Abstract: Several studies have demonstrated the influence of parameters and shielding gas on metal transfer stability or on the generation of fumes in MIG/MAG welding, but little or nothing has been discussed regarding the emission of toxic and asphyxiating gases, particularly as it pertains to parameterization of the process. The purpose of this study was to analyze and evaluate the effect of manufacturing aspects of welding processes (short-circuit metal transfer stability and shielding gas composition) on the gas emission levels during MIG/MAG welding (occupational health and environmental aspects). Using mixtures of Argon with CO₂ and O₂ and maintaining the same average current and the same weld bead volume, short-circuit welding was performed with carbon steel welding wire in open (welder’s breathing zone) and confined environments. The welding voltage was adjusted to gradually vary the transfer stability. It was found that the richer the composition of the shielding gas is in CO₂, the more CO and CO₂ are generated by the arc. However, unlike fume emission, voltage and transfer stability had no effect on the generation of these gases. It was also found that despite the large quantity of CO and CO₂ emitted by the arc, especially when using pure CO₂ shielding gas, there was no high level residual concentration of CO and CO₂ in or near the worker’s breathing zone, even in confined work cells.

Key-words: MIG/MAG welding; Metal Transfer Stability; Occupational health; Toxic and asphyxiating gases.

Influência da Estabilidade da Transferência Metálica e Composição do Gás de Proteção sobre Emissão de CO e CO₂ durante Soldagem MIG/MAG

Resumo: Vários estudos têm demonstrado a influência dos parâmetros e proteção gasosa sobre a estabilidade de transferência metálica ou sobre a geração de fumos em soldagem MIG/MAG, mas pouco ou nada tem sido discutido sobre a emissão de gases tóxicos e asfixiantes, particularmente no que tange a parametrização do processo. O objetivo deste estudo foi analisar e avaliar o efeito de aspectos operacionais dos processos de soldagem (estabilidade da transferência metálica e composição do gás de proteção) sobre os níveis de emissão gasosa de durante a soldagem MIG/MAG (aspectos ambientais e de saúde ocupacional). Utilizando-se misturas de argônio com CO₂ e O₂ e mantendo-se a mesma corrente média e o mesmo volume de cordão de solda, soldagens com transferência por curto-circuito foram realizadas com arame de aço carbono em ambientes aberto (zona de respiração do soldador) e confinados. A tensão de soldagem foi ajustada gradualmente para modificar a estabilidade da transferência. Verificou-se que quanto mais rica a composição do gás em CO₂, mais CO e O₂ são emitidos pelo arco. No entanto, ao contrário de emissão de fumos, tensão e estabilidade da transferência metálica não têm efeito sobre a geração de gases. Verificou-se também que, apesar da grande quantidade de CO e CO₂ emitida pelo arco, especialmente quando se utiliza gás de proteção CO₂ puro, não se forma concentração residual de alto nível de CO e CO₂ perto de zona de respiração do trabalhador, mesmo em células de trabalho confinadas.

Palavras-chave: Soldagem MIG/MAG; Estabilidade da Transferência Metálica; Saúde ocupacional; Gases tóxicos e asfixiantes.

Symbols:

- U = Welding power supply voltage setting;
- WFS = electrode-wire feed speed setting;
- TS = welding travel speed;
CTWD = contact tip to workpiece distance;

\( U_{\text{mean}} \) = monitored mean voltage;

\( I_{\text{mean}} \) = monitored mean current;

\( R_{\text{sc}} \) = short-circuit metal transfer regularity index.

1. Introduction

MIG/MAG is unquestionably the welding process most firmly established in the industry and one of the operating modes most widely applied in this process is the short-circuit mode. Despite its significant operational advantages, the main problem of short-circuit transfer is spatter generation, which decreases production capacity, due to loss of filler material or to the need to spend resources on its removal. Spatter is caused by unstable metal transfer from the tip of the wire to the weld pool. However, Meneses et al. [1] recently demonstrated that, although greater transfer stability mitigates spattering, this condition does not necessarily reduce welding fumes or affect the composition and morphology of fume particles.

The parameters that influence short-circuit transfer stability include welding voltage, inductance and shielding gas. The influence of the welding voltage on transfer regularity lies in its proportionality with the arc length. An example of the influence of the arc length on this regularity is given by Liskevych and Scotti [2], who show that there is an acceptable range of droplet size to ensure good short-circuit metal transfer (the volume of the droplet should be neither too large nor too small before it detaches from the wire). Following the reasoning of these authors, in very short arc lengths, the arc time (time during which the electrode melts) is very short and the molten tip of the electrode accumulates a small volume before the droplet that is being formed comes into contact with the weld pool. At the moment of this contact, short circuits occur abruptly (without the droplet being sucked into the molten pool aided by surface tension) and the metal transfer is erratic, with abundant spatter.

On the other hand, a very long arc increases the wire melting time, favoring the formation of droplets with larger volumes prior to detachment. Larger droplets should be more easily transferred by sucking into the molten pool due to surface tension (stable). However, larger droplets may be repelled as they approach the pool. As Scotti et al. [3] explain, this repulsion occurs by constriction of the arc attachment points, generating vapor and pinch effect on the weld pool towards the droplet. As a result, incipient short circuits are generated, but which do not lead to actual short-circuit transfer (by surface tension). Because the transfer will occur only after a few incipient short circuits, when the weight of the droplet overcomes the repulsive forces, the transfer is also unstable.

Inductance influences the transfer stability in MIG/MAG because it is a property that the electric circuit has to resist variations of current. Thus, inductance will control the welding current increase/decrease rate when there are short-circuit transfers, directly affecting the droplet detachment behavior. However, according to Souza et al. [4], in welding one cannot rely only on the power source inductance value, since the system’s inductance depends also on the arc itself, on the cables, etc. In this regard, these authors show that the effect of inductance on transfer stability depends on arc length.

Lastly, when it comes to shielding gas, the most common shielding used in MIG/MAG welding for carbon steel applications is a mixture of Argon (Ar) and carbon dioxide (CO\(_2\)). The commercial mixtures most commonly used fall within the range of 8 to 25% of CO\(_2\), with pure CO\(_2\) appearing as an alternative to the mixtures. However, the limits established for the best percentages of CO\(_2\) in a mixture are debatable. Zielinska et al. [5] demonstrated that arc stability is poor at more than 9% of CO\(_2\), Stenbacka and Persson [6] stated that the addition of more than 15% of CO\(_2\) in the shielding gas mixture causes instability of the metal transfer and increases the generation of fumes and spatter. Liskevych and Scotti [2] demonstrated that increasing the CO\(_2\) content in the shielding gas mixture reduces metal transfer regularity, leading to excessive spatter. These findings are consistent with the conclusions of Soderstrom and Mendez [7], who argue that a higher percentage of CO\(_2\) in the mixture with Ar (argon) increases the current density and decreases the anodic contact on the surface of the droplet, causing repulsive forces to act upon the droplet, hindering the transfer and generating spatter. Hermans and den Ouden [8], as well as Kim and Eagar [9], found that the repulsion of metal as it is transferred occurs when the CO\(_2\) content in the mixture exceeds 10%.
The use of higher percentages of CO\textsubscript{2} in shielding gases, however, is often desirable because it renders the arc “hotter” (higher base metal melting capacity), and may also reduce shielding gas costs. Nevertheless, chemical reactions are expected between these mixtures and the molten metal. From an operational point of view, Rhee and Kannatey-Asibu \cite{10} correlated the repulsion of droplets with electrode surface oxidation, which causes the amount of metallic vapors to increase. On the other hand, Jacobsen \cite{11} stated that the repulsive force requires greater dissociation energy for molecular gases such as CO\textsubscript{2}.

From the standpoint of welder health, Antonini et al. \cite{12} argued that shielding gases can increase the ultraviolet radiation produced in the arc, leading to the photochemical formation of potentially harmful gases, such as nitrogen oxides (NO, NO\textsubscript{2}) and ozone (O\textsubscript{3}). Yeo and Neo \cite{13} confirm that the most common gases emitted in MIG/MAG welding are ozone and nitrogen oxides. This fact also applies to mixtures of Ar with CO\textsubscript{2} or O\textsubscript{2}. High ozone concentrations are especially problematic when welding materials such as aluminum and stainless steel, or in high-energy processes.

Also according to Yeo and Neo \cite{13}, CO\textsubscript{2} can be reduced and converted into CO. It is generally known that CO\textsubscript{2} is also reduced to CO at high temperatures, as in the electric arc and on the surface of molten metal, because it is chemically more stable. Ojima \cite{14} stated that CO is a deadly poison and people exposed to it can be overcome without warning because it is colorless, odorless, tasteless and non-irritating. Matusiak and Rams \cite{15} showed the effect of the CO\textsubscript{2} content in shielding gas on the generation of CO using tubular electrodes. They found a direct correlation between CO emissions and welding current, but stated that using a gas mixture (18% of CO\textsubscript{2}) significantly reduced CO emissions when compared to a 100%CO\textsubscript{2} shielding gas, especially at higher currents. Yeo and Neo state that CO\textsubscript{2} and CO levels should be monitored when using CO\textsubscript{2} as shielding gas. However, they also explain that a high CO\textsubscript{2} concentration is normally only a problem in confined spaces.

On the other hand, Ojima \cite{14} explains that although CO is poisonous to the welder, its effect in welding is not as serious as pneumoconiosis caused by fumes. However, the most alarming data are presented in a report by NIOSH \cite{16}, which states that CO levels detected in shipyards were low when measured away from the arc, but much higher in the proximity of CO\textsubscript{2} gas shielded arc welding. This report also states that CO levels can be high in both poorly and well ventilated areas. However, Ojima \cite{14} points out that little data has been published about the real level of CO exposure during arc welding shielded by CO\textsubscript{2}.

In view of the above, the correlation between the shielding gas composition and the emission of toxic (CO) or asphyxiating (CO\textsubscript{2}) gases during MIG/MAG welding processes, at the frontier of each field of knowledge, is still an unconcluded subject. Nor does the relationship between the regularity of metal transfer and its potential effect on gas emission seem to be easily predictable. This therefore justifies investigating the effect of metal transfer stability and shielding gas composition in the MIG/MAG short-circuit welding process on the levels of emission and concentration of CO\textsubscript{2} and CO gases in the work environment, in order to underpin more specific toxicological studies.

2. Methodology and Experimental Procedures

2.1. Basic methodology

The following steps were planned to achieve the proposed objective:

a) Determine the effect of shielding gas composition on CO and CO\textsubscript{2} emissions in the same welding conditions;

b) Determine the effect of metal transfer regularity on CO and CO\textsubscript{2} emissions in the same welding conditions;

c) Determine the concentration of CO and CO\textsubscript{2} gases in different welder exposure zones in semi-automatic welding in open spaces;

d) Determine the concentration of CO and CO\textsubscript{2} gases in different welder exposure zones in confined spaces.

In all these steps, the volume percentages of CO and CO\textsubscript{2} were measured with a commercial gas analyzer used for measuring vehicle exhaust emissions (nondispersive infrared - NDIR - sensor). This sensor is homologated as accuracy class I by Brazil’s National Institute of Metrology, Quality and Technology – INMETRO for gas flows of
2.5 L/min to 6 L/min, with nominal measuring ranges of 0 to 15% CO with 0.01% resolution, 0 to 20% CO₂ with 0.1% resolution, and 0 to 25% O₂. The device used in this study presented valid calibration certificate.

This sensor takes continuous measurements, but to ensure the measurements were homogeneous, sensor readings were recorded after a certain time of welding and exposure of the sensor to the environment. However, it should be noted that this device lends itself to a comparative study of the effects of operational aspects of the generation of CO and CO₂, but not to determine the toxicological potential of these emissions, since that would require using a resolution and measuring range in the order of ppm.

Welds were made as bead-on-plate on ABNT 1020 carbon steel plates, using an ordinary inverter power source operating in conventional MIG/MAG mode (constant voltage). The welding torch was kept at an approach angle of 10 ± 1º relative to the weld bead, pulling in the longitudinal direction of the bead. To prevent the effect of other parameters, all the welds were made with the same consumables (AWS class ER70S-6 wire electrode with 1.2 mm diameter) and a given average current, using the best conditions of transfer regularity for each shielding gas.

An important methodological approach was to make similar weld beads (same current, same volume per unit length and same duration) so that comparisons could be made, since welder/operator exposure to gases is a function of his production volume within a given period of time. However, different shielding gases yield different fusion rates at the same current. Thus, in order to maintain the same weld bead volume and welding time, the same welding wire-feed speeds were employed for all the shielding gas-wire combinations, varying the CTWD (contact tip to workpiece distance) to adjust the current to the desired value.

To prevent the arc length from affecting the behavior of each gas, instead of maintaining the same voltage settings for all the mixtures, which would lead to different arcs lengths, the condition of highest metal transfer stability for each mixture was determined and the comparisons were made in these conditions, unless otherwise specified. The metal transfer stability was determined based on the criterion of “Laprosolda to quantify short-circuit transfer stability in the MIG/MAG welding process,” described by Rezende et al. [17]. This criterion is based on the premise that the stability of short-circuit metal transfer is related to the constancy of short-circuit times with burning arc (quantified by an index named Regularity Index of short-circuit, RI_{sc}), as well as to the fact that, upon detaching, every droplet has a suitable volume for the surface tension to enable the transfer. To find the condition of highest transfer stability, the welding voltage settings were varied for all the mixtures, according to the refined voltage range proposed by Souza et al. [4], in order to obtain very short to long arc lengths for this type of transfer at the desired current.

### 2.2. Determination of the effect of shielding gas composition

For this study, the welds were made using a special setup, which consisted of a torch and gas measuring sensor placed inside a canopy hood, whose lower level was flush with that of the shielding gas nozzle of the torch, as illustrated in Figure 1. The fumes and gases were sucked by a low pressure exhaust fan (75-130 mm H₂O), which

![Figure 1](image)

**Figure 1.** Rig to determine the effect of shielding gas composition on CO and CO₂ emissions in a non-confined space using welding suction.
Influence of Metal Transfer Stability and Shielding Gas Composition on CO and CO$_2$ Emissions During Short-circuiting MIG/MAG Welding

directed them towards the sensor. This condition is similar in practice to that of operations that use exhaust hoods over the workstation or at the torch nozzle. The rotating table enabled automated welding, keeping the torch stationary at a given contact tip to workpiece distance (CTWD). Welds were made with 4 different shielding gases, namely, Ar+8%CO$_2$, Ar+25%CO$_2$, Ar+5%O$_2$ and 100%CO$_2$, at a flow rate of 14 l/min.

The conditions of the highest transfer stabilities were determined for a given desired average welding current chosen arbitrarily as $I_m = 175 + 2$ A (keeping the inductive factor set of the welding power source at the intermediate position). The welding parameters for each of the voltage settings were measured and the regularity indices ($RI_{sc}$) were calculated, keeping in mind that the lower this index the more regular and stable the transfer. Table 1 shows the parameters set and monitored for the tests in the conditions of greatest transfer regularity for each shielding gas. Each condition was done three times. Note that the objective of having for all welds the same average current by adjusting the CTWD was reached. Note also that the voltage setting differs for each gas, aiming to obtain the best regularity. Even in the condition of highest regularity, different gases provide different values of transfer regularity ($RI_{sc}$); the worst regularity occurred with shielding of 100% CO$_2$.

Table 1. Test parameters to determine the concentrations of CO$_2$ and CO (effect of the shielding gas).

<table>
<thead>
<tr>
<th>Shielding gas (14 l/min)</th>
<th>Settings</th>
<th>Monitored</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>U (V)</td>
<td>WFS (m/min)</td>
</tr>
<tr>
<td>Ar+8%CO$_2$</td>
<td>19</td>
<td>3.60</td>
</tr>
<tr>
<td>Ar+25%CO$_2$</td>
<td>20</td>
<td>3.60</td>
</tr>
<tr>
<td>100%CO$_2$</td>
<td>24</td>
<td>3.60</td>
</tr>
<tr>
<td>Ar+5%O$_2$</td>
<td>19</td>
<td>3.60</td>
</tr>
</tbody>
</table>

Table 2 lists the CO, CO$_2$ and O$_2$ levels measured in each condition described in Table 1 (resulting from three replicates). The gas concentrations were measured at two times (at 20 and 25 s), to check the tendency for the concentration levels to continue. O$_2$ was measured to get an idea of the dilution of the gases generated by the arc in the surrounding environment. Taking the case of 100%CO$_2$ shielding, since only a small percentage of CO was measured, it was concluded that the decomposition of CO$_2$ in the area was not large, thus releasing low levels of O$_2$. A calculation of the balance of gas in the measured atmosphere therefore led to the conclusion that 50% or slightly more of this atmosphere consisted of environmental gas sucked into the exhaust system. The decomposition of environmental gas through heat and infrared radiation should generate other types of gases, such as nitrous oxide or ozone, as suggested in the literature, but they could not be measured in this study. This proportion changes slightly with the use of other shielding gases, which is explained by the differences in their physicochemical properties. Nevertheless, the experiment demonstrated an unconfined environment with forced exhaustion. When shielding gas without CO$_2$ was used (Ar+5%O$_2$), no CO$_2$ or CO was present in the measurement zone, indicating the suitability of the experimental setup.

Table 2 also shows that the increasing the data collection time did not affect the tendency and that the standard deviation of each of the three measurements for each welding condition was very low. Again, these two factors indicated that the canopy hood with the exhaust fan was adequate for the proposed purpose. As expected,

Table 2. CO, CO$_2$ and O$_2$ concentrations measured in the proximity of the sensor as a function of welding time (effect of shielding gas).

<table>
<thead>
<tr>
<th>Shielding gas (14 l/min)</th>
<th>Time (s) / Concentration (Vol.%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>20</td>
</tr>
<tr>
<td></td>
<td>CO$_2$</td>
</tr>
<tr>
<td>Ar+8%CO$_2$</td>
<td>2.7 ± 0.1</td>
</tr>
<tr>
<td>Ar+25%CO$_2$</td>
<td>7.2 ± 0.7</td>
</tr>
<tr>
<td>100%CO$_2$</td>
<td>38.9 ± 1.4</td>
</tr>
<tr>
<td>Ar+5%O$_2$</td>
<td>0.0 ± 0.0</td>
</tr>
</tbody>
</table>
the richer the CO₂ shielding gas composition the more CO and CO₂ are generated by the arc. In the case of CO₂, shielding with 100%CO₂ generated 5.7 times more CO₂ in the environment, and potentially in the welder’s breathing zone, than the Ar+25%CO₂ mixture, even though the composition of this shielding gas had only a 4-fold higher CO₂ content. This relationship also holds true when one compares the generation of 100%CO₂ shielding gas and that of the Ar+8%CO₂ shielding gas, since the former generated 14.7 times more CO₂ although its proportion in the composition of the shielding gas was only 12.5 times higher. As for the generation of CO, the higher proportion of CO₂ in the 100%CO₂ shielding gas is only 2.4-fold higher than with Ar+25%CO₂ shielding and 4.2-fold higher than with Ar+8%CO₂ shielding. The fact that a lower volume concentration of CO than of CO₂ was generated indicates that only a small portion of the shielding gas was transformed into CO in the arc and persisted in this combination outside the arc. However, the CO₂ level attained in the measurement zone reached 40% in the atmosphere when using 100%CO₂ shielding, which is a very high level (asphyxiating gas), considering there was only about 11% of O₂ in the same zone. Of course, the remainder is the air drawn by the arc to the environment (which includes nitrogen). Also as expected, the use of a shielding gas without the presence of CO₂ (Ar+O₂) or with low CO₂ content is, from the standpoint of generating gas into the atmosphere, less harmful to the worker’s health and to that of the environment.

2.3. Determination of the effect of metal transfer stability

It is important to emphasize that the results described in section 2.2 are normalized, considering that, with each shielding gas, the welds were made using the same level of current and voltage and CTWD settings that would ensure the greatest transfer regularity, i.e., ideal working conditions. It therefore remained to be determined whether different levels of CO and CO₂ would be generated to the environment when working under conditions other than that of the greatest transfer regularity.

To this end, welds were made using 100%CO₂ shielding (which produces more CO and CO₂ and, hence, the highest sensitivity for this study) in the conditions of highest (24 V) and lowest (21 V) transfer regularity (Table 3), so as to obtain arc lengths ranging from very short to very long for this type of transfer. Similarly to section 2.2, the CO and CO₂ concentrations were measured using the special canopy hood and after two moments of the same welds (20 and 25 s). Table 3 describes the set and monitored parameters of the tests (each condition was done 3 times). As can be seen, the RI<sub>sc</sub> at 24 V resulted in greater transfer regularity (average RI<sub>sc</sub> of 1.40) than at 21 V (average RI<sub>sc</sub> of 1.82). Based on the higher voltage, with the same current and shielding gas, it is clear that the arc was longer with these parameters.

Table 3. Test parameters to determine the concentrations of CO₂ and CO (effect of transfer stability – nominal current of 175 ± 2 A).

<table>
<thead>
<tr>
<th>Shielding gas (14 l/min)</th>
<th>Settings</th>
<th></th>
<th></th>
<th>Monitored</th>
<th>RI&lt;sub&gt;sc&lt;/sub&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>U (V)</td>
<td>WFS (m/min)</td>
<td>TS (cm/min)</td>
<td>CTWD (mm)</td>
<td>U&lt;sub&gt;mean&lt;/sub&gt; (V)</td>
</tr>
<tr>
<td>100%CO₂</td>
<td>21</td>
<td>3.60</td>
<td>20.8</td>
<td>9</td>
<td>20.1 ± 0.2</td>
</tr>
<tr>
<td></td>
<td>24</td>
<td>3.60</td>
<td>20.8</td>
<td>9</td>
<td>23.1 ± 0.1</td>
</tr>
</tbody>
</table>

Table 4 lists the CO, CO₂ and O₂ contents measured in each condition described in Table 3. Again, note that the data collection time did not influence the results and that the standard deviation of each measure was very low, validating the quality of the experiment. The percentage of O₂ in the measurement zone remained at about 11%, confirming that a little over 50% of the measured atmosphere was composed of environmental gases. The results of the comparison of the two conditions indicate that the voltage (or arc length or transfer stability) did not influence the generation of CO and CO₂ gases.

However, in this experiment, it was only possible to set two voltages, since a higher voltage would require the use of a CTWD too small to maintain the same desired average current. Thus, the alternative chosen was to increase the current and attempt to achieve three levels of transfer stability. To this end, another average welding current was arbitrarily chosen as I<sub>m</sub> = 190 ± 2 A (keeping the power source inductive factor set at the
intermediate position). To maintain the same volume per unit length of weld bead in the experiments with lower current, the welding speed was adjusted to maintain the same ratio of welding speed to wire feed speed. The higher welding speed (less time to produce the same deposit) would have been irrelevant because the gas was measured continuously and not cumulatively. Similarly, the respective regularity indices (RI$_{sc}$) were determined by sweeping the voltage settings from 22 to 26 V, as shown in Table 5. Table 6, in turn, lists the values RI$_{sc}$ of the volume fraction of the gases of interest in the composition of the atmosphere in which the sensor was inserted.

Table 4. CO, CO$_2$ and O$_2$ concentrations measured in the proximity of the sensor as a function of welding time (effect of transfer stability – nominal current of 175 A).

<table>
<thead>
<tr>
<th>Shielding gas (14 l/min)</th>
<th>Voltage (V)</th>
<th>Time (s) / Concentration (Vol.%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>CO$_2$</td>
</tr>
<tr>
<td>100%CO$_2$</td>
<td>21</td>
<td>38.4 ± 2.4</td>
</tr>
<tr>
<td></td>
<td>24</td>
<td>38.9 ± 1.4</td>
</tr>
</tbody>
</table>

Table 5. Test parameters to determine the concentrations of CO$_2$ and CO (effect of transfer stability – nominal current of 190 ± 2 A).

<table>
<thead>
<tr>
<th>Shielding gas (14 l/min)</th>
<th>Voltage (V)</th>
<th>Settings</th>
<th>Monitored</th>
<th>RI$_{sc}$</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>U (V)</td>
<td>WFS (m/min)</td>
<td>TS (cm/min)</td>
</tr>
<tr>
<td>100%CO$_2$</td>
<td>22</td>
<td>24</td>
<td>26</td>
<td>12</td>
</tr>
<tr>
<td></td>
<td>24</td>
<td>24</td>
<td>26</td>
<td>13</td>
</tr>
<tr>
<td></td>
<td>26</td>
<td>24</td>
<td>26</td>
<td>10</td>
</tr>
</tbody>
</table>

Table 6. CO, CO$_2$ and O$_2$ concentrations measured by the sensor as a function of welding time (effect of transfer stability – nominal current of 190 ± 2 A).

<table>
<thead>
<tr>
<th>Shielding gas (14 l/min)</th>
<th>Voltage (V)</th>
<th>Time (s) / Concentration (Vol.%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>CO$_2$</td>
</tr>
<tr>
<td>100%CO$_2$</td>
<td>22</td>
<td>32.0 ± 9.8</td>
</tr>
<tr>
<td></td>
<td>24</td>
<td>34.7 ± 5.2</td>
</tr>
<tr>
<td></td>
<td>26</td>
<td>33.0 ± 3.8</td>
</tr>
</tbody>
</table>

Tables 5 and 6, respectively, also describe the parameterization and monitoring results of new experiments and the levels of gases generated. As can be seen, the highest voltage (longest arc) led to the lowest RI$_{sc}$ (greatest regularity), ensuring the three desired levels of regularity. The fact that the uniformities were not repeated at U = 24 V between Tables 3 and 5, as they were between Tables 1 and 3, was attributed to the difference in the welding currents.

Like with lower current, the concentrations of CO$_2$ and CO did not change as a function of voltage, i.e., the voltage (or arc length or transfer stability) did not have a significant effect on the generation of CO and CO$_2$ gases by the arc, at least in the short-circuit transfer mode. Therefore, unlike fume emissions (as shown by Meneses et al. [1]), the voltage (or arc length or transfer stability) has no effect on the generation of CO and CO$_2$ gases. However, the standard deviations of the three measurements listed in Table 6 were slightly higher than those in Table 4, possibly because of the greater turbulence caused by the higher current and welding speed. Moreover, even taking into account the standard deviations upon comparing Tables 4 and 6, it is interesting...
to note that there is a tendency for CO$_2$ to decrease and O$_2$ to increase when welding with higher currents. This finding supports the hypothesis of greater turbulence of the atmosphere in the measured region, although it could also be the effect of the greater decomposition of CO$_2$ due to greater heat/radiation. Therefore, as the purpose of the experimental test was not to evaluate the effect of current, this finding is indicative rather than conclusive.

2.4. Determination of CO and CO$_2$ concentrations in different welder exposure zones in semi-automatic welding in an open environment

For this determination, the welds were made by a welder (in semi-automatic mode) and the gases were collected in three positions, namely, with the sensor inside the helmet in the welder’s breathing zone (Figure 2), with the sensor placed over the arc, ahead of welder’s helmet (Figure 3), and with the sensor positioned at the level of welded plate (Figure 4). Therefore, the idea was to evaluate the presence of CO and CO$_2$ in the key zones of the welder’s workspace, in an unconfined mode yet without forced exhaustion. The gas concentrations were measured after 30 to 60 s of continuous welding. The welds were made using 100%CO$_2$ shielding gas, with a nominal current of $I_m = 175 \pm 2$ A, in the condition of highest transfer stability (24 V). Each condition was replicated once.

Figure 2. Sensor inside the helmet in the welder’s breathing zone: (a) without arcing; (b) with arcing.

Figure 3. Sensor placed ≈ 300 mm above the plate and over the arc: (a) without arcing; (b) with arcing.
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Figure 4. Sensor placed on the plate (same plane as the arc) at \( \cong 300 \, \text{mm} \) from the torch on the plane of the plate: (a) without arcing; (b) with arcing.

Table 7 lists the set and monitored test parameters, while Table 8 shows the fraction of gases generated in the atmosphere of the measurement zone. Note some variance between the regularity indices (RI\textsubscript{ac}) of the three welding conditions, which is justified by the fact that they were semi-automatic welds (involuntary variation in the distance between torch and workpiece). Even so, the nominal current values were maintained. As in the preceding cases, the welding time for the collection of gases did not affect the measurements.

In Table 8, note that there were no significant CO and CO\textsubscript{2} concentrations in an open environment (unconfined), be it in the region of suction or close to the weld. Although these gases are generated in the arc zone, as described in sections 2.2 and 2.3, a hypothesis to explain why they disperse in the environment rather than accumulating

**Table 7.** Test parameters to determine the concentrations of CO\textsubscript{2} and CO (effect of the position of the sensor relative to the welder – 100%CO\textsubscript{2} at 14 l/min).

<table>
<thead>
<tr>
<th>Position of the sensor</th>
<th>Settings</th>
<th>Monitored</th>
<th>RI\textsubscript{ac}</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>U (V)</td>
<td>WFS (m/min)</td>
<td>TS (cm/min)</td>
</tr>
<tr>
<td>Inside Helmet</td>
<td>24</td>
<td>3.6</td>
<td>20.8</td>
</tr>
<tr>
<td>In front</td>
<td>24</td>
<td>3.6</td>
<td>20.8</td>
</tr>
<tr>
<td>Plate plane</td>
<td>24</td>
<td>3.6</td>
<td>20.8</td>
</tr>
</tbody>
</table>

Note: IM = Inside the helmet; OA = over the arc; AA = away from the arc.

**Table 8.** CO, CO\textsubscript{2} and O\textsubscript{2} concentrations measured by the sensor as a function of welding time (effect of transfer stability – nominal current of 175 A).

<table>
<thead>
<tr>
<th>Position of the sensor</th>
<th>Time (s) / Concentration (Vol.%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>30s</td>
</tr>
<tr>
<td></td>
<td>CO\textsubscript{2}</td>
</tr>
<tr>
<td>Inside Helmet</td>
<td>0.3 ± 0.4</td>
</tr>
<tr>
<td>In front</td>
<td>-</td>
</tr>
<tr>
<td>Plate plane</td>
<td>-</td>
</tr>
</tbody>
</table>
in the measurement zones is that, firstly, CO and CO\textsubscript{2} have a high capacity to diffuse in air. In other words, the diffusion coefficient of CO is 20.8 mm\textsuperscript{2}/s and that of CO\textsubscript{2} is 16.0 mm\textsuperscript{2}/s at 25 °C and under atmospheric pressure, and these coefficients increase with temperature when mixed with excess air \cite{18}. It should be kept in mind that diffusion is facilitated by higher temperatures and by the concentration gradient of such gases in the measurement zone and the environment. Secondly, when the gases are heated in the arc zone their density decreases, facilitating the effect of natural convection. The O\textsubscript{2} concentration suggests that the atmosphere in the measurement zones originated from the environment.

These findings may not be entirely in agreement with those reported by Ojima \cite{14}, who employed similar conditions to evaluate the same effect (but measured CO concentrations in ppm, reporting more refined values than those indicated here). Ojima performed MIG/MAG welding with 100%CO\textsubscript{2} shielding gas, applying currents of 100 and 300 A (the transfer mode was not mentioned). He noted that, in general, the CO\textsubscript{2} and CO concentrations varied as a function of the measurement position (much higher close to the helmet than inside the helmet) and of the welding current (higher at the higher current, although the gas flow was constant). This author also showed that forced ventilation towards the welder, even if weak, renders this difference insignificant. He pointed out that some work health standards specify that these measurements be taken at the lapel and not inside the helmet, which may not be representative. Ojima also noted that the reduction in the CO\textsubscript{2} concentration inside and outside the helmet was smaller than that of fumes. Lastly, he stated that the 8-hour Time Weighted Average (TWA) concentration during the welding period was lower than that recommended by the Japan Society for Occupational Health, but that at peaks they reach the limit specified by the National Institute of Occupational Safety and Health (NIOSH). However, a fact that Ojima did not mention directly is that fans are often used to cool the welder during welding in industrial environments.

2.5. Determination of CO and CO\textsubscript{2} concentrations in different welder exposure zones in a confined space

To define a confined environment in welding is not an easy task, especially when it comes to the generation of gases. Gases diffuse in the environment much faster than solid particles such as fumes. Moreover, gases are more sensitive to the convection effect. Therefore, welding environments are never completely confined from the standpoint of gases because, except for purposely hyperbaric chambers, they cannot be sealed due to the risk of increased pressure. This study therefore sought to simulate some of the conditions that in practice would be called confined from the standpoint of the welding fume concentrations.

2.5.1. Closed ceiling welding chamber without suction

A typical condition is welding inside tanks, vessels or containers, which are geometrically closed but not sealed. There are always doors or holes through which the gas can escape, although usually not through the ceiling. In such conditions, exhaust fans are always recommended, but extreme conditions without exhaustion are not uncommon. To simulate this condition, welds were made using a fume collector without suction, with a seal placed between the hood and the suction chamber, as illustrated in Figure 5. The gas sensor was set ± 300 mm above the plate used for welding inside the collector.

An automated short-circuit MIG/MAG weld was made using 100%CO\textsubscript{2} shielding gas, voltage set at 24 V, feed speed of 3.6 m/min, contact tip to workpiece distance of 10 mm and welding speed of 20.8 cm/min, for 120 s. These parameters resulted in an average voltage of 22.1 V, average current of 173.2 A and transfer regularity index RI\textsubscript{sc} of 1.13. Figure 6 shows a logarithmic growth in the concentration of gases in the time interval with arcing, reaching approximately 9.0% of CO\textsubscript{2} and 0.36% of CO at the end of welding. After welding, the CO\textsubscript{2} and CO concentrations fall to practically zero at the same rate. This demonstrates that during welding, if the confinement is at the ceiling (as in closed tanks), the presence of CO and CO\textsubscript{2} may be significant, but after the weld is completed, these gases are naturally diluted in ambient air.

2.5.2. Automated welding cell closed at the sides

Another typical welding environment are automated cells, which are usually open at the ceiling and closed at the bottom by the structure of the device and at the sides by safety curtains. In theory, therefore, this environment would be less confined than that described in section 2.5.1. However, since CO\textsubscript{2} is denser than air,
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Figure 5. Adaptation of a fume collector for measuring gases generated during welding in a closed chamber without exhaustion.

Figure 6. CO$_2$ and CO concentrations as a function of welding time in a closed ceiling welding chamber without suction.
unlike CO (in normal conditions of temperature and pressure, i.e., 20 °C and 1 atm, the density of air is 1,205, that of CO₂ is 1,842 and that of CO is 1,165 kg/m³), although CO₂ disperses, it could accumulate in the lower parts of the cell after long welding times before it diffuses through the environment.

Thus, long duration welding (10 minutes) using the same parameters as those described in section 2.5.1 was performed on a XYZ coordinate table. The monitored parameters showed an average voltage of 23.1 V, average current of 173.3, and transfer regularity index $R_I$ of 1.06. The gas sensor was placed in a position below the plate, as shown in Figure 7. Measurements were taken after 120 s without arcing (only the shielding gas exiting the nozzle), after 600 s of open arc, and 480 seconds after the end of welding.

Figure 7. Position of the sensor under the XYZ table of an automated welding cell (the dimensions of the zone protected by the curtain is 900 mm width × 900 mm depth × 900 mm height).

Figure 8 shows that the concentrations of CO₂ gas in the measuring zone reach up to 6.1% before the arc is struck, becoming negligible without arcing and in the arcing time intervals. The CO content showed no variations in any of the times. In other words, although large amounts of CO₂ and CO gases were generated by the arc when using pure CO₂ shielding, the CO₂ and CO gases became less dense than air when heated, rising and dispersing easily in the environment during welding and did not return after welding.

Figure 8. CO₂ and CO concentrations as a function of welding time in simulated welding in an automated welding cell protected by curtains at the sides.
Considering the possibility that gases, mainly CO\textsubscript{2}, might concentrate in the lower portions of the automated cell, but dispersed due to the non-airtight construction of the base of the welding cell, a condition was sought in which gases could not disperse through the bottom part. To this end, new welds were made inside a drum using a robot, as illustrated in Figure 9. The sheet to be welded was affixed 240 mm from the bottom of the drum and the sensor was placed on the bottom surface of the drum. The gases were measured for 120 s with arcing and for 660 s after welding. The same parameters as those described in the previous sections were used, monitoring an average voltage of 21.6 V, average current of 173.8 A and a transfer regularity index RI\textsubscript{sc} of 1.20.

Figure 10 confirms that even when prevented from dispersing through the bottom, CO and CO\textsubscript{2} gases are able to disperse upward through convection, without accumulating after welding.

![Figure 9](image1.png)

**Figure 9.** Simulation of an automated welding cell sealed at the bottom (drum with 560 mm diameter × 870 mm height).

![Figure 10](image2.png)

**Figure 10.** CO\textsubscript{2} and CO concentrations as a function of time in simulated welding in a drum with airtight bottom.
In view of the results, new measurements were taken, this time capping the drum immediately after welding. Figure 11 shows the same tendency for the non-accumulation of residual gases inside the drum during and after welding.

Finally, to evaluate the diffusion capacity of the shielding gas in a confined space, welding was simulated inside the open drum, with the release of shielding gas but without arcing. Measurements were taken over a period of 600 s, with the release of the shielding gas, and for 300 s, after the gas was shut off. The gas sensor was placed underneath the plate. Figure 12 shows that in the time interval when the shielding gas was released, without arcing, the concentration of CO$_2$ in the measurement zone reached approximately 44.5%. Of course, the presence of CO was not expected and was only measured and displayed to attest to the reliability of the measurements. When the shielding gas is shut off, the concentration of CO$_2$ gradually decreases to about 15.7%, indicating that although in theory CO$_2$ is denser than atmospheric air, it dilutes through diffusion (in this case, based on the difference in concentration) and does not accumulate in the lower regions. As, in this case, there is no heat, the diffusion is based only on the difference in concentration, leading to partial accumulation of CO$_2$ in the lower regions. A leakage of 100%CO$_2$ shielding gas, or CO$_2$ from other sources, seems to concern much more than the production of CO$_2$ during welding. In these cases, however, there is no chemical reduction of CO$_2$ to CO.

![Figure 11](image1.png)

**Figure 11.** CO$_2$ and CO concentrations as a function of time after welding simulated in a drum with airtight bottom capped immediately after welding.

![Figure 12](image2.png)

**Figure 12.** CO$_2$ and CO concentrations as a function of time after welding simulated without arcing in a drum with airtight bottom.
3. Conclusions

The results of this work lead to the conclusion that short-circuit MIG/MAG welding of carbon steel, because it allows for the use of CO\textsubscript{2} shielding gas, emits significant amounts of CO\textsubscript{2} (asphyxiating) and CO (toxic) gases in the welding region during operation. However, this emission is independent of the metal transfer stability and arc length used. The richer the shielding gas mixture is in CO\textsubscript{2} the higher this emission, reaching significant levels when using 100\% CO\textsubscript{2} as shielding gas. The use of Ar (argon) based mixtures with lower percentages of CO\textsubscript{2} or with O\textsubscript{2} proved to be an acceptable alternative also from other operational standpoints.

On the other hand, the conversion of CO in the arc in short-circuit transfer was found to be negligible, causing the emission of CO to be relatively low. Furthermore, due to the diffusion capacity of CO\textsubscript{2} and CO in atmospheric air, the emitted gases are not concentrated, at least in quantifiable levels higher than tenths and hundredths of volume percentages, after welding in the work envelope of a welder, even in spaces considered confined. Concentration of CO\textsubscript{2} in the working area can happen, however, if there is no heating to facilitate the diffusion of this gas. Other sources of CO\textsubscript{2} worry more than arc welding in confined area.

Considering these conclusions, three multidisciplinary aspects are objectives for future studies. The first would focus on workers’ health. The study would involve determining if the welder might be exposed to unacceptable levels of CO and CO\textsubscript{2} (quantified in ppm) during their emission, even considering the herein demonstrated rapid dispersion of these gases from the work area after the end of welding. The second aspect, related to the environment, would be to evaluate the contribution of the amount of CO\textsubscript{2} generated in welding processes to the enrichment of the concentration of CO\textsubscript{2} in the environment, and the consequent effects on environmental imbalance, given that welding is a widely applied manufacturing process in both industrialized and developing countries. The third aspect, of a more operational nature, would be to study the effect of other welding variables, such as current and other transfer modes, on the generation of these gases. Given the interdisciplinary nature of these studies, the natural way would be through cooperative projects supported by government and suppliers or users of the process.

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References


