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## Construction of Three-Dimensional Finite Element Model for Complicated Cathodic Protection System

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**Abstract** When condensers with carbon steel waterbox and titanium tube structure were protected by SACP in an ocean beach power plant while seawater were used to be cooling water, carbon steel was under-protected and Titanium was over-protected, because their protective potential are less than  $-0.85\text{V}$  (vs SCE) and more than  $-0.75\text{V}$  (vs SCE) separately. In order to optimize the SACP design, potential distribution of two materials must be obtained during the SACP process. In the lab, carbon steel waterbox and Titanium tube model (Q235-Ti model) were set up. A series of electrochemical techniques (steady polarization, galvanostatic charge technique) were used to obtain reasonable boundary condition and surface form of two materials during the SACP process. The experiments were carried out to validate 3-D potential distribution under one anode SACP condition by finite element method (FEM). The calculation results were consistent with measuring results. The results can be used to provide a theoretical foundation to design an optimal cathodic protection.

**Keywords** condenser; cathodic protection; FEM; potential distribution

**CLC Number**: TG 172

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### 1 Introduction

Seawater is used to be cooling water in beach power plant. Condensers, as the main part of cooling system, were consisted with carbon steel waterbox and titanium tube. Carbon steel and titanium compose galvanic couple, so galvanic corrosion always happened and caused vast economic loss to power plant.

Cathodic protection (CP) is one of the most widely used methods of protecting metallic structure against corrosion. It can prevent or reduce the corrosion of any metal or alloy exposed to an aggressive environment. Basically, this technique is accomplished by impress current cathodic protection (ICCP) or sacrificial anode cathodic protection (SACP)<sup>[1]</sup>. The

use of sacrificial anodes has also been studied to provide cathodic protection for condenser in seawater.

Carbon steel was always under-protection and titanium was over-protection in cathodic protection, because their protective potential are less than  $-0.85\text{V}$  (vs SCE) and more than  $-0.75\text{V}$  (vs SCE) respectively. Hydrogen embrittlement for titanium occurs easily under over-protection.

In order to avoid above phenomena, it is very necessary to design reasonable SACP system to obtain the reasonable current density and potential distributions across the protected surface. Mathematical model of potential distribution was built to suggest optimum SACP system, including the optimum anode location and impressing current of anodes. Numerical

method is versatile in handling problem of complex structure in complex environment<sup>[2-4]</sup>.

Q235-Ti model was designed according to condenser structure in a beach power plant. The area ratio of Q235 to Ti was 55:1 in Q235-Ti model and the size of Q235 water-box was 200 mm × 200 mm × 400 mm and Ti tube was 400 mm long.

Finite element method (FEM) was applied to calculate 3-D potential distribution of the Q235-Ti model in this paper.

## 2 Mathematical Model

When the electrolyte can be considered to be homogeneous, the calculation of potential distribution on a cathodically protected structure in seawater first requires the solution of Laplace's equation for the distribution of potential in the bulk of seawater<sup>[5-13]</sup>:

$$\Delta^2 E = 0 \tag{1}$$

In order to obtain a complete solution to this equation, the potential is required to satisfy certain conditions specified for the anode and cathode surface. Since the sacrificial anodes can be assumed to operate at a constant value, the surface condition pertaining to the anodes can be stated simply in the form:

$$E = E_{anode} \tag{2}$$

The complexity of physical, chemical and biological phenomena occurring on the cathode surface (carbon steel and titanium tube) does not permit a simple relationship of current density and potential, especially in complex structure consisted of different metals. For Q235, such a relation is governed almost entirely by the properties of calcareous deposits formed on the steel surface during cathodic protection<sup>[1]</sup>. For titanium, multilayer titanium oxides are formed easily in cathodic polarization, and configuration of surface does not change evidentially.

The current density on cathode surface can be expressed mathematically as following when in fixed pressure, temperature, velocity of medium, initial surface condition and seawater chemical property:

$$i = \kappa \left( \frac{\partial E}{\partial n} \right)_r \tag{3}$$

Where  $\kappa$  is the conductivity of seawater,  $E$  is the

potential of cathode surface measured by a reference electrode adjacent to the surface,  $n$  is normal vector in the direction of current flow.

If the polarization is at steady condition, the relation of  $i$  and  $E$  on metal/electrolyte interface can be expressed as following:

$$i = \frac{1}{R_p} (E - E_c) \tag{4}$$

Where  $R_p$  is called polarization resistance and  $E_c$  is the open circuit potential of the protected metal.

So the equation as following is obtained by two equations above:

$$-\kappa \left( \frac{\partial E}{\partial n} \right)_r = \frac{1}{R_p} (E - E_c) \tag{5}$$

To obtain mathematical model, The Laplace's equation need be solved with boundary conditions reflecting the relationship between the potential and the current density and how this relation changes with time. at same time it is necessary to retrieve polarization data compatible with actual state of the structure.

In Q235-Ti model, Q235 steel and titanium have different cathodic polarization characteristics, the relationship between the potential and the current density and how this relation changes with time are different. Two different equations must be used in boundary condition:

$$-\kappa \left( \frac{\partial E}{\partial n} \right)_{r1} = \frac{1}{R_{p1}} (E - E_{c1}) \tag{6}$$

$$-\kappa \left( \frac{\partial E}{\partial n} \right)_{r2} = \frac{1}{R_{p2}} (E - E_{c2}) \tag{7}$$

Where  $\Gamma_1$  represents surface of Q235 steel and  $\Gamma_2$  represents Titanium,  $E_{c1}$ ,  $E_{c2}$  are the open circuit potential of the Q235 steel and Titanium respectively,  $R_{p1}$ ,  $R_{p2}$  are the polarization resistance of Q235 steel and Titanium respectively.

## 3 Experimental Results

In order to achieve mathematical models of SACP, the critical aspect of solving the Laplace's equation is appropriate boundary condition by polarization curve, which is a non-linear relationship between the two main variables, polarization potential and polarization resistance.

Q235 and Ti specimens with exposed surface ar-

ea of  $1 \text{ cm}^2$  were embedded in araldite. The specimens were given a metallographic polishing prior to each experiment followed by washing with distilled water and acetone. Polarization measurements were carried out using a three-electrode configuration: saturated calomel electrode (SCE) was used as reference electrode, a platinum electrode as counter one. In the paper potential values compare to SCE. HDV-7 potentiostat was used in steady polarization experiments. Electrochemistry measurements were performed using M6e (ZAHNER, Germany) electrochemistry workstation. Cathodic polarization for the two metals was carried out in different potential scale according to practical potential in CP.

1) Boundary condition experiments about Q235<sup>[14-20]</sup> were following

In the experiments the changes in corrosion current density with time from weak polarization were obtained after potentiostatic polarization at  $-0.800 \text{ V}$ ,

$-0.850 \text{ V}$ ,  $-0.900 \text{ V}$  in seawater. Potentiodynamic measurements were performed at a potential scanning rate of  $1 \text{ mV} \cdot \text{s}^{-1}$ . The calculated  $R_p$  were list in Table 1.

2) Boundary condition experiments about Ti were following

Titanium is prone to passivation in seawater. Multilayer titanium oxides were formed easily and intermediate resistance was steady<sup>[22]</sup>. The open circuit potential moved towards less negative values continuously. Galvanostatic charge technique was used to measure electrochemical parameter of Ti after polarized at  $-0.65 \text{ V}$ ,  $-0.70 \text{ V}$  and  $-0.75 \text{ V}$  respectively. The data fitting process for galvanostatic charge curve was programmed by MATLAB computer language based on linear regression. The calculated  $R_p$  were list in Table 2.

The design protection current densities for Q235 and Ti were  $90 \text{ mA/m}^2$  and  $55 \text{ mA/m}^2$  respectively. Al-

Tab 1 Variation of  $R_p$  with polarization time for Q235 in seawater after polarized at different potential calculated by dynamic polarization curves

Polarization Time/h	Potential $-0.800 \text{ V}$								
	13	25	37	39	42.5	44.5	48.5		
$R_p / \text{k}\Omega \cdot \text{cm}^2$	19.608	15.891	10.331	9.759	9.894	9.811	6.015		
Polarization time/h	Potential $-0.850 \text{ V}$								
	6	11.5	24	30.5	35	47.5	54.5	59.5	
$R_p / \text{k}\Omega \cdot \text{cm}^2$	6.973	9.122	9.795	14.979	18.226	20.901	21.6	22.105	
Polarization time/h	Potential $-0.900 \text{ V}$								
	5	12	24	48	60	144	192	280	384
$R_p / \text{k}\Omega \cdot \text{cm}^2$	2.260	2.935	2.609	6.245	10.422	15.858	17.478	20.982	18.890

Tab 2 Variation of the  $R_p$  with polarization for Ti in seawater time after polarized at different potential calculated by the galvanostatic charge curves

Polarization time/h	Potential $-0.650 \text{ V}$										
	3	18	29	55	78	102	119	135	142		
$R_p / \text{k}\Omega \cdot \text{cm}^2$	13.6	6.5	4.3	4.7	9.6	4.5	3.7	8.3	5.7		
Polarization time/h	Potential $-0.700 \text{ V}$										
	13	19	21	37	44	61	85	97	110	120	133
$R_p / \text{k}\Omega \cdot \text{cm}^2$	14.5	13.5	7.6	2.6	3.2	2.8	3.8	3.5	2.4	3.6	2.6
Polarization time/h	Potential $-0.750 \text{ V}$										
	7	19	25	43	60	69	86	91	110		
$R_p / \text{k}\Omega \cdot \text{cm}^2$	14.3	14.4	10.5	7.7	6.2	5.9	10.8	5.9	5.1		

Zn-In sacrificial anode was used for CP system. Fig 1 depicts location of the sacrificial anode in Q235-Ti model

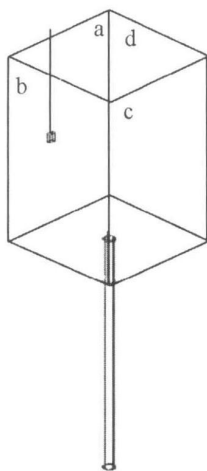


Fig 1 The sketch of sacrificial anode in Q235-Ti model (□ denote Al-Zn-In sacrificial anode)

The side which attached the sacrificial anode was defined as the a-side. And the rest sides in anticlockwise were defined as b, c, d. Fig 2 shows potential distributions by measuring in a, b, c, d sides in Q235-Ti model in turns.

In Q235-Ti model, cathode potential and current

provided by anode rapidly decrease with time at the beginning of polarization. After about 100 h, the current density decreases slowly and reaches apparent steady-state value at approximate 240 h.

### 4 Mathematic Model Analysis and Calculated Result

The boundary of the Q235-Ti model was discretized into tetrahedron unit as shown in Fig 3. Appropriate function was applied for discrete structure surface and its environment. Through analysis of each unit and synthesis in the whole area, the FEM was used to obtain the numerical solution of mathematical model.

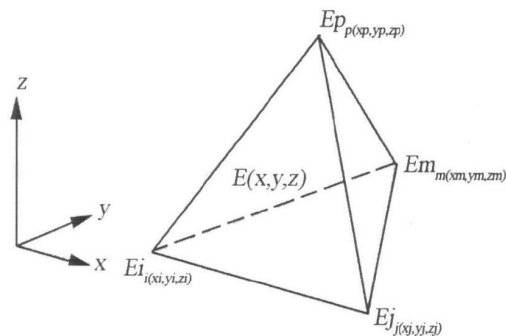


Fig 3 The sketch of tetrahedron

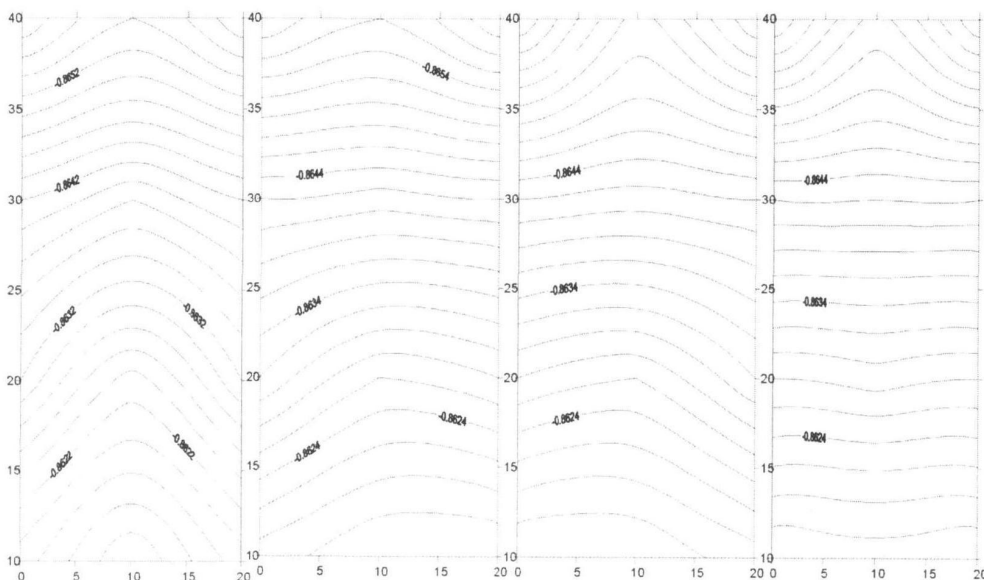


Fig 2 The distribution of potential in water box with one anode (m • V) (C)1994-2021 China Academic Journal Electronic Publishing House. All rights reserved. http://www.cnk

In the numerical procedure for calculating potential distributions the  $R_p$  can be either a constant or a variable of  $E$ . If  $R_p$  is a constant which is a good approximation in many activation controlled polarizations the constant is  $R_p$  assigned to all elements on the interfaces. The potential distribution is obtained by solving Equation (1) only once. However  $R_p$  in many corrosion systems is not a constant but is a function of  $E$ . In this case the potential distribution was obtained by iteration of  $R_p$ . After a period of protection in Q235-Ti model new  $R_p$  values were selected for individual elements on the surfaces based on the new potentials at the metal surfaces. This procedure is repeated automatically by computer until the potential closely match the previous iteration and at this point the solution has converged.

Fig 4 shows the calculated Q235 surface potential distribution by mathematical model on Q235-Ti model in SACP system in seawater. Side b is symme-

try of an equivalence relation to side d. Potential distribution of side b is same as side d. Potential distribution of underside of Q235 waterbox was depicted in Fig 4e.

Fig 5 shows the Ti-tube potential distribution. Application of the mathematical model were used to calculate the potential distribution of Q235-Ti model with one anode and two anodes in dilution seawater whose conductivity is the 1/10 of that of seawater by diluting with deioned water. the results of potential distribution in mathematical model were consist with potential distribution by measurement.

### 5 Conclusions

FEM fomulation was successfully used to calculate potential distribution in cathodic protection in complex construction based upon Laplace's equation.

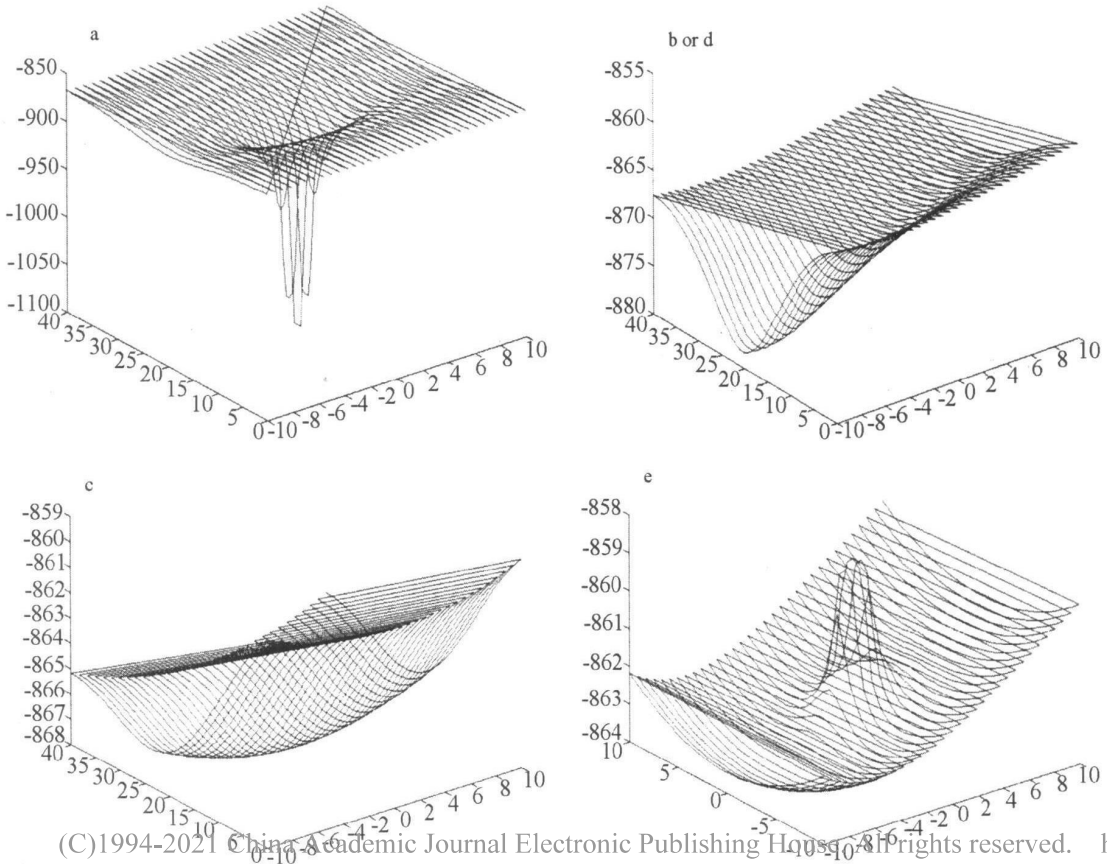


Fig 4 Potential distribution for Q235 surface by calculation (cm·mV)

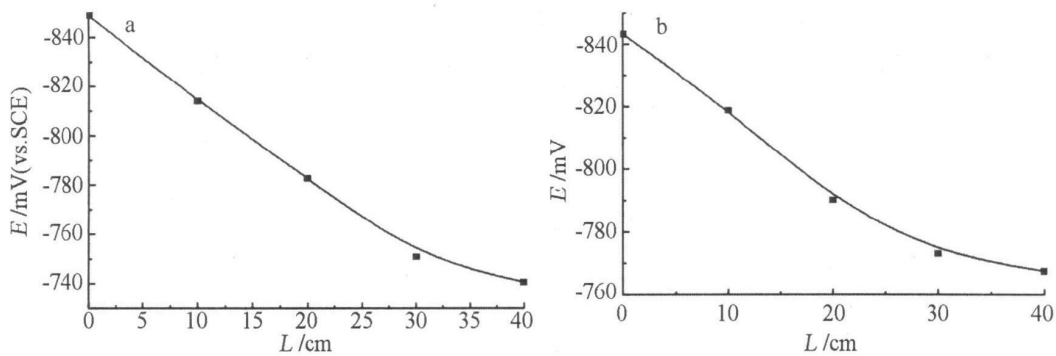


Fig 5 Potential distribution for Ti-tube a) measured data b) calculated data

Mathematical model of 3-D for water-box model was constructed. Mathematical model is feasible and available by comparing measured data and calculated data.

The use of modeling to simulate the detailed behavior of a SACP system reduces the uncertainty of the system's performance.

FEM can be applied to calculate 3-D potential distribution of the Q235-Ti model under different SACP conditions such as anode numbers, anode position etc. Computer modeling may also be used to predict potential in inaccessible part. For examples, potentials distribution for condenser in beach power plant can be predicted based on mathematical model.

## References

- [1] Chen Shaowei, Hartt W H. Steel cathodic polarization characteristic in sea water and a new approach for cathodic protection system design [J]. *Corrosion Science and Protection Technology* 1996, 8(1): 17-25.
- [2] John W Fu. A finite element analysis of corrosion cells [J]. *Corrosion* 1982, 38: 295-296.
- [3] Raymond S Munn. A mathematical model for a galvanic anode cathodic protection system [J]. *Corrosion* 1982, 38: 171-185.
- [4] Luiz C. Wrobel, Panayiotis Miliadou. Genetic algorithms for inverse cathodic protection problems [J]. *Engineering Analysis with Boundary Elements* 2004, (28): 267-277.
- [5] Ramanan V S, Muthukumar M, Gnanasekaran S et al. Green function for the Laplace equation in a 3-layer medium boundary element integrals and their application to cathodic protection [J]. *Engineering Analysis with Boundary Elements* 1999, 23: 777-786.
- [6] John W Fu, Siu-kee Chan. A finite element method for modeling localized corrosion cells [J]. *Corrosion* 1984, 40: 540-545.
- [7] Amaya K, Aoki S. Effect of boundary element method in corrosion analysis [J]. *Engineering Analysis with Boundary Element* 2003, (27): 507-519.
- [8] Cao Shengshan (曹圣山), Wang Qingzhang (王庆璋), Zhang Manping (张曼平). Numerical modeling for offshore platform cathodic protection local system [J]. *China Oceanol Limnol* 1995, 13: 247-252.
- [9] Rolf G. Kasper, Martin G. April. Electro-galvanic finite element analysis of partially protected marine structures [J]. *National Association of Corrosion Engineers* 1983, 39: 181-188.
- [10] Schwenkw. Current distribution during the electrochemical corrosion protection of pipes [J]. *Corrosion Science* 1983, 23: 871-886.
- [11] Nisancionglu K. Predicting the time dependence of polarization on cathodically protected steel in seawater [J]. *Corrosion* 1987, 43: 100-110.
- [12] Fu J W, Chow J S K. Cathodic protection designs using an integral equation numerical method [J]. *Materials Performance* 1982, 21: 8-12.
- [13] Raymond S Munn. A mathematical model for a galvanic anode cathodic protection system [J]. *Materials Performance* 1982, 21(8): 29-36.
- [14] Fu J W, Chan S K. Finite element determination of galvanic corrosion during chemical cleaning of steam generator [J]. *Materials Performance* 1986, 25: 33-40.
- [15] Hartt W H, Kunjapur M M. Influence of temperature and exposure time upon calcareous deposits [J]. *Corrosion* 1987, 43: 674-669.
- [16] Perolv G, Gartland, Roe D, Strummen. Offshore cathodic

- protection design inspection and computer modeling [J]. *Material Performance* 1993, 11, 453-460.
- [17] Brenda J little Patricia A Wanger Interrelationship between marine befouling and cathodic protection [J]. *Material Performance* 1993, 9, 369-379.
- [18] Pretice G, Holser R A, Jfarozic V, et al Cathodic protection modeling of galvanically coupled heat exchanger [J]. *Corrosion* 1990, 46, 75-84.
- [19] WEN Guomou (温国谋), ZHENG Fuyang (郑辅养). Impedance characteristics of calcareous deposit [J]. *Chin J Oceanol Limnol* 1996, 14, 277-281.
- [20] Ch Barchiche Deslouis C, Festy D, et al Characterization of calcareous deposits in artificial seawater by impedance techniques <sup>3</sup>-deposit of CaCO<sub>3</sub> in the presence of Mg (II) [J]. *Electrochimical Acta* 2003, 48, 1645-1654.
- [21] Creus J, Idrissi H, Mazille H. Corrosion behavior of Al/Ti coating elaborated by cathodic arc PVD process onto mild steel substrate thin solid films [J]. *Thin Solid Films* 1999, 346, 150-154.

## 复杂阴极保护体系三维有限元建模研究

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**摘要:** 滨海电厂常以天然海水作冷却水,其碳钢水室和钛管凝汽器在实施阴极保护过程中,要求钢电位应低于  $-0.85\text{V}$  (vs SCE), 电位过正则碳钢保护不足,而钛的电位则须高于  $-0.75\text{V}$  (vs SCE), 电位过负则易使钛管发生氢脆损伤. 研究牺牲阳极阴极保护, 需先得出两种金属在阴极保护过程中的电位分布. 本文在设计室建立碳钢水室钛管模型, 根据该实验体系阴极保护过程的 (稳态极化和恒电流极化) 测定获得两种金属有效的边界条件, 用有限元 (FEM) 计算在一个牺牲阳极阴极保护状态得到三维电位分布. 计算结果和测量结果基本一致, 为优化阴极保护设计提供了理论依据.

**关键词:** 凝汽器; 阴极保护; 有限元; 电位分布