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Katodna zaščita notranjih površin kovinskih rezervoarjev, konstrukcij, opreme in cevovodov, ki vsebujejo morsko vodo

Cathodic protection of internal surfaces of metallic tanks, structures, equipment, and piping containing seawater

Kathodischer Schutz der inneren Oberflächen von metallischen Tanks, Strukturen, Ausrüstung und Rohrleitungen die Meerwasser enthalten

Protection cathodique des surfaces internes des réservoirs, ouvrages, équipements et tuyauteries métalliques contenant de l'eau de mer

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**Cathodic protection of internal surfaces of metallic tanks,
structures, equipment, and piping containing seawater**

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COMITÉ EUROPÉEN DE NORMALISATION
EUROPÄISCHES KOMITEE FÜR NORMUNG

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European foreword

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Introduction

Metallic structures containing seawater or brackish waters are exposed to the risk of corrosion. Even when a coating is applied to reduce this risk, cathodic protection is usually used to ensure corrosion control during the design lifetime of the structure. This is especially important in the presence of galvanic couples between various metals and alloys, because corrosion is then concentrated to the less noble material.

Cathodic protection works by supplying sufficient direct current to the internal surface of the structures in contact with water in order to change the steel to electrolyte potential to values where the corrosion rate is insignificant.

The general principles and theoretical aspects of cathodic protection in seawater are detailed in EN 12473.

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1 Scope

This document specifies the requirements and recommendations for cathodic protection systems applied to the internal surfaces of metallic tanks, structures, equipment, and piping containing raw or treated seawater or brackish waters, to provide an efficient protection from corrosion.

Cathodic protection inside fresh water systems is excluded from this document. This is covered by EN 12499.

NOTE EN 12499 covers internal cathodic protection for any kind of waters, including general aspects for seawater; but excluding industrial cooling water systems. This document specifically details applications in seawater and brackish waters.

2 Normative references

The following documents are referred to in the text in such a way that some or all of their content constitutes requirements of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

EN 12473, *General principles of cathodic protection in seawater*

EN 12496, *Galvanic anodes for cathodic protection in seawater and saline mud*

EN 12499, *Internal cathodic protection of metallic structures*

EN 13509, *Cathodic protection measurement techniques*

EN ISO 8044, *Corrosion of metals and alloys - Basic terms and definitions (ISO 8044)*

EN ISO 9606-1, *Qualification testing of welders - Fusion welding - Part 1: Steels (ISO 9606-1)*

EN ISO 15607, *Specification and qualification of welding procedures for metallic materials - General rules (ISO 15607)*

EN ISO 15609-1, *Specification and qualification of welding procedures for metallic materials - Welding procedure specification - Part 1: Arc welding (ISO 15609-1)*

3 Terms and definitions

For the purpose of this document, the terms and definitions given in EN ISO 8044 and EN 12473 and the following apply.

ISO and IEC maintain terminological databases for use in standardization at the following addresses:

- IEC Electropedia: available at <http://www.electropedia.org/>
- ISO Online browsing platform: available at <http://www.iso.org/obp>

3.1

anode redundancy factor

multiplier applied to the theoretical number of anodes to allow for anode damage and failures for ensuring that protection will continue to be achieved when one or more anodes are lost, without modifying the unit weight of anodes

3.2**cathodic protection zone**

that part of the structure which can be considered independently with respect to cathodic protection design

3.3**coating breakdown factor** **f_c**

ratio of cathodic current density for a coated metallic material to the cathodic current density of the bare material

3.4**critical crevice potential**

potential under which there is no risk of initiation of crevice corrosion for a given environment and crevice geometry

3.5**driving voltage**

difference between the structure to electrolyte potential and the anode to electrolyte potential when the cathodic protection is operating

3.6**over-polarization**

occurrence in which the structure to electrolyte potentials are more negative than those required for satisfactory cathodic protection

Note 1 to entry: Over-polarization provides no useful function and might even cause damage to the structure such as cracking due to hydrogen embrittlement and coating disbondment.

Note 2 to entry: Often incorrectly referred to as over-protection.

3.7**ullage factor** **u_f**

ratio of the surface area of a ballast tank which may be in contact with water to the total surface area

Note 1 to entry: When the entire surface area may be wetted, $u_f = 1$.

3.8**wetting factor** **k**

fraction of design life when the internal surface of a tank or a structure is in contact with water.

Note 1 to entry: When the tank is permanently filled with water, $k = 1$.

Note 2 to entry: For changing conditions an average value can be considered.

Note 3 to entry : Also known as loading factor [1].

3.9**hydrogen embrittlement**

process resulting in a decrease of the toughness or ductility of a metal due to absorption of hydrogen

3.10

hydrogen stress cracking

HSC

cracking that results from the presence of hydrogen in a metal and tensile stress (residual and/or applied)

Note 1 to entry: HSC describes cracking in metals which may be embrittled by hydrogen produced by cathodic polarization without any detrimental effect caused by specific chemicals such as sulphides

4 Competence of personnel

Personnel who undertake the design, supervision of installation, commissioning, supervision of operation, measurements, monitoring and supervision of maintenance of cathodic protection systems shall have the appropriate level of competence for the tasks undertaken. This competence should be independently assessed and documented.

EN ISO 15257 constitutes a suitable method of assessing and certifying competence of cathodic protection personnel which may be utilised.

Competence of cathodic protection personnel to the appropriate level for tasks undertaken should be demonstrated by certification in accordance with EN ISO 15257 or by another equivalent prequalification procedure.

5 General considerations

5.1 Structures and equipment to be protected

This European Standard applies to internal surfaces of ballast tanks, aboveground or buried storage tanks including firewater tanks, filters such as sand filters, heat-exchangers and condensers, flooded sections of harbour and lock gates, sea defence barriers, dolphins, and offshore wind turbine foundations.

This European Standard applies to the external submerged areas of appurtenances and other independent equipment fitted within tanks, such as pumps and piping, when they are not electrically isolated from the structure. It also covers items that are integrated to the structure such as stiffeners.

This European Standard applies also to the internal cathodic protection of piping transporting seawater or brackish waters, including valves, pumps and fittings. In such applications, additional considerations are required for the design and installation of cathodic protection.

This European Standard is also applicable for temporary cathodic protection systems used to prevent corrosion by seawater or brackish waters when they are used for hydrotesting and when they are stored before commissioning and operation of tanks or other equipment.

5.2 Materials

This European Standard covers the cathodic protection of structures fabricated principally from bare or coated carbon manganese steels.

This European Standard also covers the cathodic protection of structures fabricated from stainless steels, nickel alloys, copper alloys or titanium alloys.

The requirements and the recommendations for the cathodic protection systems described in the present standard ensure control over any galvanic coupling, which could be caused by the use of various metallic materials, and minimise risks due to hydrogen embrittlement or hydrogen stress cracking (see EN 12473).

5.3 Environment

This European Standard is applicable only to structures containing seawater or brackish waters. Seawater may be either raw or treated (e.g. using chlorination systems) for preventing fouling of piping systems or for preserving biodiversity when ship ballast tanks are filled and emptied in various locations of the world [2]. Special requirements can be necessary for structures containing polluted seawater or other fluids, e.g. slop tanks.

For surfaces which are alternately immersed and exposed to the atmosphere, cathodic protection is only effective when the structure is submerged and the immersion time is sufficiently long for the steel to become polarised.

5.4 Safety and environmental protection

5.4.1 General

This European Standard does not cover routine safety and environmental protection aspects which are not especially associated with cathodic protection. The relevant national or international regulations shall apply.

Safety requirements specific to the application of cathodic protection within the scope of the present European standard are covered.

An environmental checklist is supplied in Annex A.

For impressed current systems, automatic control of applied potential should be considered and/or provision shall be made to prevent sparking if the anode is energized outside liquid level. Procedures to be adopted may include positioning of the anode so that it is always submerged and/or incorporation of emergency shutdown procedures so that if the anode is temporarily exposed all d.c. current is disabled.

The use of impressed current systems and of some galvanic anodes can be prohibited for some applications, for instance when in the vicinity of tanks containing hydrocarbons [41] (see 8.6.).

5.4.2 Evolution of dangerous gases

Cathodic protection can generate gaseous hydrogen, oxygen and chlorine. Some mixtures of oxygen and hydrogen are explosive; chlorine can be toxic and corrosive.

The design and operation of the cathodic protection systems shall take these risks into account to ensure that harmful and dangerous levels of gas accumulation are not allowed to occur. This may include the control of potential and/or the use of ventilation systems.

With galvanic anodes, the selection of alloy material should be carried out to minimise the risk.

5.4.3 Release of hydrogen gas

For impressed current systems, and with galvanic anodes (particularly magnesium anodes), the polarisation causes evolution of hydrogen gas on the protected structure. Thus, in situations such as closed tanks where hydrogen can collect, an explosion hazard can arise. To avoid this hazard, it is necessary for all designs to include venting to prevent the build-up of a significant gaseous volume of hydrogen. Gas levels should be monitored.

The rate of hydrogen evolution is related to the structure to electrolyte potential. Where hydrogen evolution can produce an explosion hazard. The structure to electrolyte potential shall be controlled and monitored.

5.4.4 Chlorine evolution

For impressed current cathodic protection systems working in seawater and brackish waters, one of the anodic reactions results in the electrolytic formation of chlorine. Such a formation can cause physical discomfort or downstream corrosion effects.

NOTE Information on toxic levels can be found in the European Union Risk Assessment Report or the US Department of Health and Human Services report [3].

In order to reduce chlorine production, the anodic current density should be minimised by a reduction in the cathodic protection current demand and by an increase in the surface area of the anodes.

5.4.5 Emptying and opening

Before opening an enclosed structure the impressed cathodic protection system shall be turned off and the enclosure gas levels declared safe.

6 Cathodic protection criteria

6.1 General

The criteria for cathodic protection generally used are given in the European Standard EN 12473 "General principles of cathodic protection in seawater". The criteria are based on the potential at the surface of the metal to be protected and the measurement techniques used should ensure that this is measured accurately (see 10.2).

When different metals and alloys are in electrical continuity, the protection potential achieved shall be the most negative one in order to prevent any galvanic corrosion of the less noble of them.

A negative limit to the potential may be required depending upon the metallic material in order to avoid coating disbondment (see ISO 15711) and/or adverse effects due to hydrogen evolution or high pH.

The potential criteria and limit values are expressed without IR errors. IR errors, due to cathodic protection current flowing through resistive electrolyte and surface films on the protected surface, are generally considered insignificant in marine applications. However, potential measurements using "Instant OFF" techniques or "coupon Instant OFF" techniques can be necessary in applications described in this European Standard in order to adequately demonstrate the achievement of the protection criteria (see EN 13509). Particular attention should be given to this in brackish waters and mud applications or close to impressed current anodes.

The criteria also apply to steel in brackish waters but the errors due to variations in salinity when using Ag/AgCl/sea water reference electrode shall be addressed (see EN 12473).

6.2 Carbon manganese steels

To ensure the protection of carbon manganese steels in aerated seawater a protection potential more negative than $-0,80$ V wrt Ag/AgCl/sea water shall be achieved. This corresponds approximately to $+0,23$ V when measured with respect to a pure zinc electrode (e.g. alloy type Z2 as defined in EN 12496) or $+0,25$ V when measured with respect to a zinc electrode made with galvanic anode alloy types Z1, Z3 or Z4 as defined in EN 12496.

When the water temperature or surface temperature of steel is higher than 60 °C, the criterion shall be $-0,90$ V with respect to (wrt) Ag/AgCl/sea water. Between 40 and 60 °C the protection potential shall be interpolated between $-0,80$ and $-0,90$ V wrt Ag/AgCl/sea water.

For tanks and structures containing seawater or brackish waters which are not frequently renewed (e.g. less than once a month) the criterion for anaerobic conditions, i.e. $-0,90$ V wrt Ag/AgCl/sea water, shall be adopted regardless of temperature.

6.3 Stainless steels and nickel alloys

In chloride-containing aerated environments such as raw seawater and brackish waters, stainless steels are known to resist uniform corrosion and the possibility for crevice corrosion and pitting remains the principal concern. To ensure the protection of such alloys in these environments, the protection potentials given in EN 12473 apply, i.e. $-0,30$ V wrt Ag/AgCl/sea water for stainless steels with $PREN \geq 40$ and $-0,50$ V wrt Ag/AgCl/sea water for stainless steels with $PREN < 40$.

However, less conservative criteria can be used provided that these are justified and documented. In this case, the selected protection criteria shall be more negative than the critical crevice potential determined for a combination of a particular alloy (PREN, microstructure, etc.), crevice parameters (geometry, surface finish, type of gaskets, sealing pressure of flanges, etc.), and operating environmental conditions (composition, temperature, velocity, etc.).

The determination of the critical crevice potential shall be carried out on the basis of demonstrated service feedback and/or on laboratory tests relevant to the design service life. In the absence of a more relevant method for a given practical situation, the method given in ISO 18070 should be used for the crevice formers applied for flat or tube specimens.

In the case of galvanic couples between parts in stainless steel or nickel alloy and parts in carbon manganese steel, the protection potential criterion of the carbon manganese steel shall be more negative than $-0,80$ V wrt Ag/AgCl/sea water.

In chlorinated seawater, the recommended protection potentials criteria for stainless steels and nickel alloys are the same as those in raw seawater, but the protection current densities are less (see 7.2.7).

6.4 Cracking risks induced by over polarization

Where there is a risk of hydrogen embrittlement or HSC of high strength steels or other metals which may be adversely affected by cathodic protection to excessively negative values, a less negative potential limit shall be defined and applied. If there is insufficient information for a given material, this specific negative potential limit shall be determined relative to the metallurgical and mechanical conditions by testing. Refer to EN 12473 for more details.

For carbon manganese steels, a potential negative limit of $-1,10$ V wrt Ag/AgCl/sea water is recommended in order to avoid crack initiation/propagation. Other potential negative limits shall be applied to prevent cathodic disbonding of coatings and HSC of vulnerable steel compositions (refer to EN 12473).

For stainless steels and nickel alloys, ferritic and martensitic microstructures can suffer from hydrogen embrittlement when the potentials are too negative. Potentials more negative than the limit values should be avoided. EN 12473 gives information on this risk and also provides recommendations for qualification of the materials.

Titanium and its alloys are prone to titanium hydride formation in cathodic protection applications. This hydride can lead to cracking when stresses reach a critical level.

Heat exchanger tube inlets are areas of high mechanical stress due to construction methods. Titanium grade 2 tubes should not be subject to a potential more negative than $-0,75$ V wrt Ag/AgCl/sea water [5], some more conservative figures being found in the literature, e.g. $-0,70$ V [6] and even $-0,65$ V [8].

This conservative approach can be replaced by other criteria provided that these are justified and documented by an assessment to ensure that the risk is acceptable. Such an assessment shall consider all load contributions causing stress and strain. In any case, potential of the titanium shall not be more negative than $-1,00$ V wrt Ag/AgCl/seawater as specified in EN 12499 and documented in the literature [5].

For other applications where stresses and strains are lower, e.g. tubular plates, a negative limit of $-1,05$ V wrt Ag/AgCl/seawater is sufficiently conservative [10].

7 Design

7.1 Objectives

The objective of a cathodic protection system is to deliver sufficient current to each part of the internal surfaces of metallic tanks and structures, including bonded equipment, to achieve the protection criteria defined in Clause 6. This current should be distributed so that the structure to electrolyte potential of each part is within the limits given by the protection criteria during its normal service conditions (see Clause 6). Permanent reference electrodes should be installed to allow measurement of the potentials at selected locations.

Uniform levels of cathodic protection may be difficult to achieve in some areas or parts of structures. In this case the use of reference electrodes installed in shielded areas is recommended, especially for assisting the commissioning. The cathodic protection system for tanks and structures is generally combined with a protective coating system, even though some equipment such as pumps and small pipes may not be coated. A good coating is especially recommended when temperature and/or velocity of the water is high, such as in pipeline systems or heat-exchanger cooling water boxes.

In the absence of design life instructions the cathodic protection system shall be designed either for the lifetime of the structure or for a period corresponding to a planned maintenance, such as dry-docking interval in the case of ballast tanks on ships or floating offshore structures. Alternatively, when it is not feasible to design the cathodic protection system for the life of the structure or if planned maintenance is not possible, the system should be designed for easy replacement of cathodic protection components. Access fittings allowing replacement of anodes and reference electrodes can be considered, e.g. for pipeline systems.

Where small current demands are anticipated over the entire life of the structure, cathodic protection can be achieved by galvanic anode systems. Impressed current systems are generally used for other instances, such as higher current demand, temporary protection of large equipment containing seawater or brackish water for hydrotesting, pipe systems or water boxes of large heat-exchangers.

Each stage of the design, the installation, the energising, the commissioning, the long-term operation and the records of all the elements of cathodic protection systems shall be documented.

Each part of the work should be undertaken in accordance with a quality plan.

NOTE EN ISO 9001 constitutes a suitable Quality Management Systems Standard which can be used.

All test instrumentation shall have valid calibration certificates traceable to national or European Standards of calibration.

This documentation shall constitute part of the permanent records.

7.2 Design parameters

7.2.1 General

The design of a cathodic protection system shall be conducted according to the following steps:

- a) Determine the coated and non-coated areas exposed to water;
- b) Determine the maximum and mean current demands for each of these areas taking into account temperature and velocity of the water;
- c) Determine the type of cathodic protection system (galvanic, impressed current, or hybrid);
- d) Determine the size, shape and number of anodes required to last the design life;
- e) Determine anode locations and installation;

- f) Determine reference electrodes types, location and installation.

The design of a cathodic protection system shall demonstrate that each structure subdivision and anode zone is supplied with the cathodic protection current necessary to provide cathodic protection to meet the criteria in Clause 6 for all service conditions.

The anode distribution shall take into account the location of interfaces between dissimilar metallic materials.

Two different values shall be considered:

- $I_{\max}$: maximum current demand, which corresponds to the most severe working conditions (e.g. end of life coating breakdown factor and maximum surface areas, maximum water temperature and velocity) and is used to design the maximum current capacity of the cathodic protection system;
- I_{mean} : mean current demand, which is used to calculate the minimum mass of galvanic anode material or life of impressed current anodes necessary to maintain cathodic protection throughout the design period.

For optimising the design, the following should be specified:

- maximum and minimum resistivity, temperature and velocity of the water;
- design life of the cathodic protection system;
- initial current density necessary to achieve initial polarization of the structure, j_i ;
- maintenance current density necessary to maintain polarization of the structure, j_m ;
- mean coating breakdown factor, f_{cm} ;
- final coating breakdown factor, f_{cf} .

NOTE As the initial polarisation period before the steady state is normally short compared to the design life, the mean current density over the lifetime of the structure is usually very close to the maintenance current density.

Indicative values for maintenance and initial protective current densities and mean and final coating breakdown factors of conventional paint systems are given in Annex B.

The units shall be consistent in the following calculations and follow the state of the art in the field of engineering.

7.2.2 Subdivision of the surfaces to be protected

Items to be protected should be divided into different cathodic protection zones to ensure a good current distribution, which can be considered independently with respect to cathodic protection design, although they may not necessarily be electrically isolated.

NOTE Complex geometries such as those encountered at the bottom of ballast tanks due to the presence of stiffeners constitute a typical example.

Each cathodic protection zone may consist of several components, the parameters of which should be identified, including material (steel, cast iron, etc.), surface area and coating characteristics if any (type, lifetime and coating breakdown factor).