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Light elements analysis using ^3He microbeam

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Abstract

Coatings containing light elements such as carbon, nitrogen or oxygen play actually an important role in surface engineering as a means to improve the tribological performance. In this paper, we have studied the wear tracks on a diamond coating after a fretting test against a Cr steel ball. The microprobe facility of the LARN has been used to determine light elements concentration in the wear tracks because the size of the tracks was less than 100 μm and the transfer layer from the Cr steel counterbody was not uniform. Quantitative analysis of oxygen has been performed using the 2.392 MeV resonance of the $^{16}\text{O}(^3\text{He}, \alpha_0)^{15}\text{O}$ reaction at 90° , while carbon has been measured simultaneously with the $^{12}\text{C}(^3\text{He}, p_1)^{14}\text{N}^*$ reaction at 175° . The cross sections and the Q -value of this reaction are so high that an absorber is not required, so we can also measure the backscattered ^3He particles in the same detector. A correlation between carbon, oxygen and iron has been made to investigate the possibility of interdiffusion between the diamond coating and the transferred material due to the fretting test.

1. Introduction

Light elements such as carbon, nitrogen and oxygen play a very important role in the surface treatment of steels, glasses or polymers [1]. Protective, decorative or adhesive coatings can be made by different techniques like DC-magnetron sputtering, ion implantation, ion beam assisted deposition (IBAD) and chemical vapor deposition (CVD). Quantitative analysis and depth profiling of light elements can be performed using nuclear reaction analysis (NRA) induced by particles giving nuclear reactions with a positive Q -value, while particle induced X-ray emission (PIXE) or Rutherford backscattering spectroscopy (RBS) are often not very convenient for light elements determination into substrate with $Z > 10$. Deuterons and ^3He incident particles are often chosen because they induce nuclear reactions with a high positive Q -value. Although the cross-sections for reactions like (d, p) or (d, α) on light elements are usually higher than the cross-sections for reactions induced by ^3He particles at the same energy, reactions like $(^3\text{He}, p)$ or $(^3\text{He}, \alpha)$ produce one hundred times less neutrons than (d, p) or (d, α) reactions at the same energy. Reactions induced by ^3He particles are also more sensitive to the near surface since the stopping power is higher for

^3He than for deuterons of the same energy. Microbeam analysis using ^3He beam is not popular because most of the nuclear microprobes in the world are connected to tandem accelerators, which are limited in the brightness of the source. In the literature, we have seen only one paper on the microbeam analysis of deuterium using the $d(^3\text{He}, p)^4\text{He}$ reaction at 980 keV [2] because this reaction has a high cross-section (70 mb/sr). The cross-sections of reactions induced by ^3He particles on light elements have been measured and reported by several authors [3–8] but a lack of data exists.

In this paper, we show that ^3He particles can be used for simultaneous quantitative micro-analysis of light elements such as carbon and oxygen. The cross-section of the nuclear reaction $^{16}\text{O}(^3\text{He}, \alpha_0)^{15}\text{O}$ shows a resonance at 2.392 MeV [9] which can be used for depth profiling even with a low current microbeam (0.5 nA). Carbon can also be analyzed in the same experimental conditions at backward angle using the $^{12}\text{C}(^3\text{He}, p_1)^{14}\text{N}$ reaction. As an example, we have treated the case of the performance of diamond coatings in unlubricated contacts subjected to low amplitude oscillatory sliding, also known as fretting wear [10]. Other wear or friction tests using a pin on disk machine give also small tracks with dimensions in the range of 50–500 μm . Micro-analysis of tracks due to such tests are very useful in order to know the composition of the transfer layer and the mechanism of wear involved.

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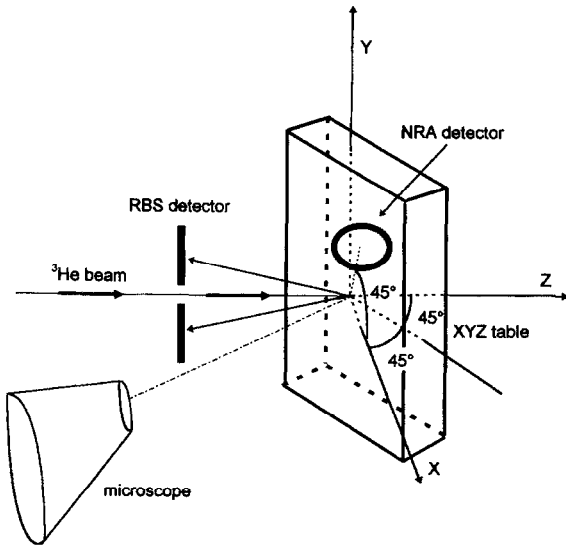


Fig. 1. 3D view of the experimental setup. The RBS detector is in the horizontal plane formed by the incident beam and the normal of the specimen, while the NRA detector is placed at 45° from the same plane and at 90° relative to the incident beam.

2. Experimental details

2.1. Experimental setup

The ^3He beam is produced with the 2.5 MV Van de Graaff accelerator located at LARN (Namur, Belgium).

The beam, typically $5\ \mu\text{m} \times 3\ \mu\text{m}$ and 0.5 nA in intensity, is focused on the sample mounted on an XYZ table, which can be moved with stepping motors in the X and Y directions. The energy of the incident particles is 2.435 MeV, just above the energy of the 2.392 MeV resonance of the $^{16}\text{O}(^3\text{He}, \alpha_0)^{15}\text{O}$ reaction and the incident angle between the impinging beam and the target is 45° . Two detectors are placed in the chamber as shown in Fig. 1. The first one, named a RBS detector, is a totally depleted annular surface barrier silicon detector located at 47 mm from the beam impact, showing a solid angle of 65 msr. The second detector is a passivated implanted planar silicon detector (PIPS), called a NRA detector, located at 90° relative to the incident beam direction and at 45° relative to the horizontal plane formed by the incident beam and the normal of the target. This detector is situated at 60 mm from the beam impact showing a solid angle of 37 msr. The bias voltage of the NRA detector was only 6 V in order to stop completely the 5.415 MeV alpha particles produced by the $^{16}\text{O}(^3\text{He}, \alpha_0)^{15}\text{O}$ reaction and to produce only a ΔE signal for the high energy protons produced by the $^{12}\text{C}(^3\text{He}, p)^{14}\text{N}$ reaction. The solid angles of both detectors have been calculated and measured by conventional RBS experiment on a known SnO_2 coating deposited on glass. An aluminum foil of $8\ \mu\text{m}$ thickness was placed in front of the NRA detector in order to stop the backscattered ^3He particles, while no absorber was required for the RBS detector. The absolute charge deposited by the incident beam on the specimen can be

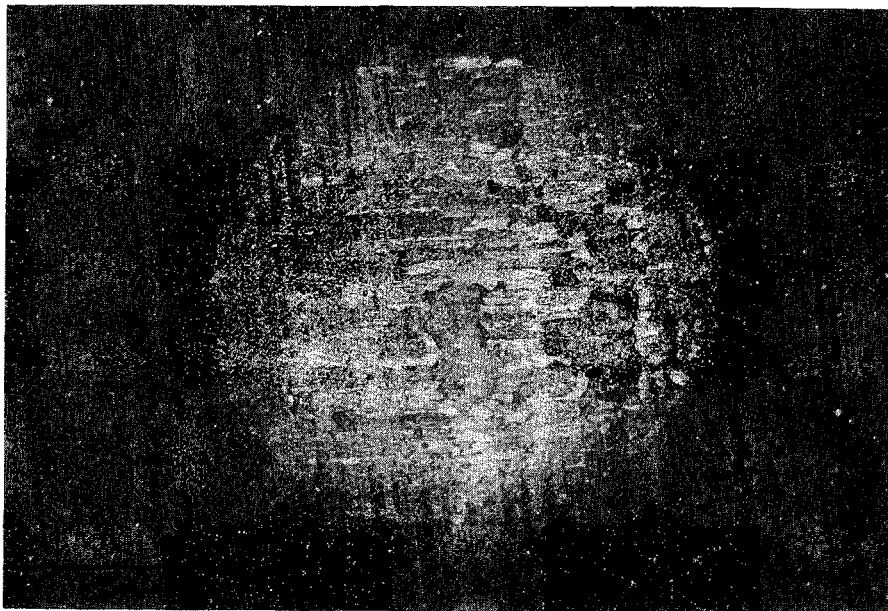


Fig. 2. Picture showing the track obtained on the diamond coating layer after the fretting wear test. The length of the index is $100\ \mu\text{m}$ and the estimated dimension of the track is about $500\ \mu\text{m}$.

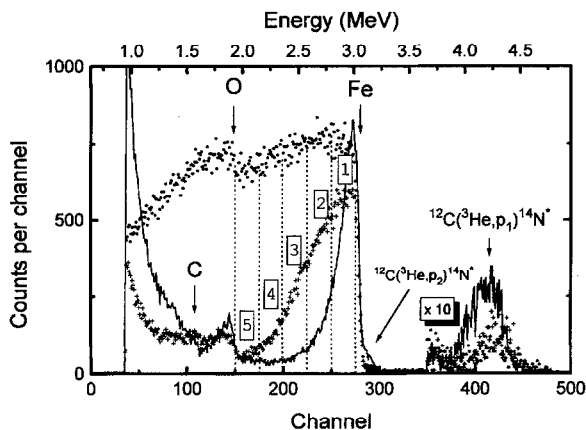


Fig. 3. RBS spectra recorded during the same incident charge (100 nCb) at three different impact points on the track showing three different thicknesses of the transferred layer. The full-line spectrum corresponds to a dark zone, the spectrum represented by crosses was taken in a semi-bright zone, while the dot spectrum was recorded in a bright area of the specimen. The regions of interest labeled 1 to 5 cover the information coming from iron and are named Fe₁ to Fe₅ in Fig. 5.

integrated using a positive bias voltage (150 V) applied to the specimen in order to attract the secondary electrons.

2.2. Samples preparation

Diamond coatings of 5 μm thickness were grown on WC-Co flat inserts (grade K10) with a hot filament chemical vapor deposition process [11]. The reactor was operated under a hydrogen/methane mixture (50 mbar) with the assistance of a W filament at about 2100°C. The substrate temperature during deposition was approximately 900°C. The coatings were then submitted to a low amplitude oscillating sliding wear test. A full description of the testing device is given in ref. [12]. The 100Cr6 steel ball counterbody (10 mm diameter) was loaded with a dead weight of 4 N against the diamond coated flat. The diamond coated sample was oscillated with a stroke of 100 μm and a frequency of 10 Hz for 100 000 cycles. Fig. 2 shows that a transferred layer has been formed [1] after the fretting test on the diamond coating in the contact zone.

3. Results and discussion

Fig. 3 shows three typical RBS spectra obtained at three different impacts with 2.435 MeV ³He particles. All spectra were recorded with the same incident charge (0.1 μCb). The full-line spectrum was recorded in a

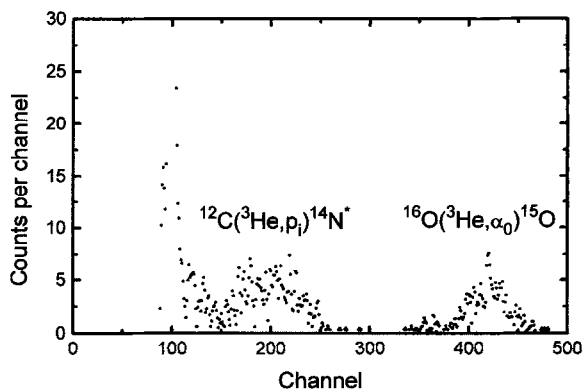


Fig. 4. NRA spectrum recorded during an incident charge of 100 nCb and simultaneously with the dotted line spectrum from Fig. 3. The bias voltage of the NRA detector was chosen in order to stop completely the alpha particles from the ¹⁶O(³He, α₀)¹⁵O reaction and to produce only a ΔE signal for the high energy protons produced by the ¹²C(³He, p)¹⁴N* reactions.

dark area of the specimen, where only a thin layer of the counterbody has been transferred on the sample. The signals coming from iron and oxygen are clearly observed, while carbon and oxygen interfere. We can also observe that information coming from the ¹²C(³He, p)¹⁴N nuclear reactions interfere with the RBS spectrum. The shape of the background under the RBS spectrum, due to nuclear reactions has been recorded outside the track and subtracted from the RBS spectra presented in order to obtain quantitative measurements. We have defined five different regions of interest covering the iron signal on the RBS spectrum. The first one, labeled 1, concerns the information coming from the surface, while the others represent the information coming deeper and deeper from the specimen. For example, there is no iron in the regions 4 and 5 for the full-line spectrum and in this case it is possible to obtain the concentration of oxygen directly from the RBS spectrum when the transferred layer is thin. The spectrum represented by crosses was recorded in a semi-bright region, where the transferred layer of the counterbody is thicker. In that case, iron is present in layers 4 and 5. The last spectrum represented by dots was obtained on a bright area where the transferred layer of the counterbody is so thick that signals coming from iron and oxygen are mixed. In that case, oxygen cannot be estimated from the RBS spectrum and we need the information recorded in the NRA detector (Fig. 4). The tail observed for the iron peak of the RBS spectra can be explained in two different ways: the first one is the possibility for the diffusion of the transferred layer in the diamond coating and the second possible explanation is the roughness of the diamond coating, which is about 200 nm. We have simulated the full-line RBS spectrum

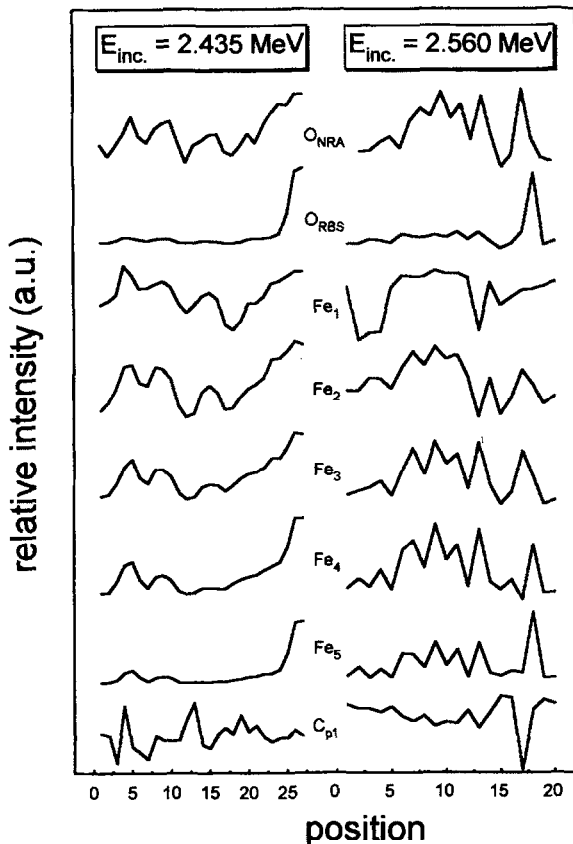


Fig. 5. Line-scans obtained at two different energies above the 2.392 MeV resonance of the $^{16}\text{O}(^3\text{He}, \alpha_0)^{15}\text{O}$ reaction. One step in the position scale represents 10 μm . The O_{NRA} curves correspond to a region of interest on the $^{16}\text{O}(^3\text{He}, \alpha_0)^{15}\text{O}$ peak (Fig. 4), while O_{RBS} , Fe_i and C_{p1} curves were obtained from a region of interest which corresponds respectively to oxygen, iron and carbon from the $^{12}\text{C}(^3\text{He}, \text{p}_1)^{14}\text{N}$ reaction in the RBS spectrum (Fig. 3).

using the RUMP code [13], taking into account the roughness of the diamond coating, and we have found a 200 nm iron oxide layer containing carbon with the stoichiometry of $\text{Fe}_{0.23}\text{O}_{0.46}\text{C}_{0.31}$.

It should be noted that for all RBS spectra, carbon coming from the diamond coating can be measured using the $^{12}\text{C}(^3\text{He}, \text{p}_1)^{14}\text{N}$ nuclear reaction and it is clearly seen that more carbon is observed when the transfer layer is thin. There is also an interference between the $^{12}\text{C}(^3\text{He}, \text{p}_2)^{14}\text{N}$ nuclear reaction and the ^3He retrodiffused by iron but in first approximation this contribution can be neglected. We can remove this contribution for absolute measurements.

Line-scans have been performed in the track shown in Fig. 2. To do this, we have introduced seven regions of interest on the RBS spectrum: the five regions in the iron signal as shown in Fig. 3, one region in the oxygen

information, labeled O_{RBS} and the last one on the peak corresponding to the $^{12}\text{C}(^3\text{He}, \text{p}_1)^{14}\text{N}$ reaction, named C_{p1} . Fig. 5 presents two line-scans of 27 and 20 steps; each step corresponds to a displacement of 10 μm . They were realized respectively at 2.435 and 2.560 MeV. For each point, the spectra (NRA and RBS) were recorded during 120 s and the total charge was also integrated during the same time for normalization of the data. The line-scans were normalized with the integrated charge and the results are presented in Fig. 5. The first energy was chosen just above the energy of the 2.392 MeV resonance of the $^{16}\text{O}(^3\text{He}, \alpha_0)^{15}\text{O}$ reaction and we can observe in Fig. 5 a correlation between the O_{NRA} and Fe_1 or Fe_2 information, while no correlation can be found between O_{NRA} and O_{RBS} or between O_{NRA} and Fe_5 . At higher incident energy (2.560 MeV), a correlation between O_{NRA} and Fe_3 or Fe_4 is evident but no correlation can be observed between O_{NRA} and Fe_1 or Fe_2 . No correlation can be found between O_{NRA} and O_{RBS} as mentioned above. This is due to the fact that the iron layer is so thick that the iron signal interferes with oxygen. We can conclude that iron and oxygen are bounded together and the stoichiometry can be measured by simulating the RBS spectrum obtained at one impact point as mentioned above. For both energies, we can observe an anticorrelation between O_{NRA} and C_{p1} , which proves that most of the information of the C_{p1} peak is coming from the diamond coating.

4. Conclusions

Quantitative microbeam analysis of light elements like C and O is possible even with a 5 $\mu\text{m} \times 3 \mu\text{m}$ ^3He beam of 0.5 nA intensity. It is more interesting to use a ^3He beam in place of deuteron particles because the reactions induced by ^3He particles are more surface sensitive and produce less neutrons. By combining RBS and NRA, it is also possible to find out the correlation between elements. Carbon and oxygen can also be measured simultaneously using two different detectors. Unfortunately, the literature concerning the cross-sections of the $(^3\text{He}, \text{p})$ and $(^3\text{He}, \alpha)$ is poor and therefore work remains to be done in this area.

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