Part of the Journal of Research of the National Bureau of Standards

# Soil-Corrosion Studies, 1946: Ferrous Metals and Alloys

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This report contains the results of measurements of corrosion made on a variety of wrought and cast ferrous materials after exposure to different soil conditions for periods up to 14 years. The steels ranged in composition from fractional percentages of nickel and chromium to the high concentrations typical of wholly austenitic steels. The soils ranged from well aerated soils deficient in soluble salts to poorly aerated soils containing high concentrations of water-soluble materials. The magnitude and progress of corrosion with respect to both weight loss and pitting are interpreted in relation to the properties of the soils at the test sites. A comparison is made of the corrosion of specimens of plain wrought materials and cast iron.

## I. Introduction

In 1932 the Underground Corrosion Section of the National Bureau of Standards undertook the investigation of the behavior of a variety of ferrous and nonferrous metals and alloys in soils representing a wide range of euvironmental conditions. The primary objective of the investigation was to ascertain whether metals and alloys that were especially resistant to corrosion in the atmosphere and in natural waters were also resistant to underground corrosion, at least in specific soil environments. It was anticipated that in the course of the investigation the specific effects on corrosion resistance of various alloying elements under different soil conditions might also be evaluated.

Originally, a sufficient number of specimens were buried at 14 test sites to provide for removal of two specimens of each material after each of five periods of exposure. Accordingly, specimens were removed from each test site in 1934, 1937, 1939, 1941, and 1946 and returned to the laboratory for measurement of weight loss and maximum depth of pits. At the time of removal of the specimens in 1937, 1939, and 1941, additional sets of specimens of some of these classes of materials were buried at the test sites and were removed in 1946. Results of exposures of 14, 9, 7, or 5 yr are, therefore, available for these materials. Corrosion data for the removals prior to 1946 have been published in a series of reports that were summarized by K. H. Logan [1].<sup>1</sup>

This paper deals only with the tests of wrought and cast ferrous materials. The results of the tests of nonferrous materials will be presented in a subsequent paper.

## II. Properties of the Soils at the Test Sites

The test sites were selected to represent a wide range of soil conditions, from slightly corrosive to highly corrosive. The considerable differences in the physical and chemical properties of the soils at the test sites are evident from the data of table 1. The resistivity ranges from 62 ohm-cm, corresponding to a high concentration of soluble salts, to 17,800 ohm-cm, indicating practically no soluble material. The water-soluble material in soil 64, Docas clay, consists of sodium and potassium chlorides, whereas the soluble material in soil 60, Rifle peat, is composed entirely of sulfates of the alkalies and of the alkaline earth metals. The hydrogen-ion concentration of the soils ranges from pH 2.6 to pH 9.4. Measurements of pH and total acidity were made in the laboratory on samples of soil shipped from the test sites in sealed containers. This precaution was taken because results of a previous study showed that aeration of certain poorly aerated soils produced marked increases in acidity because of oxidation of sulfides to sulfates [2].

#### Soil Corrosion of Ferrous Materials

 $<sup>^1\,{\</sup>rm Figures}$  in brackets indicate the literature references at the end of this paper.

TABLE 1. Properties of the soils at the test sites

	Soil			Mois-	Appar-	Resistiv-		Total acidity <sup>b</sup> (milli-		Compo	sition of	water o	extract-m .00 g of so	illigram il	equivale	nts per
No.	Type	Location	Aera- tion <sup>a</sup>	ture equiv- alent	ent specific gravity	ity at 60° F (15.6° C)	pH	gram equiv- alents per 100 g of soil)	Sulfide con- tent °	Na+K as Na	Ca	Mg	CO3	HCO3	Cl	$SO_4$
51 53 55	Acadia clay Cecil clay loam Hagerstown loam	Spindletop, Tex Atlanta, Ga	P G G	Percent 47.1 33.7 32.0	2.07 1.60 1.49	<i>Ohm-cm</i> 190 17, 800 5, 210	6.2 4.8 5.8	13.2 5.1 10.9	+	10. 27	15. 55	5.03	0.00	0. 56	5. 75	22.00
56	Lake Charles clay	El Vista, Tex	VP	28.7	2.03	406	7.1	5.1	+	3.12	0.69	0.47	0.00	0.80	1.59	3.04
58 59 60 61	Muck Carlisle muck Rifle peat Sharkey clay	New Orleans, La Kalamazoo, Mich Plymouth, Ohio New Orleans, La	P VP P P	57. 8 43. 6 43. 4 30. 8	1.43 1.28 1.78	712 1,660 218 943	$\begin{array}{c} 4.8 \\ 5.6 \\ 2.6 \\ 6.8 \end{array}$	$     15.0 \\     12.6 \\     297.4 \\     4.9   $	+ +++ ++ +	2. 03 1. 03 2. 91 0. 73	$\begin{array}{c} 2.\ 23\\ 3.\ 08\\ 10.\ 95\\ 0.\ 68\end{array}$	1. 29 2. 70 2. 86 0. 33	.00 .00 .00 .00	. 00 . 00 . 00 . 71	.47 3.47 .00 .10	2. 54 1. 04 56. 70 0. 91
	Susquehanna clay Tidal marsh Docas clay Chino silt loam	Meridian, Miss. Charleston, S. C. Cholame, Calif Wilmington, Calif.	P VP F G	34. 646. 741. 126. 4	$     1.79 \\     1.47 \\     1.88 \\     1.41 $		4.5 6.9 7.5 8.0	12.0 14.6 A A	 ++++ - -	33.60 28.10 7.65	$ \begin{array}{r}     6.85 \\     2.29 \\     12.40 \end{array} $	4.00 .76 2.20	0.00 .00 .00	0.00 .89 1.30	$12.\ 70\\28.\ 80\\6.\ 05$	36.60 . 26 16.90
66 67 70 72	Mohave fine gravelly loam Cinders Merced silt loam Papakating silty clay loam	Phcenix, Ariz Milwaukee, Wis Buttonwillow, Calif Deerfield, Ohio	$F \\ VP \\ F \\ VP$	16. 5 24. 7	1.79 1.69	232 455 278 762	8.0 7.6 9.4 7.2	A A A 17	- + - +	$\begin{array}{c} 6.55 \\ 0.77 \\ 8.38 \\ 3.11 \end{array}$	0.51 3.03 0.38 7.49	$\begin{array}{c} 0.\ 18 \\ .\ 53 \\ .\ 22 \\ .\ 95 \end{array}$	.00 .00 .02	$\begin{array}{c} 0.\ 73 \\ .\ 55 \\ 1.\ 87 \end{array}$	2.77 0.08 1.12	2. 97 2. 89 5. 57 11. 58

<sup>a</sup> Aeration of soils: G, good; F, fair; P, poor; VP, very poor.

<sup>1</sup>A indicates absence of acidity because of alkaline reaction.

- Sulfides absent; + sulfides present in low concentration; ++ sulfides present in moderate concentration; ++ + sulfides present in high concentration.

The retentiveness of the soils for moisture is indicated by the moisture equivalent, i. e., the quantity of water retained by a previously saturated soil against a centrifugal force of 1,000 times the force of gravity. Values for moisture equivalent range from 16.5 percent for soil 66, Mohave fine gravelly loam, to 57.8 percent for soil 58, muck.

 
 TABLE 2.
 Classification of soils according to type of environment

Soil No.	Soil type	Environment
51	Acadia clay	Inorganic-reducing-acid.
53	Cecil clay loam	Inorganic-oxidizing-acid.
55	Hagerstown loam	Do.
56	Lake Charles clay	Inorganic-reducing-alkaline.
58	Muck	Organic-reducing-acid.
59	Carlisle muck	Do.
60	Rifle peat	Do.
61	Sharkey elay	Inorganic-reducing-acid.
62	Susquehanna elay	Inorganic-oxidizing-acid.
63	Tidal marsh	Organic-reducing-acid.
64 65 66 67 70	Docas clay Chino silt loam Mohave fine gravelly loam Cinders Merced silt loam	Inorganic-reducing-alkaline. Inorganic-oxidizing-alkaline. Do. Inorganic-reducing-alkaline.

Values for apparent specific gravity are presented as an index of the relative porosity of the soils. Because the real specific gravities of the mineral portion of soils lie within a narrow range, the apparent specific gravity indicates the compactness and hence the relative porosity of soils that are primarily inorganic in nature.

On the basis of their properties the soils were classified according to type of environment. This classification is given in table 2.

# III. Description of the Materials

The forms, dimensions, and compositions of the specimens are given in tables 3 and 4. Although the identifying letters shown in the table are the same for a few materials, the dates of burial of these specimens are different. As comparisons will be made only between materials exposed for the same length of time, no confusion is likely to result from use of the same symbol for different samples. Specimens of some of the materials were in the form of pipe, others of plate. The ends of the pipe specimens were closed by caps which excluded moisture, thereby preventing corrosion of the interior. As an extra precaution against possible corrosion, the inner surfaces of the specimens were coated with heavy grease. The exposed area of each specimen was approximately  $0.3 \text{ ft}^2$ . All of the materials were free of mill scale except as noted in table 3.

TABLE 3.	Composition	of the	wrought	metals
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Material	Iden- tifica- tion	Year buried	Form	Nomi- nal width or di- ameter	Length	Thick- ness	C	Si	Mn	S	Р	Cr	Ni	Cu	Мо	Other elements
						WROU	GHT IR	ON								
Hand-puddled Mechanically puddled	a A a B	1932 1932	Pipe	In. 1.5 1.5	In. 12 12	In. 0. 145 . 145	Percent 0.016 .017	Percent 6.10 .125	Percent 0.029 .041	Percent 0.018 .018	Percent 0. 160 . 106	Percent	Percent	Percent	Percent	Percent Oxide+slag, 2.56. Oxide+slag, 2.681.
						CARBO	N STEI	ELS								
Low-carbon steel	N	1932	Pipe	2.3	10	0.145	0.15		0.49	0.030	0.013					
				I	OW-AL	LOY IF	ONS A	ND STI	EELS				×			· · · · · · · · · · · · · · · · · · ·
Open-hearth steel. Copper-molybdenum open-hearth iron Do Do Do Copper-nickel steel. Nickel-copper steel. Do Do * Carbon steel layers. Ni-Cu steel layers. Do Chromium-silicon-copper-phosphorus steel. 2% chromium steel with molybdenum	A MM O N H H H B J b B Y o P  D C N N	1937 1939 1937 1932 1941 1937 1937 1937 1941 1941  1932 1937 1937	Plate do Pipe Plate Plate Plate Plate Plate Plate Plate Plate	$\begin{array}{c} 2.5\\ 2.5\\ 2.5\\ 1.5\\ 1.5\\ 2.5\\ 2.5\\ 2.0\\ 2.5\\ 2.0\\ 2.5\\\\ 1.5\\ 2.5\\ 1.5\\ 1.5\\ 1.5\\ \end{array}$	$ \begin{array}{c} 12\\12\\12\\12\\12\\12\\12\\12\\12\\12\\12\\12\\14\\12\\12\\12\\12\\14.5\end{array} $	0. 188 . 250 . 243 . 250 . 145 . 145 . 265 . 248 . 210 . 172 . 036 . 100 . 145 . 188 . 145	6.033 .04 .03 .06 .04 .06 .06 .07 .12 .16 .14 .14 .075 .09	0.002 .003 .001 .05 .005 .047 .14 .12 .21 .16 .19 .84 .25	0.029 .16 .098 .32 .16 .49 .44 .58 .78 .48 .21 .20 .46	0. 017 . 027 . 032 . 029 . 027 . 025 . 025 . 025 . 025 . 030 . 009 . 018 . 010	0.006 .008 .007 .069 .016 .008 .095 .010 .07 .018 .021 .124 .015	0. 049 . 04 . 02 . 02  . 04  . 04  1. 96	0. 034 . 14 . 15 . 14 	$\begin{array}{c} 0.\ 052\\ .\ 51\\ .\ 45\\ .\ 54\\ .\ 52\\ .\ 47\\ .\ 95\\ 1.\ 01\\ 1.\ 04\\ .\ 0038\\ .\ 98\\ 1.\ 08\\ .\ 428\\ \end{array}$	0.07 .07 .13 .15 .08    	O <sub>2</sub> , 0.015; N <sub>2</sub> , .008; Sn, 0.002.
Do	KK	1937	Plate	2.5	12	•.175	. 082	. 51	. 46	.015	. 017	2.01	.07	.004	. 57	

					0	CHROM	IUM S'	FEELS								
4 to 6% chromium steel Do 4 to 6% chromium steel with molyb- denum. Do 12% chromium steel 18% chromium steel Do.	P D E H U V X	1932 1937 1937 1937 1932 1932 1932	Pipe Plate do do Pipe	2.3  2.5  2.5  2.5  4  4  1.5	$ \begin{array}{c} 10 \\ 12 \\ 12 \\ 12 \\ 6 \\ 6 \\ 12 \\ \end{array} $	$\begin{array}{c} 0.\ 154 \\ .\ 245 \\ .\ 188 \\ .\ 203 \\ .\ 063 \\ .\ 063 \\ .\ 145 \end{array}$	$\begin{array}{c} 0.13 \\ .077 \\ .074 \\ .060 \\ .065 \\ .070 \\ .12 \end{array}$	0. 43 . 41 . 39 . 28 . 34 . 277	$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$	$\begin{array}{c c} 0.\ 025\\ .\ 005\\ .\ 006\\ \hline .\ 014\\ .\ 017\\ .\ 015\\ .\ 017\\ \end{array}$	0.012 .015 .013 .021 .011 .014 .016	5.05 5.02 4.67 5.76 11.95 17.08 17.72	0.09 .09 .17 .482 .092 .287	0.008 .004 .004 .025 .021	0. 51 . 43 	Al, 0.030; Ti, 0.022. Al, 0.27.
			HIGH-CH	ROMIU	JM STH	EELS W	ITH N	 ICKEL	AND N	 IANGAI	NESE					
<ul> <li>18% chromium steel with nickel</li> <li>Do</li> <li>Do</li> <li>18% chromium steel with nickel and manganese.</li> <li>Do</li> <li>18% chromium steel with nickel, manganese, and molybdenum.</li> <li>22% chromium steel with nickel and manganese.</li> <li>22% nickel-chromium steel with manganese and molybdenum.</li> </ul>	K R W T S CM Y DT	1932 1932 1932 1932 1932 1939 1932 1939	Plate Pipe Plate do do do do do	$3 \\ 1.5 \\ 4 \\ 6 \\ 2 \\ 4 \\ 2.5 $	$ \begin{array}{c} 11\\ 12\\ 6\\ 10\\ 12.5\\ 6\\ 12 \end{array} $	$\begin{array}{c} 0.\ 025\\ .\ 145\\ .\ 063\\ .\ 063\\ .\ 25\\ .\ 063\\ .\ 25\\ .\ 25\\ .\ 25\\ \end{array}$	0.08 .05 .093 .06 .07 .144 .07	0.33 .28 .42 .40 .48 .40 .59 .91	0.44 .46 .36 6.09 9.44 1.24 1.80 1.99	0.022 .011 .017  .008 011 .012	0.015 .015 .008  .016 .015 .014	17. 20 17. 52 18. 69 17. 76 17. 78 17. 78 22. 68 19. 27	8.95 8.85 9.18 3.83 10.96 12.94 22.12	0.016 .95 .74 .021 1.07	2. 63  3. 52	

<sup>a</sup> Some mill scale on the surface at the time of burial.

b Specimens completely covered with a hard, black mill scale at the time of burial.
c Specimens composed of three layers as follows: outer layers—plain carbon steel, 0.036 in. thick; intermediate layer—2-percent nickel, 1-percent copper steel, 0.100 in. thick.

TABLE 4. Composition of the cast metals

	Tdanti	V		Nomi- nal		This		С								
Material	fication	buried	Form	width or di- ameter	Length	ness	Free	Com- bined	Total	Si	Mn	S	Р	Cr	Ni	Cu
				Inch	Inch	Inch	Per- cent	Percent	Percent	Per- cent						
Rattled cast iron a	G	1932	Pipe	1.25	13.5	0.250	2.94	0.64	3.58	1.64	0.48	0.074	0.79			
Sand-coated cast	F	1932	do	1.25	13.5	. 250	2.94	. 64	3.58	1.64	. 48	. 074	. 79			
iron.																
Plain cast iron	. A.	1941	do	1.5	13	. 250			3.22	2.19	. 91	. 12				
Low-alloy cast iron.	I	1932	do	1.25	12	. 350			2.53	1.43	. 28	. 077	. 128			0.51
Do	J	1932	do	1.25	12	. 350			2.90	2.04	. 83	. 060	. 248			. 62
Do	C	1932	do	1.5	12	. 250	3.00	. 50	3.50	2.50	. 70	. 050	. 400	0.30	0.15	
D0	В	1941	do	1.5	13	. 250			3.28	2.09	. 83	. 12			1.27	. 32
Do	C	1941	do	1.5	13	. 250			3.24	2.08	. 80	. 12			1.71	. 98
Do	NC	1941	Plate	2.5	14	. 5	1		2.80	2.03					2.08	1.10
D0	N	1941	do	2.5	14	. 5			2.75	2.00					3.10	
Do	D	1941	Pipe	1.5	13	. 250			3.21	2.11	.72	. 12			3.32	
High-alloy cast iron.	E	1932	do	1.5	10	. 250			2.98	2.13	1.00			2.61	15.00	6.58
							4									

<sup>a</sup> Ordinary iron horizontally east in green sand molds and rattled to remove sand.

# IV. Results of Exposure Tests

## 1. Effect of Composition

## (a) Low-Alloy Wrought Materials

Prior to examination the specimens were cleaned free of corrosion products by methods that have been previously described [1]. The extent of corrosion was measured by the loss in weight after exposure and by the depths of the deepest pits. These data are recorded in tables 5 and 6, respectively. Unless indicated otherwise each value is the average of measurements made on two specimens. Except as noted in tables 5 and 6, the exposure periods at the different test sites did not differ by more than 5 percent. A typical set of specimens, including ferrous, nonferrous, and miscellaneous materials, is shown in figure 1. The condition of various wrought and cast ferrous materials after exposure for 14 years in three different environments is shown in figure 2.



FIGURE 1. Specimens removed in 1946 from the test site in Hagerstown loam at Loch Raven, Md.



FIGURE 2. Corrosion of steel, cast iron, and low-alloy iron and steel in soils differing in corrosiveness—exposure 14 yr. N, Carbon steel; A, hand-puddled wrought iron; H, Cu-Mo open-hearth iron; D, 2-percent Ni, 1-percent Cu steel; G, cast iron.

## TABLE 5. Loss in weight of wrought materials

(Average of two specimens in ounces per square foot)

											Test	site nun	iber and	soil type	,					1
Identifica-	Year	D.C. doubal	п	Aver- age	51	53	55	56	58	59	60	61	62	63	64	65	66	67	70	72
tion	ied	Material	Form	expo- sure	Acadia clay	Cecil clay loam	Hag- ers- town loam	Lake Charles clay	Muck	Car- lisle muck	Rifle peat	Shar- key clay	Susque- hanna clay	Tidal marsh	Docas clay	Chino silt loam	Mo- have fine gravel- ly loam	Cin- ders	Mer- ced silt loam	Papa- kating silty clay loam
				Years																
				2.0	11.7	3.5	2.8	3.5	3.5		5.7	a 1.3	3.0	3.0	11.4	8.0	8.6	8.6		
				5.4	12.6	2.6	2.3	10.8	9.8	1.8	6.3	5.6	4.0	3.1	22.1	7.4	10.2	31.8		
A	1932	Wrought iron, hand-pud-	Pipe	7.4	15.1	3.3	3.5	17.2	11.9	2.0	5.1	6.3	6.0	3.4	34.4	9.0	11.6	29.7		
		died.		9.3	e 92 0	3.7	3.7	<sup>D</sup> 22. 8	12.6	2.4	a 14.3	6.4	7.8	d 8.5	· 16.0+	13.6	b 5.8	• 15. 2+		
				( 2.0	8 2	3 4	2.9	20.0	19.0	° 4. 0	20.1	a 1 2	8.3	9 f	° 38. 3+ 12. 2	10.4	20.3	1 D		
				5.4	13.6	3.0	2.4	7.6	10.4	1.6	6.8	4.9	4.0	2. 0	23.1	7.2	11.3	24 9		
В	1932	Wrought iron, mechani-	do	7.4	15.3	3.4	3.4	14.7	11.6	1.8	5.4	6.4	6.0	3.5	35.4	8.8	11.1	27.0		
		cally puddled.		9.3		3.7	3.8	19.5	12.7	2.3	d 16.5	5.7	9.4	4.2	• 18.4+	11.4	10.0	D		
				14.3	e 26. 7	4.8	3.7	26.5	17.4	• 4.2	28.8	11.9	7.1	d 6.8	° 36.8+	9.2	• 17. 2+	D		
				2.0	7.4	2.7	2.4	4.0	3.2		6.2	• 0.8	4.1	3.8	12.6	7.4	7.7	21.5		
N	1022	Low earbon steel	do	5.4	12.7	3.0	2.2	13.9	11.2	2.4	11.0	4.0	4.7	4.5	25.3	10.3	15.1	34.6		
19	1952	Low-carbon steel	u0	9.3	11. 5	4.2	3.2	21.0	14.1	3.0 4.7	16.7	0.0 5.9	5.3	7.1	35. 6 D	13.7	14.3	23.5		
				14.3	e 21. 9	4.4	3.1	35.2	25.5	• 3. 9	28.8	10.0	7.9	9.6	D	12.9	18.0 D	2 08.4+		
				2.1	7.5	1.8	1.8	13.8	. 5.1	1.5	4.0	2.2	3.2	2.7	8.7	4.3	9.2	40.5	4 9	
S	1937	Carbon steel	do	4.0		2.9	2.6	16.0	8.8	3.3	8.1	5.0	4.3	9.2	6.0	4.6	12.3	37.0	9.7	
				9.0	g 17.3	3.4	4.1	27.8	17.3	7.5	17.6	4.2	5.3	10.7	4.7	7.0	d 8.0	31.6	13.4	
				2	11.6	1.8	2.0	14.4	5.7	1.5	6.3	2.6	2.8	3.6	7.1	4.6	8.3	12.0	5.0	
A	1937	Open-hearth steel	Plate			3.2	2.6	18.4	9.9	4.2	9.5	5.4	3.7	<sup>b</sup> 6. 2	7.4	5.3	<sup>b</sup> 16. 8	34.3	10.6	
				1 9	g 19.1	3.9	3.8	28.0	16.9	9. 9	22.0	4.3	4.2	ь 8.9	7.5	7.2	4.6	D	17.9	
				2.0	0.2	2.8	2.0	2.3	2.9	1 7	4.9	a 0.9	3.7	2.5	16.1	7.5	8.8	7.8		
н	1932	Open-hearth iron: 0.52 Cu.	Pine	7 4	11.0	2.0	2.4	13.0	10.4	1.7	0.8	4.1	4.1	3.0	23.4	11.2	14.4	25.0		
		0.15 Mo.	1 Ipoilie	9.3	11.0	4.0	3.2	ь 18. 9	14.5	2.0	13.7	4.9 5.6	7.0	4.0	- 04.0 D	14.7	14.5	13. 8 D		
				14.3	e 21. 9	4.2	3.2	23.0	17.3	e 4. 3	25. 5	9.8	5.9	ь 8.0	31.5	14.9	D	D		
				2.1	7.5	2.1	2.0	12.4	5.8	1.3	4.1	3.0	3.0	b 2.5	8.0	5.0	9.1	33.8	5.4	
0	1937	Open-hearth iron; 0.45 Cu,	Plate	4.0		2.9	2.8	13.8	9. 9	2.9	7.4	5.6	3.6	4.7	8.0	5.0	11.6	33. 2	d 11.0	
		0.07 Mo. ,		9.0	g 19.0	3.4	4.7	28.4	17.3	7.8	16.4	4.8	4.6	7.1	6.4	6.4	5.9	25.1	13.6	
N	1027	Open hearth iners 0 54 Cur	4.	2.1	7.2	2.0	1.9	13.0	5.5	1.4	4.8	2.7	3.0	3.0	8.0	4.6	7.4	20.9	4.8	
1N	1957	0 13 Mo		4.0	σ 10 0	3.0	2.7	17.4	9.4	2.9	8.2	5.6	3.7	4.7	6.6	5.0	12.0	27.7	d 10.2	
MM	1939	Open-hearth iron: 0.51 Cu	do	[ 1 Q	B 10. 0	0.0	4.0	00.7 11.9	17.0	1.0	14.0	4.0	4.7	7.2	5.8	6.7	a 6.9	21.9	15.0	
	1000	0.07 Mo.		6.9	h 24, 1	3.6	1.8	45.8	12.7	4 6	3.4 8.4	6.4	5.0	2.1 6.7	4.1 b 20.2	5 3. 1 4 0	0.0	26.0	10.1	
Н	1941	Open-hearth iron; 0.47 Cu,	Pipe	5.0		3.7	2.0	6.3	5.0	5.5	5.3	3.0	3.4	3.5	20. 2	4. 9 5. 7	12.4	27.9 D	6.0	2.0
		0.08 Mo.		(		0.0	1.0													
т	1027	Conner-nickel stool: 0.05	Plate	$\begin{bmatrix} 2.1\\ 1.0 \end{bmatrix}$	7.0	0.8	1.2	14.0	5.2	1.6	5.3	2.3	2.4	2.5	9.0	3.4	7.3	20.3	4.2	
J	1997	Cu. 0.52 Ni.	r mue	9.0	g 18 1	1.3	1.8	19. 2 34 6	10.6	2.6	8.6	4.8	3.3	3.5	6.0	5.0	10.4	35.1	d 7.6	
		C (4) () () # 111		2.1	7.4	0.6	0.8	13.0	6.2	0.0 1.4	18.8 6.4	5.9 2.0	4.0 9.1	1.0	4.7	6.2	4 8.2 8 0	30.4	17.3	
В	1937	Nickel-copper steel; 1.96	do	4.0		1.1	1.2	19.5	11.0	2.4	9.6	3.9	2.9	2. 3	4.9	4.9 5.0	11 4	23. U 44. 3	3.9	
		Ni, 1.01 Cu.		9.0	g 20. 2	2.8	2.6	31.2	17.7	7.2	24.1	3.2	2.9	8.9	7.1	7.6	3.1	D	18.4	

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					( 2.0	4.7	2.2	1.7	3.3	3.5		4.9	a 1.2	2.7	1.6	6.5	5.0	5.1	16.4	
· .					5.4	8.3	2.2	1.4	5.1	7.7	1.7	7.8	2.6	2.3	2.3	23.6	4.2	8.4	23.5	
	D	1932	Nickel-copper steel; 2.47	Pipe	7.4	9.6	2.6	2.2	9.7	9.7	2.9	3.8	3.8	3.7	4.1	37.6	6.1	9.2	27.5	
			Ni, 1.08 Cu.		9.3		2.5	2.0	16.7	6.6	2.6	d 10.3	3.4	5.4	7.6	<sup>b</sup> 24.8	6.0	d 7.5	D	
				4 C	14.3	• 15. 9	3.4	2.4	14.5	13.1	e 3.7	17.4	6.2	3.6	10.4	25.5	6.9	14.6	D	
	Y	1941	Nickel-copper steel; 1.95	do	5.0		3.6	2.0	0.6	3.4	4.6	3. 3	2.6	3. 5	1.2	20.4	3.9	15.8	D	6.9 2.1
	P	1941	Composite steel i	Plate	5.0		4.6	4.1		6.2	5.6	5.7	3.9		ь 5.9	13.0	5.7	11.3		6.2 3.0
					2.1	9.2	1.5	1.3	12.5	4.4	1.1	5.8	2.4	1.9	2.4	4.9	4.4	6.9	17.7	5.4
	C	1937	Cr-Si-Cu-P steel, 1.02 Cr,	do	4.0		2.4	2.3	20.8	9.0	2.5	10.2	4.6	3.0	4.1	5.1	5.4	13.6	13.9	9.8
			0.42 Cu.		9.0	g 19.6	3.3	3.0	28.1	16.7	7.8	24.9	3.6	3.8	b 5.8	5.7	11.0	3.9	26.3	15.8
	NN	1020	9 percent obvornium steel	Dime	1.9		2.0	0.9	13.8	5.9	1.9	4.1	2.0	2.6	1.8	3.9	3.9	9.0	17.8	12.8
	1919	1959	2 percent chronnum steel	ripe	6.9	h 27.1	b2.9		46.0	15.6	4.0	9.8	5.8	b 4.4	5.0	15.8	7.2	18.0	23.5	12.1
			with Mo.		2.1	7.9	1.6	1.6	9.0	3.3	1.2	3.8	2.4	2.4	1.8	4.9	4.2	7.2	18.1	5.4
	KK	1937	do	Plate	4.0		2.2	2.4	14.5	· 8.4	2.5	6.0	4.7	3.0	3.1	5.4	4.5	12.3	16.4	9.7
					9.0	g 17.9	2.9	4.7	28.9	12.9	4.8	13.8	4.1	4.3	4.9	4.6	7.7	d 6.0	15.0	13.5
					( 2.0	6.4	1.4	1.2	2.5	1.8		4.3	a 0.8	2.4	1.3	14.0	7.2	7.7	7.4	
					5.4	14.6	2.0	1.2	8.2	7.4	2.0	11.6	3.3	2.6	3.6	24.1	9.7	13.9	d 27.4	
	P	1932	5.05 percent chromium	Pipe	7.4	10.7	2.4	1.6	18.0	11.7	2.3	2.9	5.0	3.4	4.2	29.6	13.4	13.0	7.5	
			steel.		9.3		2.2	1.9	23.6	13.9	2.8	15.6	5.1	4.6	d 5.8	D	10.9	18.3	27.9	
					14.3	e 19. 7	2.9	2.3	19.1	20.6	e 3.4	27.4	9.1	6.1	10.0	D	11.9	D	17.5	
				1	2.1	6.6	0.9	0.7	9.4	4.1	0.4	4.5	0.6	0.7	1.8	4.5	2.3	5.8	17.1	4.9
	D	1937	5.02 percent chromium	Plate	4.0		1.3	1.1	17.9	- 5, 9	. 9	6.6	1.6	1.2	3.8	4.4	2.2	12.1	d 18. 1	d 10. 0
			steel.		9.0	g 17.6	2.0	1.7	28.8	7.5	3.0	14.8	1.1	1.8	5.6	4.4	3.6	d 3.5	12.2	19.1
	_			1.1.1.1	2.1	7.1	0.9	0.6	9.6	3.9	0.4	4.1	0.9	0.7	1.7	4.2	2.5	7.9	12.2	5. 2
	Е	1937	4.67 percent chromium	do	4.0		1.6	1.2	16.3	7.2	. 8	6.2	1.8	1.1	5.2	5. <b>2</b>	2.1	12.0	d 11. 7	9.9
			steel with Mo.		Į 9.0	g 16. 2	2.0	1.7	28.2	6.4	2.5	13.2	1.4	2.0	5.3	4.2	3.3	d 5. 2	11.2	19.5
					2.1	<sup>b</sup> 8.3	<sup>b</sup> 0. 8	0.7	11.4	3.5	0.2	4.6	0.8	0.7	<sup>b</sup> 1.7	5.9	2.5	8.4	12.6	5.0
	Н	1937	5.76 percent chromium	do	4.0		1.4	1.1	14.3	<sup>b</sup> 6.8	. 7	6.2	1.6	1.3	5.4	5.0	2.2	14.5	d 12. 9	10.2
			steel with Mo.		l 9.0	g 16.7	1.9	2.0		6.1	2.2	14.2	<sup>b</sup> 1.0	1.9	6.4	4.0	3.7	d 5.0	8.9	17.6

<sup>a</sup> Exposed for 1.0 yr only.

<sup>b</sup> Data for 1 specimen. The other specimen was missing.

• Data for 1 specimen. The other specimen was destroyed by corrosion.

d Data for the individual specimens differed from the average by more than 50%. • Data for 4 specimens. <sup>f</sup> D, both specimens destroyed by corrosion.

g Data for 8 specimens.

<sup>h</sup> Data for 10 specimens.

 $^{\rm i}{\rm Specimens}$  composed of 3 layers as follows: outer layers—plain carbon steel, 0.036 in. thick; intermediate layer—2% nickel-1% copper steel, 0.100 in. thick.

# TABLE 6. Maximum penetration of wrought materials

(Average of	two specimens	s in mils)
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											Test	site nu i	nber and	soil type						
Identifica-	Year			Aver-	51	53	55	58	58	59	60	61	62	63	64	65	66	67	70	72
tion	bur- ied	Material	Form	expo- sure	Acadia clay	Cecil clay loam	Hag- ers- town loam	Lake Charles clay	Muck	Car- lisle muck	Rifle peat	Shar- key clay	Susque- hanna clay	Tidal marsh	Docas clay	Chino silt loam	Mo- have fine gravel- ly loam	Cin- ders	Mer- ced silt loam	Papa- kating silty clay loam
				Years																
				2.0	50	34	40	22	20		24	a 17	49	28	102	54	- 88	100		
				5.4	144	64	79	66	68	25	38	41	54	22	129	91	85	d 145+		
A	1932	Wrought ircn, hand pud-	Pipe	( 7.4	122 +	77	70	90	84	18	30	44	69	64	144+	110+	110	145 +		
		dled.		9.3		50	60	b 96	118	32	° 55	61	e 72	100	120 +	102	<sup>b</sup> 88	145+		
				(14.3	e 135+	72	76	145 +	96	e 37	78	84	74	74	145+	98	142+	145+		
	×1			2.0	60	30	42	24	18		24	a 10	70	16	118	66	82	98		
	1000			5.4	° 129+	71	84	65	64	18	37	37	56	37	110	87	140	145+		
B	1932	Wrought iron, mechani-	do	7.4	145+	76	60	106+	110	15	34	50	78	39	145+	106	140+	145+		
		cally puddled.		9.3	 0 191 I	-73	84	106	70	28	<sup>c.</sup> 04 79	° 80	6 101	° 55 80	145+	110	130+	140+		
	18 M.			( 14. 3	° 151+	00 27	41	140+	10	° 52	27	02 9 10	62	00 15	140	90 40	140 T	154		· · · · · ·
				2.0	82	57	41	20	102		94	a 10	66	10	154	40	154	110		
N	1029	Low conhon stool	do	7.4	104+	50	57	195 1	105	20	17	63	71	70	154	83	154	119+		
IN	1952	Low-carbon steel		0.3	1557	50	50	154	110	e 40	c 97	c 96	c 87	54	154+	119	154+	154+		
				14.3	e 146+	84	65	135+	154+	e 34	82	- 50	101	61	154+	86	154+	154+		
				21	52	42	33	77	29	12	15	40	40	24	80	50	145+	145+	50	
S	1937	Carbon steel	do	4.0	02	98	50	104	46	20	38	45	56	38	67	59	145 +	145 +	118+	
		Curbon Stool		9.0	f 128+	74	92	145 +	98	101	c 58	48	68	80	80	65	78	145 +	122	
				2.1	54	40	42	80	36	6	30	34	34	18	44	47	86	46	56	
A	1937	Open-hearth steel	Plate	4.0		76	54	100	61	22	40	50	47	<sup>b</sup> 26	78	51	<sup>b</sup> 188+	° 132+	77	
1				9.0	f 138+	57	90	126 +	89	98	56	90	59	ь 36	87	75	66	D	136 +	
				( 2.0	78	54	60	20	21		27	a 14	72	° 30	153	94	111	65		
				5.4	145 +	65	75	65	60	5	° 21	c 59	86	c 49	137	97	145 +	118 +		
H	1932	Open-hearth iron; 0.52 Cu,	Pipe	7.4	97	92	68	112+	145 +	° 10	16	65	78	103	145 +	117	145 +	80		
		0.15 Mo.		9.3		° 109+	93	<sup>b</sup> 145+	96	22	c 61	82	c 86	° 67	145 +	106	145 +	145 +		
				14.3	e 105+	105 +	89	108+	118+	e 38	106+	104+	84	<sup>b</sup> 74	145 +	104 +	145 +	145 +		
1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1				2.1	54	38	31	100	30	6	21	- 33	36	<sup>b</sup> 20	70	44	105	70	48	
0	1937	Open-hearth iron; 0.45 Cu,	Plate	4.0		74	44	116	48	20	28	66	38	c 48	75	65	84	55	c 97	
		0.07 Mo.		9.0	f 242+	72	85	174	97	66	38	64	66	74	102	102	63	80	89	
N	1937	Open-hearth iron: 0.54 Cu	do	2.1	66	38	29	77	32	6	20	-32	36	33	72	32	132	50	51	
	1001	C 13 Mo		4.0		° 72	51	100	44	16	° 26	54	49	47	76	e 57	98	74	c 122	
MM	1939	Open-hearth iron: 0.51 Cu.	do	9.0	f 171+	53	92	162 +	102	68	35	70	56	50	70	52	62	58	92	
		0.07 Mo.		1.9		<sup>b</sup> 42	44	60	48	b 13	16	41	40	22	47	b 43	71	128	92	
				6.9	g 124	86	49	160 +	116	26	40	62	42	76	<sup>b</sup> 84	107	144	86	80	
· H	1941	Open-hearth iron; 0.47 Cu, 0.08 Mo.	Pipe	5.0	/	67	60	40	81	40	38	30	38	114+	145+	68	145+	145+	63	24
т	1027	Copper-nickel stock 6.05	Plato	2.1	63	38	34	82	26	6	25	53	38	14	60	49	108	58	48	
J	1957	Cu 0.52 Ni	rate	4.0		57	50	- 96	64	15	40	63	60	28	84	60	85	90	82	
		Ou, 0.32 INI.		9.0	f 148+	38	77	140 +	72	33	57	84	69	40	76	58	67	102	136	
в	1937	Nickel-copper steel: 1.96	do	2.1	54	26	34	106	36	6	20	62	51	22	66	45	38	64	50	
D	1007	Ni 101 Cu		4.0		56	52	139	52	12	28	56	69	25	88	84	73	84	78	
		11, 1.01 Cu.		9.0	f 161+	45	88	135	97	32	73	80	70	37	80	75	51	D	142	

1					( 2.0	68	23	32	20	23		° 28	a İŻ	42	° 24	<u>98</u>	41	58	101		
	D	1029	Niekol coppor steel: 9.47	Dina	5.4	55	26	34	42	71	3	26	30	46	17	108	48	75	145 +		
	D	1952	Ni 108 Cu	r ipe	7.4	70	44	51	145 +	110+	14	14	51	72	41	145 +	68	141 +	145 +		
			N1, 1.08 Cu.		9.3		37	62	145 +	52	14	° 38	41	e 58	° 70	<sup>b</sup> 145+	74	° 96+	145 +		
					14.3	e 118+	42	71	145 +	106 +	e 22	92	42	68	90	145 +	73	145 +	145 +		
	Y	1941	Nickel-copper steel; 1.95 Ni, 1.04 Cu.	do	5.0		56	64	16	80	29	15	50	57	26	210 +	96	92	145 +	98	10
	P	1941	Composite steel h	Plate	5.0		. 46	42		57	43	28	53		b 33	80	42	80		71	37
	C	1097	Cr. C. Cr. D. steel: 1.02 Cr.	de	2.1	58	40	40	52	71	14	23	30	36	31	42	50	56	58	66	
	U	1937	0.42 Cu	do	4.0		64	51	77	52	42	67	41	44	41	70	44	80	47	94	
			0.42 Cu.		9.0	f 92	45	77	84	98	56	112+	76	56	b 50	60	86	58	58	148 +	
	NN	1939	2% chromium steel with	Pipe	1.9		65	59	78	46	22	18	42	40	38	40	51	145 +	124 +	130 +	
			Mo.	· · · ·	6.9	g 107	<sup>b</sup> 100		138+	100	23	48	76	<sup>b</sup> 61	48	106	98	145 +	145 +	96	
					2.1	35	40	26	38	29	22	19	31	26	20	56	56	78	53	102	
	KK	1937	do	Plate	4.0	·	52	52	60	42	27	26	35	56	24	70	55	130 +	68	94	
					9.0	f 110+	69	84	96+	67	34	57	76	58	41	70	76	80	78 +	110	
				Т	2.0	69	46	42	58	36		33	a 10	52	43	124	68	86	64	·	
-					5.4	105	56	65	154 +	70	32	67	° 37	83	87	154 +	107	133	112 +		
	P	1932	5.05% chromium steel	Pipe	7.4	106	57	88	154 +	70	° 20	62	38	125 +	89	154 +	138 +	154 +	65		
				2 X	9.3		70	84	136	111	20	110	74	70	136 +	154 +	131 +	154 +	125 +		
					14.3	e 134+	66	85	154 +	154 +	e 28	154 +	94	126	127 +	154 +	109 +	154 +	137 +		
- 1	D	1937	5.02% chromium steel	Plate	2.1	62	43	34	66	48	18	32	30	32	62	48	37	59	46	87	
		1001		1 100000000	4.0		57	48	95	44	39	51	36	52	70	60	46	99	57	121	
					9.0	f 114	66	70	104	64	66	64	42	74	108	62	63	92	48	194	
	E	1937	4.67% chromium steel with	do	2.1	50	36	34	62	39	20	26	26	26	46	46	39	81	56	79	
			Mo.		4.0		50	47	80	46	26	36	36	46	73	66	48	88	52	106	
					9.0	f 86	66	51	108+	56	55	72	30	54	88	46	52	57	52 -	154	
					2.1	<sup>b</sup> 64	<sup>b</sup> 32	30	<sup>b</sup> 60	48	18	26	24	32	<sup>b</sup> 42	46	50	72	41	88	
	Н	1937	5.76% chromium steel with	do	4.0		57	39	90	<sup>b</sup> 44	33	32	32	58	72	72	56	117	44	94	
			Mo.		(9.0	f 109	71	58		52	64	76	b 38	56	78	70	63	76	54	132 +	
				S 1 1																	

<sup>a</sup> Exposed for 1.0 yr only.

<sup>b</sup> Data for 1 specimen. The other specimen was missing.

• Data for the individual specimens differed from the average by more than 50%.

d +, one or more specimens contained holes because of corrosion.

e Data for 4 specimens.

f Data for 8 specimens.

g Data for 10 specimens.

h Specimens composed of 3 layers as follows: outer layers-plain carbon steel, 0.036 in. thick; intermediate layer-2% nickel, 1% copper, 0.100 in. thick.

In order to compare the over-all behavior of the materials and also to observe the effect of various alloying elements on corrosion, the data reported in tables 5 and 6 for each material at all of the test sites were averaged and shown graphically as weight loss and pit depth-time curves for the pipe specimens exposed 14 yr (fig. 3) and for the plate specimens exposed 9 yr (fig. 4). For comparison with the low-alloy materials in the form of plates, data for two reference materials, exposed simultaneously with the low-alloy wrought materials, are included in figure 4, namely, open-hearth iron A, and carbon steel S. Specimens of the latter material were in the form of pipe.

Although the curves shown in figure 3 indicate differences in the corrosion of certain of the materials, it is impossible to state without statistical analysis of the data whether certain of these differences at least should not be ascribed to chance causes rather to inherent differences in corrodibility of the materials. As the first step in analyzing the data statistically, the progress of weight loss and pitting of each material with time was expressed by means of equations of the form

$$y = ax^n. \tag{1}$$

Taking logarithms of both sides, this equation takes the form of the linear equation

$$\log y = \log a + n \log x. \tag{2}$$

Expressing the logarithm of the pit depth as log P, the logarithm of the time as log T, and the y-intercept as log k, the following equation is obtained:

$$\log P = \log k + n \log T. \tag{3}$$



FIGURE 3. Average loss in weight and maximum penetration-time curves of wrought pipe specimens in 13 soils.

A, Wrought iron, hand-puddled; B, wrought iron, mechanically puddled; H, open hearth iron 0.52-percent Cu, 0.15-percent Mo; N, low carbon steel; D, steel, 2.47-percent Ni, 1.01-percent Cu; P, steel, 5.05-percent Cr.

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FIGURE 4. Average loss in weight and maximum penetration-time curves of wrought plate specimens in 14 soils.

O, Open-hearth iron, 0.45-percent Cu, 0.07-percent Mo; N, open-hearth iron, 0.54-percent Cu, 0.13-percent Mo; S, plain carbon steel; A, open-hearth steel; J, steel, 0.95-percent Cu, 0.52-percent Ni; B, steel, 1.96-percent Ni, 1.01-percent Cu; C, Cr-Si-Cu-P steel—1.02-percent Cr, 0.42-percent Cu; K, steel, 5.02-percent Cr; E, steel, 4.67-percent Cr, 0.51-percent Mo; H, steel, 5.76-percent Cr, 0.43-percent Mo.

By similar substitutions, a linear equation expressing the relation between weight loss W, and time was obtained:

$$\log W = \log k_1 + u \log T. \tag{4}$$

These equations, which were derived originally by Logan, Ewing, and Denison [3] and by Martin [4] to express, respectively, the progress with time of the pitting and weight loss of wrought ferrous metals in soils, were fitted to the data shown in figure 3 by the method of least squares. The mean values of weight loss and pit depth given in table 7 were computed by eq 3 and 4, respectively, for the period of 14 yr. Values for the standard error of these means, shown also in the table, were calculated by the method of Ezekiel [5]. In order to determine whether the various wrought materials corroded at significantly different rates from plain steel, the mean values of weight loss and pitting for each material at 14 yr were compared with the corresponding values for plain steel, taking into account the respective standard errors. The standard errors of the difference between the means for plain steel and for each material with which it was compared were calculated by the following equation to be found in standard texts on statistical methods [6]:

$$\sigma_D = \sqrt{\sigma_{\overline{x}_1}^2 + \sigma_{\overline{x}_2}^2} \tag{5}$$

$$=\sqrt{\frac{\sigma_1^2}{N_1} + \frac{\sigma_2^2}{N_2}},$$
 (6)

## Soil Corrosion of Ferrous Materials

			Loss in	ı weight (o	z/ft <sup>2</sup> )	9 		Maximu	m pit deptl	n (mils)	
Identi- fica- tion	Material	$\begin{array}{c} \operatorname{Mean} \\ \overline{X} \\ (W_T =_{14} y_r) \end{array}$	$\begin{array}{c} \text{Stand}\\ \text{ard}\\ \text{error}\\ \overline{\sigma} \end{array}$	Differ- ence of the mean from material N	Standard error of the dif- ference a $\sigma_D$	$2\sigma_D$	$\begin{array}{c} \text{Mean} \\ \overline{X} \\ (P_T =_{14 \text{ yr}}) \end{array}$	$\begin{array}{c} \text{Stand}\\ \text{ard}\\ \text{error}\\ \overline{\sigma} \end{array}$	Differ- ence of the mean from material N	Standard error of the dif- ference a $\sigma_D$	$2\sigma_D$
		· · · · · · ·		D	-	·			D		
N	Disir starl	10.7	1.4				110	1.0			
IN .	Plain steel	10.7	1.4				113	4.0			
A	Hand-puddled wrought iron	14.8	1.2	-1.9	1.8	3.6	103	4.1	-10	5.7	11.4
В	Mechanically puddled wrought iron	14.3	1.2	-2.4	1.8	3.6	107	5.5	-6	6.9	13.8
H	Open-hearth iron	13.6	1.0	-3.1	1.7	3.4	109	1.5	-4	4.3	8.6
Р	5% Cr steel	13.5	1.2	-3.2	1.8	3.6	125	5.0	+12	6.4	12.8
D	2% Ni-1% Cu steel	10.7	1.1	-6.0	1.8	3.6	92.2	8.3	-20.8	9.1	18.2

TABLE 7. Comparison of average corrosion at all test sites of wrought materials exposed 14 years

$$\sigma_{D} = \sqrt{\frac{\sigma_{1}^{2} + \sigma_{2}^{2}}{N_{1} + N_{2}}}$$
 (see text).

where

 $\sigma_1$  = standard deviation of first sample,

 $\sigma_2$ =standard deviation of second sample,

 $N_1$  = number of observations of first sample,

 $N_2$ =number of observations of second sample.

By comparing the standard error of the difference  $\sigma_D$  with the difference between the two means  $\overline{X}_1$  and  $\overline{X}_2$ , the probability that the observed difference is significant can be found by reference to tables of probability corresponding to the number of times the observed difference exceeds the standard error. For the present study, a difference between means of at least twice the standard error was selected. A difference of this magnitude . corresponds to a probability of 21 to 1 that the observed difference is significant.

On the basis of this criterion, only material D, containing 2 percent of nickel and 1 percent of copper, can be considered definitely to be more resistant to weight loss and to pitting than the plain steel, N. However, there is some probability that material P, containing 5 percent of chromium, is slightly more resistant than plain steel with respect to weight loss, but less resistant to pitting. The small differences shown by the other materials as compared with plain steel are probably to be ascribed to chance causes.

The weight loss data shown in figure 4 for the plate specimens exposed for a maximum of 9 yr indicate that all of the materials except those containing 2 percent or more of chromium, namely, materials K, D, E, and H, corroded essentially alike. The beneficial effect of chromium in higher concentration is indicated by the lower average weight losses for the 5-percent chromium steel D.

The effect of molybdenum in reducing the corrosion of chromium steels is indicated by the curves for materials E and H, both of which contain approximately 0.5 percent of molybdenum in addition to 4.67 and 5.76 percent of chromium, respectively.

The benefit conferred by chromium in reducing weight loss apparently does not extend to maximum penetration, because the 5-percent chromium steel D, which showed definitely lower weight losses than the materials with smaller amounts of chromium, showed no improvement with respect to pitting. On the other hand, the effect of molybdenum in reducing the depth of the deepest pits as well as weight loss is indicated by the fact that the materials containing approximately 0.5 percent of molybdenum, E, H, and K, also show relatively low values for maximum penetration. As a matter of fact, the average depth of the deepest pits on material K, which contained only 2 percent of chromium was no greater than the corresponding values for materials E and H, both of which contained from two to three times as much chromium.

Although the specific effect of chromium on the maximum penetration of the irons and steels under consideration is somewhat difficult to evaluate because most of the materials also contain molybdenum, there was no reason to predict that chromium in the range of 4 to 6 percent might accelerate pitting, such as is indicated by the data for material P shown in figure 3 and in table 7. In view of the possibility that chromium carbides precipitated along grain boundaries might have served as local centers of attack and so might have



FIGURE 5. Longitudinal sections of 4-6 percent chromium steel specimens. × 500. D, 5.02-percent chromium; E, 4.67-percent chromium with 0.51-percent molybdenum; P, 5.05-percent chromium.

accelerated pitting, microphotographs of material P were prepared for comparison with chromium steels that were considered to show normal behavior.

Microscopical examination of steel P (fig. 5) reveals that the carbides are in lamellar form, partly spheroidized. By contrast, the carbides of steel D are completely spheroidized and are completely dispersed throughout the matrix. Because neither of these steels contains molybdenum, the carbides are probably chromium carbides. As the microstructure of material P is considered to be that of a normal 4- to 6-percent chromium steel, the apparent accelerated pitting of this material cannot at present be accounted for.

The carbides of steel E (fig. 5) are localized around the grain boundaries. Because this steel contains an appreciable amount of molybdenum most of the carbon was probably combined with molbydenum, and the chromium remained in solution in the ferrite. This may account for the slightly improved resistance of steel E as compared with steel D.

The results of the tests of steels B and D, containing 2 percent of nickel and 1 percent of copper, are conflicting. Material D is definitely superior to plain steel with respect to weight loss and pitting (fig. 3), whereas material B, having the same nominal composition, shows no advantage from its content of nickel and copper. A probable explanation for this difference in behavior is the acceleration of corrosion from the presence of mill scale on material B, which unlike D, had not been pickled.

Comparison of the average corrosion resistance of the different materials in all of the soils reveals nothing concerning the behavior of these materials in any one soil or environment. Consequently, superior corrosion resistance of a material in any one environment might be obscured by inferior resistance in another. In order to compare the behavior of the materials under specific environmental conditions, the corrosion data for the soils classified according to environment (table 2) were calculated on a relative basis for each period of exposure, the standards of reference, carbon steel (N) and open-hearth steel (A), being taken as 100 percent. The averages for all periods and for all soils in the same environmental group are given in tables 8 and 9.

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 TABLE 8.
 Effect of composition on the corrosion of low-alloy iron and steel pipe specimens in different soil environments—maximum exposure 14 years

	Comp	osition of	steel (pe	ercent)	Environment									
Identification					Organic		Inor	rganic						
Identification	$\mathbf{Cr}$	Ni	Cu	Mo	Reducing	Red	lucing	Oxid	lizing	Cinders				
					acid	Acid	Alkaline	Acid	Alkaline					
AVERAGE I	LOSS IN	WEIG	HT FO	R 5 PEI	RIODS OF	EXPOS	URE-REI	LATIVE E	BASIS (PE)	RCENT)				
N D		2.47	1.08		100 70	$\frac{100}{76}$	$\begin{array}{c} 100 \\ 63 \end{array}$	$\begin{array}{c} 100 \\ 67 \end{array}$	$     100 \\     53   $	100 100				
H P	5.05		0. 52	0.15	78 75	96 94	96 78 94 81		$\begin{array}{c ccc} 94 & 104 \\ 62 & 94 \end{array}$					
AVERAGE	MAXI	MUM ]	PENET	RATIO	N FOR 5 P (PERCEN	PERIODS T)	S OF EXP	OSURE—I	RELATIVE	BASIS				
N D H		2.47	$1.08 \\ 0.52$	 0. 15	100 85 103	100 66 97	100 90 94	100 75 132	100 78 127	100 99 80				
Р	5.05					82	132	118	124	74				

 TABLE 9.
 Effect of composition on the corrosion of low-alloy iron and steel plate specimens in different soil environments—maximum exposure 9 years

	Comp	osition of	f steel (pe	ercent)			Envir			
Identification					Organic		Inor	rganic		
ruentineation	Cr	Ni	Cu ,	Мо	Reducing	Red	lucing	Oxio	lizing	Cinders
					acid	Acid	Alkaline	Acid	Alkaline	
AVERAGE I	LOSS IN	WEIG	HT FO	R 3 PE	RIODS OF	EXPOS	URE-REI	LATIVE 1	BASIS (PE)	RCENT)
A	0.049	0.034	0.052		100	100	100	100	100	100
0	. 02	.15	. 45	0.07	83	98	94	104	93	133
· N	.02	.14	. 54	. 13	82	94	96	102	92	102
J		. 52	. 95		79	87	96	73	87	124
B		1.96	1.01		90	82	93	59	91	145
C	1.02	. 22	. 428		85	90	88	80	102	84
K	2.01	.07	. 004	. 57	62	88	80	89	91	71
D	5.02	. 09	. 008		52	53	85	41	60	68
Ε	4.67	. 09	. 004	. 51	51	54	85	43	61	51
Н	5.76	. 17	. 004	. 43	52	54	76	42	72	50
AVERAGE	MAXII	MUM P	ENETF	RATION	FOR 3 P (PERCEN	ERIODS NT)	OF EXPC	SURE-R	ELATIVE	BASIS
A	0.049	0.034	0,052		100	100	100	100	100	100
0	. 02	. 15	. 45	0.07	102	123	110	97	97	62
Ν	. 02	. 14	. 54	.13	97	106	104	95	84	56
J		. 52	. 95		81	113	102	93	86	76
В		1.96	1.01		85	113	111	100	82	91
C	1.02	0.22	0.428		130	81	88	91	80	50
К	2.01	.07	. 004	. 57	82	78	90	93	96	60
D	5.02	. 09	. 008		149	77	105	97	76	46
Е	4.67	. 09	. 004	. 51	130	62	94	83	72	49
Н	5.76	.17	.004	. 43	128	72	94	88	87	42

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Additions of 0.5 percent of copper and 0.15 percent of molybdenum are seen from the data reported in table 8 (material H) and in table 9, materials O and N, to have had no significant influence on the rate of corrosion. As a matter of interest, it may be noted that the addition of copper tends to increase slightly the loss in weight of steels in sea water, but has little or no effect on pitting [7]. The inconsistent behavior of materials B and D, each containing 2 percent of nickel and 1 percent of copper, has been previously commented upon.

As was previously noted, the effects of additions of chromium are difficult to evaluate because most of these steels also contain molybdenum. All of the chromium steels lost less weight than the reference material A (table 9). The maximum penetration is also less for all environments except the organic-reducing. This benefit of chromium is especially marked in cinders. In general, the maximum resistance to corrosion is shown by steel



FIGURE 6. Effect of chromium content on loss in weight and pitting of steels in three poorly aerated soils, exposure 9 yr.

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E, containing slightly less than 5 percent of chromium and 0.5 percent of molybdenum.

The curves shown in figure 6 indicate that the advantage gained in reduction of pitting by the addition of small amounts of chromium may be lost if the amount added exceeds a certain optimum value. In fact, the pit depth curves for Acadia clay and Lake Charles clay in figure 6<sup>2</sup> indicate that steels containing 18 percent of chromium show a slight acceleration of pitting over plain steel. The tendency for chromium to accelerate the pitting of steels in natural waters has previously been noted; LaQue [7] cited the work of Speller on the corrosion of steels in river water. and on the basis of his own observations in sea water, concluded that addition of chromium to steels for the purpose of reducing weight losses in natural waters should probably not exceed 3 percent, because larger amounts may accelerate pitting.

#### (b) High-Alloy Steels

Corrosion data for the high-chromium and chromium-nickel steels, some exposed for 7 yr and some for 14 yr, are given in table 10; and the effects of composition on the corrosion of three steels in three soils are illustrated in figure 7. It is unfortunate that complete information concerning the rolling and heat treatment of the various steels is not available, because these factors undoubtedly played a part in the corrosion of the materials. It is shown quite conclusively, however, that plain chromium steels are subject to severe pitting, and that steels containing at least 18 percent of chromium and 8 percent of nickel are considerably more resistant than the straight chromium steels in these environments. Steels containing larger amounts of chromium and nickel, with and without molybdenum, were wholly resistant to corrosion, but two of these steels were exposed for 7 yr only, as compared with 14 vr for the other materials.

Because high-chromium-uickel steels are susceptible to pitting where oxygen is excluded locally as, for example, by adherent deposits in sea water, the corrosion resistance shown by some of the steels in soils deficient in oxygen and high in chlorides is noteworthy. Docas clay (soil 64)

 $<sup>^2</sup>$  The composition of the steel containing 18% of chromium was not recorded in table 3 because the complete analysis of this steel is not available.



FIGURE 7. Specimens of high-alloy steels after exposure for 14 yr. U, 12-percent-chromium steel; V, 18-percent-chromium steel; W, 18-percent-chromium, 9-percent-nickel steel.

M, Shallow metal attack, roughening of surfaces, but no definite pitting.

P, Definite pitting, but no pits greater than 6 mils.

+, One or more specimens contained holes because of corrosion.

Specimen	U	ſb	V	c	2	ĸ	ł	ζ	I	R	W	ď	Y	d	C	М	D	т
AISI Type No	4	10	43	30	43	30	30	04	30	04	3	02	30	)9	31	16		
Çomposition <sup>.</sup> C, % Mn, % Cu, % Cr, % Ni, % Mo, %	e ( 0. 111. 0.	5) 065 38  95 48	(( 0. 17. 0.	5) 070 36 02 08 09	(; 0 17 0	2) . 12 . 42 . 72 . 29	(: 0 17 8	2) . 08 . 44 . 20 . 95	() 0 17 8	2) . 05 . 46 . 52 . 85	( 0.	5) 093 36 02 69 18	(4 0. 1. 22. 12.	5) 144 80 02 68 94	() 0 1 17 10 2	2) . 07 . 24 . 78 . 96 . 63	(3 0, 1, 19 .22 3	2) . 07 . 99 . 07 . 27 . 12 . 52
Soil No. <sup>t</sup>	Loss in weight	Max- imum pene- tration	Loss in weight	Max- imum pene- tration	Loss in weight	Max- imum pene- tration	Loss in weight	Max- imum pene- tration	Loss in weight	Max- imum pene- tration	Loss in weight	Max- imum pene- tration	Loss in weight	Max- imum pene- tration	Loss in weight	Max- imum pene- tration	Loss in weight	Max- imum pene- tration
51	0 <i>z</i> / <i>ft</i> <sup>2</sup> 0.0002	Mils	oz/ft <sup>2</sup>	Mils	oz/ft <sup>2</sup>	Mils	0 2/ft <sup>2</sup> 0. 03 . 0009 . 20 . 002	Mils  29+ 0 $32+0$	oz/ft 2	Mils	0 <i>z</i> / <i>ft</i> <sup>2</sup>  0. 0005	Mils	oz/ft <sup>2</sup>  0. 0003 	Mils  0 	oz/ft <sup>2</sup> 0.0004 .03 .002 .002 .0003	Mils M 0 0 8 0	<i>oz/ft</i> <sup>2</sup> 0.001 .03 .0007 0 0	<i>Mils</i> 0 0 0 0 0
59	. 0002 5. 4	M 37+	. 001 . 17 	M 16+ 	.03	 6 13	.0007 .0006 .0004 .0004 .06	0 0 P M P	. 003	 6 13	. 002 . 003	P M 	. 001 . 001	P M 	.0009 .002 0 .0007 .002	0 0 0 0	. 0006 . 0003 . 004 . 0007 . 0004	0 0 0 0
64 65 66 67 70	5.7 0.46 3.2	63+62+62+	1.4 0.74 2.3	63+ 61+ 62+ 	. 34	48	. 003 . 01 . 0009 . 0008	P 16+ P 12	. 04	7	.002 0 0	0 M M	.0007 .00004 0	0 0 M	. 0004 . 002 0 . 002 . 002 . 0005	M 0 M 0	.0004 .001 0 .002 0	0 0 0 0 0

\* Specimens "CM" and "DT" were exposed to the soils for 7 yrs. The remaining specimens were exposed 14 yrs.

<sup>b</sup> Annealed at 1,400° F, pickled, lightly cold-rolled.

• Annealed at 1,600 F, pickled.

<sup>d</sup> Heated to 1,850° F, air quenched, pickled.

• The numbers in parentheses indicate the number of specimens removed from each test site.

! See table 1 for names and location of soils.

TABLE 11. Results of exposure of high-alloy steels to a poorly aerated soil containing sodium chloride (Docas clay, soil 64)

M, Shallow metal attack, roughening at surface, but no definite pitting.

+, One or more specimens contained holes because of corrosion. The thickness of the specimen has been used as the maximum pit in this case. D, Destroyed by corrosion.

Specimen	U	v	x	К	R	W	Y	СМ	DT	Low car- bon steel
AISI Type No	410	430	430	304	304	302	309	316		
Composition: C, % Mn, %	0.065 .38	0.070 .36	0.12 .42	$\begin{array}{c} 0.08 \\ .44 \end{array}$	0.05.46	0.093	0.144 1.80	0.07 1.24	$0.07 \\ 1.99$	
Cu, % Cr, % Ni, % Mo, %	11.95 0.48	.02 17.08 0.09	17. 72 0. 29	17. 20 8. 95	17.52 8.85	. 02 18. 69 9. 18	0.02 22.68 12.94	$     17.78 \\     10.96 \\     2.63 $	$ \begin{array}{c} 1.07 \\ 19.27 \\ 22.12 \\ 3.52 \end{array} $	
Period of exposure		l	1		Loss in we	ight (oz/ft²)		1	1	
Years										
2 5	0.67 1.0	0. 53 . 28	0.50 .30	0.01 .002	0.01	0.002 .002	0.003 .002	0.006	0.002	12. 6 25. 3
9	1.7 3.2	. 44 1. 2	(a) (a)	. 003	.02	.002	. 005	. 004	. 004	35.6 D
14	5. 1	1.4	0.34	. 003	. 04	.002	.0007			
-				M	aximum per	netration (m	ils)			
2 5 7 9	63+63+63+63+63+	63+54+63+63+63+	(a) 10 21 96 48		8 0 0 36 7		5 < 6 < 6 < 6 < 6 0	M M	0	130 154+ 154+ 154+ 154+
14	00 T	0.01	40	~0		0	U			104-

» Data not used because of corrosion caused by asphalt on the ends of the specimens.

is a very corrosive soil because of its low permeability to air and its high content of sodium chloride. The results of the field tests in this soil, given in table 11 for all periods, indicate the beneficial effect of nickel in promoting corrosion resistance of these steels. Steels containing chromium but less than 0.5 percent of nickel pitted deeply, perforations occurring within 2 yrs; but steels containing more than 9 percent of nickel developed only shallow pits, which showed no tendency to increase with time.

#### (c) Cast Materials

The weight losses and maximum depths of pits of cast iron after exposure are given in tables 12 and 13, respectively. Following the procedure employed in studying the corrosion of the wrought specimens, the data for weight loss and pitting for each of the cast irons were averaged for all of the soils for each period of exposure. From the results for the pipe specimens buried in 1932, shown as weight loss and pit depth-time curves in figure 8, it is evident that weight loss and maximum penetration for all materials were approximately the same for each period except for alloy E, which corroded considerably less than the other materials. The corrosion data shown graphically in figure 9 for the series of nickel and nickel-copper irons exposed in 1941 does not indicate any significant improvement from the additions of these alloying elements.

#### (2) Effect of Environment

Photographs of corroded specimens of unalloyed steel pipe N and of plain cast iron pipe G after exposure for 14 yr in the various soils are shown in figures 10 and 11, respectively. Photographs of open-hearth iron plates A exposed for 9 yr at 15 test sites are shown in figure 12.

Typical corrosion-time curves for the wrought materials that behaved essentially like plain steel are shown in figure 13. Each point on these curves is the average weight loss or maximum penetration of two specimens of each of the following materials: low carbon steel N, hand-puddled wrought iron A, mechanically puddled wrought iron B, and

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FIGURE 8. Average loss in weight and maximum penetration-time curves of cast iron pipe specimens exposed in 14 soils. G, Rattled cast iron; F, sand-coated cast iron; I, cast iron, 0.51-precent Cu, 0.28-percent Mn; J, cast iron-0.62-percent Cu, 0.83-percent Mn: C, cast iron-0.30-percent Cr, 0.15-percent Ni; E, cast iron-15.0-percent Ni, 6.6-percent Cu. 2.6-percent Cr.

copper-molybdenum open-hearth iron H. Values are given for five periods of exposure except for those soils in which the number of perforations rendered the corrosion data meaningless.

It is evident from the corrosion-time curves shown in figure 13 that the rate of corrosion with respect to both weight loss and maximum penetration varies from zero after a short period of exposure to a rate that is proportional to time.



FIGURE 9. Average loss in weight and maximum penetration of specimens of cast irons exposed for 5 yr in 14 soils.

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In general, it may be said that curves showing a marked diminution of the rate of corrosion with time (soils 53 and 55) are typical of corrosion in well aerated soils, whereas curves indicating a linear rate (soil 56) are typical of poorly aerated soils.

Within the limits of the accuracy of the data, the curves obtained by plotting the values for maximum penetration and weight loss against the length of exposure conformed to the following equations, which were employed in comparing the corrosion of the low-alloy irons and steels:

$$P = kT^n$$

$$W = k' T^u$$

where P = maximum pit depth at time T

W=weight loss at time T.

As will be recalled, these equations are linear on logarithmic coordinates, the constant k (k') being the *y*-intercept and n (u) the slope. The values of

## TABLE 12. Loss in weight of specimens of cast iron

(Average of **2** specimens in ounces per square foot)

					ри .						Test	site nur	nber and	soil type	)					
Idontifiao	Year			Aver-	51	53	55	56	58	59	60	61	62	63	64	65	66	67	70	72
tion	bur- ied	Material	Form	expo- sure	Acadia clay	Cecil clay loam	Hag- ers- town loam	Lake Charles clay	Muck	Car- lisle muck	Rifle peat	Shar- key clay	Susque- hanna clay	Tidal marsh	Docas clay	Chino silt loam	Mo- have fine gravel- ly loam	Cin- ders	Mer- ced silt loam	Papa- kating silty clay loam
				Years											· · · · · · · ·					
				2.0	11.9	4.7	2.7	9.5	5.8		10.6	a 1.1	6.7	3.8	12.6	10.3	6.0	26.8		
~				5.4	18.4	2.0	2.4	14.8	12.1	2.6	7.2	4.8	5.6	3.2	21.8	6.2	6.2	48.2		
G	1932	Rattled cast iron b	Pipe	57.4	20.8	2.6	3.1	22.0	19.0	° 3.9	4.9	4.5	5.0	1.4	35.5	7.1	5.6	23.0		
				9.3		3.2	3.6	d D 4 D	20. I D	° 3. 6	20.8	7.1	8.1	6.1		7.6	4.0 D			
	1041	Diain cost inco	4.	14.3	e D	3.3	2.4	1 D		° 0. 8	28.6	12.3	8.0	° 13. 1 2 0	D 7	9.0	6.9	р. Д	7 5	0.8
A	1941	Plain cast iron	do	5.0	14.0	3.2	2.2	0. D 5 1	0.8	14.4	0.5	0.8	3.0	3.8	g 19.7	8.0	0. 2 5. 1	30.3	1.0	0.0
				2.0	14.9	3.0	2.8	0.5	4.0	9.2	9.8	×0.9 5.6	0.0	2.8	10.0	81	5.4	48.9		
т	1039	Low-alloy: 0.51 Cu. 0.28	do	7 4	24.0	1.7	2.4	94 7	20.6	03.0	7.2	5.0	4.2	2.0	39.4	8.0	4.7	29.8		
4	1352	Mn	•uo	9.3	21.1	2.8	2.1	37.3	20.0	4 1	0 18 2	7.5	6.6	o 10, 0	46.8	11.0	8.2	h 61.4+		
				14.3	1 27 6+	2.3	2.3	42.0	37.1	e 5. 1	33.6	11.0	6.7	7.4	49.6	12.3	18.6	D		
				1 2.0	14.7	3.1	3.9	5.1	3.7		7.8	a 1.0	6.8	4.6	15.9	11.0	6.0	24.6		
				5.4	22.9	1.8	1.5	12.5	11.5	2.5	7.2	6.2	4.6	3.6	26.7	9.2	7.0	51.4		
J	1932	Low-alloy; 0.62 Cu, 0.83	do	7.4	21.0	1.8	2.4	22.7	20.0	3.2	5.7	5.3	4.8	3.0	44.1	9.0	7.6	36.5		
		Mn		9.3		2.6	2.1	33.7	27.1	2.9	° 17.0	6.9	6.4	° 11. 4	44.6	10.3	12.2	h 65.0+		
				14.3	i 31.1+	3.4	2.6	f 43.0	h 39.6+	e 6.2	30.7	10.7	7.5	8.5	57.7	12.5	22.6	h 51.5+		
				( 2.0	15.1	3.9	3.4	7.2	9.0		9.0	■ 1.2	6.4	3.1	14.4	12.9	8.3	21.5		
				5.4	29.3	1.9	1.5	10.4	14.3	1.9	6.7	4.7	4.6	1.8	31.2	12.2	11.2	35.9	• •	
C	1932	Low-alloy; 0.30 Cr, 0.15	do	7.4	23.7	1.6	2.0	19.6	18.0	2.2	4.3	5.0	2.5	f 2.1	44.7	11.1	6.0	26.7		
		Ni		9.3		2.2	2.0	29.7	21.9	° 2.4	° 13. 1	7.2	6.9	2.6	f 41. 9	14.6	10.7	45.7		
				14.3	e D	3.5	3.3	h 34.0+	D	e 5.0	25.3	10.8	6.6	° 4. 5	h 36.1+	17.5	D	D		
B	1941	Low-alloy; 1.27 Ni, 0.32	do	5.0		3.9	2.5	3.9	4.4	13.3	2.0	5.7	3.5	2.4	20.4	7.7	8.2	D	4.8	1.0
		Cu																		
C	1941	Low-alloy; 1.71 Ni, 0.98	do	5.0		3.5	2.6	4.1	3.5	11.1	1.9	5.6	3.8	2.3	18.7	9.9	9.3	D	4.2	0.9
		Cu																	-	
NC	1941	Low-alloy; 2.08 Ni, 1.10 Cu	Plate	5.0		3.3	1.6	4.8	3.6	5.4	4.4	3.4	3.1	f 4.3	15.5	7.9	6.6	D	7.0	1.2
N	1941	Low-alloy; 3.10 Ni	do	5.0		3.0	2.1	7.3	3.2	5.4	3.6	3.4	2.5	f 3.5	15.8	8.2	f 10.2	D	7.6	1.2
D	1941	Low-alloy; 3.32 Ni	Pipe	5.0		3.5	2.6	2.9	1.4	7.6	3.5	5.2	2.6	9.6	14.4	7.0	8.3	D	3.7	0.8
				1 2.0	4.6	1.2	0.8	3.3	0.8		4.2	a 0.3	1.7	. 6	4.4	1.5	3.1	17.6		
				5.4	4.1	0.7	. 6	4.6	4.9	0.4	4.9	1.8	1.1	. 6	3.1	3.0	3.0	38.4		
E	1932	High-alloy; 15.0 Ni, 6.6	do	7.4	3.9	. 7	.7	9.4	8.6	°.6	f 1.3	1.7	1.0	. 7	5.8	2.0	3.5	24.3		
		Cu, 2.6 Cr		9.3		1.4	. 7	14.6	9.9	. 7	° 10.0	2.3	2.7	1.6	12.8	° 2. 6	c 3. 3	52.3		
				14.3	e 9.7	1.2	1.1	17.3	10.0	• 1.3	11.6	5.2	2.3	1.5	4.8	3.1	7.6	D		

<sup>a</sup> Exposed for 1.0 yr only.

<sup>b</sup> Ordinary cast iron horizontally cast in green-sand molds and rattled to remove sand.

° Data for the individual specimens differed from the average by more than 50 percent.

<sup>d</sup> D, both specimens destroyed by corrosion.

e Data for 4 specimens.

<sup>f</sup> Data for 1 specimen. The other specimen was missing.

g Data for 3 specimens.

 $^{\rm h}$  Data for 1 specimen. The other specimen was destroyed by corrosion.

i Data for 3 specimens. The other specimen was destroyed by corrosion.



FIGURE 10. Corrosion of unalloyed steel exposed 14 yr at 14 test sites. See table 1 for identification and properties of soils.

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FIGURE 11. Corrosion of plain cast iron exposed 14 yr at 14 test sites. See table 1 for identification and properties of soils.



FIGURE 12. Corrosion of open-hearth iron plates exposed 9 yr at 15 test sites. See table 1 for identification and properties of soils

# Soil Corrosion of Ferrous Materials



FIGURE 13. Weight loss and pit depth-time curves for wrought materials in typical soils

these constants, calculated according to the method of least squares, are recorded for the individual soils in table 14. The standard errors of the constants, calculated according to the method of Ezekiel [6] are included in the table. Values for the constants k and n for a much larger group of soils than those under consideration at present have been reported [3].

Because the constants k and k' represent, respectively, maximum penetration and weight loss at 1 yr., the values of these constants may be considered to measure approximately the initial rate of corrosion and consequently the inherent corrosiveness of the soils. The relatively large values for the constant k for the group of six soils of fair to good aeration, No. 64, 66, 62, 65, 55, and 53 (table 14) are to be ascribed largely to the depolarization of the cathodic areas by the oxygen of the soil atmosphere. The low values of k for the remaining poorly or very poorly aerated soils are a consequence of cathodic polarization because of the deficiency of oxygen in these soils.

Comparison of the values for k and n shows that there is a good inverse correlation between these two constants, from which it follows that the higher the initial rate of pitting the more rapidly does the rate of pitting decrease with time. A necessary corollary of this conclusion is that measurements of the initial rate of pitting of wrought ferrous materials in soils cannot be used for predicting corrosion over a long period unless the measurements are accompanied by some expression for the change in corrosion with time for the particular environmental conditions.

A probable explanation for the inverse correlation noted between the constants k and n is as follows: In poorly aerated soils, characterized by low values for the constant k, ferrous ions, migrating and diffusing from the local anodes have a negligible effect on the rate of corrosion, the rate being determined entirely by the depolarization of hydrogen at the cathode. However, in soils that contain oxygen in excess of that required for cathodic depolarization, ferrous ions are oxidized and precipitated in close proximity to the local anodes. The tubercles formed in this manner prevent the further migration and diffusion of ferrous ions with consequent reduction in the rate of corrosion by anodic polarization. The mechanism of tubercle formation and its significance in soil corrosion have been considered in previous papers [8, 9].

## TABLE 13. Maximum penetration of specimens of cast iron

#### (Average of 2 specimens in mils)

	2	*	211								Test	site nur	nber and	soil type	е					
Identifica-	Year	Matarial	Farm	Aver-	51	53	55	56	58	59	60	61	62	63	64	65	66	67	70	72
tion	ied	Material	Form	expo- sure	Acadia clay	Cecil clay loam	Hag- ers- town loam	Lake Charles clay	Muck	Car- lisle muck	Rifle peat	Shar- key clay	Susque- hanna clay	Tidal marsh	Docas clay	Chino silt loam	Mo- have fine gravel- ly loam	Cin- ders	Mer- ced silt loam	Papa- kating silty clay loam
				Years											-					-
				2.0	90	48	48	41	34		50	a 24	70	38	131	46	56	129		
E	1000	Gen d sected cost inco	D!	5.4	° 250+	49	66	119	104	52	60	48	66	43	116	74	71	250+		
r r	1932	Sand-coated cast iron	Pipe	7.4	250+	102 b 75	126	250+	200+	46	35	56	106	52	150	86	122	250+		
				14 3	d 250∔	128	135	250+ g 250+	250+ 250+	44 d 64	149	18	100	131 b 164 l	250+	145	250 1	250+		
			~	14.5	118	42	48	s 230∓ 36	250+	~ 0±	52	3 43	64	94	194	182+	250+	250+		
				5.4	250+	51	50	109	179+	33	68	55	95	74	124	94	79	250+		
G	1932	Rattled cast iron e	do	7.4	250 +	71	126	250 +	250+	52	26	76	118	61	122	112	181+	210+		
				9.3		57	123	250+	250+	52	b 175+	78	b 81	114	250 +	131	152	250+		
			×	14.3	d 250+	138	149	250+	250+	d 70	197+	98	140	156	250+	146	250+	250+		
A	1941	Plain cast iron	do	5.0		179 +	110	124	106	89	52	78	53	99	f 151	113	67	250+	104	39
	1.1			2.0	102	30	28	22	41		38	a 30	100	20	136	56	57	128		
· · ·	1020	Low allow 0.51 Cu. 0.99	4.	5.4	305+	47	56	98	107	43	50	83	71	60	116	95	60	ь 217+		
1	1932	Low-anoy; 0.51 Cu, 0.28	do	$) \frac{7.4}{0.2}$	304+	50	108	191	192	44	80	90	94	b 90	146	110	149	240		
		14114.		14 3	d 317⊥	08	108	249	240	d 03	0 140	113	102	104	317+	172	214	327+		
				(2.0)	106	54	36	30	238	- 55	38	a 26	68	51	1201+	58	350+ 54	350+ 108		
				5.4	309+	45	-59	- 93	109	35	57	60	80	83	104	100	70	b 210+		
J	1932	Low-alloy; 0.62 Cu, 0.83	do	7.4	260	60	97	184	179	57	25	78	84	72	156	118	200	276+		
		Mn.		9.3		62	ь 96	215	233	49	b 130	80	b 63	85	246	156	193	266 +		
				14.3	d 350+	57	108	g 230	314+	d 57	152	124	112	g 78	256+	150	350+	314 +		
				2.0	112	39	26	32	44		34	a 15	80	10	142	58	68	119		
0	1029	Low allow 0.20 Cn. 0.15 Ni	1	5.4	250+	40	58	101	124	11	72	44	76	12	146	91	120	250+		
C	1952	Low-alloy, 0.30 Cr, 0.15 MI_		1.4	250+	88 b 66	95	150	176	D 20	18	53	83	g 132	143	128	161	250+		
				14.3	d 250+	171+	123	210+	250+ 250+	24 d 56	0 93 250⊥	119	194	40	g 250+	101	141	195+		
в	1941	Low-alloy: 1.27 Ni, 0.32 Cu	do	5.0	200 1	124	114	96	140	- 50	200-	197	77	102	187	195+	250+	250+	100	
C	1941	Low-alloy; 1.71 Ni, 0.98 Cu_	do	5.0		144	123	64	109	111	32	112	112	86	210+	129	80	250 + 250 +	00	32 59
NC	1941	Low-alloy; 2.08 Ni, 1.10 Cu.	Plate	5.0		133	110	110	118	78	56	84	94	g 110	138	149	88	250 +	101	32
N	1941	Low-alloy; 3.10 Ni	do	5.0		120	113	120	119	80	66	96	76	g 106	138	100	g 80	250+	107	29
D	1941	Low-alloy; 3.32 Ni	Pipe	5.0		136	96	50	144	87	94	104	104	24	226+	157	80	250+	93	34
				2.0	30	30	30	42	50		24	a 36	b 43	22	28	26	26	94		
E	1020	High allow 15.0 Ni C.C.	d e	5.4	50	36	37	32	34	14	<sup>b</sup> 80	39	49	16	27	42	36	250+		
E	1932	High-alloy; 15.0 N1, 6.6	do	7.4	35	51	41	53	b 58	28	g 22	30	37	55	40	42	38	208 +		
		Ou, 2.0 Or.		9.3	d 79	33	34	58	53	26 d 20	D 45	0 36	53	<sup>D</sup> 32	74	35	40	250+		
11 - 2 <sup>4</sup> 2				( 14. 3	a 78	44	51	12	54	a 30	12	58	60	30	34	59	54	250+		

<sup>a</sup> Exposed for 1.0 yr only.

<sup>b</sup> Data for the individual specimens differed from the average by more than 50%.

• +, one or more specimens contained holes because of corrosion.

d Data for 4 specimens.

• Ordinary cast iron horizontally cast in green-sand molds and rattled to remove sand.

f Data for 3 specimens.

# Data for 1 specimen. The other specimen was missing.

 TABLE 14.
 Corrosion of wrought and cast materials at 14 years and constants of equations connecting weight loss and maximum penetration with duration of exposure \*

	Soil				Weight	loss			Maximum penetration						
No.	Туре	Aeration	W <sub>T=14</sub> yr	σ <sub>P<sub>T=14</sub> yr</sub>	k'	$\overline{\sigma}_{k}'$	u	$\overline{\sigma}_u$	PT=14 yr	σP <sub>T=14</sub> yr	k	$\overline{\sigma}_k$	n	$\overline{\sigma}_n$	
			W	ROUG	НТ МА	TERIA	LS								
			$oz/ft^2$	$oz/ft^2$	oz/ft <sup>2</sup>	$oz/ft^2$			Mils	Mils	Mils	Mils			
64	Docas clay	Fair	50.9	9.0	8.41	1.71	0.68	0.13	(b)						
66	Mohave fine gravelly loam	do	15.0	1.6	7.12	1.12	. 28	. 09	(b)						
65	Chino silt loam	Good	13.1	1.1	6.04	0.89	. 29	. 08	108	9	57.9	8.9	. 24	0.08	
62	Susquehanna clay	Fair	6.9	1.0	3.18	. 87	. 29	. 14	83	5	55.7	6.4	. 15	.06	
55	Hagerstown loam	Good	3.4	0.4	2.08	. 52	. 19	. 13	82	6	40.8	5.6	. 27	.08	
53	Cecil clay loam	do	4.2	.4	2.46	. 44	. 20	. 09	77	8	34.1	6.3	. 31	. 09	
60	Rifle peat	Poor	17.8	6.2	2.61	1.78	. 73	. 33	56	16	16.4	9.0	. 46	. 27	
63	Tidal marsh	Very poor	7.7	1.5	1.58	0.64	. 60	. 20	81	11	13.8	3.8	. 67	. 14	
61	Sharkey clay	Poor	9.3	0.8	1.12	. 14	. 80	.07	92	8	13.5	1.7	. 72	.06	
56	Lake Charles clay	Very poor	30.5	2.3	1.72	. 25	1.09	.08	(b)						
58	Muck	Poor	21.0	1.8	1.85	. 31	0.92	.08	194	54	9.1	2.8	1.16	. 17	
59	Carlisle muck	Very poor	4.1	0.1	0.49	. 02	. 81	. 02	36	2	4.4	0.7	0.80	.07	
51	Acadia clay	Poor	20.9	. 2	5.59	. 10	. 50	. 09	(b)						
67	Cinders	Very poor	41.0	3.0	8.67	. 76	. 59	. 06	(b)						
			-	<u> </u>											
				CAST	MATE	RIALS									
64	Doess elsy	Fair	58.0	5.8	0.12	1.4	0.70	0.09	140	20	120	20	0.06	0.10	
62	Susquebanna clay	do	5.0	1.1	5 53	1.1	0.10	16	98	10	70.5	14 0	13	10	
65	Chino silt loam	Good	10.7	1.1	9.93	2.0	.03	. 10	163	10	40.5	6.4	. 15 53	08	
66	Mohave fine gravelly losm	Fair	8.0	1.1	5 78	0.69	.03	. 12	294	58	34 5	13.3	71	.03	
53	Cocil clay loam	Good	9.0	0.5	3 13	1.4	- 11	. 10	81	10	28 5	6.8	30	12	
61	Sharkey clay	Poor	10.1	1.0	1 12	0.17	84	. 22	110	8	28.1	3.6	. 53	05	
60	Bifle nest	do	17.0	8.0	4 45	4.3	51	43	72	68	23.7	29.0	42	59	
55	Hagerstown loam	Good	2.3	0.4	2.82	0.96	- 07	17	146	17	20.5	4.3	74	. 11	
58	Muck	Poor	32.0	2.4	3.05	35	80	06	369	26	14 9	1.3	1.22	05	
63	Tidal marsh	Very poor	6.1	2.4	1.96	1.5	70	.00	126	7	14.8	1.0	0.81	06	
56	Lake Charles clay	do	40.5	6.8	3.12	1.0	97	16	369	51	12.1	2.0	1.30	. 11	
59	Carlisle muck	do	5.4	0.5	0.51	0.16	89	14	67	11	10.5	4.3	0.70	26	
51	Acadia clay	Poor	29.6	6.0	11.4	. 96	36	. 16	(b)		10.0		00		
67	Cinders	Very poor	60.3	14.0	19.4	7.0	43	. 20	(b)						
		, or y post	00.0	1											

•  $W_{=k'T^u}$ ,  $P_{=kT^n}$ , where  $W_{=}$  weight loss at the time, T; and  $P_{=}$  depth of the deepest pit at the time, T.

<sup>b</sup> Specimens perforated at an early period. See table 6 for data on pit depths.

Although the permeability of the soil to air appears to be the major factor in determining the value of the constant n, it should be noted that any property of the soil that tends to increase the solubility of ferrous ions at the local anodes, such as high contents of chloride, sulfate, or hydrogen ions, would also have the effect of increasing the magnitude of n and consequently the depths of the deepest pits.

The inverse correlation between the values of the constants for maximum penetration, k and n, is also to be noted between the constants for weight loss, k' and u, showing that weight loss and pitting follow the same general tendencies with respect to the soil environment.

The average values of the constants k, k', n, and u for the cast materials that corroded at approximately the same rates are also presented in table 14. The materials represented are plain cast iron G, and the low-alloy cast irons C, I, and J. In this table the same general tendencies noted in the case of the wrought materials are to be observed, but with more numerous exceptions.

#### 3. Comparison of Wrought and Cast Materials

Because of dissimilarities in the dimensions of the wrought and cast ferrous specimens in the previous field tests, no comparison could be made of the relative corrodibility of the two classes of materials. However, in the present series of tests, the dimensions of the cast and wrought specimens were similar, so that comparisons between the two materials are possible. The cast and wrought materials were compared by the method previously described in connection with the comparison with plain steel of the low-alloy wrought materials exposed 14 yr. The differences between the values of weight loss and pitting for cast iron and steel at 14 yr recorded in table 14 for the different soils were calculated, and the standard errors of these differences were computed by means of eq 6. These data are given in table 15.

Because the differences between the mean values for weight loss of the cast and wrought materials were, with few exceptions, less than twice the respective standard errors of the differences, it is necessary to conclude that no generally significant difference between the weight losses of the two materials can be detected. With respect to maximum penetration, differences less than twice the standard error of the difference were noted in four of the nine soils for which the data could be statistically treated. In the remaining five soils, the greater depth of pits on the cast specimens should be considered significant.

# V. Summary

This report contains the results of measurements of corrosion made on a variety of wrought and cast ferrous materials after exposure to different soil conditions for periods up to 14 vr. Steels containing small amounts of nickel and chromium showed increased resistance to soil corrosion, but the resulting improvement was small. However, certain wholly austenitic steels containing high percentages of chromium and nickel were completely resistant to corrosion. In those soils in which there was a significant difference in the pitting of cast iron and wrought materials, cast iron usually pitted at somewhat higher rates. High sustained rates of corrosion occurred generally in poorly aerated soils high in soluble salts or in acidity. In well aerated soils low in soluble salts, corrosion virtually ceased after a relatively short period because of the formation of layers of corrosion products close to the metal surface.

			Loss	in weigh	at (oz/ft 2)			Maximum pit depth (mils)									
Soil a	Wro mate	ught rials	Cast m	aterials	Differ- ence	Standard error		Wrot mate	ught rials	Cast m	aterials	Differ- ence	Standard error				
5011 -	Mean	Stand- ard error	Mean	Stand- ard error	between the means	of the differ- ence <sup>b</sup>		Mean	Stand- ard error	Mean	Stand- ard error	between the means	of the differ- ence <sup>b</sup>				
	$\overline{X}_1$	$\overline{\sigma}_1$	$\overline{X}_2$	$\overline{\sigma}_2$	$\overline{X}_1\!-\!\overline{X}_2$	$\overline{\sigma}_D$	$2\sigma_D$	$X_1$	$m{ar{\sigma}}_1$	$\overline{X}_2$	$\overline{\sigma}_2$	$\overline{X}_1 \!-\! \overline{X}_2$	$\overline{\sigma}_D$	$2\sigma_D$			
51	20.9	0.2	29.6	6.0	-8.7	6.0	12.0										
53	4.2	.4	2.4	0.5	+1.8	0.7	1.4	77	8	81	10	-4	12	24			
55	3.4	.4	2.3	.4	+1.1	. 6	1.2	82	6	146	17	-64	18	36			
56	30.5	2.3	40.5	6.8	-10.0	7.2	14.4										
58	21.0	1.8	32.0	2.4	-9.0	3.0	6.0	194	54	369	26	-175	60	120			
59	4.1	0.1	5.4	0.5	-1.2	0.5	1.0	36	2	67	11	-30	11	22			
60	17.8	6.2	17.0	8.0	+0.8	10.1	20.2	56	16	72	68	-16	70	140			
61	9.3	0.8	10.1	1.0	8	1.3	2.6	92	8	110	8	-18	11	22			
62	6.9	1.0	5.9	1.1	+.9	1.5	3.0	83	5	98	10	-16	11	22			
63	7.7	1.5	6.1	2.3	+1.6	2.8	5.6	81	11	126	7	-45	13	26			
64	50.9	9.0	58.0	5.8	-7.1	10.7	21.4					· · · · · · · · · · · ·					
65	13.1	1.1	10.7	1.4	+2.4	1.8	3.6	108	9	163	14	-55	17	33			
66	15.0	1.6	8.0	1.5	+7.0	2.2	4.4										
67	41.0	3.0	60.3	14.0	-19.0	14.3	28.6										

TABLE 15. Comparison of wrought and cast materials at 14 years

\* See table 1 for type, location, and properties of soils.

$$T_D = \sqrt{\frac{\sigma_1^2}{N_1} + \frac{\sigma_2^2}{N_2}}$$
 (see text)

The field tests described in this paper were planned and installed, and until 1946 were conducted under the supervision of K. H. Logan. The measurements of corrosion were made chiefly by Warren P. Dettmers, who assisted in other capacities.

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WASHINGTON, June 10, 1949.