

<b>COMPONENTS:</b>  (1) Eicosane; C <sub>20</sub> H <sub>42</sub> ; [112-95-8] (2) Water; H <sub>2</sub> O; [7732-18-5]	<b>ORIGINAL MEASUREMENTS:</b>  Sutton, C.; Calder, J.A.  <i>Environ. Sci. Technol.</i> <u>1974</u> , 8, 654-7.
<b>VARIABLES:</b>  One temperature: 25°C	<b>PREPARED BY:</b>  M.C. Haulait-Pirson
<b>EXPERIMENTAL VALUES:</b>  The solubility of eicosane in water at 25°C was reported to be $1.9 \times 10^{-7}$ g(1)/100 g(2) corresponding to a mole fraction $x_1$ , of $1.1 \times 10^{-10}$ .	
<b>AUXILIARY INFORMATION</b>	
<b>METHOD/APPARATUS/PROCEDURE:</b>  175 mg (1) were equilibrated with 700 mL (2) in closed flasks by shaking on a constant temperature bath for 12 hours. The flasks were then allowed to stand for 24 hours. Aliquots of 100 mL were removed, filtered through a 0.45 μm Millipore filter, then extracted three times with 10-mL portions of hexane containing an internal standard. The concentration of (1) was determined by injection of the hexane extract into a dual column gas chromatograph equipped with flame ionization detectors.	<b>SOURCE AND PURITY OF MATERIALS:</b>  (1) Analabs Inc., 99+%. (2) doubly distilled.  <b>ESTIMATED ERROR:</b>  temp. ± 0.1°C soly. ± 16%  <b>REFERENCES:</b>

<b>COMPONENTS:</b>  (1) Eicosane; C <sub>20</sub> H <sub>42</sub> ; [112-95-8]  (2) Seawater	<b>ORIGINAL MEASUREMENTS:</b>  Sutton, C.; Calder, J.A.  <i>Environ. Sci. Technol.</i> <u>1974</u> , 8, 654-7.
<b>VARIABLES:</b> One temperature: 25°C One salinity: 35 g salts/kg sln	<b>PREPARED BY:</b>  P.A. Meyers and D. Shaw
<b>EXPERIMENTAL VALUES:</b>  The solubility of eicosane in seawater was reported to be $8 \times 10^{-8}$ g(1)/100 g sln and $x_1 = 5 \times 10^{-11}$ .	
<b>AUXILIARY INFORMATION</b>	
<b>METHOD/APPARATUS/PROCEDURE:</b>  (1) and (2) were placed in a glass stoppered flask fitted with a Teflon stopcock near the bottom. The components were equilibrated by gentle shaking for 12 hrs at 25.0 ± 0.1°C. The mixture was then allowed to stand for 24 hrs. Samples removed via the stopcock were filtered with suction through 0.45 μm membrane filters to remove any hydrocarbon droplets. The filtrate was extracted three times with hexane and analyzed by gas chromatography.	<b>SOURCE AND PURITY OF MATERIALS:</b>  Analabs, Inc., North Haven, Conn. 99 + % pure hydrocarbons.  Seawater collected from 25 m depth in Gulf of Mexico, poisoned with HgCl <sub>2</sub> sln to prevent bacterial growth, and filtered through Gelman glass fiber filter. Natural n-alkane levels too low to cause interference.  <b>ESTIMATED ERROR:</b> Eight replications were made. The average of the deviations of the mean gave an experimental error of ± 16%, yet some accommodation may have occurred due to presence of natural dissolved organic matter.  <b>REFERENCES:</b>