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TABLE I

FIRST-ORDER REACTION RATE CONSTANTS FOR AQUATION; $ML + H_2O \rightarrow MH_2O + L$

Metal Complex	k_{1-1} (sec ⁻¹)	μ (M)	T (°C)	pH	Reference
$Mn(CN)_6^{-3}$	4.4×10^{-4}	---	8	9-8	1
$Fe(H_2O)_5NCS^+$	8.8×10^{-1}	0.4	14.5	1.5	2
$Fe(H_2O)_5SO_3^+$	1.1	1.0	8.4	1.7	3
$Cr(OX)_2(AC)H_2O^{-2}$	6.2×10^{-4}	0.1	16.3	2-5	4
$Cr(OX)_2(AC)OH^{-3}$	2.7×10^{-2}	0.1	8.4	9-12	4
$Co(en)_2(H_2O)Cl^{+2}$	1.6×10^{-6}	---	---	2-3	5
$Co(en)_2(OH)Cl^+$	1.3×10^{-2}	---	---	7-9	6
$Mn(H_2O)_6^{+2}$	3.1×10^7	0.1	25	1	7
$Fe(H_2O)_6^{+2}$	3.2×10^6	0.1	25	1	7
$Ni(H_2O)_6^{+2}$	3.0×10^4	0.1	25	1	7
$Cu(H_2O)_6^{+2}$	8.0×10^9	0.1	25	1	8
$Fe(H_2O)_6^{+3}$	3.0×10^3	0.1	25	1	8
$Co(cyclam)(OH)Cl^+$	1.2×10^{-2}	0.1	18.8	7-12	9
$CaEDTA^{-2}$	1.3	0.5	25	6-8	10
$Ca(PDTA)^{-2}$	6.6×10^{-2}	0.5	25	7-8	10
$Ni(dipy)^{+2}$	5.3×10^{-5}	---	25	7	11
$Ni(AC)^+$	1.0×10^3	1.0	20	5	12
$CO_2(aq) + H_2O \longrightarrow$					
$HCO_3^- + H^+$	2.6×10^{-2}	0.1	25	8.7	13
$H^+ + FeS + 2.25 O_2 \longrightarrow$					
$Fe^{+3} + SO_4^{=2-} + H_2O$	$3.0 \times 10^{-3}^*$.001	20	7.0	14

*pseudo first-order constant

TABLE II
SECOND-ORDER REACTION RATE CONSTANTS FOR REDOX PROCESSES/ELECTRON TRANSFERS

REACTION	k_2 (M ⁻¹ sec ⁻¹)	μ (M)	T (°C)	pH	Reference
Fe ⁺² + Mn ⁺³	1.7x10 ⁴	3.1	25	0	15
MnTPP ⁻³ + HO-quinoline	7.9x10 ⁵	0.1	16	8	16
FeEDTA ⁻² + ClO ₂ ⁻	3.0x10 ⁴	1.0	20	9	17
Fe(phen) ₃ ⁺² + ClO ₂ ⁻	3.0x10 ⁵	1.0	20	9	17
Fe(CN) ₆ ⁻³ + ascorbate	1.3x10 ²	0.24	25	2.0	18
Fe(CN) ₆ ⁻³ p-NH ₂ -φ-NH ₂	4.0x10 ¹	0.00	30	7.5	19
MnNTA ⁻ + HO-quinoline	1.9x10 ⁷	0.1	16	8	16
MnATP ⁻² + HO-quinoline	1.0x10 ⁶	0.1	16	8	16
Fe(phen) ₃ ⁺² + MnO ₄ ⁻	6.0x10 ³	0.45	25	1	20
Fe ⁺³ + penicillamine	4.5x10 ⁻¹	0.1	20	9	21
Fe(phen) ₃ ⁺² + HOCl	2.2x10 ⁻²	1.0	25	7.2	22
Fe ⁺³ + ascorbate	30	1.0	0	1	23
Fe ⁺³ + I ⁻	16	0.2	25	1	24
Fe(phen) ₃ ⁺³ + S ₂ O ₅ ⁻²	7.2x10 ³	0.5	25	1	25
Fe(phen) ₃ ⁺³ + SO ₃ ⁻²	4.2	0.5	25	1	25
Cytochrome C + FeEDTA ⁻²	2.6x10 ⁴	0.1	25	6-9	26
Fe ⁺³ + HS-succinic acid	1.0	1.0	25	1	27
Mn ⁺³ + 1,2-(pH) ₂ -benzene	6.5x10 ³	3.0	25	0	28
Cu ⁺² + ascorbate	3.1x10 ⁴	0.1	30	4-6	29
Mn ⁺³ + thiourea	2.6x10 ⁴	---	25	0	30
Cu(DMP) ₂ ⁺² + H ₂ O ₂	1.2x10 ⁶	0.1	25	6-9	31
Fe ⁺² + benzoquinone	9.1x10 ⁻⁴	0.5	25	0	32
Cytochrome C + Fe(CN) ₆ ⁻⁴	1.3x10 ⁴	0.2	12	7	33

TABLE III
SECOND-ORDER REACTION RATE CONSTANTS FOR COMPLEXATION

REACTION	k_2 (M ⁻¹ sec ⁻¹)	μ (M)	T (°C)	pH	Reference
Fe(OH) ⁺ + chloride	1.1×10^4	1.0	25	1.2	34
Fe(OH) ⁺ + thiocyanate	1.1×10^4	0.4	25	1.5	34
Fe(OH) ⁺ + acetate	2.8×10^3	0.5	25	1.5	35
Fe(OH) ⁺² + H ₂ -salicylate	5.5×10^3	1.0	25	2.0	36
Fe(OH) ⁺² + H-salicylate	1.4×10^4	1.0	25	2.0	36
Fe ⁺³ + oxalate	1.4×10^2	0.5	25	0.5	37
Fe ⁺³ + sulfate	6.4×10^3	0.5	25	1.3	38
Fe ⁺³ + sulphosalicylate	1.2×10^4	1.0	25	2	39
Fe ⁺³ + salicylate	1.4×10^4	1.0	25	2	39
Fe(OH) ⁺ + HIDA ⁻	8.8×10^3	1.0	25	2	40
Fe(OH) ⁺ + H ₂ NTA ⁻	5.6×10^4	1.0	25	2	40
Fe(OH) ⁺ + H ₃ EDTA ⁻	1.1×10^5	1.0	25	2	41
Fe ⁺² + sulfate	1.0×10^6	0.1	20	1	42
Fe ⁺² + phen	7.9×10^5	1.0	25	1	43
Fe(dipy) ₂ ⁺² + dipy	4.0×10^5		25		44
Mn ⁺² + NTA ⁻³	5.0×10^8		25		45
Mn ⁺² + HNTA ⁻²	2.0×10^5		25		45
Mn ⁺² + sulfate	4.0×10^6	0.1	20	1	42
Mn ⁺² + chloride	1.6×10^7	0.1	20	1	46
Cu ⁺² + H ₂ ATP ⁻²	8.8×10^8	0.1	25	1-5	47
Cu ⁺² + serinate	1.8×10^9	0.1	25	1-5	48
Co ⁺² + α -alanine	1.3×10^9	0.1	20	9	49
Co ⁺² + histidine	1.3×10^7	0.1	20	9	49

TABLE IV
SECOND-ORDER REACTION RATE CONSTANTS FOR AUTOXIDATIONS

REACTION	k_2 ($M^{-1} sec^{-1}$)	T (°C)	pH	Reference
$H^+ + Fe^{+2} + 1/4 O_2 \rightarrow Fe^{+3} + 1/2 H_2O$	$9.6 \times 10^{-1}^*$	20	7	50
$Mn^{+2} + H_2O + 1/2 O_2 \rightarrow MnO_2 + 2H^+$	$9.1 \times 10^{-2}^*$	25	9	51
$Fe^{+2} + O_2$	2.0×10^{-4}	25	5	52
$Fe^{+2} + O_2$	7.9×10^{-4}	30	2-3	53
$(M^{-2} atm^{-1} sec^{-1})$				
$Fe^{+2} + O_2$	7.1×10^{-1}	30	2-3	54
$(M^{-2} sec^{-1})$				
$Co(hist)_2^{-2} + O_2$	3.5×10^3	25	8-11	55
$Co(glygly)_2(OH)_2^{-2} + O_2$	1.0×10^3	25	8-12	56
$Cu(dipy)_2^+ + O_2$	6.5×10^3	25	5	57
$Fe(HTPP)_2^{-2}$	4.0×10^1	25	4.5	58
$Fe(cyst) + O_2$	5.0×10^3	25	6	59
$Fe(cyst)_2^{-2} + O_2$	2.0×10^4	25	9.7	59
$Co(trien)(H_2O)_2^{+2}$	2.5×10^4	25	7-12	60
$Co(trien)(OH)(H_2O)^+ + O_2$	2.8×10^5	25	7-12	60

* Calculated from data presented in reference.

TABLE V
SECOND-ORDER REACTION RATE CONSTANTS FOR PROTON TRANSFER

REACTION	k_2 (M ⁻¹ sec ⁻¹)	μ (M)	T (°C)	pH	Reference
H ₃ O ⁺ + OH ⁻	1.4x10 ¹¹	0	25	15.75	61
NH ₄ ⁺ + OH ⁻	3.4x10 ¹⁰	0	20	9.25	61
imidazole ⁺ + OH ⁻	1.3x10 ¹⁰	0	25	6.95	61
C ₆ H ₅ OH + OH ⁻	1.3x10 ¹⁰	0	25	4.20	61
H ₃ O ⁺ + HS ⁻	7.5x10 ¹⁰	0	25	7.24	61
H ₃ O ⁺ + NH ₃	4.0x10 ¹⁰	0	25	9.25	61
H ₃ O ⁺ + HCO ₃ ⁻	5.0x10 ¹⁰	0	25	3.77	61
H ₃ O ⁺ + CH ₃ CO ₂ ⁻	3.2x10 ¹⁰	0	25	4.20	61
CH ₃ CO ₂ H + formate	5.0x10 ⁸	1	20	3.75	62
CH ₃ CO ₂ H + propionate	4.0x10 ⁸	1	20	4.87	62
CH ₃ CO ₂ H + malonate	5.6x10 ⁸	1	20	5.70	62
CH ₃ CO ₂ H + imidazole	1.3x10 ⁹	1	20	7.0	62
CH ₃ CO ₂ H + HPO ₄ ⁻	4.8x10 ⁹	1	20	7.2	62
HCO ₃ ⁻ + H ₃ O ⁺ → CO ₂ (H ₂ O)	5.6x10 ⁴	0	25	6.35	62

TABLE VI
SECOND-ORDER RATE CONSTANTS FOR Ni(II) COMPLEXATION

LIGAND	$k_2 (M^{-1} sec^{-1})$	log K	$\mu (M)$	T (°C)	pH	Reference
histidine	2.2×10^3	8.7	0.1	24	5-7	63
phenanthroline	4.1×10^3	8.6	0.1	25	6-8	64
dipyridyl	1.6×10^3	7.0	---	---	---	65
salicylicate	5.3×10^3	7.0	0.1			65
sulfosalicylate	4.7×10^3	6.4	0.1			66
proline	3.4×10^4	5.9	0.1	25	6-8	67
α -alanine	2.0×10^4	5.4	0.1	20	9	68
ethylmalonate	7.6×10^3	3.2	0.1	25	3-5	69
HEDTA ⁻³	2.0×10^5	3.2	0.1	25		70
phthalate	2.6×10^5	2.2	0.1	25	4-5	69
lactate	2.6×10^4	1.6	0	25	5.8	71
acetate	2.7×10^4	1.5	0	25	3-7	72
sulfate	1.6×10^4	1.0		25		42
oxalate	7.9×10^4	52	0.1	25	7	73
imidazole	3.2×10^3	3.0	0.1	24	6	74
pyridine	4.6×10^3	1.9	---	25	7	75
thiocyanate	5.0×10^3	1.2	0.1	25	7	76
methionine	$\sim 1 \times 10^{-3}$	5.2	0.15	24	6-7	77
glutathione	$\sim 1 \times 10^{-3}$		0.15	24	6-7	77
m-tyrosine	1.2×10^{-1}	5.1	0.15	24	6-7	77
cysteine	8.5×10^2	9.8	0.15	24	6-7	77

TABLE VII
KINETIC DATA FOR AUTOXIDATION OF SULFIDE; COMPARISON OF RESULTS

#	REACTION	WATER	pH	$t_{1/2}(\text{min})^{**}$	T(°C)	$[\text{HS}^-]^n$	$[\text{O}_2]^m$	pH Range	R
1	$\text{HS}^- + \text{O}_2$	sea water	8.0	280	23.0	1	---	8-8.5	78
2	$\text{HS}^- + \text{O}_2$	distilled [†]	7.9	3,000	25.0	1.34	0.56	6-12	79
3	$\text{HS}^- + \text{O}_2$	sea water	8.2	24	25.0	1	1	---	80
4	$\text{HS}^- + \text{O}_2$	sea water	7.8	175	9.8	1	---	---	81
5	$\text{HS}^- + \text{O}_2$	fresh ^{††}	7.6	880	25.0	1	1	4-10	82
6	$\text{HS}^- + \text{O}_2$	distilled	11.0	130	---	1	---	11-14	83
7	$\text{HS}^- + \text{O}_2$	fresh	8.6	2,200	25.0	1	1	7-8.6	84
8	$\text{HS}^- + \text{O}_2$	fresh	8.0	114	25.0	1	---	6-10	85
9	$\text{HS}^- + \text{O}_2$	sea water	7.7	600	9.0	---	---	7-8	86
10	$\text{HS}^- + \text{O}_2$	distilled [†]	8.3	816	25.0	1	1	5-12	87
11	$\text{HS}^- + \text{O}_2 + \text{CoTSP}^*$	distilled	8.3	7	25.0	1	1	5-12	88

** Calculated from reported data

† EDTA added

†† simulated

* CoTSP = Co(II)-4,4',4'',4'''-tetrasulphophthalocyanine, $[\text{CoTSP}] = 2 \times 10^{-9} \text{ M}$

TABLE VIII: KINETIC AND EQUILIBRIUM DATA FOR Fe(II) OXIDATIONS

#	OXIDANT	$k_2 (M^{-1} sec^{-1})^a$	ΔG^\ddagger b	ΔG° b	$\mu (M)$	T (°C)	R
1	Fe ⁺³	4.0	69.7	0	0.5	25	89
2	Fe(EDTA) ⁻	4.0×10^{-4}	92.5	61.3	0.5	25	90
3	FeCl ⁺²	3.73×10^1	64.1	- 2.5	0.5	25	89
4	Fe(terpy) ₃ ⁺³	8.5×10^4	44.9	-18.3	0.5	25	91
5	Fe(dipy) ₃ ⁺³	2.7×10^4	47.7	-22.2	0.5	25	91
6	Fe(5-nitrophen) ₃ ⁺³	1.1×10^5	38.5	-49.2	0.5	25	91
7	Fe(5-chlorophen) ₃ ⁺³	2.1×10^5	42.7	-36.7	0.5	25	91
8	Fe(2,5-dimethylphen) ₃ ⁺³	7.8×10^3	50.8	-22.2	0.5	25	91
9	Fe(tetramethylphen) ₃ ⁺³	2.3×10^2	59.5	- 6.7	0.5	25	91
10	benzosemiquinone ^c	7.19×10^{-1}	73.9	52.3	0.5	25	92
11	tolusemiquinone ^c	2.98×10^{-1}	76.1	62.3	0.5	25	92
12	durosemiquinone ^c	2.66×10^{-4}	93.6	85.8	0.5	25	92
13	2,6-dichlorosemiquinone ^c	3.58	70.0	38.9	0.5	25	92
14	FeSO ₄ ⁺	6.77×10^2	56.9	6.5	0.5	25	93

a. $[H^+] = 1.0 M$ b. ΔG in kJ/molec. Calculated from K and k_f (Marcus, R. A., J. Chem. Phys., 26, 872 (1957)).

References

1. Adamson, A. W., Welker, J. P., Wright, W. B., J. Am. Chem. Soc., 73, 4786 (1951).
2. Below, J. F., Connick, R. E., Coppel, C. P., J. Am. Chem. Soc., 80, 2961 (1958).
3. Carlyle, D. W., Inorg. Chem., 10, 761 (1971).
4. Casula, M., Illuminati, G., Ortaggi, G., Inorg. Chem., 11, 1062 (1972).
5. Chan, S. C., J. Chem. Soc., 5137 (1963).
6. Pearson, R. G., Meeker, R. E., Basolo, F., J. Am. Chem. Soc., 78, 2673 (1956).
7. Swift, T. J., R. E. Connick, J. Chem. Phys., 37, 307 (1962).
8. Connick, R. L., "Symposium on Relaxation Techniques," Buffalo, N.Y. June (1965). ,
9. Poon, C. K., Tobe, M. L., J. Chem. Soc. (A) 2069 (1967).
10. Carr, J. D., Swartzfager, J. Am. Chem. Soc., 97, 315 (1975).
11. Melson, G. A., Wilkins, R. G., J. Chem. Soc., 4208 (1962).
12. Fronaeus, S., Acta Chem. Scand., 6, 1200 (1952).
13. Pinsent, B. R. W., Pearson, L., Roughton, F. J. W., Trans. Faraday Soc., 52, 512 (1956).
14. Nelson, M. B., Davis, J. A., Benjamin, M. M., Leckie, J. O., "The Role of Iron Sulfides in Controlling Trace Metals in Anaerobic Sediments," AFOSR report No. CEED0-TR-77-13, Environ. Sci. Res. Div., Tyndall AFB, FL, 398 pages (1977).
15. Sutin, N., J. Phys. Chem., 68, 174 (1964).
16. Hague, D. N., Zetter, M. S., Trans. Faraday Soc., 66, 1176 (1970).
17. Burke, R. L., Marzluff, W. L., J. Am. Chem. Soc., 91, 3481 (1969).
18. Mehrota, U.S., Agrawal, M. C., Mushran, S. P., J. Phys. Chem., 73, 1996 (1969).
19. Corbett, J. F., J. Chem. Soc. (B), 207, 213, 818 (1969).
20. Hicks, K. W., Sutter, J. R., J. Phys. Chem., 75, 1107 (1971).
21. Bell, C. M., McKenzie, E. D., Orton, J., Inorg. Chimica Acta, 5, 109 (1971).

22. Ondrus, M. G., Gordon, G., Inorg. Chem., 10, 474 (1971).
23. Lawrence, G. S., Ellis, K. J., J. C. S. Dalton 1667 (1972).
24. ibid., 2229 (1972).
25. Carlyle, D. W., J. Am. Chem. Soc., 94, 4525 (1972).
26. Hodges, H. L., Holwerda, R. A., Gray, H. B., J. Am. Chem. Soc., 96, 3132 (1974).
27. Ellis, K. J., McAuley, A., J. C. S. Dalton, 1533 (1973).
28. Mentasti, E., Pelizzetti, Pramanro, E., Giraudi, Inorg. Chim. Acta, 12, 61 (1975); Gazzetta 105, 551 (1975); Inorg. Chim. Acta, 15, L1 (1975).
29. Hayakawa, K., Minami, Nakamura, S., Bull. Chem. Soc. Japan, 46, 2788 (1973).
30. Davies, G., Inorg. Chim. Acta, 14, L13-14 (1975).
31. Davies, G., Higgins, R., Loose, D. J., Inorg. Chem., 15, 700 (1976).
32. Baxendale, J. H., Hardy, H. R., Trans. Faraday Soc., 50, 808 (1964).
33. Sutin, N., Christman, R., J. Am. Chem. Soc., 83, 1773 (1961).
34. Connick, R. E., Coppel, C. P., J. Am. Chem. Soc., 81, 6389 (1959).
35. Pandey, R. N., Smith, W. F., Can. J. Chem., 50, 194 (1972).
36. Saini, G., Mentasti, E., Inorg. Chim. Acta, 4, 585 (1970).
37. Bauer, R. F., Smith, W. F., Can. J. Chem., 43, 2763 (1965).
38. Davis, G. G., Smith, W. F., Can. J. Chem., 40, 1836 (1962).
39. Gonger, S., Stueher, J., Inorg. Chem., 13, 379 (1974).
40. Mentasti, E., Pellizzetta, E., Saini, Gazzetta, 104, 201 (1974).
41. ibid, J. C. S. Dalton, 1944 (1974).
42. Eigen, M., Tamm, K. Z., Electrochem., 66, 107 (1962).
43. Bell, R. S., Sutin, N., Inorg. Chem., 1, 359 (1962).
44. Irving, H., Mellor, D. H., J. Chem. Soc., 5222 (1962).
45. Koryta, J., Z. Electrochem., 61, 423 (1957).

46. Hayes, R. G., Myers, R. J., J. Chem. Phys., 40, 877 (1964).
47. Brundage, R. S., Karpel, R. L., Kuotin, K., Weisel, J., Biochim. Biophys. Acta, 267, 258 (1972).
48. Cobb, M. A., Hague, D. N., J. C. S. Faraday (I) 68, 932 (1972).
49. Pasternak, R. F., Justin, K., Weinstock, E. I., J. Am. Chem. Soc., 88, 4610 (1966).
50. Stumm, W., Lee, G. F., Ind. Eng. Chem., 53, 143 (1961).
51. Morgan, J. J., "Chemical Equilibria and Kinetic Properties of Manganese in Natural Waters," in Principles and Applications in Water Chemistry, S. D. Faust and J. V. Hunter, eds., Wiley, N.Y., p. 561 (1967).
52. Tamura, H., Goto, K., Nagayama, M., J. Inorg. Nucl. Chem., 38, 113 (1976).
53. Cher, M., Davidson, N., J. Am. Chem. Soc., 77, 793 (1955).
54. Utsumi, S., Murosima, K., J. Chem. Soc., Japan, Pure Chem. Sect., 86, 593 (1965).
55. Simplicio, J., Wilkins, R. G., J. Am. Chem. Soc., 84, 6092 (1967).
56. Tanford, C., Kirk, D. C., Chantoni, M. K., J. Am. Chem. Soc., 76, 5325 (1954).
57. Pecht, J., Anbar, M., J. Chem. Soc., A, 1902 (1968).
58. Kurimura, Y., Kuriyama, H., Bull. Chem. Soc. 42, 2239 (1969).
59. Gilmour, A. D., McAuley, J. Chem. Soc. (A), 1006 (1970).
60. Miller, F., Wilkins, R. G., J. Am. Chem. Soc., 92, 2687 (1970).
61. Eigen, M., Angew. Chem., 75, 489 (1963); Internat. Edt., 3, 1 (1964); Pure Appl. Chem., 6, 97 (1963).
62. Ahrens, M. L. Maass, G., Angew. Chem., Internat. Edt., 7, 818 (1968).
63. Letter, J. E., Jordan, R. B., Inorg. Chem., 10, 2692 (1971).
64. Kustin, K., Liu, S. T., J. C. S. Dalton, 278 (1973).
65. Bennetto, H. P., Caldin, E. F., Chem. Comm., 599 (1969).
66. Williams, J., Petrucci, S., J. Am. Chem. Soc., 95, 7619 (1973).
67. Wilkins, R. G., J. Am. Chem. Soc., 94, 8399 (1972).

68. Pasternak, R. F., Kustin, K., Weinstock, E. M., J. Am. Chem. Soc., 88, 4610 (1966).
69. Cavasino, F. P., Di Dio, E., Locanto, G., J. C. S. Dalton, 2419 (1973).
70. Margerum, D. W., Zabin, B. A., J. Phys. Chem., 66, 2214 (1962).
71. Harada, S., Tanabe, H., Yasunaga, T., Bull. Chem. Soc. Japan, 46, 3125 (1973).
72. Harada, S., Yasunaga, T., Tamura, K., Tatsumoto, N., J. Phys. Chem., 80, 313 (1976).
73. Nancollas, G. H., Sutin, N., Inorg. Chem., 3, 360 (1964).
74. Hammes, G. G., Stinfeld, J. I., J. Phys. Chem., 67, 528 (1963).
75. Melson, G. A., Wilkins, R. G., J. Chem. Soc., 4208 (1962).
76. Davies, A. G., Mac F. Smith, W., Proc. Chem. Soc., 380 (1961).
77. Letter, J. E., Jordan, R. B., J. Am. Chem. Soc., 97, 2381 (1975).
78. Almgren, T., Hagstrom, I., Water Res., 8, 395 (1974).
79. Chen, K. Y., Morris, J. C., Environ. Sci. Tech., 6, 529 (1972).
80. Östlund, H. G., Alexander, J., J. Geophys. Res., 68, 3395 (1963).
81. Cline, J. D., Richards, F. A., Environ. Sci. Tech., 3, 838 (1969).
82. O'Brien, D. J., Birkner, F. B., Environ. Sci., Tech., 12, 1114 (1977).
83. Avrahami, M., Golding, R. M., J. Chem. Soc., (A), 647 (1968).
84. Demirjian, Y. A., "The Electrochemical Characterization of the Rates of Oxidation of Soluble Sulfides by Molecular Oxygen," Ph.D. Thesis, University of Michigan, Ann Arbor, 1971.
85. Moussavi, M., "Autoxidation of Aqueous Sulfide in the Presence of Phenols-Kinetics and Products," Ph.D. Thesis, University of Southern California, Los Angeles, 1974.
86. Sorokin, Y. I., Okeanologiya, 11, 423 (1971).
87. Lim, B. C. H., "Kinetics and Mechanism of the Oxidation of Sulfide by Oxygen Catalyzed by Homogeneous Organometallic Complexes," M.S. Thesis, University of Minnesota (1978).

88. Hoffmann, M. R., Lim, B. C. H., (1979) to appear, Environ. Sci. Tech., 14, 1406.
89. Silverman, J., Dodson, R. W., J. Phys. Chem., 56, 846 (1952).
90. Reynolds, W. L., J. Am. Chem. Soc., 83, 1078 (1961).
91. Ford-Smith, M. H., Sutin, N., J. Am. Chem. Soc., 83, 1830 (1961).
92. Baxendale, J. H., Hardy, H. R., Trans. Faraday Soc., 50, 808 (1954).
93. Reynolds, W. L., Fukushima, S., Inorg. Chem., 2, 176 (1963).