

COMPONENTS:		EVALUATOR:	
(1) <i>sec</i> -butylbenzene; C ₁₀ H ₁₄ ; [135-98-8]		A. Maczynski Institute of Physical Chemistry of the Polish Academy of Sciences Warszawa, Poland.	
(2) Water; H ₂ O; [7732-18-5]		November 1981	
CRITICAL EVALUATION:			
The solubility of <i>sec</i> -butylbenzene (1) in water (2) has been reported in three works listed below:			
<u>Authors</u>	<u>Method</u>	<u>T/K</u>	<u>g(1)/100 g sln</u>
Andrews and Keefer (ref 1)	spectrophotometric	298.15	0.0309
Sutton and Calder (ref 3)	GLC	298.15	0.00176
Krzyzanowska and Szeliga (ref 4)	GLC	298.15	0.00101
The data of Andrews and Keefer seem to be high. The mean value of the remaining data is tentative.			
The solubility of water (2) in <i>sec</i> -butylbenzene (1) has been reported in one work listed below:			
<u>Authors</u>	<u>Method</u>	<u>T/K</u>	
Englin <i>et al.</i> (ref 2)	gasometric	273.15-323.15	
Since these data are from one source only, they are regarded as tentative.			
<u>SOLUBILITY OF <i>sec</i>-BUTYLBENZENE (1) IN WATER (2)</u>			
<u>TENTATIVE VALUE</u>			
<u>T/K</u>	<u>g(1)/100 g sln</u>	<u>x₁</u>	
298	0.0014	1.9 x 10 ⁻⁶	
<u>SOLUBILITY OF WATER (2) IN <i>sec</i>-BUTYLBENZENE (1)</u>			
<u>TENTATIVE VALUES</u>			
<u>T/K</u>	<u>g(2)/100 g sln</u>	<u>x₂</u>	
283	0.020	0.0020	
293	0.029	0.0029	
303	0.039	0.0038	
REFERENCES			
1. Andrews, L.J.; Keefer, R.M. <i>J. Amer. Chem. Soc.</i> <u>1950</u> , <i>72</i> , 5034-7.			
2. Englin, B.A.; Plate, A.F.; Tugolukov, V.M.; Pryanishnikova, M.A. <i>Khim. Tekhnol. Topl. Masel</i> <u>1965</u> , <i>10</i> , 42-6.			
3. Sutton, C.; Calder, J.A. <i>J. Chem. Eng. Data</i> <u>1975</u> , <i>20</i> , 320-2.			
4. Krzyzanowska, T.; Szeliga, J. <i>Nafta (Katowice)</i> <u>1978</u> , <i>12</i> , 413-7.			

COMPONENTS: (1) sec-Butylbenzene; C ₁₀ H ₁₄ ; [135-98-8] (2) Water; H ₂ O; [7732-18-5]	ORIGINAL MEASUREMENTS: Andrews, L.J.; Keefer, R.M. <i>J. Am. Chem. Soc.</i> <u>1950</u> , <i>72</i> , 5034-7.
VARIABLES: One temperature: 25°C	PREPARED BY: A. Maczynski and Z. Maczynska
EXPERIMENTAL VALUES: <p>The solubility of sec-butylbenzene in water at 25°C was reported to be 0.0309 g(1)/100 g sln.</p> <p>The corresponding mole fraction, x_1 calculated by the compilers is 4.15×10^{-6}.</p>	
AUXILIARY INFORMATION	
METHOD/APPARATUS/PROCEDURE: A mixture of (1) and (2) was rotated for twenty hours in a constant temperature bath at 25°C. A sample (5-20 mL) of the aqueous phase was withdrawn and extracted with a measured volume of hexane (10-50 mL) by shaking in a glass-stoppered Erlenmeyer flask. Next, the absorbance of the hexane phase was measured against a hexane blank on the Beckman spectrophotometer.	SOURCE AND PURITY OF MATERIALS: (1) Eastman Kodak Co. white label; fractionally distilled; b.p. range 175.0-175.8°C. (2) not specified. ESTIMATED ERROR: not specified. REFERENCES:

COMPONENTS: (1) sec-Butylbenzene; C ₁₀ H ₁₄ ; [135-98-8] (2) Water; H ₂ O; [7732-18-5]	ORIGINAL MEASUREMENTS: Englin, B.A.; Plate, A.F.; Tugolukov, V.M.; Pryanishnikova, M.A. <i>Khim. Tekhnol. Topl. i Masel</i> <u>1965</u> , 10, 42-6.												
VARIABLES: Temperature: 10-30°C	PREPARED BY: A. Maczynski and Z. Maczynska												
EXPERIMENTAL VALUES: <p style="text-align: center;">Solubility of water in sec-butylbenzene</p> <table border="1" style="margin-left: auto; margin-right: auto;"> <thead> <tr> <th style="text-align: center;"><u>t/°C</u></th> <th style="text-align: center;"><u>g(2)/100 g sln</u></th> <th style="text-align: center;"><u>x₂ (compiler)</u></th> </tr> </thead> <tbody> <tr> <td style="text-align: center;">10</td> <td style="text-align: center;">0.0226</td> <td style="text-align: center;">0.00223</td> </tr> <tr> <td style="text-align: center;">20</td> <td style="text-align: center;">0.0317</td> <td style="text-align: center;">0.00313</td> </tr> <tr> <td style="text-align: center;">30</td> <td style="text-align: center;">0.0426</td> <td style="text-align: center;">0.00420</td> </tr> </tbody> </table>		<u>t/°C</u>	<u>g(2)/100 g sln</u>	<u>x₂ (compiler)</u>	10	0.0226	0.00223	20	0.0317	0.00313	30	0.0426	0.00420
<u>t/°C</u>	<u>g(2)/100 g sln</u>	<u>x₂ (compiler)</u>											
10	0.0226	0.00223											
20	0.0317	0.00313											
30	0.0426	0.00420											
AUXILIARY INFORMATION													
METHOD/APPARATUS/PROCEDURE: Component (1) was introduced into a thermostatted flask and saturated for 5 hr. with (2). Next, calcium hydride was added and the evolving hydrogen volume measured and hence the concentration of (2) in (1) was evaluated.	SOURCE AND PURITY OF MATERIALS: (1) not specified. (2) not specified. ESTIMATED ERROR: Not specified. REFERENCES:												

COMPONENTS: (1) sec-Butylbenzene; C ₁₀ H ₁₄ ; [135-98-8] (2) Water; H ₂ O; [7732-18-5]	ORIGINAL MEASUREMENTS: Sutton, C.; Calder, J.A. <i>J. Chem. Eng. Data</i> 1975, 20, 320-2.
VARIABLES: One temperature: 25°C	PREPARED BY: A. Maczynski and Z. Maczynska
EXPERIMENTAL VALUES: <p>The solubility of sec-butylbenzene in water at 25°C was reported to be 17.6 mg(1)/kg(2). The corresponding mass percent and mole fraction, x_1, calculated by the compilers are 0.00176 g(1)/100 g sln and 2.36×10^{-6}.</p>	
AUXILIARY INFORMATION	
METHOD/APPARATUS/PROCEDURE: The concentration of (1) in (2) was determined by gas chromatography.	SOURCE AND PURITY OF MATERIALS: (1) Aldrich Chemical Co. or Matheson Coleman and Bell 99+%. (2) distilled. ESTIMATED ERROR: temp. $\pm 0.1^\circ\text{C}$ soly. 0.2 (the standard deviation of the mean for six replicates). REFERENCES:

COMPONENTS: (1) sec-Butylbenzene; C ₁₀ H ₁₄ ; [135-98-8] (2) Water; H ₂ O; [7732-18-5]	ORIGINAL MEASUREMENTS: Krzyzanowska, T.; Szeliga, J. <i>Nafta (Katowice)</i> , <u>1978</u> , <i>12</i> , 413-7.
VARIABLES: One temperature: 25°C	PREPARED BY: M.C. Haulait-Pirson
EXPERIMENTAL VALUES: <p>The solubility of sec-butylbenzene in water at 25°C was reported to be 10.1 mg(1)/kg(2).</p> <p>The corresponding mass percent and mole fraction, x_1, calculated by compiler are 0.00101 g(1)/100 g sln and 1.36×10^{-6}.</p>	
AUXILIARY INFORMATION	
METHOD/APPARATUS/PROCEDURE: <p>The saturated solutions of (1) in (2) were prepared in two ways. First, 200 μL of (1) was injected into 20 mL of (2) and thermostatted at 25°C. Second, the mixture of (1) and (2) as above was thermostatted at 70°C and then cooled to 25°C. The time required to obtain equilibrium was three weeks. The solubility of (1) in (2) was measured by glc. A Perkin-Elmer model F-11 gas chromatograph equipped with a 100-150 mesh Porasil column (70°C) and a flame ionization detector was used. Saturated solutions of heptane in (2) were used as standard solutions.</p>	SOURCE AND PURITY OF MATERIALS: (1) not specified. (2) not specified. <hr/> ESTIMATED ERROR: soly. 0.3 mg(1)/kg(2) (standard deviation from 7-9 determinations). <hr/> REFERENCES:

COMPONENTS: (1) sec-Butylbenzene; C ₁₀ H ₁₄ ; [135-98-8] (2) Artificial seawater (ref 1)	ORIGINAL MEASUREMENTS: Sutton, C.; Calder, J.A. <i>J. Chem. Eng. Data</i> <u>1975</u> , 20, 320-2.
VARIABLES: One temperature: 25.0°C One salinity: 34.5 g salts/kg sln	PREPARED BY: M. Kleinschmidt and W. Shiu
EXPERIMENTAL VALUES: <p>The solubility of sec-butylbenzene in artificial seawater is reported to be 11.9 mg(1)/kg sln. The corresponding mass percent and mole fraction, x_1 calculated by the compiler are 1.19×10^{-3} g(1)/100 g sln and 1.64×10^{-6} assuming the artificial seawater composition of ref 1.</p>	
AUXILIARY INFORMATION	
METHOD/APPARATUS/PROCEDURE: A test tube containing (1) was placed in a flask containing (2) thus allowing for equilibration through the vapor phase. The saturated solution was extracted with hexane and analyzed by gas chromatography.	SOURCE AND PURITY OF MATERIALS: (1) from either Aldrich Chemical Co. or Matheson Coleman and Bell, 99+% pure. (2) made from doubly distilled water and salts 99+% pure. ESTIMATED ERROR: temp. \pm 0.1°C soly. 0.2 (std. dev.) REFERENCES: 1. Lyman, J.; Fleming, R.H.; <i>J. Mar. Res.</i> <u>1940</u> , 3, 135.