Ti_2AlC . The shape of the obtained isotherms suggested the presence of slit pores which are also characteristic for graphenes [7, 8].

The results of the physical properties of layered Ti₂AlC and expanded Ti₂C were summarized in Table 3. As expected, the BET specific surface area of expanded Ti₂C was ca. 42% higher in comparison with layered Ti₂AlC. In particular, the significant increase in the surface area is related to the formation of large number of slit-pores in Ti₂C as a result of expanding process. The results of the total volume of pores (V_{BJH}) also confirmed this assumption. The value of V_{BJH} obtained for Ti₂C was ca. 0,007 cm³/g larger in comparison with Ti₂AlC (Tab. 3).

Table 3. Physical properties of the Ti₂AlC and expanded Ti₂C.

sample name	S_{BET} [m ² /g]	S_{pores} [m ² /g]	V_{pores} [cm ³ /g]	D_{pore} [nm]
Ti ₂ AlC	3,51	3,01	0,014	4
Ti ₂ C	6,09	4,74	0,021	4

The pore size distributions obtained for Ti₂AlC and Ti₂C are presented in Fig. 7. The obtained BJH results indicate that, in all of our materials, there were mainly nanopores (in the range of 3–5 nm) present. As can be also observed, the mesopores (in the range of 5–50 nm) were present in great minority in comparison to the general amount of nanopores.

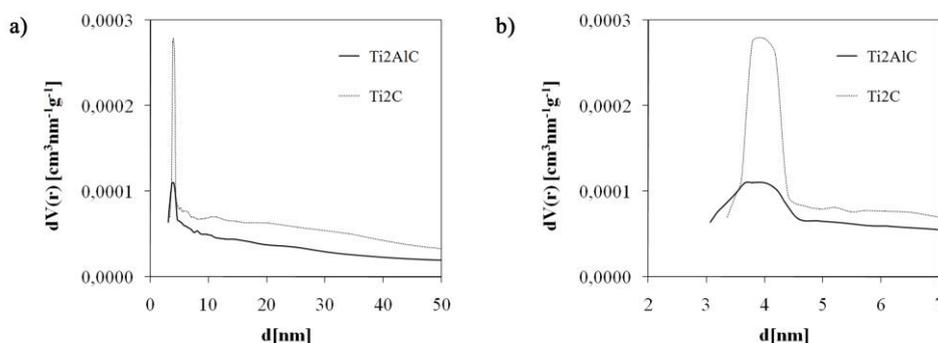


Figure 7. The distributions of pore diameter obtained using BJH method for the Ti₂AlC a) and expanded Ti₂C b).

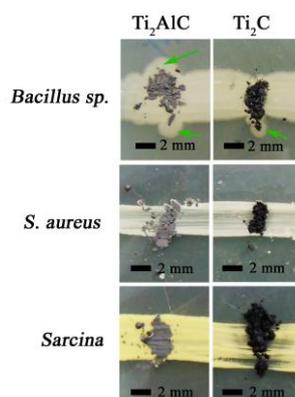


Figure 8. Photographs of the surface of Petri plates, presenting the growth of: *Bacillus sp.*, *S. aureus* and *Sarcina* bacteria in presence of Ti₂AlC and expanded Ti₂C. Green arrows mark regions where microorganisms exceed their primary inoculation line.

The present study also investigates the bioactive properties of Ti_2C , which is the expanded form of layered titanium carbide (Ti_2AlC). The investigations of the bioactive properties of the samples were performed using the classical culture method. Due to enable comparison, the layered Ti_2AlC was also subjected for microbiological analysis. As can be seen in Fig. 8, no inhibition of bacterial growth was observed in the presence of the tested samples. However, in case of *Bacillus sp.* bacteria the intensified growth was observed in the area directly surrounding the applied sample. As a result, the *Bacillus sp.* bacteria exceeded their primary growth lines. This effect may be related to the presence of the specific adhesion mechanisms between microorganisms and a specific surface properties of the tested material. For further studies, the expanded form of Ti_2C was used. The results of the qualitative analysis of preferential sites for adsorption of bacteria cells on the surface of expanded Ti_2C partially confirmed the described observations of the specific adhesion mechanisms.

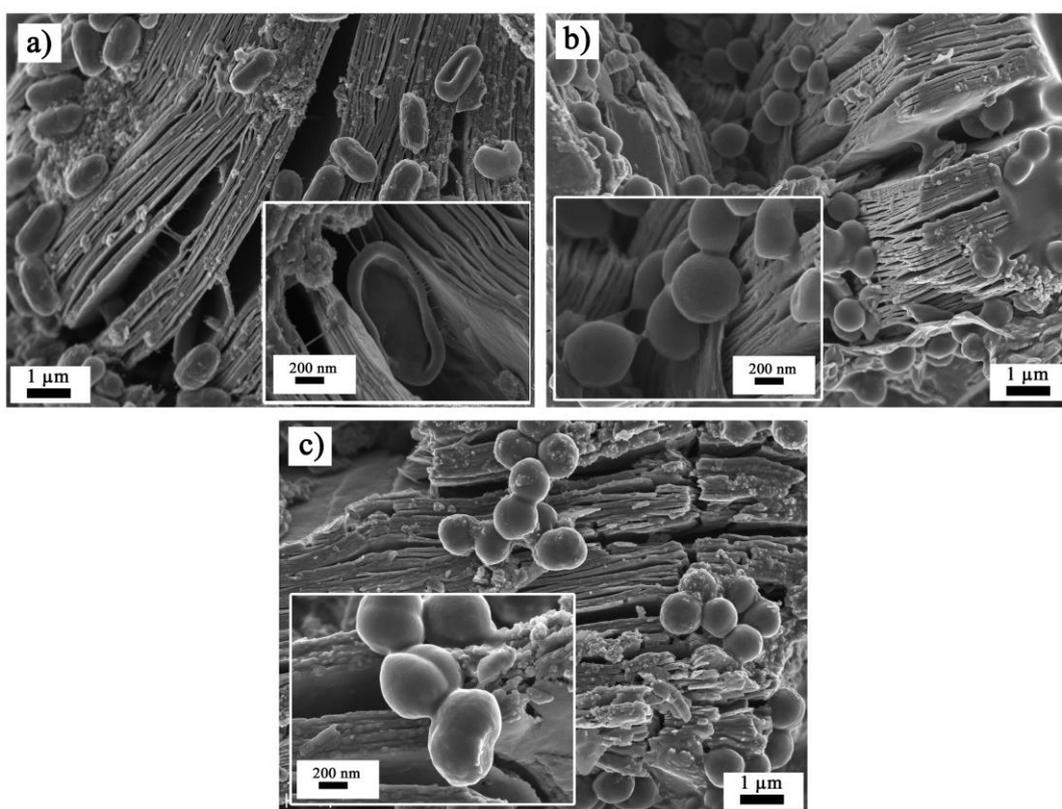


Figure 9. SEM images of the Ti_2C sheets with adsorbed bacteria cells such as: *Bacillus sp.* a), *S. aureus* b), and *Sarcina* c). The high magnification insets present in detail the adsorbed cells morphology.

The aim of this study was also to describe the preferential sites for adsorption of the bacteria cells on the surface of expanded Ti_2C using scanning electron microscope. The Fig. 9 shows the representative SEM images of the Ti_2C expanded sheets with the adsorbed bacteria cells of *Bacillus sp.*, *S. aureus*, and *Sarcina*. The morphology of the analysed cells was presented in detail on the high magnification insets. In case of *Bacillus sp.* species (Fig. 9a), the cells were randomly attached to the surface of the expanded walls of Ti_2C . However, one can notice that the cells also presented the specific ability to fit into the larger slit-shaped pores. The cells located between Ti_2C sheets were in

some cases highly disrupted. This effect was not seen for other bacteria species. The *S. aureus* and *Sarcina* cells were also found successfully attached to the surface of expanded Ti_2C sheets as can be seen in Fig. 8b and c, respectively. Their morphology was not significantly disrupted. However, in case of *S. aureus* some areas of the biofilm formed between individual Ti_2C sheets are present.

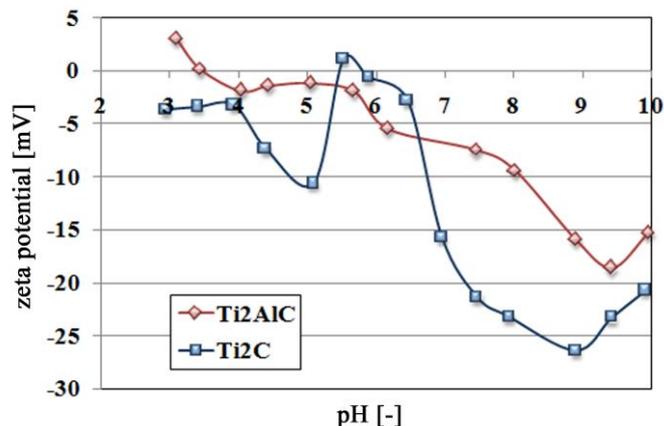


Figure 10. Zeta potential curves obtained for Ti_2AlC and expanded Ti_2C suspended in distilled water environment. The isoelectric points (IEPs) obtained for Ti_2AlC was 3,47 and for Ti_2C was 5,48 and 5,76.

In the next part of the study, the analysis of changes of zeta potential of Ti_2C as a result of bacteria cells adsorption was investigated. Such an analysis is very useful when there is a need for information on how the adsorption of bacteria cells changes the stability of a suspension and how much it influences the electrical charge formed on the surface of the analysed material or bacteria cells [7, 9, 10]. The methodology used for zeta potential investigations was similar to our previous reports [7, 11]. It should be also noted, that the Ti_2AlC was used as a reference only for the purpose of comparison of zeta potential of pure suspensions.

The zeta potential curves were recorded for the expanded Ti_2C and were presented in Fig. 10. It should be noted, that the Ti_2AlC sample was used in our investigations as a reference. It can be seen, that the zeta potential of Ti_2C was generally decreasing ca. -2,5 mV up to -26 mV when pH increased. However, it can be also noticed one clearly visible peak located around pH=6. Consequently, zeta potential curve two times crossed the zero zeta potential value. As a result, two IEPs (isoelectric points) were obtained i.e. 5,48 and 5,76. Moreover, at the highly basic environment (pH=ca. 9) the trend of the curve reversed and the zeta potential constantly increased with pH increase. When compare the shape of the zeta-curve to that obtained for Ti_2AlC , one can notice that it is closer to zero line without any peaks around pH=6. The IEP value obtained for Ti_2AlC was 3,47.

It should be however noted, that before our research there were no detailed reports on zeta potential curves of the expanded Ti_2C . To the best of our knowledge, there are only few reports available on other types of MXenes. Ren *et al.* [12] reported zeta potential of -29 mV at pH=7 for Ti_3C_2 (IEP= 2.4). Ying *et al.* [13] found that the Ti_3C_2 was characterized by a highly negative surface charge and the zeta potential of as-produced Ti_3C_2 suspension was -39.5 mV. Authors assumed that the negative surface charge originated from the =O, -OH and -F terminating functional groups present

on the surface of Ti_3C_2 sheets. Others researchers demonstrated that the V_2C sheets were negatively charged within the entire range of pH (from 0.97 to 11.3). At pH=7, the V_2C was found to possess highly negative surface charge (in the range from -46 mV to -58 mV) [14]. In contrast to those studies, the Ti_2C synthesized by us was characterized by a highly negative zeta potential only in the basic environment (in the 7-10 pH range). However, when the pH reached the value of 7, the rapid drop of zeta potential was observed.

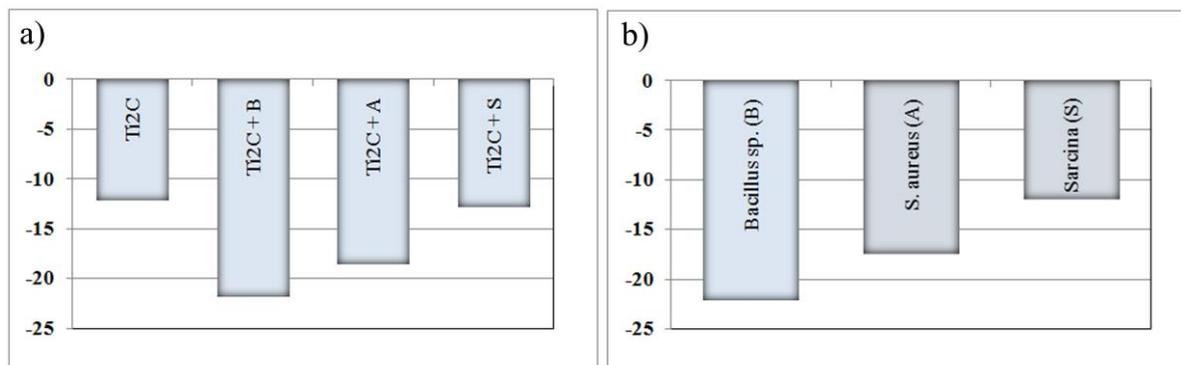


Figure 11. Zeta potentials obtained for the expanded Ti_2C with the presence of different bacteria cells a) as well as for pure bacteria cells suspensions in drinking water environment (pH=7,3) c). The bacteria strains were marked as: B for *Bacillus sp.*, A for *S. aureus* and S for *Sarcina*.

The zeta potentials obtained for Ti_2C suspensions in drinking water without and with the presence of bacteria cells were presented in Figure 11. The Ti_2C suspension was used in this study as a reference (Fig. 11 a). The pure suspensions of particular bacteria cells of strains such as: *Bacillus sp.*, *S. aureus* and *Sarcina* were presented in Fig. 11 b. We expected that the adsorption of bacteria cells onto the surface of Ti_2C should change its surface charge to similar to adsorbed cells by simple adjustment of the potential as a result of covering of the adsorbent surface [7, 9]. Our results indicate, that the adsorption of bacteria cells on the surface of Ti_2C resulted in adjustment of its zeta potential to that of bacteria cells. Thus, our results are in agreement with other data [7, 9]. It is known that during bacterial adhesion the electric double layer interactions lead to the redistribution of charges and the heterogeneous distribution of charges on the surface of bacteria reduces the effect of double layer repulsion during bacterial adhesion without affecting of bacteria viability [15]. Also the ion-penetrable layer of surface polymeric groups that are known to exist on the surface of bacteria may increase the attractive steric interactions promoting the adhesion by bridging the gap between two surfaces that repel each other (the ‘bridging’ interactions) [15]. All the mentioned phenomena could interplay on the surface of tested bacteria due to achieve the adsorption on the surface of Ti_2C .

4. SUMMARY

Recently, the chemically expanded structures of early transition metal carbides, known as MXenes have been introduced together with their specific properties which can be described as characteristic for both metals and ceramics. It made them attractive in the high-tech area of science. However, the potential of the biological activity of MXenes have not yet been fully discovered, highly

limiting the possibility of their application in many fields of bioscience and biotechnology. The present study, for the first time, investigates the biological activity of Ti_2C , which is the expanded form of layered titanium carbide (Ti_2AlC) against the gram-positive bacteria commonly occurring in the natural environment such as: *Bacillus sp.*, *S. aureus* and *Sarcina*. The aim of this study was also to describe the preferential sites for adsorption of the bacteria cells using scanning electron microscope examinations as well as analysis of changes of zeta potential of expanded Ti_2C MXene as a result of bacteria cells adsorption.

The SEM investigations results have shown that the layers of the dense Ti_2AlC grains were successfully split into individual sheets of the expanded Ti_2C characterized by the extended widths of the grains located perpendicular to the surface of the sheets. The diameters of the individual sheets varied between 26 and 61 nm. The sheets were also connected to each other forming the specific stack of nano-sized sheets with the pores of slit-type. The EDS results revealed that the amount of aluminium was reduced from c.a. 17 wt% in Ti_2AlC to almost 0 wt% in Ti_2C which confirmed the efficiency of the chemical process of aluminium extraction from Ti_2AlC . The presence of significant amounts of oxygen and fluorine was also confirmed. The shape of the isotherms obtained for the expanded Ti_2C suggested the presence of slit-shaped pores in the size range of 3–5 nm. As expected, the BET specific surface area of expanded Ti_2C was ca. 42% higher in comparison with layered Ti_2AlC .

It was also revealed, that both Ti_2AlC and Ti_2C did not influence negatively the analyzed gram-positive bacteria. Moreover, the slightly intensified growth of gram-positive *Bacillus sp.* strain was observed in the vicinity of tested Ti_2AlC and Ti_2C . What is interesting, this effect was not revealed for other gram-positive *S. aureus* and *Sarcina* strains. It should be noted, that the SEM investigations of the preferential sites for bacteria adsorption indicated the presence of minor apoptosis only for *Bacillus sp.* strain, especially when the cells were located between individual sheets of the expanded Ti_2C . The analysis of changes of zeta potential of Ti_2C in the presence of bacteria has shown that adsorption of bacteria cells on the surface of Ti_2C resulted in changing of its zeta potential to that of bacteria cells.

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The authors declare that there is no conflict of interest regarding the publication of this paper.

References

1. M. W. Barsoum, *Prog. Solid State Chem.* 28 (2000) 201.
2. M. Naguib, O. Mashtalir, J. Carle, V. Presser, J. Lu, L. Hultman, *ACS Nano* 6 (2012) 1322.
3. M. Naguib, V. N. Mochalin, M. W. Barsoum, Y. Gogotsi, *Adv. Mat.* 26 (2014) 992.
4. O. Mashtalir, M. Naguib, B. Dyatkin, Y. Gogotsi, M. W. Barsoum, *Mat. Chem. Phys.* 139 (2013) 147-152.
5. F. Wang, C. H. Yang, C. Y. Duan, D. Xiao, Y. Tang, J. F. Zhu, *J. Electrochem. Soc.* 162 (2015) B16.
6. X. Xie, S. Chen, W. Ding, Y. Nie, Z. Wei, *Chem. Commun.* 49 (2013) 10112.

7. A. M. Jastrzębska, E. Karwowska, P. Kurtycz, A. R. Olszyna, A. Kunicki, *Surf. Coat. Technol.* 271 (2015) 225.
8. A. M. Jastrzębska, J. Jureczko, A. R. Kunicki, A. R. Olszyna, *Int. J. Appl. Ceram. Technol.* 12 (2015) 522.
9. A. Jastrzębska, E. Karwowska, A. Olszyna, Chapter XI - Bio-Assessment and Toxicology: Influence of the Staphylococcus Aureus Bacteria Cells on the Zeta Potential of Graphene Oxide Modified with Alumina Nanoparticles in Electrolyte and Drinking Water Environment. In *Springer Proceedings in Energy, 2nd International Congress on Energy Efficiency and Energy Related Materials (ENEFM 2014)*, Oludeniz, Fethiye/Mugla, Turkey, 2014, pp 245-250.
10. P. Kurtycz, A. M. Jastrzębska, E. Karwowska, A. Olszyna, A.; Comparative assessment of the Graphene oxide and reduced Graphene oxide influence on the zeta potential of selected bacteria in drinking water environment. *Proceedings of the 9th International Conference on Composite Science and Technology (ICCST) – 2020 – Scientific and Industrial Challenges*, Naples, Italy, 2013, pp 383-393.
11. A. M. Jastrzębska, E. Karwowska, P. Kurtycz, E. Miałkiewicz-Pęska, D. Basiak, A. Olszyna, M. Załęska-Radziwiłł, N. Doskocz, *Int. J. Appl. Ceram. Technol.* 12 (2015) 1157.
12. C. E. Ren, K. B. Hatzell, M. Alhabeab, Z. Ling, K. A. Mahmoud, Y. Gogotsi, *J. Phys. Chem. Lett.* 6 (2015) 4026.
13. Y. Ying, Y. Liu, X. Wang, Y. Mao, W. Cao, P. Hu, X. Peng, *ACS Appl. Mat. Interfaces* 7 (2015) 1795.
14. M. Naguib, R. R. Unocic, B. L. Armstrong, J. Nanda, *Dalton Trans.* 44 (2015) 9353.
15. A. T. Poortiga, R. Bos, W. Norde, H. J. Busscher, *Surf. Sci. Rep.* 47 (2002) 1.

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