

<b>COMPONENTS:</b> 1. Manganese(II) selenite; $MnSeO_3$ ; [15702-34-8] 2a. Sulfuric acid; $H_2SO_4$ ; [7664-93-9] 2b. Hydrochloric acid; $HCl$ ; [7647-01-0] 3. Water; $H_2O$ ; [7732-18-5]	<b>ORIGINAL MEASUREMENTS:</b> Chukhlantsev, V.G.; Tomashevsky, G.P. <i>Zh. Anal. Khim.</i> 1957, 12, 296-301; * <i>J. Anal. Chem. USSR</i> 1957, 12, 303-9.																																													
<b>VARIABLES:</b> Sulfuric and hydrochloric acid concentrations One temperature: 293 K	<b>PREPARED BY:</b> Mary R. Masson																																													
<b>EXPERIMENTAL VALUES:</b> All concentrations are expressed in units of $mol\ dm^{-3}$ . <table border="1" data-bbox="185 520 1138 747"> <thead> <tr> <th>Soln.</th> <th>Initial pH</th> <th>Final pH</th> <th><math>[Mn^{2+}]</math></th> <th>pMn</th> <th><math>\log \alpha_{L(H)}</math></th> <th><math>p[SeO_3^{2-}]</math></th> <th><math>pK_{SO}</math></th> </tr> </thead> <tbody> <tr> <td rowspan="2"><math>H_2SO_4</math></td> <td>2.69</td> <td>6.07</td> <td><math>3.1 \times 10^{-3}</math></td> <td>2.51</td> <td>1.94</td> <td>4.45</td> <td>6.96</td> </tr> <tr> <td>2.06</td> <td>4.35</td> <td><math>2.8 \times 10^{-2}</math></td> <td>1.55</td> <td>3.63</td> <td>5.18</td> <td>6.73</td> </tr> <tr> <td rowspan="3"><math>HCl</math></td> <td>2.79</td> <td>6.24</td> <td><math>2.2 \times 10^{-3}</math></td> <td>2.66</td> <td>1.77</td> <td>4.43</td> <td>7.09</td> </tr> <tr> <td>2.41</td> <td>5.37</td> <td><math>6.8 \times 10^{-3}</math></td> <td>2.17</td> <td>2.60</td> <td>4.77</td> <td>6.94</td> </tr> <tr> <td>2.05</td> <td>4.42</td> <td><math>2.1 \times 10^{-2}</math></td> <td>1.68</td> <td>3.58</td> <td>5.26</td> <td>6.94</td> </tr> </tbody> </table> <p>The average value is <math>K_{SO} = 1.2 \times 10^{-7} mol^2 dm^{-6}</math>.                      (<math>pK_{SO} = 6.9</math>)</p> <p><b>Notes.</b>  <math>[Se_{tot}] = [Mn^{2+}]</math> and <math>[SeO_3^{2-}] = [Se_{tot}]/\alpha_{L(H)}</math>                      where <math>\alpha_{L(H)} = (1 + [H^+]/K_2 + [H^+]^2/K_1K_2)</math>                      and the acid dissociation constants have the values <math>K_1 = 4 \times 10^{-3}</math> and <math>K_2 = 1.0 \times 10^{-8}</math> (ref. 1).</p>		Soln.	Initial pH	Final pH	$[Mn^{2+}]$	pMn	$\log \alpha_{L(H)}$	$p[SeO_3^{2-}]$	$pK_{SO}$	$H_2SO_4$	2.69	6.07	$3.1 \times 10^{-3}$	2.51	1.94	4.45	6.96	2.06	4.35	$2.8 \times 10^{-2}$	1.55	3.63	5.18	6.73	$HCl$	2.79	6.24	$2.2 \times 10^{-3}$	2.66	1.77	4.43	7.09	2.41	5.37	$6.8 \times 10^{-3}$	2.17	2.60	4.77	6.94	2.05	4.42	$2.1 \times 10^{-2}$	1.68	3.58	5.26	6.94
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<b>AUXILIARY INFORMATION</b>																																														
<b>METHOD APPARATUS/PROCEDURE:</b> Solutions of sulfuric and hydrochloric acid were saturated with manganese(II) selenite by shaking in a thermostat at $20 \pm 0.05^\circ C$ for 8 hr. The remaining solid phase was removed by centrifugation, then the pH was measured ("Moskip" pH meter, to 0.01 pH unit) and the manganese concentration was determined spectrophotometrically, after oxidation to Mn(VII) with ammonium persulfate.	<b>SOURCE AND PURITY OF MATERIALS:</b> C.P.-grade reagents were used. Manganese selenite was prepared by mixing stoichiometric amounts of 0.2N manganese(II) sulfate and 0.2N sodium selenite, and heating the solution to 50 - 70°C. After prolonged standing, the precipitate was washed with water and dried at 40°C. Manganese was determined gravimetrically as the sulfate, and selenium as the element.																																													
<b>ESTIMATED ERROR:</b> $\pm 0.4 \times 10^{-7}$ . (The spread in the results is 0.36 of a log unit.) Temperature: $\pm 0.05$ K																																														
<b>REFERENCES:</b> 1. Rumpf, P. <i>Compt. Rendu</i> 1933, 197, 686.																																														