

Comprehensive Review on Carbon Steels Corrosion in Chloride-Rich Media

Abstract

Corrosion is a serious engineering and economic problem that affects metals and alloys, particularly carbon steel, which is widely used in many industrial applications. This review provides a comprehensive overview of the corrosion of carbon steels in rich chloride media, starting with corrosion mechanisms and factors that influence corrosion, types of corrosion, and types of the main corrosion products (rust). It also explores the main methods used in studying and monitoring corrosion (non-electrochemical and electrochemical techniques), starting from laboratory experiments to advanced analytical techniques. Among the strategies to mitigate corrosion, the use of corrosion inhibitors plays a crucial role in reducing the corrosion rate of carbon steels. This is especially true in harsh environments rich in chloride, known for accelerating metal degradation. This review also presents previous studies on corrosion in chloride-rich waters and evaluates the effectiveness of different inhibitors in controlling corrosion, contributing to the development of efficient protective solutions for metals under harsh and aggressive conditions.

Keywords Corrosion, Carbon Steels, Chlorides Ions, Corrosion Products, Inhibitors.

1.Introduction

Metal corrosion creates a significant risk to environmental and human health. Corrosion may release hazardous metals, including lead, cadmium, and chromium, into soil and water, resulting in pollution that impacts ecosystems and penetrates the food chain. Moreover, infrastructure deterioration can lead to dangerous leaks, impacting water quality and presenting immediate health hazards. Corrosion is not only a matter of material degradation; it also poses significant environmental and public health risks.

The corrosion of metals significantly affects several industrial sectors [1–5]. Metals, such as steels and alloys, have been widely utilized in industrial and technical applications. Moreover, metals are utilized in the offshore petroleum, power generation, nuclear energy, aerospace, shipping, and building industries [6–11]. Corrosion is the irreversible deterioration of a metal surface as a result of chemical reactions, whereby pure metal is converted into more chemically stable forms such as metal oxide, hydroxide or oxyhydroxide within a corrosive environment. These environments can exist in solid, liquid or gaseous states and are commonly referred to as electrolytes [12]. Corrosion is a surface process that occurs on the outer layer of the metal and manifests itself in several forms: An initial attack on the surface may cause general corrosion, leading to a gradual reduction in thickness, or it may cause localized damage, leaving only specific areas corroded, as sometimes seen at grain boundaries or in areas of weakness resulting from differences in the metal's resistance to the corrosive environment [13]. This process is slow, but over time it causes the deterioration of metal machinery and equipment, reducing their production efficiency. The annual economic losses resulting from various forms of corrosion in India are estimated at US\$6.5 billion, while direct waste in the United States amounts to about 3.2% of gross domestic product [14]. A good understanding of corrosion and the application of appropriate and timely solutions are essential for controlling it [15]. Furthermore, a special type of corrosion can occur in relatively pure water at temperatures reaching around 1300°C on the product side, known as classic hot spot corrosion [16]. Corrosion is widely recognized as a universal phenomenon. Carbon steels (CS), due to their excellent mechanical properties [17–19]. In addition to the possibilities for processing (welded

[20], chipped [21], deformed [22]), are an appropriate choice for use in the manufacture of different parts of machines [23], accessories of fall arrest systems such as carabiners, hooks, and pythons [24,25], vehicle bodies [26], shipbuilding [27], or in use in buildings, bridges, rails, water, gas, and industrial pipes, cooling tower parts [28-31].

This study focuses on carbon steels with carbon at or below ~0.35% to facilitate welding. A subsequent differentiation can be established based on carbon content. Low carbon steels (containing fewer than 0.15% carbon) possess insufficient carbon to undergo hardening and are commonly utilized in hot-working or to achieve maximum ductility in the annealed state. Steels containing less than 0.25% carbon, commonly known as mild-steel, exhibit increased strength as they approach the maximum limit of carbon content. Medium carbon steels (0.25–0.55% carbon) are frequently subjected to heat treatment (quenching and tempering) to attain greater strength; however, this review focuses mainly on compositions below 0.35% carbon [32]. CS may get corroded during industrial operations; thus, corrosion inhibition sometimes is crucial for safe and economical operations for prolonged use of CS. This review examines CS as the most suitable alloy for constructing pipelines that transport water, oil, and gas. The majority of corrosion problems in the water, the oil and gas industries are related to pipelines and their exposure environments of CS, which determine the appropriate kind of CS selection. The present article offers an extensive examination of the corrosion characteristics of CS in saline solutions, emphasizing the fundamental electrochemical processes, significant influencing variables, and different experimental methodologies employed to evaluate corrosion rates. Furthermore, it analyzes various mitigation measures, including the use of corrosion inhibitors. This study examines recent research to identify contemporary difficulties, knowledge deficiencies, and emerging trends in corrosion science. The insights provided attempt to enhance the effectiveness as well as sustainability of corrosion prevention techniques, especially in shipping, industrial, and oil and gas sectors.

2. Corrosion mechanism

Corrosion is characteristics due to chemical (dry corrosion) or electrochemical (wet corrosion) interactions with its surroundings. Nonmetals are excluded from the current definition [32]. The breaking down of reactions into partial oxidation and reduction processes is referred to as electrochemical [33]. For corrosion, these reactions must be performed [34]:

Corrosion happens at the anode. At the point of the anode, oxidation or the loss of electrons from the metal occurs. The metal with the greater reduction potential is often referred to as the anode. The anodic metal is transformed into its equivalent ion by the loss of electrons.



At the cathode, reduction occurs, wherein the metal receives electrons from the anode. This may occur through the evolution of hydrogen and/or the absorption of oxygen.

2.1 By hydrogen evolution

It is the corroding process which hydrogen is released during it. The process often occurs in an acidic or alkaline media, when (H^{+}) ions absorb anodic electrons and release hydrogen.



For example, Fe metal reaction at anode:



And reaction at cathode:



Overall cell reaction:



2.2 By absorption of oxygen

It is the corrosion process involving the absorption of oxygen. This typically occurs in a neutral liquid, with NaCl being employed as the conducting medium, as seen in Fig. 1 in which (O₂) is absorbed to give (OH⁻) ions.



For example, at anode the carbon steel reaction is:



And at cathode is:



Overall reaction:

Formation of Fe²⁺ ions at anode and OH⁻ ions released at cathode which come to form iron hydroxide [Fe(OH)₂] [35].

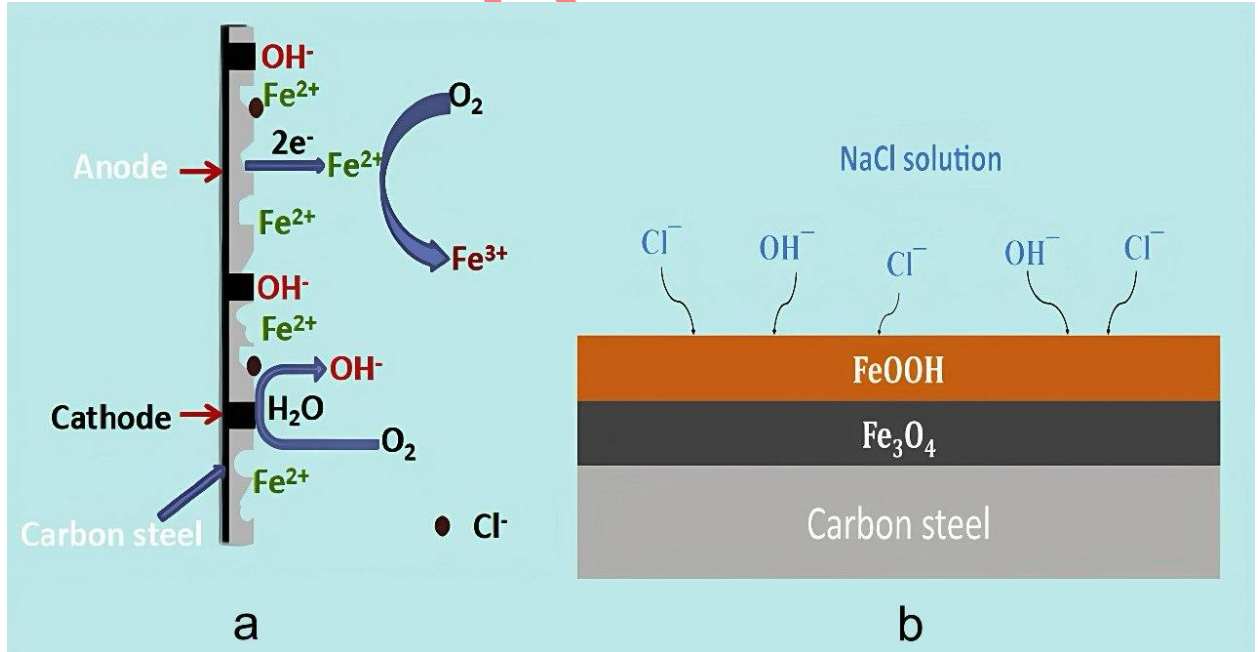


Fig. 1. Schematic representation of carbon steel immersed in NaCl solution
(a) Mechanism [36], (b) Corrosion layer formation on the steels surface

3. Factors affecting corrosion process

Numerous internal and environmental variables directly or indirectly influence corrosion; some of these elements are illustrated in Fig. 2. We will focus on the nature of the metal and natural corrosion environment. Additional elements influencing corrosion include metal purity, characteristics of the surface coating, properties of the corrosive product, temperature, concentration of dissolved salts in the a medium, air humidity, and the pH of the electrolyte.

3-1 Nature of alloy or metal

It further depends upon:

3-1-1 Galvanic series position: When different metals are electrically connected in a specific electrolytic medium, the metal with higher oxidation potential, or the one higher in the electrochemical series, corrodes, while the other metal remains protected. The rate of corrosion depends directly on the electrochemical difference between the two metals; the greater the difference, the faster the corrosion. The susceptibility of metals to corrosion is shown in Fig 3. For example, in the presence of copper and cast iron in the same electrolytic medium, the cast iron corrodes first while the copper is protected from corrosion.

3-1-2 Purity of alloy or metal: The corrosion rate typically increases with rising impurity levels. This occurs because impurities create small electrochemical cells in which the anodic component undergoes corrosion. For instance: Zinc with impurities such as iron or lead exhibits accelerated corrosion.

3-1-3 Nature of the surface film: A thin layer of oxides forms on the surface of all metals in aerated environments. The effectiveness of this layer is determined by the volume ratio between the metal oxide and the original metal, known as the 'specific volume ratio.' In light of this, the higher this ratio, the lower the rate of metal oxidation.

3-1-4 Nature of corrosion product: The rate of corrosion accelerates when the corrosion product is soluble in the corrosive medium, as this allows the reaction to continue on the metal surface. The volatile product evaporates immediately after formation, exposing the metal to further attack and thus increasing rust formation.

3-1-5 Alloy or metal grain size: The mechanical characteristics of low carbon steel are predominantly influenced by ferrite particle size, with finer grains being advantageous for mechanical characteristics [37-39]. Nevertheless, the high energy and chemical reactivity of grain boundaries result in a larger density of these boundaries, which enhances surface reactivity through improved electron activity and diffusion, so influencing corrosion resistance. Enhanced corrosion resistance leads to prolonged durability of steel buildings [40–45]. Consequently, examining the effects of grain size on the corrosion resistance of low carbon steel is a significant topic.

3-2 Nature of corrosive environment

It further requires:

- *High temperature:* The rate of corrosion increases sharply with rising temperature, doubling with every 10 °C increase, provided that other biological and chemical factors remain constant [46]. This behavior is often depicted as an exponential curve showing the direct effect of heat, but the reality is more complex, as thermal changes also affect the effectiveness of other factors such as ion transport, gas dissolution (especially oxygen), and the properties of the protective layer formed on the metal surface. The effect of increased temperature is evident in two main ways: First, by accelerating electrochemical reactions thanks to increased molecular energy and more frequent effective collisions, and second, by modifying intermediate factors, such as solubility, ion transport, and microbial activity adding further depth and complexity to the underlying exponential curve.
- *Media humidity:* The rate of corrosion increases significantly when a certain level of relative humidity, known as critical humidity, is exceeded, as the ability of the iron oxide layer to absorb water from atmospheric humidity enhances electrochemical corrosion. The available moisture provides the metal surface with the electrolyte necessary to form an effective electrochemical cell, which accelerates oxidation and recovery reactions at the metal interface.
- *Effect of dissolved salts in media:* The rate of corrosion initially increases with increasing salt content, then gradually decreases until it becomes lower than the rate of corrosion in distilled water once the concentration reaches saturation. Different types of salts such as sodium chloride, alkali metal salts, alkaline earth metal salts, and acid salts affect the corrosion behavior of iron and steel in various ways related to their ability to alter the properties of the oxide layer and the conductivity of ions in solution [47].
- *Effect of pH in media:* pH is the most important factor in determining the rate of corrosion; a decrease in pH is usually accompanied by a significant increase in the rate of corrosion, indicating that acidic media (with a pH below seven) are more corrosive than neutral or alkaline media [48].
- *Effect of dissolved oxygen in media:*
Dissolved oxygen plays a critical and complicated function in the corrosion of metals. Oxygen participates in cathodic processes on the metal surface in neutral, alkaline, and acidic environments. Therefore, the presence of corrosion is necessary for its occurrence. In the absence of dissolved oxygen, corrosion in neutral and alkaline solutions diminishes to nearly nothing. An increase in dissolved oxygen content, due to its participation in cathodic processes, promotes corrosion. What would occur if we were to inject increasing amounts of water infused with oxygen? It was established that oxygen, under specific conditions (in high-purity water) and elevated temperatures, may lead to the production of a thick passive protective coating of metal oxides on the metal surface, hence reducing corrosion [49].

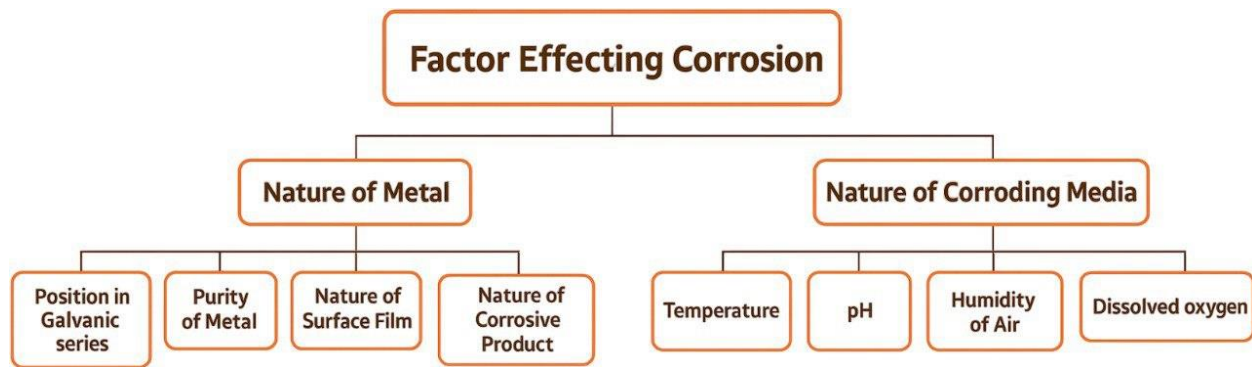


Fig. 2. Some of the factors affecting corrosion process

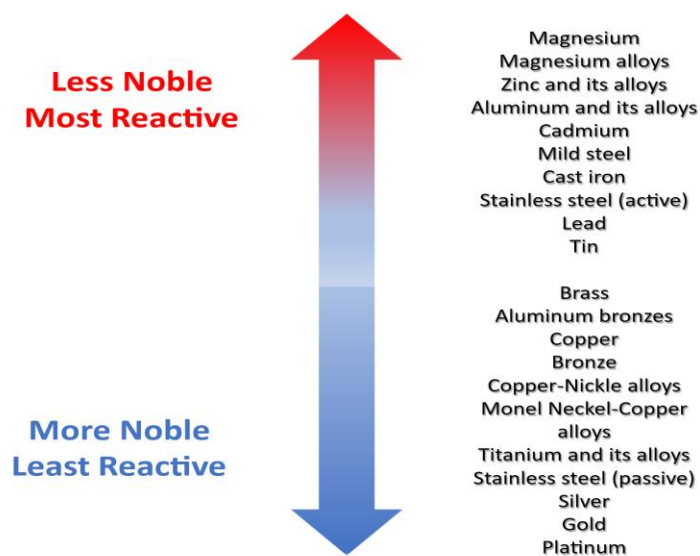


Fig. 3. Corrosion susceptibility of the metal

4- Methods for studying corrosion

4-1 Non-electrochemical technique (weight loss method)

This is a gravimetric technique where the weight of a carbon steel sample is measured before and after exposure to a corroding environment (e.g., saline solution). The difference in weight is used to calculate the corrosion rate according to the following equation [50]:

$$\text{Corrosion rate} = \frac{K W}{D \times T \times A} \quad (10)$$

The corrosion rate can be characterized as an increase per unit in the depth of corrosion over time, measured in mils per year (mpy) or as weight loss per unit area per time, often in milligrams per square decimeter per day (mdd) or as the corrosion current ($\text{mA} \cdot \text{cm}^{-2}$). The standard SI metric for denoting the corrosion rate is millimeter per year (mm/y) or inch per year (inch/y). Table 1 illustrates the resistance of corrosion classification of CS.

Table 1. The comparison resistance of corrosion classification of CS [51]

Relative corrosion resistance	Approximate corrosion rate	
	mpy	mm/y
Outstanding	More than 1	More than 0.02
Excellent	1-5	0.002-0.1
Good	5-20	0.1-0.5
Fair	20-50	0.5-1
Poor	50-200	1-5
Unacceptable	Less than 200	Less than 5

Many researchers (52-58) have been measured the corrosion rate of different types of carbon steel in different chloride media at 25°C of immersion as shown in Table 2.

Table 2. Corrosion rates of various types of carbon steels in different chloride media

Types of carbon steel	Type of salt solution	Total Immersion Time	Corrosion Rate	Reference
Low carbon steel	Marine water	24 months	500 mm/y	[52]
Low carbon steel	Seawater	28 days	0.037 mm/y	[53]
Low carbon steel	Lake Water	28 days	0.045 mm/y	[53]
Low carbon steel	Tap Water	28 days	0.04 mm/y	[53]
Medium carbon steel	Banda Aceh's	12 months	0.015 mpy	[54]
Carbon steel	Seawater	288 hours	0.18 mm/h	[55]
Carbon steel	Beibu Gulf Tidal Zone	360 days	600 mm/day	[56]
Mild Steel	1% NaCl	28 days	0.788 mpy	[57]
Mild Steel	3% NaCl	28 days	0.687 mpy	[57]
Mild Steel	5% NaCl	28 days	0.360 mpy	[57]
Mild Steel	7% NaCl	28 days	0.779 mpy	[57]
Mild Steel	10% NaCl	28 days	0.315 mpy	[57]
Mild Steel	Salt water	5 weeks	1.0622 mm/y	[58]

4-2 Electrochemical Techniques for Studying Corrosion of Carbon Steel

The most common employed electrochemical techniques are the potentiodynamic polarization technology (Tafel) combined with electrochemical impedance spectroscopy (EIS) [59,60]. The primary benefit of electrochemical methods exceed weight loss techniques is the ability to investigate additional information relating to corrosion mechanisms. Electrochemical procedures are typically conducted using a three electrode established consisting of a working electrode, a reference electrode, and a counter electrode [61]. The impact of corrosion inhibitors has been investigated by modifying the extract concentration, testing temperature, and liquid flowrate. Electrochemical analysis is efficient, rapid, and simple to use [62]. Potentiodynamic polarization yields many electrochemical characteristics, including corrosion potential ($E_{corr.}$), current density of corrosion ($I_{corr.}$), anodic slope (β_a), and cathodic slope (β_c). Electrochemical impedance spectroscopy is capable of accurately estimating the rate of corrosion with minimal impact on the electrode. Different electrochemical techniques exist to assess the process and corrosion degree of the metal, as well as the efficiency and mechanisms of corrosion inhibitors to a certain extent.

4-2-1 Potentiodynamic Polarization (Tafel)

The potentiodynamic polarization technique is widely employed and consists of polarization current density and electrode potential [63,64]. Polarization curves (Anodic and cathodic) represent the relationship between polarization electrical current density and electrode potential. However, when the potentiodynamic polarization utilizing the potentiodynamic polarization curve technique is excessively high, significant curve deviations arise, preventing from getting hold of a standard potentiodynamic polarization curve. Fig. 4 illustrates a diagram of the potentiodynamic polarization curve.

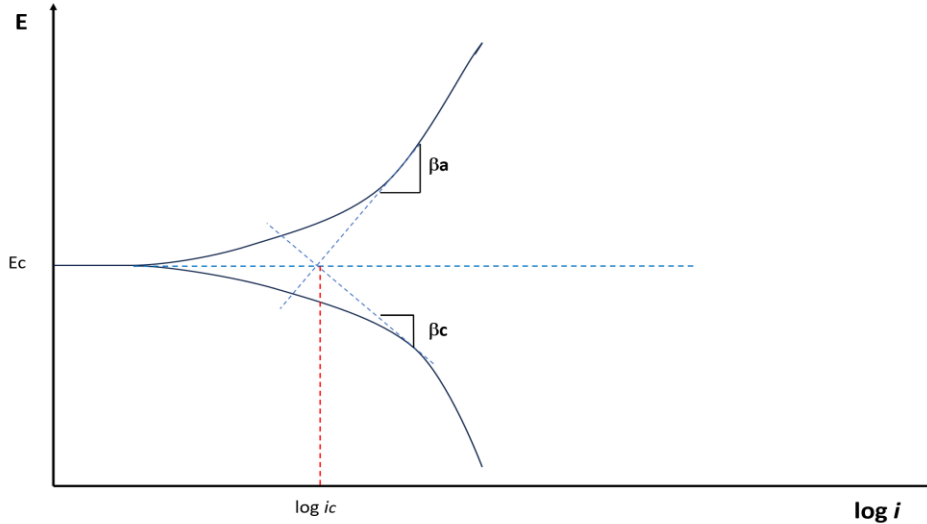


Fig. 4. Potentiodynamic polarization curve diagram

4-2-2 Electrochemical Impedance Spectroscopy (EIS)

Electrochemical Impedance Spectroscopy (EIS) quantifies the impedance spectra of an electrode system throughout an extensive frequency range, employing a small amplitude sinusoidal potential as the perturbation signal to provide a nearly linear correlation between the system's responses. EIS is a prevalent testing method that little affects the electrode surface condition. Consequently, it is extensively utilized in scientific research and constitutes a significant methodology in electrochemical testing approaches [65,66]. In the analogous circuit, R_s denotes the resistance of electrolyte solution between the working electrode and reference electrode, R_{ct} signifies the charge transfer resistance associated with the corrosion process at the metal-electrolyte interface, and CPE represent the constant phase element. In corrosion science, EIS provides data on corrosion resistance, variations surface roughness of electrodes, adsorption of corrosion inhibitors, and the generation of corrosion products. In contrast to other conventional electrochemical techniques, such as potentiodynamic polarization curves, EIS offers enhanced insights into corrosion kinetics as seen in Fig. 5.

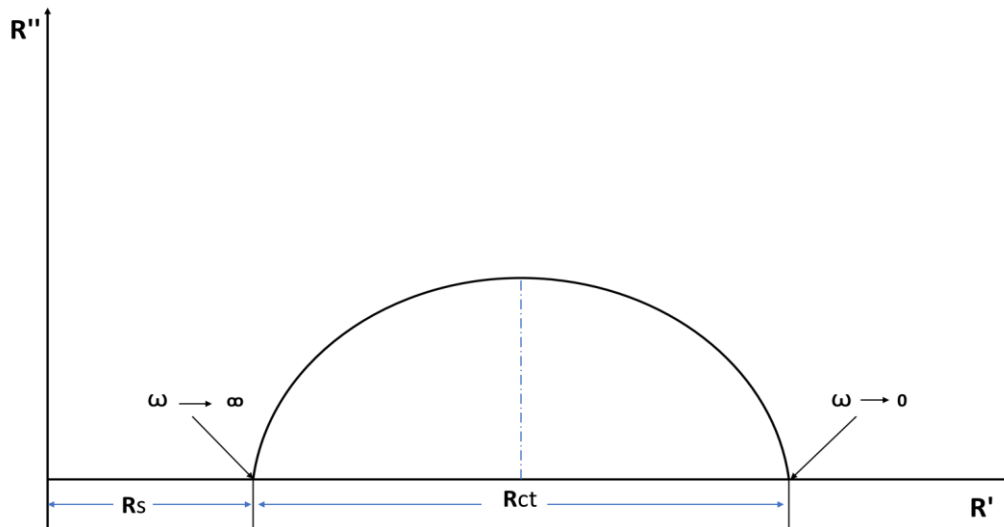


Fig. 5. Schematic diagram of electrochemical impedance spectroscopy

4-3 Surface characterization techniques

This portion of the article examines surface analytical approaches for characterizing carbon steel surfaces, both treated and untreated (e.g., corrosion inhibitors) in corrosive environments. The discussion addresses aspects affecting the use of certain approaches in carbon-steel corrosion research, including high surface roughness under corrosive circumstances [67].

4.3.1. Scanning electrochemical microscopy (SEM)

Employing scanning electron microscopy (SEM), an accurate imaging technique capable of giving high-resolution images of metallic surfaces, enables a comprehensive examination of corrosion, protective films, and the underlying substrates of metals [68,69]. This approach involves directing an electron beam at a sample, resulting in the production of secondary electrons by surface contact. The electrons are subsequently collected to provide a high-quality image [70]. A key benefit of scanning electron microscopy is its flexibility, allowing application in environmental chambers, both high and low vacuum environments, and the investigation of corrosion processes under different conditions [71]. SEM may be utilized with various techniques, such as X-ray spectroscopy, for the chemical evaluation of corrosion products to determine the kind of corrosion and its source. SEM is an essential tool for examining corrosion; it may potentially identify localized corrosion sites and yield data on the concentration of active corrosion species, hence advancing study into localized corrosion at particular areas. It also offers helpful understanding into inhibitors to improve protective methods and enhance the useful lifetime of metal structures. Research evaluated the performance of a newly developed organic corrosion inhibitor by SEM testing.

4.3.2 Energy dispersive X-ray spectroscopy (EDX)

EDX is a chemical analysis method that employs a high-energy electron beam to target substances, excite electrons from inner shells, and concurrently emit X-ray photons. The energy content of these photons is characteristic to the emitting element, facilitating the examination of the sample's chemical structure. SEM-EDX is a high magnification, stereoscopic microscopy technique employed for the analysis of healing products. It employs a concentrated high energy

electron beam to examine materials, revealing physical information on the surface topography of the specimen. This technique is frequently employed to ascertain crystal structure, shape, dimensions, and distribution. The interface composition is assessed by backscattered SEM images. SEM/EDX is a method that offers an optical representation of a surface together with the elemental composition of selected regions [72-76].

4-3-3 Atomic force microscopy (AFM)

Atomic force microscopy facilitates the examination of surface topography, involving measures of surface roughness. Assessing the adhesion of the inhibitor or corrosive coating on the substrate (metal or alloys) only using topographical imaging has consistently posed challenges. Moreover, capturing topographical photos might be challenging due to the irregular film development. Consequently, AFM current pictures and force versus distance curves are employed for providing further insights into the characteristics of surface layers, such as the adherence of corrosion inhibitors and coatings [77,78]. AFM offers the capability to function in different liquid conditions and facilitates real-time surface monitoring in corroding environments [79,80].

4-3-4 X-ray diffraction spectroscopy (synchrotron-sourced)

Grating-induced X-ray diffraction (GI-XRD) is an effective method for studying interfacial chemistry at the solid-liquid interface, but its traditional application suffers from a low signal-to-noise ratio that can lead to inaccurate conclusions about phase evolution. This challenge is exacerbated when analysing thin films—such as corrosion inhibitor layers—on carbon steel, as surface roughness in corrosive environments contributes to a reduction in the received signal. Synchrotron-radiation-induced X-ray diffraction (SR-XRD) addresses these limitations by providing a high signal-to-noise ratio sufficient for examining thin films on carbon steel even under aggressive corrosion conditions. SR-XRD has high resolution due to the use of a parallel beam that reduces peak broadening at low angles, as well as high brightness, high intensity, tight collimation, and low emission polarisation, making it the preferred choice of crystallographers [81].

It is also necessary to monitor the transformations undergone by corrosion products under different conditions and to conduct in situ corrosion tests. In carbon dioxide environments, where oxygen is consumed, it has been observed that carbon steel produces iron carbonate as a by-product, which may oxidise when subsequently exposed to oxygen-rich conditions. It has also been demonstrated that combining field SR-GI-XRD with additional electrochemical methods such as cyclic voltammetry and electrochemical impedance spectroscopy provides a deeper mechanistic understanding of electrochemical reactions on copper surfaces [82,83].

4-3-5 X-ray photoelectron spectroscopy (XPS)

XPS is a quantitative spectroscopic method that provides information on the elemental composition at parts-per-thousand levels depending on the electronic and chemical state of the surface being examined. This technique has been widely used in corrosion research to describe the composition of metal surfaces, particularly in the study of corrosion products. For example, researchers applied it to carbon steel exposed to seawater, and three-dimensional analyses revealed the absence of an iron signal in the spectrum, indicating that the surface was completely covered with a layer of corrosion products [84]. The presence of iron in its metallic and oxidised states was associated with the production of oxides, and techniques such as Raman spectroscopy and X-ray diffraction confirmed these results. In addition, XPS was used in the qualitative analysis of surface

films to measure the effectiveness of protective layers, such as studying the formation of iron carbonates on carbon steel surfaces [85] and examining the presence of hydroxide ions within the layer to assess its cohesion and protective properties [86].

4-3-6 Raman spectroscopy (IR)

Raman spectroscopy (IR) has been acknowledged as a crucial surface characterization technique for corrosion investigations, applicable for assessing both single layer and multilayer surface characteristics. Raman spectroscopy offers insights into surface connecting (vibrational data), hence giving structural information regarding the contact between the metal substrate and corrosion inhibitor [87].

4-3-7 Fourier transform infrared spectroscopy (FTIR)

Unique molecular fingerprints can be formed based on the absorption and transmission of infrared light. The quality and quantity of individual components in samples may be determined using FTIR. IR recognized as the dispersive IR method, was employed in previous investigations. Currently, FTIR is widely utilized because to its several benefits, including accurate measurement, rapid data collection, high sensitivity, non-destructive analysis, and the absence of a need for external calibration. Conventional FTIR, usually examined in the mid IR wavelength range, has been effectively utilized in the analysis of surface coatings on CS substrates. The existence or nonexistence of certain bands of the corrosion inhibitor molecule correlates with the molecule's orientation on the CS surface and/or inside the film of inhibitor. This helps in understanding how corrosion inhibitors adsorbed on steel surfaces [88].

To evaluate and examine the corrosion of metals and alloys, monitoring alterations in the color of electrolyte solutions may serve as a simple way, but alternative approaches such as the Standard Immersion Test (SIT) should also be studied. Fig. 6 illustrates the flowchart of the experimental approach based on SIT. At the conclusion of the SIT, the corrosion behavior of metals subjected to corrosive solutions is assessed by measuring the corrosion rate (mm/y).

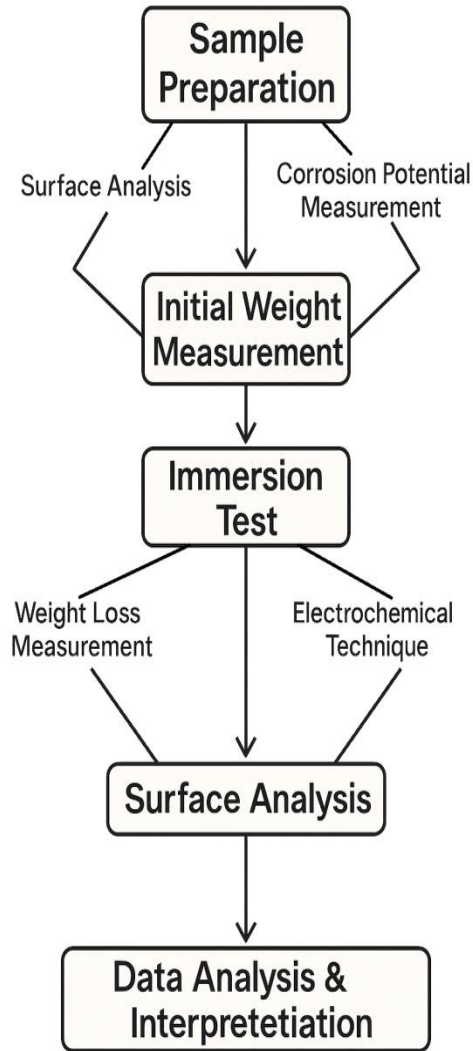


Fig. 6. Flowchart of an experimental methodology employed to examine corrosion behavior of steels

5. Types of corrosion

Corrosion interactions are classified into two types based on the characteristics of the corrosive environments: wet corrosion and dry corrosion. Corrosion can be categorized into several types [89-97] as shown in Fig. 7, they depend on the environment, the substrate type, or the chemical procedure.

5-1 General Corrosion: It is also known as uniform corrosion, is distinct from dry corrosion; this type of corrosion degrades the metal's untreated surface very consistently. It may also be characterized as a form of corrosion that progresses uniformly throughout the entire exposed surface. Oxygen serves as the primary catalyst for this corrosion. The main materials exhibiting general corrosion are steel and cast iron. When subjected to a humid environment, they exhibit a rust-like appearance.

5-2 Intergranular Corrosion: Intergranular corrosion is highly pertinent to the brewing industries. It involves a localized attack, wherein a restricted pathway is preferentially corroded along the grain boundaries of a metal. This form of corrosion can significantly impact mechanical qualities, leading to decreases in strength and ductility.

5-3 Galvanic Corrosion: Galvanic corrosion often arises when different conductive materials are electrically interconnected and subjected to an electrolyte. Consequently, the subsequent essential criteria must be satisfied for galvanic corrosion:

1. Various metals or alternative conductors, such as graphite.
2. Electrical contact between dissimilar conductive materials, which may occur by direct contact or an alternative connection, such as a shared grounding pathway.
3. Electrolyte (the corrosive media) in contact with different conductive materials.

5-4 Pitting Corrosion: Pitting corrosion is considered more hazardous than uniform corrosion damage because to its challenging detection, prediction, and design reduction. Pitting corrosion can result in pits that are either open or enveloped by a semipermeable barrier of corrosion byproducts. Pits can be either hemispherical or cup-shaped [97].

5-5 Crevice Corrosion: This type of corrosion is represented a localized type. Crevice corrosion happens in small voids or interstices between two metallic surfaces or between metals and non-metallic surfaces. A concentration cell is established with the crevice lacking oxygen. The differential aeration between the cracking (micro-environment) and the exterior surface (bulk environment) imparts an anodic aspect to the crevice. This may lead to a very corrosive environment within the cracks. Instances of crevices involve flanges, deposits, washers, ends of rolled tube, and threaded connections as shown in Fig. 8 [98].

5-6 Erosion Corrosion: Erosion Corrosion is the accelerated degradation or attack on a metal resulting from the relative motion between a corrosive liquid and the metal surface.

5-7 Stress Corrosion Cracking (SCC): It denotes cracking resulting from the simultaneous existence of tensile stress and a corroding environment. The influence of SCC on a material often occurs between dry cracking and the material's fatigue threshold [99].

5-8 Selective Leaching: Also referred to as "parting" or "dealloying," it is the specific elimination of one element from a steel alloy, resulting in a modified residual structure. A common instance is the selective extraction of zinc from brass alloy, known as dezincification as shown in Fig. 8 [100].



Fig. 7. Different corrosion patterns in carbon steels manufactured parts as observed at the thermal power station and refinery of Al-Dora-Baghdad, Iraq



Fig. 8. Crevice corrosion and selective leaching [98,100]

6-The types of corrosion product producing on CS

The corrosion products that develop on the steel alloy surface are the principal results of the metal's dissolution. They can significantly influence the current corrosion process. Primarily, they establish a physical barrier between the alloy and the environment, therefore protecting the metal by preventing the transfer of dissolved oxygen from saltwater to the metal surface [101,102]. Secondly, their porous nature provides a unique home for micro-organisms that may exist in a particular environment [103-107]. Thirdly, some phases are used as electronic conductors, such as magnetite [108,109] and iron sulfides [110,111], which can enhance galvanic cells. The

composition of the corrosion product layer varies according to the exposure zone and may alter over time. The layers, developed on steel alloy surfaces during laboratory tests or at seaport exposure locations, illustrate the complex nature of iron chemistry in natural saltwater. Their composition varies in regions (anodic and cathodic), hence actively contributing to the maintenance of corrosion cells and promoting localized corrosion processes [112].

Many research investigations have been performed on corrosion products that developed in different atmospheres utilizing XRD and IR. The corrosion product of CS typically consists of crystalline compounds of iron oxide, hydroxide, and oxide-hydroxides [113,114]. The shape and dimensions of the crystals created are depending upon the varying conditions under which they create. The chemical composition [115-121] and morphologies of solid corrosion products were examined according to characteristics given in Table 3. Some of crystals are shown in Fig. 9.

Table 3. Chemical composition and morphology of corrosion products [54,122-129]

Corrosion product	Composition	Crystal structure	Morphology
Goethite	α -FeOOH	Orthorhombic	Cloudy-shaped, flat and thin sheet, Needle-shaped, filiform, whiskers, bipyramids, cubes, thin rods cotton balls, tiny rods, nest-like, star-like, cotton ball.
Lepidocrocite	γ -FeOOH	Orthorhombic	Dense plates, granular, thick sheet Laminar, spherical, sandy granules, worm burrow, birds nest, plumage or shattered glass, floral, sandy mixture.
Hematite	α -Fe ₂ O ₃	Hexagonal	-
Feroxyhyte	δ -FeOOH	Hexagonal	Flowery, bent plates.
Akageneite	β -FeOOH	Monoclinic	cylinder, tube, cigar-shaped, plate-like morphology.
maghemite	γ -Fe ₂ O ₃	Cubic	Dark flat layer, circular grain, donuts-like, Black circular rings.
Magnetite	Fe ₃ O ₄	Cubic	-
Wustite	FeO	Cubic	-

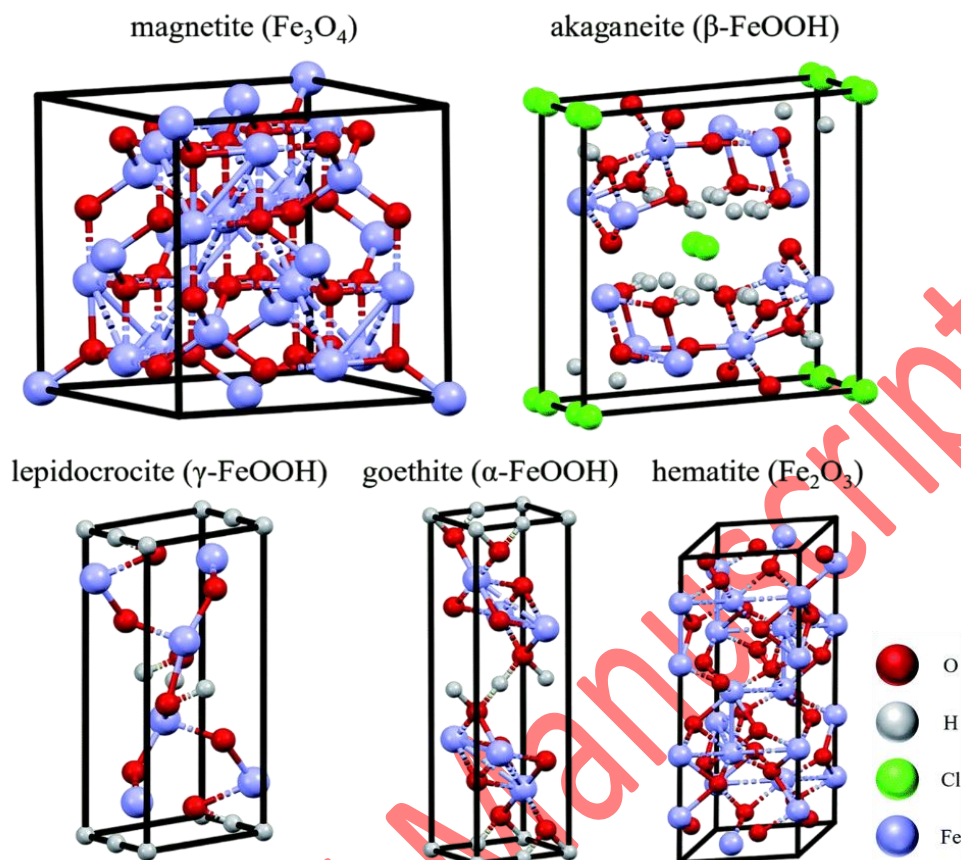


Fig. 9. Some of crystals cell of carbon steel corrosion [128]

7- Corrosion inhibition

The solubility of metal in the solution affects weight loss, corrosion, and metal instability. The addition of inhibitors results in a linear decrease in the weight loss in the specimens proportional to the corrosion rate [129].

Corrosion inhibition is the most cost-effective, practical, and easy method for controlling corrosion on metals in saltwater environments [130]. Corrosion inhibitors control metal dissolution and acid consumption. Inhibitors are adsorbed onto the metallic surface; Inhibitors inhibit the solubility of metals. Inhibitors adsorb onto the metal surface, creating a physical protective barrier, and interact with anodic and/or cathodic reaction sites to prevent oxidation and/or reduction of corrosion reactions [131].

Corrosion inhibitors are natural or chemical substances added in low concentrations into corrosive environments to prevent or minimize (control) corrosion without significantly reacting with surrounding components [132]. Concentration ranges from (1 to 15,000) ppm [133]. Corrosion inhibitors are crucial in many chemical industries such as oil extraction, processing. The use of corrosion inhibitors into the system minimizes the corrosion process or inhibits the metal oxidation rate [134]. Inhibition is the method that prevents the damage caused by of corrosive substances by adsorbing inhibitors onto a whole metallic substrate [135,136].

The inhibitors can be categorized into both inorganic and organic types [51]. Inorganic inhibitors indicate either anodic or cathodic behaviors. The organic inhibitors have mixed cathodic and

anodic activity together with adsorption properties. Figure 10 illustrates how to classify of the inhibitors. Inorganic corrosion inhibitors have superior effectiveness over an additional temperature spectrum and for longer periods of time in comparison to organic corrosion inhibitors. Organic corrosion inhibitors, while more expensive than their inorganic different forms, indicate less toxicity.

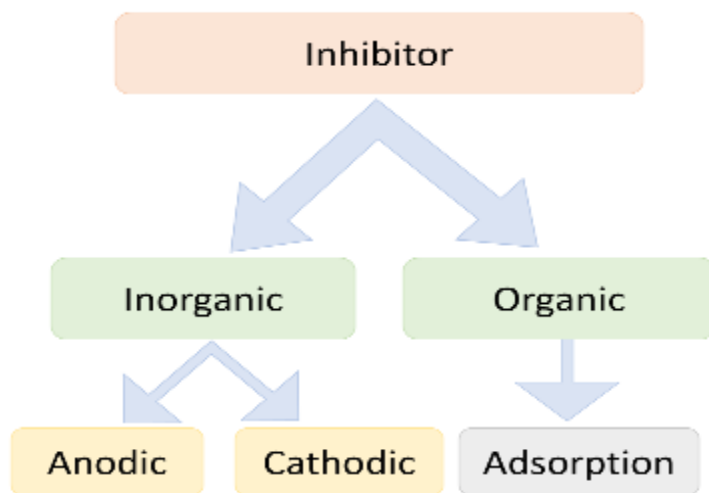


Fig. 10. Classification of Inhibitors

The inhibition efficiency of inhibitors is determined by the structure and chemical properties of the film formed over the substrate surface [137]. The efficiency of the inhibitor is affected by the state of the substrate surface, the type of corrosive media, the steel composition, and the chemical structure of the inhibitor [138]. Physical-chemical properties induced by the functional group the and strength of the inhibitor-metal bond in the molecule affect the adsorption of organic inhibitors [139]. The variety of organic and inorganic corrosion inhibitors as well as some green eco-friendly, some biobased corrosion inhibitors are explored for the carbon steel application.

Recent research conducted by many authors on the efficiency of inhibition of different types of carbon steels in rich chloride solutions are shown in Table 4.

Table 4. Efficiency of some corrosion inhibitors for different types of carbon steel in salt water at 25°C.

Corrosion Inhibitor	Concentration of inhibitors	Type of Media	Steel type	Efficiency %	References
4-[6-bromo-benzothiazolylazo] thymol BTAT	70 ppm	seawater	Low Carbon Steel	94.6	[140]
Syzygium cumini fruit	500 ppm	seawater	Low Carbon Steel	92.99	[141]
Coriandrum sativum	10 μ L 30 μ L	138 and 200 mg/L	Mild Steel	83.6 78.5	[142]

Aloe vera	50 µL 300 ppm	seawater	API 5L Carbon steel	86.6 83.75	[143]
Silicate	1.2 g was dissolved in 100 ml of 3 mol/L NaOH	Saline solution	Carbon steel	94	[144]
Alkanol ammonium salts	From $[5 \times 10^{-4}]$ to $5 \times 10^{-3}]$ mol/L	3 % NaCl	Carbon steel	96	[145]
Amino Acid L-Histidine	250 ppm 500 ppm 1000 ppm	3.5 %NaCl	AISI 1018 carbon steel	Up to 89	[146]
New organic ammonium salts	Different concentration s	3.5 % Saline solution	Mild steel	High	[147]
Morinda citrifolia	300 ppm	Saline environme nt	Mild steel	85.1	[148]
plum tree gum	25 mg/L 50 mg/L 100 mg/L 250 mg/L 500 mg/L	3.5% saline water	Carbon steel	94 95 96 97 97	[149]
4-(dimethylamino) -1-nonylpyridin-1 -ium Bromide((4DMN)	Different concentration s	3.5 %NaCl	Carbon steel	94	[150]
4-(dimethylamino)-1 -(prop-2-yn-1-yl)pyridine -1-ium Iodide (4DMP)	Different concentration s	Saline medium	Carbon steel	92	[150]
Sodium silicate	0.1, 0.5, 1, 5, 10 and 20 mmol/L	3 % NaCl	Carbon steel	Up to 99.8	[151]
Cassava leaf (DNA)	20 ppm	3.5% NaCl	AISI 1015 low carbon steel	96.4	[152]
Ficus (FIC) leaf	2 ppm	Saline medium	Carbon steel	87	[153]
Bis(2-ethylhexyl) phosphate (BEP)	500 ppm	1 % NaCl	Carbon steel	93.07	[154]

Curcuma longa	200 ppm	Seawater	Mild steel	98	[155]
Octylsilanol and Ce(III) ions	400 ppm	0.1 NaCl mol/L	Carbon steel	>96	[156]
(1-[[5-(2-Chloro-phenylazo)-2-hydroxy-benzylidene]-amino]-4,7-dimethyl -6-nitro- 1H quinolin-2-one (CPHAQ2O))	5 ppm 10 ppm 20 ppm	3.5% NaCl	carbon steel	86.82 87.05 90.55	[157]
Imidazole and Benzimidazole	50 ppm	3% NaCl	AISI 1010 carbon steel	73	[158]
coconut oil-modified imidazoline	20 ppm	3% NaCl	Carbon steel	85	[159]

8- Conclusions and Future Prospects

Corrosion of carbon steel in salty environments is a serious concern due to its extensive use in oceanic building materials, pipelines, and industrial applications. Temperature, pH, chloride concentrations, and exposure length have significant effects on corrosion. In order to limit the degradation of materials, it is necessary to have a thorough understanding of these variables. A variety of approaches have been used to investigate corrosion behavior, including weight loss, electrochemical techniques (potentiodynamic polarization, electrochemical impedance spectroscopy) and surface analysis instruments (SEM, EDX, XRD, AFM). The weight loss method allows for direct measurement of corrosion rate, electrochemical methods can provide real-time information, and surface analysis methods can provide insight into morphological and chemical changes in corrosion-damaged materials. The prevention of corrosion has been achieved by employing various protective strategies, including corrosion inhibitors, protective coatings, cathodic protection, and nanotechnology-based materials. It is becoming increasingly essential for early detection of corrosion symptoms and averting structural failures to employ real-time corrosion monitoring techniques, such as electrochemical sensors and acoustic emission techniques. It should be noted, however, that progress in corrosion research continues, but obstacles remain in creating eco-friendly inhibitors, increasing coating longevity, and developing real-time monitoring systems. By developing environmentally friendly corrosion inhibitors and self-healing coatings, by implementing intelligent monitoring systems that utilize artificial intelligence and Internet of Things-based sensors, and by developing alloys and nanotechnology-based coatings, future corrosion research must focus on sustainable corrosion mitigation. With the incorporation of emerging technologies and sustainable materials, carbon steel structures are able to last longer, require fewer repairs, and provide increased operational safety in adverse conditions. Industrial applications benefit from improved material durability and environmental sustainability as a result.

Nomenclature

CR	Average corrosion rate (mm/y)
K	Conversion factor (3650) of the corrosion rate (mm/y)
W	Weight loss (g)
D	Density (g/cm ³)
T	Corrosion exposure time (Day)
A	Surface area of the sample (cm ²)

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