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## [CONTRIBUTION FROM THE JOHN HARRISON LABORATORY OF CHEMISTRY.] THE EFFECT OF SULPHURIC ACID ON THE DEPOSITION OF METALS WHEN USING A MERCURY CATHODE AND ROTATING ANODE.

By LILY G. KOLLOCK AND EDGAR F. SMITH. Received March 19, 1907.

In the course of a study of the precipitation of metals with the help of a mercury cathode and rotating anode, observations were made which led to a review of the experiments to ascertain more especially what effect varying amounts of acid would have upon the electrolytic decomposition of certain metallic sulphates.



The metals studied under these conditions were zinc, copper, cadmium,

iron, cobalt and nickel. The volume of the solution of the salt was, in nearly all cases, 10 c.c. containing about 0.25 gram of metal. The current was maintained at two amperes, while the pressure varied from 2 to 4.5 volts. The sulphuric acid was concentrated. Curves were plotted from the results obtained.

#### Zinc

When one cubic centimeter of sulphuric acid was present in a solution of zinc sulphate, the zinc was completely deposited in ten minutes. The solution contained 0.25 gram of metal, the volume of the solution being 11 cc. The following observations were made :

In 2.5 minutes, 0.1040 gram of zinc was deposited.

` 5	**	0.1974	" "	" "	٤,
7.5	••	0.2400	• •		" "
8	4.4	0.2428	" "	4.	٤٠
, 10	ъi	0.2494	"	•••	٠.

#### (Figure 1, Curve 1.)

When two cubic centimeters of sulphuric acid were present, the solution which was siphoned from the cup, showed the presence of a trace of zinc after it had been electrolyzed for ten minutes. Eleven minutes were necessary for the complete removal of the zinc. The volume of the solution in this case was 12 cc. The following rate of deposition was observed :

In 2.5 minutes, 0.0953 gram of zine was deposited.

``5	•••	0.1805	• •	••	••
	٠,	0.2306	••	• -	•••
" 10	÷ •	0.2467	••	1 .	• •
ʻ'I1	۰.	0.2488	• •	• •	" "

#### (Figure 1, Curve 2.)

In the presence of three cubic centimeters of acid, twelve minutes were required for the deposition of the zinc. The volume of the solution in these experiments was 13 cc. Under these conditions

In 2.5 minutes, 0.0815 gram of zinc was deposited.

·· 5	•••	0.1633	••	••	••
·· 7.5	* *	2.200	<b>、</b> (	4.	÷+
·· 10	۰.	0.2428	• 6	••	••
`` 12	٠.	0.2493	٠.	• •	••

#### (Figure 1, Curve 3.)

When 4 cc. of acid were present, considerable retardation in the decomposition was observed; for at the end of the fifteen minute period, there was still some undeposited zinc. The test at the end of eighteen minutes showed that no zinc was present in the solution. The rate of precipitation was as follows:

In 2.5 minutes, 0.0668 gram of zinc was deposited.

ʻ` 5	۴۰	0.1506	••		• •
		0.2050	**		i 1
ʻʻ 10	<b>6</b> v	0.2315	6.6	• •	• •
"12	4.4	0.2423	÷ •	÷ •	• •
<b>** 1</b> 5	4.1	0.2480	6 F	* *	* *
·· 18	" "	0.2495	4.4	" "	4.4
		( Fig	gure 1	, Curve 4.)	

That the presence of 5 c.c. of acid greatly retards the precipitation was shown when a solution, containing 0.2110 gram of zinc was electrolyzed. It was thirty minutes before the last traces of metal were thrown out.

In 2.5	minutes,	<b>o.o</b> 668	gram	of zinc was	deposited
" 5	"	0.1306	••	" "	" "
"10	**	0.1980	" "	"	" "
ʻʻ 20	" "	0.2077	"	"	" "
ʻʻ 30	**	0.2110	" "	**	" "
		(Fi	gure 1	, Curve 5.)	

In these experiments the volume of the solution was increased, in each case, by the addition of the acid. Thinking that, perhaps, the increased period of time might be due to the greater volume of solution, it was observed that the retardation was due principally to the acid present. From a solution of 14 cc. volume, containing 1 cc. of sulphuric acid, all the zinc was deposited in twelve minutes. In the same volume of solution, containing 4 cc. of acid, eighteen minutes were necessary for complete precipitation.

Copper

The volume of the copper sulphate solution in these experiments, as



well as in the solutions of the sulphates of the succeeding metals, was 10 cc. including the volume of the acid added.

The amount of copper present in the solution was 0.2485 gram. In

the presence of 1 cc. of sulphuric acid, all the copper was deposited in ten minutes. The current registered two amperes and showed a pressure of from 3 to 3.5 volts. The solution became colorless in seven minutes. Three minutes additional were necessary to deposit the remaining 0.007 gram of copper. The following rate of precipitation was observed :

I 11	2.5	minutes,	0.0985	gram	of copper	was deposited.
6.6	5	"	0.1887	Ŭ ,,	6.5	- ,,
" "	7.5	• •	0.2418	i 1	4 <del>.</del>	
6 4	io	* *	0.2437	* *	• •	÷ ,
			(Fi	gure 2	, Curve 1.	)

When 2 cc. of acid were present and the current was maintained at two amperes, the pressure varying from 4 to 4.5 volts, it was found that the copper was completely deposited in ten minutes. It will be observed that the pressure in this case was half a volt to a volt greater than in the previous experiment. The two curves almost coincide.

> In 2.5 minutes, 0.1157 gram of copper was deposited. '' 5 '' 0.1897 '' '' '' '' '' '' '' 7.5 '' 0.2385 '' '' '' '' '' '' 10 '' 0.2489 '' '' '' '' '' (Figure 2, Curve 2.)

The presence of 3 cc. of acid affects the curve but slightly. The departure is noticed in the deposition of the last traces of the metal. The copper was completely removed in twelve minutes.

[n	2.5	minutes,	0.1079	gram	of	copper	was	deposited.
44	5	6 6	0.1807	- í,		17		·
" "	7.5	• •	0.2311	٤٠		4.4		• •
" "	io T	• •	0.2450	٠.				* *
<b>''</b> 1	12	" "	0.2488	" "		" "		" "

#### (Figure 2, Curve 3.)

The effect of a large amount of sulphuric acid was noticed again in the latter part of the decomposition, when 4 cc. of acid were added. It was then observed that it took six minutes longer than it did when but 3 cc. were present to remove all copper.

In 2.5 minutes, 0.1060 gram of copper was deposited.

·· 5	* *	0.1684	<b>6</b> 4	* *	**
·· 7.5	" "	0.2078	* *	• •	
" ío"	"	0.2402	4 4		÷ 1
ʻ`12	"	0.2458	* *	6.6	. 4
ʻʻ 18	"	0.2487	• •	" "	۴۰
		( 11:-		0	

#### (Figure 2, Curve 4.)

In the presence of 5 cc. of acid, twenty-five minutes were necessary in order that the last traces of copper might be removed. The solution did not become colorless until after eighteen minutes.

In 5 minutes, 0.1422 gram of copper was deposited.

" 10	* *	0.1943	•••	- 7,	• .,
"12	÷ 6	0.2140	"	4 4	÷
ʻʻ 15	" "	0.2334	" "	÷ 4	* *
" 20	6.6	0.2451	" "	• •	• •
·· 23	* *	0.2476	• •	6 <del>.</del>	••
·' 25	* *	0.2484	" "	" "	4.4

800

From a neutral solution of the salt, under the same conditions, the copper was completely precipitated in eight minutes.

#### Cadmium

In the presence of 1 c.c. of sulphuric acid 0.25 gram of cadmium was deposited in ten minutes. The current equalled two amperes, and the pressure from 4 to 4.5 volts.



FIG. 3. Cadmium

In 2.5 minutes, 0.1720 gram of cadmium was deposited.

'' 5		0.2418	" "	* *	" "
" 7.5	"	0.2480		" "	**
''10	" "	0.2500	4 G	• 4	" "
		(Fig	ure 3, C	urve 1,)	

When 2 cc. of acid were present, hydrogen sulphide, after fifteen minutes gave a faint yellow color to the solution from the cell. In eighteen minutes there was no color on testing the liquid in the same way.

In 2.5 minutes, 0.1250 gram of cadmium was deposited.

" 5	6 F	0.2260	"	* *	
·· 7·5	• •	0.2398	" "	* *	" "
" 10	" "	0.2439	* *		" "
" 15		0.2480	" "	* *	**
·· 18		0.2501	" "	" "	4.4
		(Trime			

(Figure 3, Curve 2.)

Twenty-two minutes were required to precipitate all of the cadmium in the presence of 3 cc. of acid.

I 11	5	minutes,	0.2028	gram	$_{\rm of}$	cadmium	was deposited
••	7.5		0,2260	· · ·		• •	
" "	10	* 6	0.2433	4.6		٠,	• 1
61	15	• 4	0.2475	<del>7</del> 4		۰,	14
1.	26	* *	0.2486	• •		* *	
۰،	22	••	0.2498	• •		• •	۰.
			/ * **		~		

(Figure 3, Curve 3.)

When 4 cc. of acid were added to the electrolyte, twenty-seven minutes were necessary for the complete precipitation of the cadmium.

In 5 minutes, 0.2023 gram of cadmium was deposited.

' <b>1</b> 0	• •	0.2392	۴۰	••	,
15	" "	0.2491	5.6	• •	•
' 20	" "	0.2488	* *	**	,
6 25	" "	0.2495	14	**	,
· 27	"	0.2505	" "	" "	,
		(F	oure	2 Curve A	

When 5 cc. of acid were present in the solution, at the end of thirty-five minutes there was considerable cadminin undeposited (0.003).

In 5 minutes, 0.1974 gram of cadminu was deposited.

~					
' IO	• •	0.2354	"	11	•
<b>2</b> 0		0.2419	• •	**	•
' 30	٤,	0.2454	••	÷ 1	•
35	••	0.2475	<b>6</b> 1	• •	•
		(Fig	gure 3	, Curve 5.)	

Iron

Sulphuric acid retards the decomposition of iron sulphate to a far



Fig. 4. from

greater extent. Even after forty minutes, in the presence of 1 cc. of acid, a trace of iron was held in solution.

In	5	minutes,	0.0858	gram	of iron	was	deposited.
" "	IO	÷ 4	0.1204	· · ·	4.4		
6.6	15	• •	0.16:9	" "	* *		٠.
**	20	" "	0.2196	" "			• •
4.4	30	• •	0.2439	" "	**		" "
• •	35	" "	0.2481	" "	" "		**

#### (Figure 4.)

In the presence of 3 cc. of acid, in twenty-five minutes, 0.069 gram of iron was found in the mercury. 4 cc. of acid were not enough to completely hold up the iron.

#### Nickel

In fifteen minutes the nickel solution, containing I cc. of sulphuric acid, became colorless. It required twenty minutes, however, to remove the last traces of metal.

[n	5	minutes,	0.1404	gram	of nickel	was deposited.
"	10		0.2070	С <i>.</i> ,	* *	
"	15		0.2370	4.4		" "
۴.	20	* *	0.2476	" "	**	
""	25	* *	0.2511		" "	* *
			$(\mathbf{F})$	ioure	Curve I	)

In the presence of 2 cc. of acid, after twenty minutes, the solution gave a brown color after adding ammonium hydroxide and ammonium sulphide. In thirty minutes the test showed that there was still a small amount of nickel unprecipitated. In thirty-two minutes nickel was found in the solution siphoned from the cup.



I 11	5	minutes,	0.1265	gram	of nickel	was deposited.
* *	10	" "	0.2053	- · ·	* *	
۴۰	15	• •	0,2290	÷ 6	£ 1	••
4.4	20	• •	0.2383	44		* 6
4.4	25	••	0,2445	* *	• •	
"	30	**	0.2485	( <b>i</b>	• •	4.1
"	32	" "	0.2500	4.7	* *	

(Figure 5, Curve 2.)

#### Cobalt

The solution of cobalt sulphate contained 0.1700 gram of cobalt. Twenty-five minutes were required for its complete precipitation in the presence of 1 cc. of sulphuric acid.

In	5	minutes,	0.1010	gram	of cobalt	was	deposited.
	10	" "	0.1539		* *		
" "	15	• •	0.1663	" "	* *		
" "	20	**	0.1681	÷ í	4.5		
"	25	" "	0.1700	63	6.7		* *
(Figure 6, Curve 1.)							

When 2 cc. of sulphuric acid were present, ten minutes additional were necessary for the removal of the cobalt.



(Figure 6, Curve 2.)

The ease and rapidity with which metals are precipitated upon the mercury cathode naturally suggested the inquiry as to the separations which were possible in this manner. The following lines communicate a few results obtained in this direction. The idea will, however, be pursued in detail as far as it can be carried out.

## Cadmium from Aluminium

The aluminium salt appears to retard the complete deposition of cadmium. In the presence of a few (4) drops of concentrated sulphuric acid with a current of two amperes and seven volts, considerable cadmium remained undeposited after the current had acted fifteen minutes. A higher current was, therefore, employed. The separation was complete in twenty minutes, using a current of three amperes and seven volts. The cadmium in solution was 0.25 gram, while the aluminium sulphate was equivalent to 0.1 gram of aluminium. The total volume of the solution was 10 cc. The increase in the weight of the mercury in the first experiment was 0.2502 gram and in the second 0.2495 gram.

# Cadmium from Magnesium

This separation was accomplished under the same conditions as the previous separation (cadmium from aluminium).

#### CONDITIONS

Volume of Solution =	10 cubic centimeters.
Magnesium sulphate =	0.1 grain magnesium.
Cadmium sulphate	0.25 gram cadmium.
Sulphuric acid =	4 drops (40 drops = $1 \text{ cc.}$ )
Current =	2–3 amperes.
Pressure =	11–9 volts.
Time = :	25 minutes.
Cadmium found (first experiment) ···· . = a	0.2505 gram.
Cadmium found (second experiment) = a	0.2497 gram.

### Cadmium from Chromium

Inasmuch as chromium is deposited from its sulphate from slightly acidulated solution, the separation of chromium from cadmium was carried out in the presence of 3 cc. of concentrated acid. The conditions were as follows:

Volume $\dots \dots \dots$
Chromium sulphate e= 0.1 gram chromium.
Cadmium sulphate
Sulphuric acid 3 cubic centimeters.
Current $\dots = 2-3$ amperes.
Pressure $\ldots = 3.5-4$ volts.
Time = 25 minutes.
Cadmium found (first experiment) = 0.2499 gram.
Cadmium found (second experiment) = 0.2492 gram.

#### Copper from Aluminium

This separation was accomplished in the presence of 0.5 cc. of sul-

phuric acid (1.1). The current registered 1 ampere and 4 volts. In four minutes the solution was colorless. The current was allowed to act for ten minutes.

Volume of the solution ==	to cubic centimeters.
Copper sulpliate	o.1150 gram coppe <b>r</b> .
Aluminium sulphate	o.1 gram aluminium.
Sulphuric acid	0.5 cubic centimeter.
Current	1–1.6 ampere.
Pressure	4-4.5 volts.
Time	10 minutes.
Copper found	0.1150 gram, 0.1153 gra

Copper found ...... 0.1150 gram, 0.1153 gram, 0.1152 gram. A low current (0.01 ampere and 2 volts) was passed through zinc sulpliate in the presence of 0.25 cc. of sulpluric acid. In twenty minutes the niercury had increased 0.0006 gram in weight.

Iron sulphate containing 0.1110 gram of iron, acidulated with 0.5 cc. of acid, with a current of 0.01 ampere and a pressure of 2 volts gave in thirty minutes 0.0434 gram of iron. When 1 cc. of acid was present, a current of 0.1 ampere with a pressure of 2 volts gave 0.0911 gram of metal.

The separation of copper from zinc was attempted but it was not successful. A copper solution containing 0.1150 gram of copper in the presence of 2.5 cc. of acid and a current of 0.6 ampere and three volts, caused the mercury to increase in weight 0.1360 gram, showing that 0.0212 gram of zinc had also been deposited.

The separation of copper from iron was tried, but this, too, failed. With a current (like that given in the preceding paragraph) iron was detected in the mercury, and in thirty minutes the solution still showed the presence of copper. 0.25 cc. of acid was present. The iron seemed to hold back the copper.

When 3 cc. of acid were present in a solution of iron sulphate, containing 0.1 gram of iron and it was electrolyzed with a current of two amperes and four volts, 0.069 gram of iron went into the mercury. The experiment was repeated after the addition of 4 cc. of acid. Even this quantity failed to hold up all the iron. The separation, therefore, of iron from copper and of iron from zinc was unsuccessful.

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[CONTRIBUTION FROM THE JOHN HARRISON LABORATORY OF CHEMISTRY.] THE USE OF A ROTATING ANODE IN THE ELECTROLYTIC PRECIPI-TATION OF URANIUM AND MOLYBDENUM

> By EDGAR T. WHERRY AND EDGAR F. SMITH. Received March 19, 1967.

#### I. URANIUM

The early suggestion of Smith<sup>1</sup> that uranium could be completely precipitated by the current from an acetate electrolyte has been amply veri-

<sup>1</sup> Am. Ch. J., 1, 329.