Effects of boron concentration on the microstructure, mechanical and tribological properties of powder-pack borided AISI 4140 steel

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Research Article

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Abstract

Three different boron amounts were used in this study to control the microstructure, mechanical, and tribological properties of borided cases on AISI 4140 steel. A single Fe$_2$B phase was observed for the lowest boron amount, while both Fe$_2$B and FeB phases were present on the other two boron compositions. The highest surface hardness was observed in the highest boron amount. The scratch tests revealed different cohesive failure mechanisms on the samples, such as arc and chevron tensile cracks, and even chipping was observed for all three-studied boron amounts; it appeared at lower applied loads for the highest boron amount, indicating a likely detachment of the fragile FeB layer. The borided layers presented abrasive wear during the ball-on-flat wear tests. Despite the lower specific wear rate, the surface damage was more severe in the sample with the highest boron amount, due to its brittleness, resulting in lower resistance to surface fatigue.

1 Introduction

Wear is one of the most common metal surface degradation mechanisms; there are several ways to promote wear mitigation, such as increasing the surface hardness. Boriding is a very effective surface hardening treatment, which is applied to various ferrous and non-ferrous materials [1]. The boriding process in steels involves heating the material in a range between 700 and 1000°C, for 1 to 12 hours, in contact with a boriding medium such as a solid powder, slurry, liquid, or gas [1]. Boriding in steels generates layers with excellent adhesion [2], mainly due to its columnar morphology [3]. Besides, borided layers present high hardness and, consequently, good wear resistance [4–13]. One of the boriding challenges is to set the appropriate process parameters to obtain the proper microstructure to withstand a given wear environment.

Pack boriding methods present the simplest experimental setup, compared to the other boriding methods, but higher temperatures and longer treatment times are required [14]. Pack boriding has been recently studied in the boriding of titanium-based alloys [14–19], steel alloys [6–8, 10, 20–26], cobalt-base alloys [9, 27–31], and nickel-base alloys [32–34], and is usually performed using commercial powder mixtures, however, the pack boriding of steel and iron with different powder compositions was previously investigated [2, 10, 17, 19, 26, 35, 36]. The borided layers in carbon and alloyed steels usually have the Fe$_2$B single-phase or a two-phase morphology: FeB and Fe$_2$B. The FeB phase is richer in boron; however, a single Fe$_2$B phase is more desirable since the FeB, of higher hardness, is brittle [1].

One highly used commercial powder mixture consists of B$_4$C, KBF$_4$, and SiC [1]. The mechanism of pack boriding of steels using B$_4$C (boron source), KBF$_4$ (activator), and SiC (diluent) occurs according to the following reactions [37]:

$$KBF_4(s) \xrightarrow{530°C} KF(s) + BF_3(g)$$
According to Martini et al. (2004) [38], the thermochemical growth of borides on iron can be described as a three-step mechanism. The first step is characterized by Fe₂B crystals growing on the metal surface, creating a layer with randomly oriented or locally-oriented (110) or (200) crystals. The second step is the Fe₂B crystals' growth towards the metal bulk, creating an inner and compact region, showing a (002) preferred orientation. A (002) Fe₂B texture is established during the third step, and the needles grow perpendicular to the specimen's free surface [38]. The Fe₂B and FeB layers can form, but the free boron concentration on the steel surface dictate which of the following physicochemical processes will occur [37, 39]:

\[ \begin{align*}
B + Fe &\rightarrow FeB, \text{ at the B}/\text{Fe}_2B \text{ interface} \\
B + 2Fe &\rightarrow Fe_2B, \text{ at the B}/\text{Fe}_2B \text{ interface} \\
\text{Fe}_2B + B &\rightarrow 2FeB, \text{ at the FeB}/\text{Fe}_2B \text{ interface} \\
2FeB + B &\rightarrow Fe_2B + B, \text{ phase transformation (8)}
\end{align*} \]

A lower boron concentration favors the Fe₂B formation, while a higher boron concentration enables FeB to form as well [37], as shown in Equations (4) and (6). The FeB → Fe₂B phase transformation (Equation 8) occurs by prolonging the boriding process under a boron depletion condition [37].

In the present work, the authors sought to perform the pack boriding of AISI 4140 steel with new easy to manufacture and repeatable boriding mixtures, to study the feasibility of controlling the microstructure, and, consequently, the mechanical properties and wear resistance of the material.

### 2 Materials And Methods

#### 2.1 Materials

AISI 4140 steel (0.43 wt% C, 1.00 wt% Mn, 0.03 wt% P, 0.04 wt% S, 0.35 wt% Si, 1.10 wt% Cr, and 0.25 wt% Mo) specimens were sectioned into 5 mm thick disc samples. These disc samples were prepared with 320 to 1200 grit SiC abrasive paper and rinsed with ethanol before the boriding process. A commercial boron mixture (Ekabor®) consisting of 5 wt% of B₄C, 5 wt% of KBF₄, and 90 wt% of SiC was used as a basis for the development of two new boriding mixtures. The boron amount was varied by setting three amounts of B₄C: 1, 5, and 10 wt% yielding in 1.21, 4.35, and 8.26 wt% of boron amount. Each of the three
power mixtures contained 650 g, and the amount of each reactant and the amount of boron in each powder mixture is shown in Table 1. Boron carbide ($B_4C$) acts as a boron source, potassium tetrafluoroborate ($KBF_4$) as an activator of the boriding process, and silicon carbide ($SiC$) as a diluent, not taking part in the chemical reaction of boriding.

Table 1
Powder composition of the boriding mixtures (wt%).

| Sample | $B_4C$ | KBF$_4$ | SiC | B (wt%) |
|--------|--------|---------|-----|---------|
|        | wt%    | g B(g)  | wt% | g B(g)  |
| 1.21B  | 1      | 6.5     | 5   | 32.5    |
| 4.35B  | 5      | 32.5    | 5   | 32.5    |
| 8.26B  | 10     | 65      | 5   | 32.5    |

The $B_4C$ and SiC were supplied by Fiven Brazil, while KBF$_4$ was acquired from Sigma Aldrich. The reactants of each boriding mixture, with a granulometry below 100 µm, were mixed in a Y-mixer for 2 hours at 31.5 rpm on a lathe; and then mixed with 25 mm diameter alumina balls for 1 hour. Subsequently, a portion of each powder mixture was placed in a steel container, enough to fill the container's bottom and create a 25 mm thick powder layer. Afterward, four-steel disc samples were placed in the container, then the rest of the container was filled with the powder mixture, after which the steel container was closed and placed in a furnace at 900°C for 2 h. After the boriding process, the samples were air-cooled.

To facilitate the understanding of the manuscript, the AISI 4140 steel samples were identified according to the respective amount of boron in the powder mixture used in the pack boriding, as follows: 1.21B, 4.35B, and 8.26B.

2.2 Characterization

The borided specimens were prepared with 320 to 1200 grit SiC abrasive paper and polished with 0.25 µm diamond paste to a surface roughness of $Ra < 0.1$ µm. Subsequently, the samples were etched with Nital 2%. The microstructural features of the borided layers were determined through a specimens' cross-section inspection in a scanning electron microscope – SEM (Tescan Vega 3), allowing the examination of the morphology and the determination of the thicknesses of the borided cases. The average thickness of the FeB and Fe$_2$B layers was obtained by measuring 20 points in an SEM image with 2000x magnification. The thickness measurements consisted of establishing straight lines from the free surface to the ends of the peaks and valleys of the saw-tooth structure.

The identification of the phases in the borided layers was obtained through X-ray diffraction, XRD (Shimadzu XRD-7000) with CuK$_\alpha$ radiation ($\lambda = 1.54$ nm), with thin-film setting using 2° incidence angle, a scan speed of 0.5°/min and 0.02° step, with a 2θ scan range from 35° to 100°.
The surface hardness (H) was obtained by Vickers indentation, using a 250 mN load and 15 seconds of indentation time. The reduced Young's moduli (E) of the borided cases were extracted from the load-displacement curves using the method proposed by Oliver and Pharr \[40\], by instrumented indentation using a Berkovich indenter, following ASTM E2546-15 Standard \[41\], applying 24 steps up to a maximum load of 400 mN.

### 2.3 Scratch adhesion tests

The scratches were done and interpreted based on the ASTM C1624-05 Standard \[42\]. The scratch grooves were obtained in a Revetest Scratch Tester (Anton-Paar Instruments), with a Rockwell C diamond indenter with a tip radius of 200 µm. The test parameters were 398 N/min loading rate, 6 mm/min speed, progressive load from 1 to 200 N, and a scratch length of 3 mm. The scratch grooves were examined in a Tescan Vega 3 SEM to determine the failure mechanisms and to establish the critical loads (\(L_C\)).

### 2.4 Tribology tests

The tribological tests were conducted using a ball-on-flat configuration of an Anton-Paar Instrument according to the ASTM G99-17 Standard \[43\]. The tribological pair counterpart was a 6 mm diameter alumina (\(\text{Al}_2\text{O}_3\)) sphere. A sliding speed of 5 cm/s, a sliding track radius of 9 mm, a constant load of 20 N, and a sliding distance of 500 m were used for the tests. The coefficient of friction (CoF) was continuously recorded during the tests. After the tests, the wear tracks and their debris were analyzed by SEM/EDS (Tescan Vega 3) to determine the wear mechanisms during the tribotests and the chemical composition of the formed debris. An optical profiler (Taylor-Robson CCI Lite) was used to quantify the disk specimen wear, by measuring the cross-sectional area of the disk wear track. The average wear track area was multiplied by the wear track perimeter to obtain the total volumetric wear. The following equation determined the specific wear rate (\(\kappa\)):

\[
\kappa = \frac{V}{F \cdot L} \quad (9)
\]

where \(V \ [\text{mm}^3]\) is the total volumetric wear, \(F \ [\text{N}]\) is the normal load, and \(L \ [\text{m}]\) is the sliding distance \[44\].

The wear of the alumina ball counter body was inspected by SEM/EDS (Tescan Vega 3) and optical profiler (Taylor-Robson CCI Lite).

### 3 Results And Discussions

#### 3.1 Morphology and surface properties

The XRD diffraction patterns, shown in Fig. 1, reveal that the lowest boron amount (1.21B specimen) presented only peaks corresponding to the \(\text{Fe}_2\text{B}\) phase. In contrast, for the 4.35B and 8.26B specimens, the diffraction peaks of the \(\text{FeB}\) phase appear along with the diffraction peaks of the \(\text{Fe}_2\text{B}\), due to a higher concentration of free boron that enabled the \(\text{FeB}\) formation \[37\] by the physicochemical processes shown in Equations (4) and (6) \[39\].
The SEM images of the three borided conditions are in Fig. 2. The microstructures show a typical saw-tooth morphology. The Fe$_2$B needles grow toward the substrate in the three conditions, with a (002) texture. The preferred orientation for all samples follows a path of minimum resistance in the [001] direction, which is normal to the substrate surface, corresponding to the third step of thermochemical growth of iron borides proposed by Martini et al. (2004) [38]. The SEM images, Fig. 2 (a) and (d), confirm the Fe$_2$B single-phase layer for the 1.21B sample. The outermost FeB and the inner Fe$_2$B layers in the 8.26B specimen are visible in SEM images Fig. 2 (c) and (f). Although the FeB phase peaks appeared in the XRD pattern (Fig. 1), the SEM images, in Fig. 2 (b) and (e), do not show the presence of FeB in the 4.35B sample, which indicates that the FeB layer on this sample was not continuous, but a series of FeB crystals dispersed in the Fe$_2$B phase.

Also, cracks are present in the three conditions, the cracks in the 8.26B specimen outnumber the other two boron amounts most likely due to the difference in thermal expansion coefficients between FeB and Fe$_2$B phases ($\alpha_{\text{FeB}} = 23 \times 10^{-3} / ^{\circ}\text{C}$, $\alpha_{\text{Fe}_2\text{B}} = 7.85 \times 10^{-3} / ^{\circ}\text{C}$) [1].

The thicknesses of the 1.21B and 4.35B samples are 39 µm and 52 µm, respectively (Table 2). While the 8.26B sample showed a total thickness of 62 µm, being 11 µm of the outermost FeB layer.

| Sample | Phases     | Thickness (µm) | Hardness - H (GPa) | Reduced Young’s Modulus – E (GPa) |
|--------|------------|----------------|---------------------|-----------------------------------|
| 1.21B  | Fe$_2$B    | 39 ± 2         | 12.6 ± 0.7          | 318 ± 4                           |
| 4.35B  | FeB + Fe$_2$B | 52 ± 2     | 13.7 ± 1.0          | 319 ± 4                           |
| 8.26B  | FeB + Fe$_2$B | 11 ± 2 (FeB) | 16.7 ± 1.7          | 344 ± 6                           |
|        |            |                |                     |                                    |
|        | AISI 4140  | –              | 3.5 ± 0.2           | 222 ± 4                           |

The borided layers (Table 2) were three to four times harder than the untreated AISI 4140 steel substrate, which presents a hardness of 3.5 GPa. The surface hardness increased with the boron amount since the 1.21B sample showed the lowest (1282 HV or 12.6 GPa) while the 8.26B sample showed the highest surface hardness (1705 HV or 16.7 GPa). The higher surface hardness of the 8.26B specimen is related to the higher hardness of the outer FeB phase. The 4.35B sample showed an intermediate hardness (1395 HV or 13.7 GPa), closer to the 1.21B sample since the harder FeB layer was not homogeneous in the 4.35B sample.
The reduced elastic moduli (Table 2) significantly differ among the samples. All borided samples presented a higher reduced elastic modulus than the untreated steel substrate (222 GPa). The boron diffusion in the material's crystal lattice leads to higher surface rigidity [45]. The dual-phase boride layer on the 8.26B sample showed the highest modulus (344 GPa) since the outer FeB layer, with a massive amount of boron, results in a higher surface rigidity. The 1.21B and 4.35B samples showed similar elastic modulus (318 and 319 GPa, respectively) to the one found by da Costa Aichholz et al. (2018) [46] in samples with the Fe$_2$B phase (313 GPa).

### 3.2 Adhesion

The scratch tests were performed; consequently, the failure modes were identified, and the critical loads were assigned for each boriding condition. SEM images were taken to accurately determine the critical loads to reveal the scratch groove details, as shown in Fig. 3. The critical loads were correlated to the defined and repeated failure mechanisms, as shown in Table 3. Examining the scratch grooves, in Fig. 3, three failure modes can be identified in the borided cases: arc tensile cracks, forward chevron tensile cracks, and chipping. The first failure to occur is arc tensile cracking, which occurs due to a tensile stress field generated behind the indenter caused by its penetration [47]. The arc tensile cracks were observed at 37 N, 33 N, and 7 N, for the 1.21B, 4.35B, and 8.26B samples, respectively. Arc tensile cracks correlate with Young's moduli (E) difference between the borided layer and steel substrate (Table 2). The borided layer of the 8.26B sample showed a much higher surface reduced Young's modulus due to the FeB layer, which results in a smaller capacity to deform under tensile stresses, consequently leading to the lowest critical load for this failure mode.

| Sample | Failure modes                     | Critical damage load (N) |
|--------|-----------------------------------|--------------------------|
| 1.21B  | Arc Tensile Cracks                | 37                       |
|        | Forward Chevron Tensile Cracks    | 37                       |
|        | Chipping                          | 145                      |
| 4.35B  | Arc Tensile Cracks                | 33                       |
|        | Forward Chevron Tensile Cracks    | 44                       |
|        | Chipping                          | 146                      |
| 8.26B  | Arc Tensile Cracks                | 7                        |
|        | Forward Chevron Tensile Cracks    | 53                       |
|        | Chipping                          | 126                      |

Furthermore, the borided layers present different fracture toughness; the FeB phase has almost a quarter of the fracture toughness of the Fe$_2$B phase [48], which directly impacts the material's ability to deform without the formation of cracks. Then, as the scratch test progresses, forward chevron tensile cracks...
form on the borided samples, initiating inside the scratch groove and propagating away from it. This failure mechanism occurs due to the plastic deformation of the substrate [49]. Its onset seems to depend mainly on the borided layer's thickness since thinner layers tend to have less resistance to accommodate such deformation; consequently, it takes place at 37 N, 44 N, and 53 N for the 1.21B, 4.35B, and 8.26B samples, respectively. Finally, the borided layer's chipping takes place at 145 N, 146 N, and 126 N for the 1.21B, 4.35B, and 8.26B specimens. Notice that the borided layer with the FeB phase, namely sample 8.26B, shows lower chipping critical load, revealing the fragile nature of the FeB layer.

### 3.3 Wear behavior

Figure 4 shows the CoF behavior versus the sliding distance. The CoF behavior of the borided cases shows two stages: running-in and steady-state. The running stage varied among the borided cases; the shortest running-in corresponds to the 1.21B sample, and the longest to the 8.26B sample. The variation in the running-in is due to the difference in hardness among the samples since the running-in occurs when the asperities of the Al₂O₃ ball and the tested borided surfaces are deformed, and the surfaces are brought together. Then, the 1.21B sample has the lowest surface hardness, which resulted in faster deformation of the surface by the alumina counterpart, therefore the shorter running-in. In contrast, the 8.26B sample, with the higher hardness, exhibited more resistance to the asperity's deformation process, and consequently, a longer running-in is observed. In contrast, the steady-state period showed no difference among the three borided cases, and small fluctuations are related to oxide formation and its breakage [50], as well as material removal and formation of debris.

The optical profiler images and cross-sectional views of the tracks (Fig. 5) were used to obtain the total volumetric wear (V) and then the specific wear rate (κ) of the tested samples (Table 4). The wear tracks did not exceed the thickness of the Fe₂B (for the 1.21B and 4.35B samples) and FeB (for the 8.26B sample) layers. Since borided layers are useful for tribological applications if κ < 10⁻⁶ mm³/Nm [21], the samples of the three conditions presented mild wear. The 1.21B sample, with the higher wear rate (κ=1.3 x 10⁻⁶ mm³/Nm), presented the widest wear track, Fig. 5 (a) and (b), with a lateral agglomeration of material (tribofilms and smearing). The wear track of the 4.35B sample (κ=1.1 x 10⁻⁶ mm³/Nm), Fig. 5 (c) and (d), is narrower and has a significant material agglomeration, mostly at the sides of the track. The 8.26B sample had the lowest wear rate among the samples (κ=0.9 x 10⁻⁶ mm³/Nm), and the narrowest wear track, as observed in Fig. 5 (e) and (f), which is expected, since the wear rate is usually related to the hardness of the material [51, 52], and sample 8.26B, containing the FeB phase, showed the higher surfaces hardness among the studied conditions. Previous results also showed higher wear abrasion resistance of the FeB phase over the Fe₂B phase [53].
Table 4  
Coefficient of friction (CoF), wear volumes, and specific wear rate of the borided samples.

| Sample | Wear volume of the borided disk specimens (V), $10^{-3}$ [mm³] | Specific wear rate of borided disk specimens (k), $10^{-6}$ [mm³/Nm] |
|--------|-------------------------------------------------------------|--------------------------------------------------------|
| 1.21B  | 12.5 ± 1.6                                                 | 1.3 ± 0.2                                              |
| 4.35B  | 11.1 ± 1.8                                                 | 1.1 ± 0.2                                              |
| 8.26B  | 9.4 ± 1.4                                                  | 0.9 ± 0.1                                              |

The main wear mechanisms of the borided specimens are shown in Fig. 6 (a), (c), and (e). The first wear mechanism is grooving, resulting from the plowing effect due to hard particles, which are removed from the borided surface during the tests, and act as an abrasive surface, as previously observed [54], resulting in micro-abrasion of the borided layers. It can be identified as scratches parallel to the sliding direction. This mechanism is more severe on the surface of the 1.21B sample but is also present in the 4.35B sample, as revealed in Fig. 6 (a) and (c).

The 1.21B sample showed higher material removal during the tests, resulting in higher grooving levels on its surface, consequently a higher wear rate. The 8.26B sample has a thicker borided layer than the other two samples (Table 2), which decreases the wear loss, as supported by previous results [54–56]. The 8.26B sample surface has a large portion of polished regions visible in smooth areas and occurs due to lower levels of material removal, with no visible scratching [21]. The higher surface hardness improved resistance to the plowing effect, which was also observed previously [54].

The repeated loading and unloading of the surfaces can lead to surface fatigue [21], which results in flaking and pitting of the surface, the latter occurring when portions of material are torn out, creating irregular craters on the surface namely pits. Although the 8.26B sample has the lowest wear rate, its very hard surface could not sustain large plastic deformation, leading to larger flaking of the surface, followed by pitting, as can be seen in Fig. 6 (e). Tribofilms can be observed at the surface of the samples. These films result from material agglomeration after debris adhere and cluster due to mechanical contacts at the surface, as previously observed [21, 51]. Heavily oxidized tribofilms, identified as smearing, can be seen inside and on the sides of the wear tracks, mostly in the 8.26B specimen.

The aspect of the Al₂O₃ balls after sliding is shown in Fig. 7. The images reveal that the sliding contact resulted in the flattening of the Al₂O₃ balls. The wear scar of the alumina ball against 1.21B samples, Fig. 7 (a), shows plowing lines. It indicates grooving as the main wear mechanism, which occurred due to the three-body abrasive wear on the ball and disk, resulting in higher wear of 1.21B disk (borided specimen) (Table 4). The wear scar of the alumina ball against the 4.35B sample, Fig. 7 (b), shows less grooving than for 1.21B-ball, which indicates a lower formation of debris and less borided specimen wear (Table 4). In the case of the alumina ball against the 8.26B samples, the wear scar was smooth, with no grooving lines. The higher surface hardness of the 8.26B sample resulted in higher alumina's ball wear.
The EDS analysis of the wear scars of the alumina balls shows material adhesion, Fig. 7 (b), (d), and (f). they indicate the presence of mainly iron-oxides. The iron-oxides found in the pair 8.26B sample- ball (Fig. 6 (f) and 7 (f)), in higher amount, may have reduced the wear in the borided sample. Because the iron-oxides films can act as a lubricant, reducing the CoF; consequently, mitigating wear [21, 51]. On the other hand, it was observed high levels of flaking and pitting of its surface, probably due to the high brittleness of the FeB phase, resulting in less surface resistance to fatigue.

4 Conclusions

- The mixtures manufactured in this work can be easily reproduced, and new mixtures can be easily obtained in an industrial environment. Furthermore, by controlling the amount of boron in the mixture, the microstructure resulting from the boriding process can also be manipulated.
- A FeB single-phase was obtained by boriding with 1.21 wt% of B, and the FeB phase appeared in both 4.35 B and 8.26 B samples. However, the FeB phase on the 4.35B sample was not homogeneous, and this sample showed similar surface properties to the 1.21 B sample.
- The scratch test revealed the same failure modes regardless of the boron amount: arc tensile cracks, forward chevron tensile cracks, and chipping. On the other hand, the critical loads are different, revealing that microstructure and, consequently, surface hardness, Young's modulus, thickness affect the critical load. The 8.26 wt% B specimens have the most fragile behavior due to the outer FeB phase.
- The sliding contact between the borided cases and alumina balls generated abrasive wear. The average friction coefficients are practically the same for the three borided specimens. Although the running-in stage of the friction coefficient behavior is the longest for the 8.26B specimen, which is related to the highest hardness of this borided condition, the higher the hardness, the more difficult is the breakage of asperities.
- The specific wear rate for the 8.26B sample was the lowest among the three investigated conditions due to its higher surface hardness and a larger amount of iron-oxides formation during the tribological tests. On the other hand, the FeB phase increases the brittleness of the borided case causing several different surface fatigue mechanisms.

Declarations

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Competing Interests

The authors have no relevant financial or non-financial interests to disclose.

Availability of data and material

All data generated or analysed during this study are included in this published article (and its supplementary information files).

Code availability

Not applicable

Ethics approval

Not applicable

Consent to participate

Informed consent was obtained from all individual participants included in the study.

Consent for publication

The Authors hereby consent to publication of the Work in any and all JAMT publications.

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Figures
Figure 1

XRD patterns of boride samples: (a) 1.21B, (b) 4.35B, and (c) 8.26B.
Figure 2

SEM cross-section images of borided samples: (a, d) 1.21B, (b, e) 4.35B, and (c, f) 8.26B, etched with Nital 2%. 
Figure 3

SEM images of the scratch grooves, with detail of the failure mechanisms: (a) 1.21B, (b) 4.35B, and (c) 8.26B samples.
Figure 4

Friction behavior of the borided samples sliding against alumina balls.
Figure 5

Optical profiler images and cross-sections of the wear tracks: (a, b) 1.21B, (c, d) 4.35B, and (e, f) 8.26B samples.
Figure 6

SEM images of the wear tracks and the corresponding EDS analysis: (a, b) 1.21B, (c, d) 4.35B, and (e, f) 8.26B samples.
Figure 7

SEM images and EDS analysis of worn surfaces of Al$_2$O$_3$ balls against: (a, b) 1.21B, (c, d) 4.35B, and (e, f) 8.26B samples.