COMPONENTS: (1) Cadmium oxide; CdO; [1306-19-0]	ORIGINAL MEASUREMENTS:
(1) Cadmium oxide; CdO; [1306-19-0]	
	Piater, J. Z. anorg. allgem. Chem. <u>1928</u> , 174, 321-41.
(2) Cadmium hydroxide; Cd(OH) ₂ ; [21041-95-2]	
(3) Sodium hydroxide; NaOH; [1310-73-2]	
(4) Water; H ₂ 0; [7732-18-5]	
VARIABLES:	PREPARED BY:
Method of preparation of Cd(OH) ₂ and the NaOH concentration	T. P. Dirkse
EXPERIMENTAL VALUES: Table	1
Solubility of CdO or Cd(OH)	2 in water at 25°C.
Preparation mol dm ⁻³	Solubility Product (mol ³ dm ⁻⁹)
Cd0 1.79 x 10 ⁻⁵	2.3×10^{-14}
$Cd(0H)_2^a$ 1.30 x 10 ⁻⁵	8.8×10^{-15}
$Cd(0H)_2^{b}$ 1.14 x 10 ⁻⁵	5.9×10^{-15}
$Cd(0H)_{2}^{c}$ 1.11 x 10 ⁻⁵	5.5×10^{-15}
^a prepared by the method of De Schulten (2)	
	kept under water at least 3 months before use
AUXILIARY	INFORMATION
AUXILIARY METHOD/APPARATUS/PROCEDURE: Apparently the solid was added to the solvent and shaken. The specific conductance of the solution was measured and the validity of the method was judged by the conductance behavior. Corrections were made for dissolved CO ₂ (1) and the solubility was calculated from the specific conductance and accepted values for ionic conductances.	SOURCE AND PURITY OF MATERIALS: Analysis of each preparation showed about 0.5% impurities. Conductivity water was

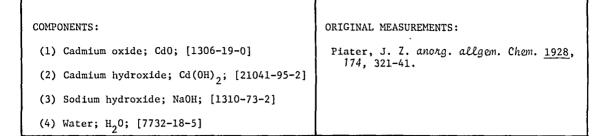


Table 2

Solubility of Cd(0H) $_{\rm 2}$ in NaOH solutions at 25°C.

mol NaOH dm ⁻³	mol Cd(OH) ₂ dm ⁻³
0.00 0.01 0.1 1.0 5.0	1.2×10^{-5} 1.3×10^{-6} 1.3×10^{-6} 1.3×10^{-5} 0.7×10^{-5} 9.0×10^{-5}

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COMPONENTS:			ORIGINAL MEAS	IIDENENTC .	
	ide; Cd(OH) ₂ ; [21041-				F 7. аполо.
(2) Sodium hydroxide; NaOH; [1310-73-2] (3) Water; H ₂ O; [7732-18-5]		Scholder, R.; Staufenbiel, E. Z. anorg. allgem. Chem. <u>1941</u> , 247, 259-76			
(3) water, $n_2 0$, $[7]$, 32-10-)]				
		(
VARIABLES:			PREPARED BY:		
Concentration of Na	OH and temperature.		T. P. Dirks	se	
EXPERIMENTAL VALUES	:	Table	2 1		
Solubility	of Cd(OH), in NaOH	solutio	ons at 100.0°	с.	
	mg CdO/100 g soln			$(mo1 \ Cd0 \ dm^{-3})$ x 10 ³	solid phase
22.7 ^a _b	18.3^{a}_{b} 18.4^{b}_{c}				ĺ
22.7^{b} 30.0 ^a _b	18.4 ⁰ 37.8 _b		6.8	1.72	Cd (OH) 2
30.0_{a}^{b} 30.3_{b}^{a}	37.7		9.6	3.74	п
36.3 ^{°°} 36.3	65.9 ^a 65.6 ^b		12.1	6.84	**
39.4^{a}_{b}	02 0 ^a				
39.4° 40.4,	92.5 ^b 100.0 ^a		13.5	9.86	"
40.4 ^b	100.7		13.9	10.81	**
42.3^{a}_{b} 42.3^{a}_{b}	89.2 ^a 89.5 ^b		14.8	9.73	Na, [Cd(OH)]]
46.3 ⁶ 46.3	51.2 ^a 51.5 ^b		16.6	5.75	2 7
42.3 ^b 46.3 ^a 46.3 ^b 50.0 ^a 50.0 ^b	25.2 ^a 26.5 ^b		18.4	3.03	
					nat temperature.
<u> </u>	AUX	ILIARY	INFORMATION		
METHOD/APPARATUS/PI	ROCEDURE :		SOURCE AND PI	URITY OF MATERIALS	5:
in a silver vessel In the solid phase, dissolving the solid ing the Cd by treat thiocyanate (1). I was determined by a CdS, changing it to	added to the NaOH sol and refluxed for $1-2$, Cd was determined b d in H_2SO_4 , and dete ment with dipyridine in the liquid phase C adding H_2SO_4 , precipi o CdSO ₄ and weighing. Ined by titration. I use was identified	days. y rmin- d ²⁺ tating NaOH	of cadmium a KOH solution	prepared by addin acetate to boiling n. Presumably rea ere used but this y stated.	g concentrated agent grade
			controlled to values appea about 3-5%. REFERENCES: 1. Spacu,	ROR: stated. The tempo to ± 0.1°C. The s ar to have an unce G.; Dick, I. Z. a 73, 279.	solubility ertainty of

COMPONENTS:	ORIGINAL MEASUREMENTS:
(1) Cadmium hydroxide; Cd(OH) ₂ ; [21041-95-2]	Akselrud, N. V.; Fialkov, Ya. A. Ukr. Khim. Zh. <u>1950</u> , 16, 283-95.
(2) Cadmium sulfate; CdSO ₄ ; [10124-36-4]	<u> </u>
(3) Water; H ₂ 0; [7732-18-5]	
(), mater, m ₂ 0, [//32 10 3]	
VARIABLES: Concentration of CdSO ₄ at 18.0°C	PREPARED BY: T. Michalowski
	1. MICHAIOWSKI
EXPERIMENTAL VALUES:	
Values of -log K _s o for Cd(OH) ₂ at	= 18.0°C.
mol $Cd^{2+} dm^{-3}$	
in liquid phase pH	a -log K _S o
0	14.5820 ^b
0.1020 7.42	
0.1382 7.37	
0.3583 7.19	
0.5115 7.05	
1.1298 6.54 1.4411 6.28	15.1198 15.5338
 ^a -log K_so = -2log K_w -log [Cd²⁺] - 2_F ^b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. 	
b this value was obtained by graphical	
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0.</pre>	extrapolation of the
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN AUXILIAN</pre>	extrapolation of the
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN METHOD/APPARATUS/PROCEDURE:</pre>	extrapolation of the NY INFORMATION SOURCE AND PURITY OF MATERIALS:
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN</pre>	extrapolation of the Y INFORMATION SOURCE AND PURITY OF MATERIALS: he CdSO ₄ was recrystallized twice from water. The Cd(OH) ₂ was precipitated from a CdSO ₄ solution and then washed with hot water.
^b this value was obtained by graphical corresponding curve to [Cd ²⁺] = 0. AUXILIAN METHOD/APPARATUS/PROCEDURE: Cd(OH) ₂ was added to solutions of CdSO ₄ . T mixtures were shaken in a thermostat until equilibrium was attained. There is no	extrapolation of the Y INFORMATION SOURCE AND PURITY OF MATERIALS: he CdSO ₄ was recrystallized twice from water. The Cd(OH) ₂ was precipitated from a CdSO ₄ solution and then washed with hot water.
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN METHOD/APPARATUS/PROCEDURE: Cd(OH)₂was added to solutions of CdSO₄. T mixtures were shaken in a thermostat until equilibrium was attained. There is no</pre>	extrapolation of the Y INFORMATION SOURCE AND PURITY OF MATERIALS: he CdSO ₄ was recrystallized twice from water. The Cd(OH) ₂ was precipitated from a CdSO ₄ solution and then washed with hot water.
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN METHOD/APPARATUS/PROCEDURE: Cd(OH)₂was added to solutions of CdSO₄. T mixtures were shaken in a thermostat until equilibrium was attained. There is no</pre>	extrapolation of the Y INFORMATION SOURCE AND PURITY OF MATERIALS: he CdSO, was recrystallized twice from water. The Cd(OH) ₂ was precipitated from a CdSO ₄ solution and then washed with hot water. s.
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN METHOD/APPARATUS/PROCEDURE: Cd(OH)₂was added to solutions of CdSO₄. T mixtures were shaken in a thermostat until equilibrium was attained. There is no</pre>	extrapolation of the Y INFORMATION SOURCE AND PURITY OF MATERIALS: he CdSO ₄ was recrystallized twice from water. The Cd(OH) ₂ was precipitated from a CdSO ₄ solution and then washed with hot water. ESTIMATED ERROR:
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN METHOD/APPARATUS/PROCEDURE: Cd(OH)₂was added to solutions of CdSO₄. T mixtures were shaken in a thermostat until equilibrium was attained. There is no</pre>	extrapolation of the Y INFORMATION SOURCE AND PURITY OF MATERIALS: he CdSO, was recrystallized twice from water. The Cd(OH) ₂ was precipitated from a CdSO ₄ solution and then washed with hot water. s.
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN METHOD/APPARATUS/PROCEDURE: Cd(OH)₂was added to solutions of CdSO₄. T mixtures were shaken in a thermostat until equilibrium was attained. There is no</pre>	extrapolation of the Y INFORMATION SOURCE AND PURITY OF MATERIALS: the CdSO, was recrystallized twice from water. The Cd(OH), was precipitated from a CdSO, solution and then washed with hot water. S. ESTIMATED ERROR: There is insufficient information in the article to allow the possible error to be
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN METHOD/APPARATUS/PROCEDURE: Cd(OH)₂was added to solutions of CdSO₄. T mixtures were shaken in a thermostat until equilibrium was attained. There is no</pre>	extrapolation of the Y INFORMATION SOURCE AND PURITY OF MATERIALS: CdSO ₄ was recrystallized twice from water. The Cd(OH) ₂ was precipitated from a CdSO ₄ solution and then washed with hot water. ESTIMATED ERROR: There is insufficient information in the article to allow the possible error to be estimated.
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN METHOD/APPARATUS/PROCEDURE: Cd(OH)₂was added to solutions of CdSO₄. T mixtures were shaken in a thermostat until equilibrium was attained. There is no</pre>	extrapolation of the Y INFORMATION SOURCE AND PURITY OF MATERIALS: CdSO ₄ was recrystallized twice from water. The Cd(OH) ₂ was precipitated from a CdSO ₄ solution and then washed with hot water. ESTIMATED ERROR: There is insufficient information in the article to allow the possible error to be estimated.
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN METHOD/APPARATUS/PROCEDURE: Cd(OH)₂was added to solutions of CdSO₄. T mixtures were shaken in a thermostat until equilibrium was attained. There is no</pre>	extrapolation of the Y INFORMATION SOURCE AND PURITY OF MATERIALS: CdSO ₄ was recrystallized twice from water. The Cd(OH) ₂ was precipitated from a CdSO ₄ solution and then washed with hot water. ESTIMATED ERROR: There is insufficient information in the article to allow the possible error to be estimated.
<pre>b this value was obtained by graphical corresponding curve to [Cd²⁺] = 0. AUXILIAN METHOD/APPARATUS/PROCEDURE: Cd(OH)₂was added to solutions of CdSO₄. T mixtures were shaken in a thermostat until equilibrium was attained. There is no</pre>	extrapolation of the Y INFORMATION SOURCE AND PURITY OF MATERIALS: CdSO ₄ was recrystallized twice from water. The Cd(OH) ₂ was precipitated from a CdSO ₄ solution and then washed with hot water. ESTIMATED ERROR: There is insufficient information in the article to allow the possible error to be estimated.

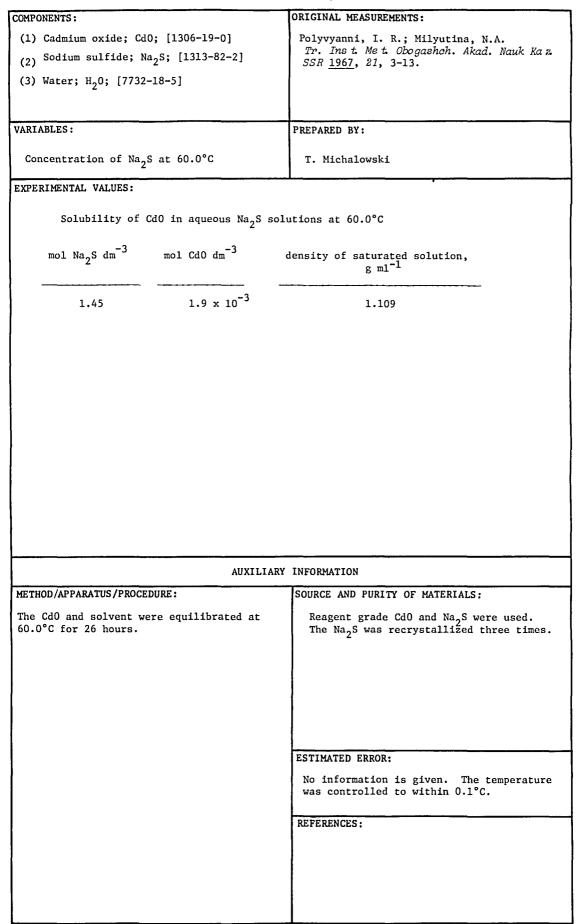
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COMPONENTS: (1) Cadmium hydroxide; Cd(OH) ₂ ; [21041-95-2]	ORIGINAL MEASUREMENTS: Schindler, P. Helv. Chim. Ac tz 1959,42,
(2) Sodium perchlorate; NaClO ₄ ; [7601-89-0]	2736-42.
(3) Water; H ₂ 0; [7732-18-5]	
VARIABLES:	PREPARED BY:
Method of reaching equilibirum at 25° C and at a constant total ionic strength of 3 mol dm ⁻³ .	T. P. Dirkse
EXPERIMENTAL VALUES: log [H ⁺] For solutions from which Cd(OH) ₂	$\frac{-\log [H^+]^2 / [Cd^{2+}]}{\text{has precipitated.}}$
8.09 ± 0.01	14.04 ± 0.02
8.29 + 0.01	14.03 ± 0.03
7.77 ±_0.01	14.03 ± 0.02
8.16 ± 0.01	14.06 ± 0.03
8.15 ± 0.01	14.04 ± 0.03
8.69 ± 0.01	14.04 ± 0.04
8.67 ± 0.01	14.00 ± 0.04
For solutions into which Cd(OH)	(s) dissolves.
7.80 ± 0.01	13.99 ± 0.02
8.56 ± 0.01	14.04 ± 0.04
7.60 ± 0.01	14.02 ± 0.02
Analysis of the solution was carried out for gradual change over the first 100 hours. Aft	about 450 hours. The results showed a cer that, the values remained constant.
The above values were converted to log K_{s}^{o} values $\log K_{w}^{o} = -14.22 \pm 0.02$ at 25°C in 3	alues using the following value. mol dm ⁻³ NaClO ₄ (1).
	INFORMATION
METHOD/APPARATUS/PROCEDURE: Two solutionsone containing Cd^{2+} and the other containing OH ⁻ were mixed to precipi- tate $Cd(OH)_2$. The hydrogen ion concentration (determined by an e.m.f. measurement) of the solution was measured over a period of about 20 days. The Cd^{2+} ion concentration was calculated from the volume and the concentra- tions of the solutions. The second method involved adding $Cd(OH)_2$ to a 3 mol dm ⁻³ aqueous solution of NaClO ₄ and again measur- ing the hydrogen ion concentration over a period of time.	
	ESTIMATED ERROR:
	The calculated solubility product (concen- tration product) has an uncertainty of about 10%.
	REFERENCES:
	1. Ingri, H.; Lagerstrom, G.; Fryaman, M.; Sillen, L. G. Ac ta Chem. S cand. <u>1957</u> , 11, 1034.
	L

COMPONEN	IS:			ORIGINAL M	ASUREMENT	S:	
(1) Cadmium hydroxide; Cd(OH) ₂ ; [21041-95-2]			Ladeinova, L. V.; Lozhkina, L. G.;				
(2) Hydrogen peroxide; H ₂ O ₂ ; [7722-84-1]			neva, A. M. n. Nauk 19	. I zv. Akad. Na 61, 12-16.	uk SSR ,		
(3) Water; H ₂ 0; [7732-18-5]							
(3) Wat	er; H ₂ 0;	[7732-18-5]					
VARIABLE			- 1	PREPARED B	Y:		
tempera		e concentration a	and	T. P. Dir	kse		
EXPERIME	NTAL VALU	ES:					
		Data on the Cd(O		le 1	۰° <i>C</i>		
		Data on the Cd(OH	¹ ² ⁻ⁿ ² ⁰ ² ⁻ⁿ ² ⁰	system at t			
		sition of hase in wt %		-	sition of phase in v	at %	
	TTINTO P		aa		P.1.000 211 1	73	a a
active 0 ₂	H ₂ O ₂	(CdO) x 10 ³	Solid ^a phase	active 0 ₂	H ₂ 0 ₂	(CdO) x 10 ³	solid ^a phase
				<u> </u>	<u> </u>		
		1.6	A	21.18	45.03	7.03	C + D
0.81 2.54	1.72 5.40	4.19 7.83	" A + B	22.14 22.85	47.0 48.60	5.56 1.87	D ''
4.44	9.44	5.29	B	24.27	51.60	1.86	,,
5.42	11.52	5.63	**	26.43	56.19	1.76	11
5.54	11.78 15.60	5.52 5.07	11	27.44 27.47	58.34 58.40	5.16 4.85	D + E E
9.35	19.88	4.23	**	28.10	59.74	4.43	15
10.06	21.73	5.96	**	28.53	60.65	4.41	"
11.21	23.83	7.28	B + C	32.41	68.90	1.43	11
12.59	26.76 28.50	4.52 4.85	C	35.66 40.80	75.81 86.74	0.99 1.04	11
15.45	32.85	4.40		44.17		2.10	11
18.31	38.93	5.18	**				
$a_{A} = C$	1(OH) : B	= Cd0 ₂ ·2H ₂ 0; C =	= CdO. •1.5H.C	D = Cd0	•H 0• E = 1	Cd0 •0.5H 0	
	2, 2, 2	³⁴⁰ 2 ² ² 2 ³ , ³	0002 110020	, 2 0002	¹¹ 2 ⁰ , ¹	2 0.5.20	
		· · · · · · · · · · · · · · · · · · ·	AUXILIARY	INFORMATIO	N		
METHOD //	PPARATIIS	PROCEDURE :		SOURCE AND	PURTTY OF	MATERIALS:	
		were mixed and st	irred until	1		(OH), and disti	11ed H 0
	-	reached. Equili		without s	stabilizers	s were used.	1100 1202
1	-	onstancy of activ					
		h solid and liqu: reached in 1.5 –					
Active	oxygen co	ontent was determ	ined by				
		0.1 N KMn04 in th					
imetric	ally with	content was deter n dithizone in th	mined color-	se			
		ally by weighing		ľ			
pyropho	osphate in	n the solid phase	•				
				ESTIMATED	ERROR:		
						s controlled to ation is given	
						the results.	
				REFERENCES	S:		
	· · · .		· · · · · · · · · · · · · · · · · · ·	<u>, A</u>			

292	Cadmium Oxide and Hydroxide				
COMPONENTS: (1) Cadmium h (2) Hydrogen (3) Water; H ₂	peroxide; H	d(OH) ₂ ; [21041-9 2 ⁰ 2; [7722-84-1 -5]	ORIGINAL MEASUREMENTS: Ladeinova, L. V.; Lozhkina, L. G.; Chernysheva, A. M. I w. Akad. Nauk SSR, O td Khim. Nauk <u>1961</u> , 12-16.		
EXPERIMENTAL	RESULTS, c	ontinued			
			Table		
	Data	on the Cd(OH) ₂ -H	2 ⁰ 2 ^{-H} 2	0 system at 20°C.	
		ition of ase as wt %			
active 02	H202 ^a	(CdO) x 10 ³	So:	lid phase ^b	
$\begin{array}{c} 3.55\\ 5.62\\ 6.10\\ 6.32\\ 6.80\\ 8.47\\ 9.61\\ 10.97\\ 12.27\\ 13.20\\ 13.52\\ 13.37\\ 13.98\\ 14.82\\ 15.56\\ 16.47\\ 18.12\\ 18.64\\ 21.38\\ 21.74\\ 23.43\\ 24.23\\ 25.08\\ 25.50\\ 25.83\\ 26.79\\ 27.98\\ 31.15\\ 34.21\\ 35.18\\ 37.41\\ 40.48\\ \end{array}$	7.55 11.95 12.97 13.44 14.47 18.01 20.43 23.32 26.08 28.74 28.42 29.72 31.51 33.08 35.02 38.52 39.63 45.45 46.22 49.81 51.51 53.32 54.21 54.92 56.95 59.48 66.22 72.73 74.80 79.53 86.06	1.78 2.91 3.66 3.58 1.60 2.88 0.26 2.80 3.33 3.41 3.11 2.84 3.00 2.85 2.83 2.20 1.07 0.63 1.03 2.16 2.42 2.61 2.73 3.34 3.24 2.92 2.18 0.93 0.96 2.56 1.09 0.69 0.69 0.8		$ \begin{array}{c} A \\ H \\ B \\ B \\ H \\ B \\ H \\ C \\ C \\ H $	
41.91 ^a recalculated	89.10 by compile	1.58 r	; D=Cd	" 0 ₂ ·H ₂ 0; E=Cd0 ₂ ·0.5H ₂ 0.	



COMPONENTS:	ORIGINAL MEASUREMENTS:
(1) Cadmium oxide; CdO; [1306-19-0]	Danilov, V. V.; Martinson, I.G.; Ravdel, A. A. Zh. Prik ad Khim. (Leningrad)
(2) Ammonia; NH ₃ ; [7664-41-7]	<u>1971</u> , 44, 3-6.
(3) Water; H ₂ 0; [7732-18-5]	
VARIABLES:	PREPARED BY:
Temperature and concentration of ammonia.	T. P. Dirkse
EXPERIMENTAL VALUES:	
Solubility of CdO in aqueous ammon	la solutions.
2	(mol Cd dm ⁻³) x 10^2
mol NH ₃ dm ⁻³	<u>10°C 20°C 30°C</u>
2.18	0.65 0.68 0.70
3.40	1.52 1.59 1.70
4.40	2.61 2.70 2.77
7.10	5.22 4.94 4.46
9.45	5.76 5.54 5.06
11.45	6.20 6.02 5.42
AUXILIARY	INFORMATION
METHOD/APPARATUS/PROCEDURE:	SOURCE AND PURITY OF MATERIALS:
Solutions were equilibrated for 30 days with intermittent stirring. Equilibrium was approached from both high and low tempera- tures. Cadmium content was determined by titration with Trilon B at a pH of about 10, using Eriochrome Black T as indicator.	
	ESTIMATED ERROR: No information is given as to how closely the temperature was controlled, nor to how reproducible the analyses were.
	REFERENCES:

	ORTOTIVAL AND AND THE
COMPONENTS:	ORIGINAL MEASUREMENTS:
(1) Cadmium oxide; CdO; [1306-19-0]	Kudryaytsev, N. T.; Selivanova, G. A. Izv. Vyssh, Ucheb. Zaved., Khim. Tekhnol. <u>1971</u> ,
(2) Ammonium sulfate; (NH ₄) ₂ SO ₄ ; [7783-20-2]	14, 835-8.
(3) Water; H ₂ O; [7732-18-5]	
(3) "accr, "20, (7) 22 10 3]	
VARIABLES:	PREPARED BY:
Concentration of ammonium sulfate.	T. Michalowski
EXPERIMENTAL VALUES:	
Solubility of CdO in aqueo	us (NH,),SO, at 25°C.
	. – .
$c_{(NH_4)_2} s_{0_4}^{mo1 dm^{-3}} c_{cd}^{mo}$	1 dm ⁻⁵ pH
0 9 x 1 0.5 0.2	
1.0 0.6	5 9.5
1.5 0.9	0 8.9
2.0 1.4 2.5 1.8	
3.0 2.0	
or NH ₄ OH). In the 1 mol dm ⁻³ soluti mol dm ⁻³ at a pH = 7-10. In the 2 m mol dm ⁻³ at a pH = 6-7. The solid phase in equilibrium with is described as a compound of the ty Using the water solubility value in is obtained for the value of $K_{s}o = C$	the solutions in the above Table pe $CdSO_4 \cdot nCd(OH)_2 \cdot mNH_3 \cdot xH_2O$. the above Tąble, a value of 2.26 x 10^{-14}
AUXILIARY	INFORMATION
METHOD/APPARATUS/PROCEDURE:	SOURCE AND PURITY OF MATERIALS:
No information is given about the preparation and mixing of the solutions. Cadmium content was determined by a compleximetric titration or photometrically using dithizone. The pH of the solutions was measured by using a glass electrode.	
	FORTMATED EDDOD.
	ESTIMATED ERROR:
	No information is given.
	REFERENCES:
	<u> </u>

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COMPONENTS:		ORIGINAL MEASUREMENTS:		
 (1) Cadmium oxide; Cd0; [1306-19-0] (2) Sodium hydroxide; NaOH; [1310-73-2] (3) Water; H₂0; [7732-18-5] 		Soloveva, V. D.; Svirchevskaya, E. G.; Bobrova, V. V.; Eltsov, N. M. Tr. Inst. Metal. Obogashch. AN KazSSR, <u>1973</u> , 49, 37-44.		
VARIABLES:		PREPARED BY:		
NaOH concentration and	temperature	T. Michalowski		
EXPERIMENTAL VALUES:		· · · · · · · · · · · · · · · · · · ·		
Solubility of	CdO in NaOH solutions.			
Temperature, <u>°C</u>	mol NaOH dm ⁻³	Cd solubility, (mol dm ⁻³) x 10^5	Solid phase	
20 20	2.75 5.10	4.45	Cd (0H) 2	
20	10.25	8.9 162.0	11	
20	13.08	177.0	11	
40	3.0	15.0	Cd (0H) 2	
40	5.65	57.0	"2	
40	8.85	131.0	11	
40	10.84	320.0	11	
105	3.5	71.0	Cd (0H) 2	
105 105	7.5	177.0	n 2 n	
105	10.0 12.0	500.0 970.0	**	
200	2.75	134.0	Cd (OH) 2	
200	5.0	223.0		
200 200	7.5 10.0	339.0 900.0	"	
200	12.0	1070.0		
200	14.0	1340.0	11	
200 200	16.5 18.3	1080.0 _366.0	Na2Cd (OH)4	
	AUXILIARY	INFORMATION		
METHOD/APPARATUS/PROCE	DURE:	SOURCE AND PURITY OF MAT	TERIALS:	
Apparently the mixture at the temperature rec description of the pro	orded, but no explicit	Analytical grade CdO wa is said about the other were used.		
		ESTIMATED ERROR:		
		No information is giver		
		REFERENCES :		
L				

COMPONENTS:		ORIGINAL MEASUREMENTS:		
(1) Cadmium hydroxide; Cd(OH)	; [21041-95-2]	Soloveva, V. D.; Bobrova, V. V.; Orld	va	
(=) =====,===,===; ==(;==;2]	, [22012)5 2]	L. F.; Adeyschvili, E. U. Tr. Inst.	va,	
2) Sodium hydroxide; NaOH; [1310-73-2]		Met. Obogashch. AN. Kaz SSR <u>1973</u> , 49,		
		45-8.	-	
(3) Water; H ₂ 0; [7732-18-5]				
ARIABLES:	<u></u>	PREPARED BY:		
NaOH concentration and temperat	ture.	T. Michalowski		
XPERIMENTAL VALUES:				
Solubility of Cd(OH) ₂ at	different tem	eratures		
(Concentration of	f cadmium in the solution,		
		dm^{-3} x 10 ⁵		
mol NaOH dm ⁻³	20°	105° 200°		
······································				
3.75	4.46	44.6 44.5		
5.62	9.00	71.4 125.0		
7.50 10.00	71.4 90.0	178.0 250.0		
11.50	180.0	357.0 543.0 455.0 580.0		
13.35	340.0	803.0 759.0		
14.00	338.0	770.0 1160.0		
15.65		455.0 990.0		
16.85		277.0 786.0		
18.75		149.0 290.0		
	AUXILIARY	INFORMATION		
ETHOD / APPARATUS / PROCEDURE :		SOURCE AND PURITY OF MATERIALS:		
Apparentlythe mixtures were equ	dilibrated at			
the temperature recorded but no description of the procedure is	explicit	Cd(OH), was precipitated by the addi of a NãOH solution to a solution of The precipitate was washed and dried	CdSO,.	
	8-10-11	its purity was checked by X-ray diff		
		diagrams.		
		ESTIMATED ERROR:		
		No information is given.		
		DEDEDENOVO.		
		REFERENCES:		

Cadmium Oxide and Hydroxide

COMDONENTS .	· · · · · · · · · · · · · · · · · · ·		OPTOTNAL MEASUPENE	NTC .		
COMPONENTS: (1) Cadmium oxide; CdO; [1306-19-0]				ORIGINAL MEASUREMENTS: Omezzine, B. K.; Ariguib-Kbir, N. Compt.		
			rend. <u>1978</u> , 2			
(2) Arsenic(V) oxide; As ₂ 0 ₅ ; [1303-28-2]						
(3) Water; H ₂ 0;	[7732-18-5]					
	••••••					
VARIABLES:			PREPARED BY:			
Solution compo	osition at 25°C.		T. P. Dirkse			
EXPERIMENTAL VAL Solubilit	UES: y of CdO in aqueo	ous solution	s of As ₂ 0 ₅ at 25°C.	a		
^{As20} 5,	^{As} 2 ⁰ 5	CdO,	CdO,			
··· -	mol/kg H ₂ 0 ^b			Solid ^C		
mass %			mol/kg H ₂ 0 ^{,b}	phase		
0.08	0.0035	trace				
3.85	0.17	0.40	0.033	Α		
7.85	0.37	0.75	0.064			
11.80	0.59	1.70	0.15			
11.80 11.85	0.59 0.60	1.72 1.75	0.15 0.16	A + B A + B		
11.85	0.59	1.70	0.15	A + B		
14.22	0.74	2.20	0.21	В		
17.75	0.97	2.42	0.24	B		
22.00	1.28	3.45	0.36			
22.20	1.30	3.35	0.35			
22.15	1.30	3.42	0.36	B + C		
22.10	1.29	3.39	0.35	B + C		
22.10	1.29	3.30	0.34	B + C		
28.40	1.81	3.40	0.39			
32.60	2.22	3.60	0.44			
39.80	3.06	3.65	0.50	<u>_</u>		
46.20	4.01	3.70	0.58	С		
53.20	5.26	2.80	0.50			
59.80 65.40	6.81 8.68	2.00 1.82	0.41 0.43			
65.52	8.74	1.85	0.44	C + D		
65.50	8.74	1.87	0.45	C + D		
66.20	8.84	1.20	0.29			
		AUXILIARY	INFORMATION			
METHOD/APPARATUS			SOURCE AND PURITY			
	placed in polyeth		s No information :	is given.		
	a thermostat. Ed	•				
	l by analysis. An l iodometrically,	senic conte	114			
cadmium conter						
cadmium conter pleximetric ti	tration. The con	position of				
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COMPONENTS:

(1) Cadmium oxide; CdO; [1306-19-0]

ORIGINAL MEASUREMENTS:

Omezzine, B. K.; Ariguib-Kbir, N. Compt. rend. <u>1978</u>, 286, 197-9.

- (2) Arsenic(V) oxide; As₂0₅; [1303-28-2]
- (3) Water; H₂0; [7732-18-5]

EXPERIMENTAL VALUES, CONTD:

Solubility of CdO in aqueous solutions of $As_20_5^{0}$ at 25°C. ^a

As2 ⁰ 5,	^{As} 2 ⁰ 5	CdO,	CdO,	· · · C
mass %	mol/kg H ₂ 0 ^b	mass %	mol/kg H ₂ 0 ^b	Solid ^C phase
····				_
68.00	9.48	0.80	0.20	D
70.20	10.25	trace		
71.60	10.97	trace		

^a The article contains only a phase diagram. These numerical data were given in a personal communication of Sept. 11, 1982 from Prof. N. Kbir-Arguib.

^b calculated by the compiler.

^c A = $5Cd0 \cdot 2As_20_5 \cdot 5H_20$; B = $2Cd0 \cdot As_20_5 \cdot 2H_20$; C = $Cd0 \cdot As_20_5 \cdot 4H_20$; D = $Cd0 \cdot 2As_20_5 \cdot 5H_20$.