

NOTE ON THE FREEZING POINT OF "ISO-OCTANE" (2, 2, 4-TRIMETHYLPENTANE)¹

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ABSTRACT

A high-grade sample of commercial iso-octane was purified by equilibrium melting. The freezing point of the pure hydrocarbon was found to be -107.41°C . The purity of an iso-octane sample may be calculated from the equation: Mole per cent purity $= 3.86_2 t_F + 514.8$ in which t_F is the initial freezing point of the sample in $^{\circ}\text{C}$.

"Iso-octane" (2, 2, 4-trimethylpentane) is used as the upper reference standard for antidetonation tests of motor fuels. As a

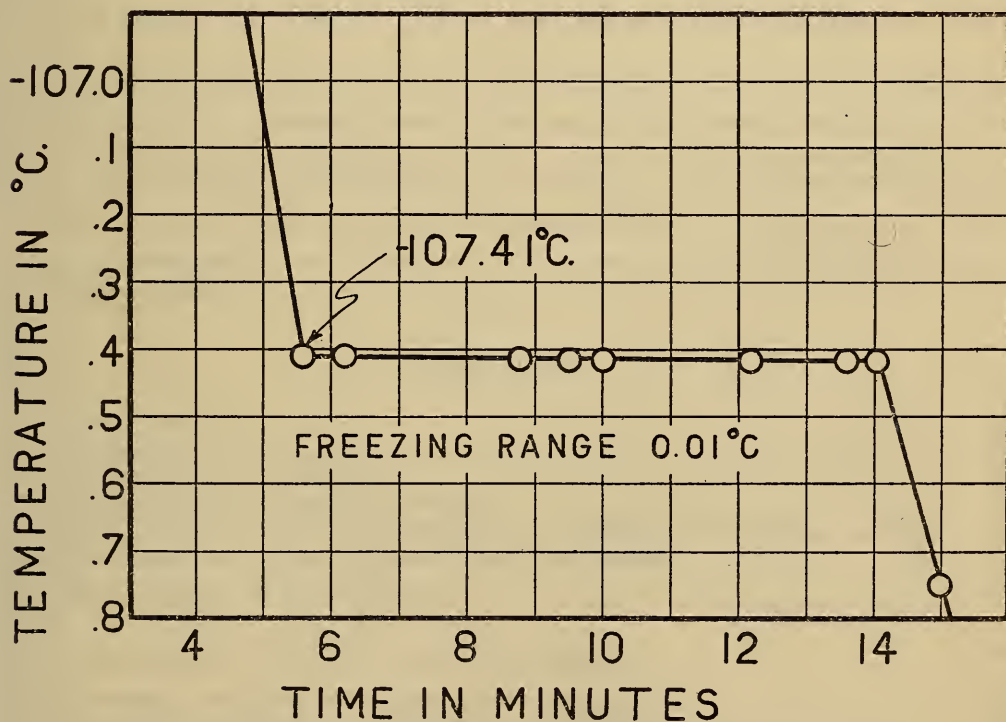


FIGURE 1.—Time-temperature cooling curve of iso-octane

criterion for the purity of commercial "iso-octane," it is desirable to have a reliable value for the freezing point of the pure hydrocarbon. The value (-107.8°C .) reported in the literature³ is apparently too low as many commercial samples were found to freeze at temperatures above this value.

¹ Financial assistance has been received from the research fund of the American Petroleum Institute. This work is part of Project No. 6, "The Separation, Identification, and Determination of the Constituents of Petroleum."

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³ G. M. Parks and H. M. Huffman, Ind. Eng. Chem., vol. 23, p. 1139, 1931.

A sample of high-grade commercial "iso-octane" was purified further by subjecting it to a number of fractionations by equilibrium melting in a centrifuge.⁴ As a result of these fractionations four different fractions with freezing points ranging from -107.7° to -107.4° C. were obtained. The time-temperature cooling curve of the purest fraction of "iso-octane" was determined, and is shown in Figure 1.

From the value, -107.41° C., found for the freezing point of pure iso-octane and from Parks and Huffman's value ⁵ (18.9 cal./g) for the heat of fusion, the purity of the iso-octane may be calculated from the laws of ideal solutions and is expressed by the equation: $P = 3.86_2 t_F + 514.8$ in which t_F is the initial freezing point of the sample in $^{\circ}$ C., and P is the purity in mole per cent.

Temperatures were determined by means of a platinum-resistance thermometer calibrated at this bureau in accordance with the International Temperature Scale⁶ as adopted in 1927.

WASHINGTON, May 4, 1932.

⁴ For method see M. M. Hicks-Bruun and J. H. Bruun, B. S. Jour. Research, vol. 8, p. 527, 1932.

⁵ See footnote 3, p. 269.

⁶ B. S. Jour. Research, vol. 1, p. 635, 1928.

