# **Study on Safety Measures To Reduce The Corrosion of Reinforcement In The Concrete**

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*Abstract- This paper deals with the consumption of strengthening steel can be issue in solid structures. Corrosion of steel produces hydrated iron oxide or rust which is broad. This extension develops inner weight until the solid flops as spalling.This is a main source of solid disintegration and various examinations have been performed to avert corrosion of fortifying steel. Strengthened cement has demonstrated to be very fruitful material regarding both auxiliary execution and sturdiness Epoxy-covered fortifying steel is the most widely recognized corrosion security strategy is utilized. Hardened steel fortification demonstrates that it might stay free of consumption in chloride defiled cement for over 75 years. While erosion of strengthening steel can be issue in solid it very well may be decreased utilizing legitimate detail and fast recognizable proof of vulnerability asphalts. Solid blends that are intended for administration presentation condition sick decrease the open doors for consumption*

*Keywords-* spalling,corrosion

#### **I. INTRODUCTION**

The disintegration of strengthened solid structures is a noteworthy issue. The expense of fixing or supplanting disintegrated structures has turned into a noteworthy risk for interstate offices, assessed to be more than \$20 billion and to increment at \$500 million every year. The essential driver of this weakening (breaking, delamination, and spalling) is the consumption of steel strengthening bars because of chlorides. The two principle wellsprings of chlorides are de-icing synthetic substances and seawater. The exposed asphalt arrangements of numerous roadway organizations for winter snow and ice expulsion have brought about broad utilization of salt-based de-icing synthetic substances. The most widely recognized compound utilized has been sodium chloride. Numerous scaffolds have additionally been worked in beach front territories and are presented to seawater. Extensions worked with dark fortifying steel are indicating dynamic solid crumbling as the centralization of chloride particles increments. As indicated by a May 1997 report, The Status of the Nation's Highway Bridges: Highway Bridge Replacement and Rehabilitation Program and National Bridge Inventory, Thirteenth Report to the United States Congress, around

80,000 scaffolds on the Federal-guide framework and 103,000 extensions off the Federal-guide framework are insufficient somehow or another. This incorporates both fundamentally lacking and practically out of date spans. The normal scaffold deck situated in a snow-belt State with dark fortifying steel and 40 mm (1.5 in) of solid spread hasshown spalling in around 7 to 10 years after development and has required recovery in around 20 years after construction.The increment in the sum and seriousness of extension deck weakening in the late 1960's and mid 1970's frightened the State Highway Agencies and presented genuine security risks to the voyaging driver. Accordingly a few measures have been created and actualized to forestall the chloride-incited erosion of steel fortifying bars and the subsequent crumbling. A portion of the early estimates utilized included bringing down the waterbond proportion of the solid and expanding the solid spread over the steel fortifying bars. Solid penetrability can likewise be decreased by the utilization of admixtures. Consumption inhibitors are additionally being utilized. Epoxy-covered fortifying steel (ECR) was presented in the mid 1970's as a security framework for new extension decks. Another security measure is the utilization of consumption safe strong fortifying bars or clad dark strengthening bars. The utilization of waterproof films related to asphaltic cement overlays as a defensive framework has delivered variable outcomes. A different insurance methodology is the synchronous utilization of at least two assurance measures, for example, epoxycovered fortifying steel and a consumption inhibitor.The utilization of prestressed solid individuals in scaffolds is a generally new practice and thus most extension applications are moderately youthful. Thus, consumption incited weakening of these individuals just ended up obvious in the mid 1980's. In spite of the fact that prestressed solid individuals are commonly made with a higher quality cement under better quality control, they are liable to indistinguishable impacts of erosion from is customarily strengthened cement. In any case, on account of the high worries in the prestressing strands, the consumption procedure is quickened. Indeed, even little consumption pits could make a strand crack, contrasted with customary strengthening that may rust to for all intents and purposes nothing before breaking. There have been reported instances of prestressing strands breaking because of consumption. This is a major issue as prestressed solid

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individuals depend on the elasticity of the strands to oppose connected loads.Protective frameworks for prestressed solid scaffold superstructure individuals (braces and shafts) and substructure individuals (heaps, docks, and so on.) are still under scrutiny and are being improved and grown further. These individuals are commonly worked with dark prestressing steel. Ebb and flow research shows guarantee for the utilization of epoxy-covered prestressing strands for prestressed solid individuals. Starting at now, there is no for the most part acknowledged defensive framework for solid scaffold substructure individuals or prestressed solid extension individuals. In any case, on account of the utilization of uncoated prestressing steel, the utilization of consumption inhibitors in prestressed, solid individuals has picked up acknowledgment. For most consumption insurance measures, the fundamental standard is to keep the chloride particles from responding with the steel surface and furthermore to expand the time required for the chloride particles to penetrate through the solid spread. While these measures for the most part don't prevent consumption from in the end starting, they do build the administration life of fortified solid structures by moderating the erosion procedure. Cathodic assurance, notwithstanding, has demonstrated to be a fruitful consumption security measure for expectedly fortified and pretensioned, prestressed solid extension individuals.

#### **II. LITERATURE REVIEW**

Overview of Corrosion-Induced Concrete Deterioration Processes General Although concrete has evolved to become the most widely used structural material in the world, the fact that its capacity for plastic deformation (so its ability to absorb mechanically imparted energy is essentially nil) imposes major practical service limitations. This shortcoming most commonly is overcome by incorporating steel reinforcement into specific locations in the concrete where tensile stresses are anticipated. Consequently, concerns regarding performance must not only focus upon properties of the concrete but also of the embedded steel and, in addition, the manner in which these two components interact. In this regard, steel and concrete are, in most aspects, mutually compatible, as exemplified by the fact that the coefficient of thermal expansion for each is approximately the same. Also, while boldly exposed steel corrodes actively in most natural environments at a rate that requires instituting extrinsic corrosion control measures (for example, protective coatings for atmospheric exposures and cathodic protection in submerged and buried situations), the relatively high pH of concrete pore water (pH  $\approx$  13.0–13.8) helps form a protective oxide (passive) film about 10 nanometres' thick. This film effectively insulates the metal and electrolytes so that the corrosion rate is negligible, allowing decades of relatively low maintenance.

Corrosion Mechanism Disrupting the passive film

upon embedded reinforcement and onset of active corrosion can arise in conjunction with either of two causes: carbonation or chloride intrusion (or a combination of the two). In the case of carbonation, atmospheric carbon dioxide (CO2) reacts with pore water alkali according to the generalized reaction, Ca( ) OH  $2 + CO2 \rightarrow CaCO3 + H2O$ , (1) which consumes reserve alkalinity and reduces pore water pH to the 8–9 range, where steel is no longer passive. For dense, high-quality concrete (for example, high cement factor, low water-cement ratio, and pozzolanic admixture), carbonation rates are typically on the order of 1 mm per decade or less; loss of passivity from this cause within a normal design life is not generally a concern. Carbonation must be anticipated at concrete cracks, however, where air essentially has direct access to the reinforcement, irrespective of concrete cover and quality. Older structures are also at issue because of their age, because earlier generation concretes were typically more permeable when compared to more recent concretes, and because of relatively low concrete cover. Chlorides, on the other hand, arise in conjunction with deicing activities upon northern roadways or from coastal exposure, as noted above. While this species (Cl- ) has only a small influence on pore water pH, concentrations as low as 0.6 kilograms per cubic meter (kg/m3 ) (concrete weight basis) have been projected to compromise steel passivity. In actuality, it probably is not the concentration of chlorides that governs loss of passivity but rather the ratio of chlorides-to-hydroxides ([Cl-]/[OH-]), because the latter species (OH- ) acts as an inhibitor. This has been demonstrated by aqueous solution experiments from which it is apparent that the Cl- threshold for loss-of-steel passivity increased with increasing pH.[4–5] However, in cementitious materials, this interrelationship is more complex due to Cl- binding and the dependence of such binding upon pH.[6] Thus, Cl- binding evidently decreases with increasing OH- above pH 12.6, such that a decrease in pH can result in decreasing[Cl-]/[OH- Considerable research efforts have focused on identifying a chloride threshold; however, a unique value for this parameter has remained elusive, presumably because of the numerous influential variables, including type of cement, cement alkalinity, concrete mix design, environmental factors, potential, and reinforcement composition and micro structure. Because Cl- , not carbonation-induced loss of passivity, is of primary concern for bridge structures, subsequent focus is placed upon this cause of corrosion alone.

#### **III. METHODOLOGY**

The corrosion of steel reinforcing bars is an electrochemical process that requires a flow of electric current and several chemical reactions

The corrosion of steel in concrete in the presence of oxygen, but without chlorides, takes place in several steps:

1. At the anode, iron is oxidized to the ferrous state and releases electrons.

$$
Fe \parallel Fe^{++} + 2e-
$$

2. These electrons migrate to the cathode where they combine with water and oxygen to form hydroxyl ions.

# **2e- +H2O+1/2O2|2OH-**

3. The hydroxyl ions combine with the ferrous ions to form ferrous hydroxide.

$$
Fe^{++}+2OH^{\cdot} \mid Fe(OH)_2
$$

4. In the presence of water and oxygen, the ferrous hydroxide is further oxidized to form

# $4Fe(OH)<sub>2</sub> + O<sub>2</sub> + H<sub>2</sub>O$  |  $4Fe(OH)<sub>3</sub>$ . **2Fe(OH)3 | Fe2O3@2H2O**

The corrosion of steel in concrete in the presence of chlorides, but with no oxygen (at the anode), takes place in several steps:

1. At the anode, iron reacts with chloride ions to form an intermediate soluble iron-chloride complex.

 $Fe + 2cl$  |  $(Fe^{++} + 2cl) + 2e^-$ 

4. When the iron-chloride complex diffuses away from the bar to an area with a higher pH and concentration of oxygen, it reacts with hydroxyl ions to form  $Fe(OH)$ . This complex reacts with water to form ferrous hydroxide.

 $(\text{Fe}^{++} + 2\text{cl}) + 2\text{H}_2\text{O} + 2\text{e}^{\cdot}$  |  $\text{Fe}(\text{OH})_2 + 2\text{H}^{\text{+}} + 2\text{cl}$ 

5. The hydrogen ions then combine with electrons to form hydrogen gas.

 $2H^{+} + 2e^{-}$  |  $H_{2}8$ .

As in the case of the corrosion of steel without chlorides, the ferrous hydroxide, in the presence of water and oxygen, is further oxidized to form  $Fe<sub>2</sub>O<sub>3</sub>$ .

#### $4Fe(OH)<sub>2</sub> + O<sub>2</sub> + H<sub>2</sub>O$  |  $4Fe(OH)<sub>3</sub>$ **2Fe(OH)<sup>3</sup> |Fe2O3@2H2O**

The corrosion products resulting from the corrosion of steel reinforcing bars occupy a volume equal to three to six times that of the original steel. This increase in volume induces stresses in the concrete that result in cracks, delaminations, and spalls. This accelerates the corrosion process by providing an easy pathway for the water and chlorides to reach the steel.There are two main types of chloride contents that are tested for and reported – acidsoluble chlorides, sometimes referred to as total chlorides, and water-soluble chlorides. Acid-soluble chlorides are the chlorides extracted from a concrete sample using an acid. Water-soluble chlorides are those chlorides present that can dissolve in water. The amount of water-soluble chlorides is less than the total or acid-soluble amount of chlorides present in a concrete sample.

## **3.1. FACTORS INFLUENCING CORROSION RATE:-**

Once a sufficient amount of chlorides has reached the steel reinforcing bars to depassivate the bars and initiate corrosion, factors influencing the corrosion rate of steel reinforcing bars embedded in concrete include:

->Availability of water and oxygen.

->Ratio of the steel surface area at the anode to that at the cathode.

- ->Amount of chloride ions in the pore water.
- ->Resistivity of the concrete.
- ->Temperature.
- ->Relative humidity (both internal and external).

->Concrete microstructure.

The availability of oxygen is a function of its rate of diffusion through the concrete, which is affected by how saturated the concrete is with water. When totally submerged, the diffusion rate is slowed because the oxygen must diffuse through the porewater. When the concrete is dry, the oxygen can freely move through the pores. Alternating wet-dry cycles accelerates the corrosion process. Wet concrete has a lower resistivity than dry concrete due to the presence of water as an electrolyte. When the ratio of the area at the cathode to the area at the anode increases, the current density at the anode increases. The current density is the amount of electrical current passing through a unit area at the anode. An increase in current density results in an increase in the corrosion rate.

# **3.2. FACTORS INFLUENCING DETERIORATION RATE**

To date, not much research has been done in this area. The main focus has been on the depth of the concrete cover and permeability. High-strength concretes generally have low water-cement ratios, low porosity, and a relatively high modulus of elasticity. Because of its low porosity, highstrength concrete may have less ability to absorb corrosion products (i.e., they have fewer voids where corrosion products may accumulate without exerting any internal pressure on the concrete). In general, high-strength concretes have a higher modulus of elasticity and are less forgiving than concretes with a lower modulus. Concretes with a lower modulus may deflect without cracking, while in concretes with a higher modulus, stresses may build up and cause fracturing.On the other hand, higher strength concretes generally have lower permeability and therefore it takes longer for chlorides to accumulate at the reinforcement level compared to lower strength concretes.

## **3.3. CORROSION-CONTROL MEASURES**

Corrosion-induced deterioration of reinforced concrete structures occurs when the environmental loading on the structure is greater than the ability of the structure to resist the environmental loading (environmental resistance). One can either decrease the loading or increase the resistance or do a combination of both. The main deterioration mechanisms (chloride-induced corrosion of rebar) focus on the reinforcement and its protection.

Corrosion can also occur as a result of other deterioration processes: freeze-thaw cycles, expansive reactions, excessive deflections, fatigue, etc. These processes cause the concrete to crack, which subsequently allows water and chlorides easy access to the interior of the concrete and the steel reinforcing bars. These other deterioration mechanisms create conditions more conducive to the corrosion of the embedded steel reinforcing bars, which leads to further deterioration of the concrete

# **3.4. SELECTION OF PROTECTION SYSTEMS**

The proper corrosion-protection strategy will vary from structure to structure. Some factors to be considered during the design of a structure include:

->Intended design life of the structure.

->Effects of corrosion and corrosion-induced deterioration – This includes the costs due to closure (either permanent or temporary) for repair. Bridges on major roads are more critical than bridges on local roads.

->Quality of workmanship in construction – The quality of construction entails good consolidation, proper rebar placement, sufficient concrete cover over the steel reinforcing bars, and other measures.

->Possible rehabilitation methods – The design of structures should include provisions for the possible future rehabilitation of corrosion-induced deterioration.

->Initial costs – May need to consider more than just initial costs (i.e., life-cycle costs). As the rehabilitation and replacement costs increase, corrosion-control measures become more cost-effective.

->Multiple protection strategies may be cost-effective for long-term corrosion protection.

One such strategy is the use of epoxy-coated rebar in combination with a durable concrete containing corrosion inhibitors, having a low permeability, and adequate concrete cover. Silica fume and fly ash can be added to the concrete to reduce permeability and provide additional corrosion control. However, there is a need to balance the costs of the additional control measures against how much additional service life can be expected as a result of the added control measures. The additional costs can usually be justified based on a life-cycle cost analysis.

Some factors to be considered when choosing a corrosioncontrol measure include:

- ->Reliability and effectiveness of the measure.
- ->Risk of unintended side effects.
- ->Possibility of future installation of other control measures.
- ->Life expectancy of the measure.
- ->Any incremental costs over the "do nothing" option.
- ->Any impacts on the cost of other elements in the structure.

->How aggressive the environment is where the structure will be located.

Corrosion-protection strategies for steel reinforcing bars embedded in concrete can be grouped into four general categories: design, concrete, corrosion inhibitors, and reinforcement type.

The design category includes such items as:

->Concrete cover.

- ->Maximum allowable crack widths in service.
- ->Reinforcement distribution (crack control provisions).

->Rigid overlays (silica fume concrete, latex-modified concrete, dense concrete, polymer concrete).

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The concrete category includes such items as:

->Water-cement ratio.

-> Pozzolans (silica fume, fly ash, slag).

->Latex, epoxy, and polymer admixtures.

->Cement type.

->Aggregate gradation.

The inhibitor category includes such items as:

->Organic corrosion inhibitors.

->Inorganic corrosion inhibitors.

->Mixed corrosion inhibitors.

The reinforcement category includes such items as:

->Epoxy-coated bars.

- ->Galvanized bars.
- ->Nickel-clad bars.
- ->Copper-clad bars.
- ->Stainless steel-clad bars.
- ->Stainless steel bars.
- ->Corrosion-resistant alloyed bars.

->Non-metallic bars.

#### **3.3GENERAL DESIGN PROVISIONS**

It is generally the design details that influence the overall performance and durability of bridge decks and other bridge components, both conventionally reinforced concrete and prestressed concrete. Some design factors that affect the durability of concrete structures include:

- ->Construction type.
- ->Expansion joints.
- ->Construction joints.
- ->Tendency of concrete to crack.
- ->Duct and anchorage layout in post-tensioned concrete.
- ->Drainage details.
- ->Access for inspection and maintenance.
- ->Proximity to seawater.
- ->Exposure to deicing chemicals.

The effectiveness and lifespan of expansion joints depend on how well they are installed. Whenever possible, construct continuous structures and integral abutments to eliminate expansion joints. However, when joints are used, provide adequate and proper drainage so that water does not reach the anchorages or bearings and does not pond. Include provisions for the inspection of the joints and structural components under the joint. Even well-constructed joints leak. Whenever possible, locate deck construction joints away from

critical areas (prestressed anchorages in particular).Cracks may be thermal or shrinkage. Cracking may also be due to creep or to the high modulus of elasticity of the hardened concrete. Exercise proper care in the layout and sequencing of concrete pours to minimize the risk of cracking. For posttensioned structures, provide an adequate amount and distribution of reinforcement in the anchorage areas.In posttensioned concrete structures, the ease of grouting will influence the quality of the completed grouting operation. Both tendon profiles and duct size affect the ease of grouting. The location of anchorages affects the ease of stressing and inspection, as well as susceptibility to the ingress of water. Anchorages located in the top surfaces of decks are easy to construct, stress, and grout. However, due to their location, it is easier for chloride-contaminated water to penetrate and reach tendons.

#### **IV.CONCLUSION**

Security measures to reduce the corrosion of support in the solid structure by utilizing post strain and pre tensioned strategies are to be utilized for decrease the consumption in solid structures. The assurance of anchorages relies upon the presentation of the encompassing cement

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