

COMPONENTS: (1) Benzenesulfonamide, 4-amino-, benzene-sulfonamide, 4-amino-N-2-thiazolyl- (molecular compound); $C_6H_8N_2O_2S \cdot C_9H_9N_3O_2S_2$; [1704-78-3] (2) Water; H_2O ; [7732-18-5]	ORIGINAL MEASUREMENTS: Ito, K.; Sekiguchi, K. <i>Chem. Pharm. Bull.</i> <u>1967</u> , <i>15</i> (4), 420-6.																	
VARIABLES: Temperature	PREPARED BY: R. Piekos																	
EXPERIMENTAL VALUES: <table border="1" data-bbox="454 554 944 826"> <thead> <tr> <th rowspan="2">t/°C</th> <th colspan="2">Solubility</th> </tr> <tr> <th>10^3 mol dm^{-3} solution</th> <th>g dm^{-3a}</th> </tr> </thead> <tbody> <tr> <td>20</td> <td>0.173</td> <td>0.074</td> </tr> <tr> <td>25</td> <td>0.231</td> <td>0.099</td> </tr> <tr> <td>30</td> <td>0.320</td> <td>0.137</td> </tr> <tr> <td>35</td> <td>0.412</td> <td>0.176</td> </tr> </tbody> </table> <p data-bbox="482 856 791 897">^a Calculated by compiler.</p>		t/°C	Solubility		10^3 mol dm^{-3} solution	g dm^{-3a}	20	0.173	0.074	25	0.231	0.099	30	0.320	0.137	35	0.412	0.176
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METHOD/APPARATUS/PROCEDURE: The earlier described method (1) was used: in a 200-ml egg-plant type flask immersed in a thermostat, an excess (0.6 and 0.8 g per 100 ml of water at expts at 20°C and 30°C, resp) of the mol compd was placed with 100 ml of distd water which was previously kept at appropriate temp. Immediately after addn of water the mixt was vigorously agitated with an elec stirrer. The soly equilibrium at 20°C and 30°C was attained after 15 and about 9 h, resp. Aliquots were withdrawn at certain time intervals with a pipet equipped with a filter and the concn of solute was detd spectrophotometrically in the uv region.	SOURCE AND PURITY OF MATERIALS: The source and purity of the mol compd was not specified. Distd water was used. ESTIMATED ERROR: Nothing specified. REFERENCES: 1. Sekiguchi, K.; Ito, K. <i>Chem. Pharm. Bull.</i> <u>1965</u> , <i>13</i> (4), 405.																	