

# 8th Internation Conference on Alkali–Aggregate Reaction

# SUPPRESSION OF ALKALI-AGGREGATE REACTION BY CONCRETE SURFACE COATING

Tohru Taki\*, Michio Takeyoshi\*\* Kazuaki Noda\*\*\*, and Kiyoshi Katawaki\*\*\*\*

- \* Fundamental Research Laboratory, Dai Nippon Toryo Co., Ltd., 1382-12 Shimoishigami, Ohtawara, Tochigi-Pref., 329-26, Japan
- \*\* Protective Coatings Dept., Nippon Paint Co., Ltd., 4-1-15 Minamishinagawa, Sinagawa-Ku, Tokyo, 140, Japan
- \*\*\* Marine & Protective Coatings R & D Dept., Kansai Paint Co., Ltd., 4-17-1 Higashi-yawata, Hiratsuka, Kanagawa Pref., 254, Japan
- \*\*\*\* Public Works Research Institute, Ministry of Construction, Asahi-1, Tsukuba, Ibaragi Pref., 305, Japan

# 1.INTRODUCTION

One of the causes of concrete deterioration is alkali-aggregate reaction. In this study, concrete specimens containing alkali reactive aggregate were coated with various organic and inorganic coatings to examine suppressing effect of cracking caused by alkali-aggregate reaction (AAR). Some of the specimens were coated prior to AAR acceleration test, and the others were coated after cracking has appeared. In addition, transmission rate of AAR accelerating substances and extension ratio at break for the coating films to discuss relationship between these basic properties of coating films and crack formation.

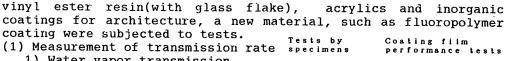
### 2. OUTLINE OF THE STUDY

### 2.1 Objective of the study

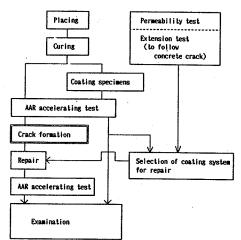
The study was conducted to make following points clear. (1) Suppressing effect of coating films on concrete stability, before and after cracking starts. (2) Which properties of coating films affect to suppress cracking? Flow diagram of the study is shown in FIGURE-1. The experiment mainly consists of two parts; the stability test using concrete specimens, and the measurements of coating film perfomances.

#### 2.2 Procedures of the experiment

2.2.1 <u>Performance tests of coating films</u> Thirty two coatings, i.e., heavy duty types, such as epoxy resins and a



- 1) Water vapor transmission rate
  - a) JIS-Z-0208 (dish method)
  - b) Lyssy method (electrical method, using humidity sensor)
  - 2) Water transmission rate
    - a) JIS-A-6900-5-10
  - b) Gravimetrical method (using coated specimens)
  - 3) Ion transmission rate Transmission rate of Na\* and Cl was determined after a month's immersion test using double cells (3%NaCl soln./distilled water) which were divided by a free film.



(2) Extension ratio at break of coating films to over coat surface cracking concrete

FIGURE-1 The flow diagram of the study

Extension ratio at break was determined using Instron type tester with crosshead speed 5mm/min. at 20 C. 2.2.2 Preperation of concrete specimens and the procedure of

AAR acceleration test

(1)Mix proportion of concrete

Mix proportion of concrete is shown in TABLE-1. Specimens were subjected to the test after 28 days' curing in water.

No.1 and No.3 were subjected to the test as uncoated, No.2 both coated and uncoated. Specimens were cubic, having as dimension of 100 mm on each side.

No.	(1)	V kg/m³	C kg/m³	S kg/m³	G kg/∎³	(2)	NaCi kg/m³	₩/C ¥	S/A X	Air ¥	Slump cm	(3)	(4)	(5)
Stand	lard	181	362	686	993	0.025	•	50	40	5.0	8.0	-	-	-
1	т	180	362	687	993	0.025	-	50	40	5.2	11.3	0.1	502	3.27
2	T	176	362	691	993	0.025	8.8(6)	50	40	5.4	13.8	0.7	481	3.09
3	ĸ	180	362	687	993	0.025	-	50	40	5.2	11.3	0.1	459	3.30

TABLE-1 Mix proportion of concrete

(1) Sort of aggregate

T: Judged potentially reactive through ASTM C 289. Rc=183, Sc=639

K: Judged nonreactive through ASTM C 289. Rc=34.3, Sc=39.6

(2) A E Agent (%)

(3) Surface moisture in fine aggregate (%)

(4) Compression strength (kgf/cm<sup>2</sup>)

(5) Modulus of elasticity  $(10^5 \text{kgf/cm}^2)$ (6) As total R<sub>2</sub>0(Na<sub>2</sub>0+0.658k<sub>2</sub>0)=8kg/m<sup>2</sup>

- 828 ---

(2) Coating of specimens

1) Coating before crack appeared

Specimens were treated with disc sander (cc#50 paper) prior to coat after 28 days' curing in water. They were coated all of the surfaces. Targets of dry film thickness were 100  $\mu$ m as standard, 350  $\mu$ m as specified. Coatings contained neither primer nor putty inorder to evaluate their properties directly. They were dried for 7 days at 20 °C prior to subject to the test.

2) Repair coating after crack appeared

Target of dry film thickness was specified value of each coating system. Coating films were evaluated as film system (each contains primer, putty, top coat). Pretreatment and coating method were as same as mentioned above.

(3) AAR acceleration test

Specimens were subjected to the wet/dry cycle exposure test. They were placed under the sun periodically being sprayed water.

Spraying period was 30 minutes, four times a day. It was obvious that the wet/dry cycle test accelerated AAR more effectively than immersion or high temperature humidity test.

## 3. RESULTS AND DISCUSSION

#### 3.1 Performance tests of coatings

Test results were shown in TABLE-2. Vinyl ester, epoxy and polyurethane resins were highly resistant to water or water vapor transmission, on the other hand, inorganic and acrylic coatings were highly transmittable. Inorganic and acrylic coatings were highly transmittable to sodium ions, while vinyl ester, epoxy and polyurethane coatings hardly transmitted sodium ions.

1) film thickness

File thickness
 Faint maker
 Vaint maker
 Vair vapor transmission rate (g/m<sup>2</sup> + 24hrs.) a: Heasured through JIS-2-0208 (20 °C) b: Heasured through Dr.Lyssy method (40°C) 4) Vater transmission rate (m1/m<sup>2</sup> + 24hrs.) Vater transmission rate (m2/m<sup>2</sup> + 24hrs.) b) A' transmission rate (m2/m<sup>2</sup> + 24hrs.) Cl transmission rate (m2/m<sup>2</sup> + 24hrs.) B) Cl transmission rate (m2/m<sup>2</sup> + 24hrs.) B) Extension ratio at break point (1)

TABLE-2 Test results of coatings

TADL	L-Z lest	resu	1173	S 01	coa	lings				
Coaling	1) f.T.	2) H	٧V	T 3)	4) ¥T(1)	5) VT(2)	6) Na*	(1 <sup>-</sup>	8) E.R.	
		(µ)			b					
	Epoxy		A				0.017			
	(primer)	-	B			23.0	0.079			
		-	C			78.0	0.024	1	1	
Ероху	Ероху	100	A	3.4		7.0	0.000	+	6.96	0.93
	(int.)	100	R		1.7		0.007			1.2
1		100	C	1.4		2.1	0.003	*	*	4.1
	Epoxy(top)	100	A	2.7		6.9		*	0.10	2.2
	Epoxy	100	A	5.8		10.0	0.015	* [	1.38	130
	(flexible)	100	C	2.8			a ine	* 1	*	25
	Epoxy	350	A	1.4	2.5	1.7	0.025	.*	<b>*</b>	1.8
	(high-build)	350 350	BC	0.2	2.5			*		1.5
			Ľ.							
	Ur.(primer)	-	(B	1		1,2	0.004			
Urelhane	Ur.(top)	100	8		35			*		41
		100	A	2.6		21		+		77
	Ur.(flex.)	100	B	1	52			+	1.38	68
	Vater		A	81		60	0.10	900	400	
In-	repeilent		ß		Į	20	0.14		100	
огдаліс		-	İč			60	0.16			
	Inorganic	350	A	ļ		4900	1.40	31000	53700	55
	-	100	c	240	1	• ·	1		944	55
	P.cement	350	18	j	35	7.0		6.88		94
	(flexible)			ţ			[			
V.Es.	Glass flake	350	B		0.6	+		<b>\$</b> .		1.5
	Acrylic	100	1	13.0		10.0	0.018	0.17	2.43	171
Others	(solvent)	100	lî.	1.3.0	160	,	0.010	2490	2.13	28
0.001.3		100	le.	13.0	1			20.5	12.6	
	Acrylic(em.)	100	B		1960	1560	1	21000		8.3
	Acrylic(em.)	100	1c	200	1	1			4330	940
	(flexible)		1			i i	1			
	Polybulailien	100	C	29	1		ļ	\$	*	260
	(flexible)	1			1	1				Ì
	Vinyl	100	B	· ·	34	]	1	1		3.0
	Chid.rubber	100	C	2.9		ļ		19.3	14.6	250
	(flexible)	1	1	1	1	j				1
	Fluoric	100	B		26			+		44

--- 829 ----

#### 3.2 AAR acceleration test of uncoated or originally coated specimens

All of the No.1 and No.3 specimens in TABLE-1 were subjected to the test as uncoated. Thirty in 86 of No.2 specimens were subjected to the test after being coated, while 56 as uncoated.

3.2.1 Crack formation in uncoated specimens As shown in FIGURE-2, cracking was observed in 90 per cent of No.2 specimens after 12 months, 100 per cent after 24 months. While, No.1 specimens (potentially reactive, but without NaCl) were observed no cracking after 17 months, and 7 in 54 were observed to have been cracked (fine crack; under 0.05mm) after 24 months. The crack was considered to originate in AAR, because white gel was observed and it was remarkably accelerated to appear by addition of NaCl.

3.2.2 Crack formation	in	specime	ens	which	have	be	en	coat	ted
before crack appeared		Crack	was	obset	rved	in	33	per	cen
originally coated									
awarimawa often 16									

specimens after 16 months, and in 48 per cent after 23 months. Crack suppressing effect of coatings was obvious.

Relationship between crack formation and water vapor transmission rate or extension ratio is shown in TABLE-3 and FIGURE-2. As shown in FIGURE-2, cracking formation ratio was low in A type, and it formed in early time, no increace was noticed by passage of time. C type showed low ratio of crack

formation too, but it increaced by

TABLE-3 Number of cracked specimens

	Coating prope	rties		Number o	Number of specimens					
	Transmission Extension		Crack	11months	16months	23months				
A			Finded	1	1	1				
	Low	Small	Non	4	4	4				
			(XFinded)	20	20	20				
			Finded	4	6	8				
8	Mediuma	Small	Non	4	6 5	8 3				
•			(%Finded)	36	55	73				
			Finded	0	1	4				
С	Medium	Large	Non	07	_1 6	3				
•			(%Finded)	0	14	57				
			Finded	0	0	1				
D	High	Large	Non	1	1	0				
•			(XFinded)	0	0	100				
E			Finded	2	2	2				
	High	Small	Non	1	1	1				
			(%Finded)	67	67	67				

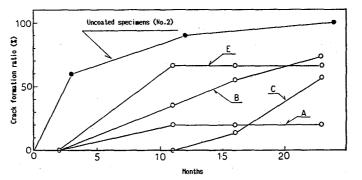
cent of

1 - 100 Medium 100 < High Small 0 - 10 % Extension

Large 20 % < ratio

rate

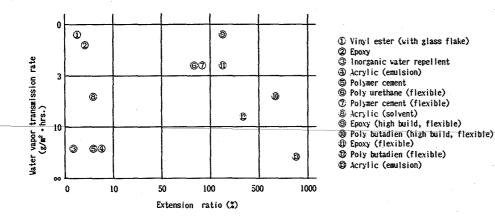
passage of time. The reason was considered as follows. As the specimens were coated under well dried condition, if they were coated with nontransmmittible coatings, expansion from AAR was considered to be suppressed.



In case that a specimen is FIGURE-2 Crack formation ratio with time

coated with a coating which is comparatively low water vapor transmittible and having high extension ratio, it is considered that the coating may yet suppress cracking of the specimen after it expands to a certain extent, because the coating film will extend following to expantion of the specimen and may keep a certain extent of transmission resistance. On the other hand, in case that a coating film has comparatively high transmission resistance, but poor in extending property, micro-fault might occur in it according to the specimen expands, as a result, transmission resistance of the film might decrease and water transmission into the specimen will occur, resulting in further expantion of the specimen and crack formation on it.

3.2.3 <u>Crack formation in repair coated specimens</u> After 4 months' acceleration test, cracked specimens of No.2 were repaired with 13 types of coating systems shown in FIGURE-3 and subjected to the acceleration test again.



- 831 -

FIGURE-3 Properties of selected repair systems

Each system was coated on two specimens, having small crack and large one. No crack growth was observed on the coating surface after 20 months' acceleration test. Then coating films were removed to examine crack growth under the films, but no growth of crack was observed.

# 4. CONCLUSION

- Cracking due to AAR of concrete which contains alkali reactive aggregate is expected to be suppressed by surface coatings.
- (2) As shown in this study, if all of the surface are coated with a coating of low transmission rate towards water, cracking of concrete is considered to be suppressed effectively.
- (3) Concrete cracking could not be prevented perfectly by coating concrete surface before crack formation, while crack effectively suppressed by coating growth was after former case too large crack formed, because in the extension ratio of the coating film would be required to cover crack width, on the other hand requirement for extension ratio of the coating film would be much smaller than that of latter case.
- (4) In the actual concrete structures which might be inevitable to have uncoated portion from where water can permeate, other coating system might be required. (The authors are conducting an further experiment, in which coated concrete specimens having uncoated portion are exposed under the sun being buried uncoated portion in the earth.)

#### 5. ACKNOWLEDGMENT

This report is the result of the study which was performed by M.Takeyoshi, K.Noda, and T.Taki when engaging as outside researcher in Public Works Research Institute in 1985-1986. The authors are grateful to the participants in the Institute who gave guidance and advice.

#### REFERENCE

[1] K.Noda, M.Takeyoshi, T.Taki, K.Katawaki, Proceedings of JCI Vol.9, No.1, p591-596, (1987)