

<p>COMPONENTS:</p> <p>(1) Lanthanum bromide; LaBr_3; [13536-79-3]</p> <p>(2) 1,4-Dioxane; $\text{C}_4\text{H}_8\text{O}_2$; [123-91-1]</p>	<p>ORIGINAL MEASUREMENTS:</p> <p>Kirmse, E.M.; Zwietasch, K.J.; Tirschmann, J.; Oelsner, L.; Niedergesass, U.</p> <p>Z. Chem. <u>1968</u>, <u>8</u>, 472-3.</p>
<p>VARIABLES:</p> <p>Room temperature: T/K around 298</p>	<p>PREPARED BY:</p> <p>Mark Salomon and Tomasz Mioduski</p>
<p>EXPERIMENTAL VALUES:</p> <p>The solubility of LaBr_3 in p-dioxane at about 25°C was given as</p> <p style="text-align: center;">0.2 mass %</p> <p>The corresponding molality calculated by the compilers is</p> <p style="text-align: center;">$0.005_3 \text{ mol kg}^{-1}$</p> <p>The nature of the solid phase was not specified.</p>	
<p>AUXILIARY INFORMATION</p>	
<p>METHOD/APPARATUS/PROCEDURE:</p> <p>The solute-solvent mixture was isothermally agitated at 25°C or room temperature. The authors state that the difference found for the solubility was within experimental error limits.</p> <p>La was determined by complexometric titration.</p> <p>No other details given.</p>	<p>SOURCE AND PURITY OF MATERIALS:</p> <p>The anhydrous salt was prepared by the method of Taylor and Carter (1).</p> <p>No other information given.</p> <hr/> <p>ESTIMATED ERROR:</p> <p>Nothing specified.</p> <hr/> <p>REFERENCES:</p> <p>1. Taylor, M.D.; Carter, C.P. <i>J. Inorg. Nucl. Chem.</i> <u>1962</u>, <u>24</u>, 387.</p>

COMPONENTS: (1) Lanthanum bromide; LaBr_3 ; [13536-79-3] (2) Alcohols; ROH (3) 1,4-Dioxane; $\text{C}_4\text{H}_8\text{O}_2$; [123-91-1]	ORIGINAL MEASUREMENTS: Golub, A.M.; Yankovich, V. N. <i>Ukr. Khim. Zh.</i> <u>1977</u> , 43 , 1139-42; Ukr. J. Chem. (Engl. Transl.) <u>1977</u> , 43 , 16-20.																
VARIABLES: Concentration of ROH T/K = 295	PREPARED BY: M. Salomon and T. Mioduski																
EXPERIMENTAL VALUES: Numerical data were not given, but results were presented graphically and in the form of the equation $K = [\text{LaCl}_3 \cdot n\text{ROH}] / [\text{ROH}]^n \quad [1]$ In this equation $[\text{LaBr}_3 \cdot n\text{ROH}]$ is the solubility in units of mol dm^{-3} , $[\text{ROH}]$ is the total alcohol concentration in units of mol dm^{-3} , and n is the solvate number in solution (see ref. 1). According to this equation, n is calculated from the slope of a plot of the logarithm of the solubility, $\log [\text{LaBr}_3 \cdot n\text{ROH}]$, against $\log [\text{ROH}]$. Thus the solubility of LaBr_3 can be calculated as a function of ROH concentration using the reported values of n and K (see table below). The alcohol concentrations were varied from 1-5 mol dm^{-3} . <table border="1" data-bbox="144 745 1166 944"> <thead> <tr> <th>alcohol</th> <th>n</th> <th>$-\log K$</th> <th>nature of the solid phase</th> </tr> </thead> <tbody> <tr> <td rowspan="2">methanol; CH_4O; [67-56-1]</td> <td>1</td> <td>0.89</td> <td rowspan="2">$\text{LaBr}_3 \cdot 2\text{CH}_3\text{OH}$</td> </tr> <tr> <td>2</td> <td>1.16</td> </tr> <tr> <td rowspan="2">1-propanol; $\text{C}_3\text{H}_8\text{O}$; [71-23-8]</td> <td>1</td> <td>1.71</td> <td rowspan="2">$\text{LaBr}_3 \cdot 2\text{C}_3\text{H}_7\text{OH}$</td> </tr> <tr> <td>2</td> <td>2.16</td> </tr> </tbody> </table> In the above systems, two values of n and K are reported, and the overall solubility of LaBr_3 is obtained by using the values for n - K in eq. [1] which give the greater solubility.		alcohol	n	$-\log K$	nature of the solid phase	methanol; CH_4O ; [67-56-1]	1	0.89	$\text{LaBr}_3 \cdot 2\text{CH}_3\text{OH}$	2	1.16	1-propanol; $\text{C}_3\text{H}_8\text{O}$; [71-23-8]	1	1.71	$\text{LaBr}_3 \cdot 2\text{C}_3\text{H}_7\text{OH}$	2	2.16
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METHOD/APPARATUS/PROCEDURE: Isothermal method used as described in (1). Solvent mixtures of known alcohol concentration were saturated with anhydrous LaBr_3 at $22 \pm 1^\circ\text{C}$. Equilibrium was confirmed from constancy of the rare earth metal concentration upon repeated analyses. Liquid phases were analysed for rare earth metal concentration (method not specified). At least 3 separate experiments were carried out for each system studied. In addition, the solid phases were analysed for several arbitrary points of each series of experiments (method not specified).	SOURCE AND PURITY OF MATERIALS: Source and purity of LaBr_3 not specified. Anhydrous LaBr_3 prepared by method described in (2). C.p. grade organic solvents were purified by "known" methods (3). ESTIMATED ERROR: Soly: nothing specified. Temp: precision ± 1 K REFERENCES: 1. Golub, A.M.; Golovorushkin, V.I. <i>Zh. Neorg. Khim.</i> <u>1968</u> , 13 , 3194. 2. Spedding, F. H.; Doan, A. H. <i>J. Am. Chem. Soc.</i> <u>1952</u> , 74 , 2783. 3. Kolotyrkin, Ya.M. (ed). <i>Electrochemistry of Metals in Nonaqueous Solutions</i> . Khimiya Press. Moscow. <u>1974</u> . p 440.																