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<td>Author(s)</td>
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Kyoto University
Fluorometric Determination of Europium with Benzoyltrifluoroacetone

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and Takayuki Sumida.*

Received September 16, 1968

Determination of traces of europium as benzoyltrifluoroacetone chelate was studied. Trioctylphosphine oxide adduct of the chelate, which has a fluorescence maximum at 612 nm, was formed in n-hexane solution by utilizing synergistic effect in solvent extraction. The effect of various factors on the fluorescence intensity at 612 nm was examined. This method enabled us to detect europium in a concentration as low as $2 \times 10^{-8}$ M (or 3.04 x $10^{-2}$ µg per 10 ml). Of the other rare earth ions, only samarium gave a positive error. Other common divalent cations had little effect except ferric ion which caused a negative error.

I. INTRODUCTION

Recently many works have been done on the spectrofluorometric determination of rare earth ions because of its high sensitivity and selectivity. The simultaneous determination of dysprosium, europium, samarium and terbium was investigated in 0.6M sodium tungstate solution. Europium and terbium were determined fluorometrically in an o-phenanthroline-salicylic acid system, by the extraction of them into benzene. These methods are interfered by some of the other rare earth ions. Taketatsu et al. have investigated the fluorometric analysis of terbium and europium in potassium carbonate solution. Dagnall, Smith and West examined EDTA-sulphosalicylic acid system and made a specific determination of terbium in the concentration ranging from $6.4 \times 10^{-3}$ to 3.2 µg of terbium per ml. McCarthy and Wineforder studied the possibilities of determining dysprosium, europium, samarium and terbium fluorometrically in nonaqueous solutions of aromatic carbonyl compounds.

Some β-diketone chelates of europium are also known to fluoresce. They have attracted attention as possible laser materials. These compounds have strong and broad absorption bands in the near ultraviolet which were caused by the $\pi \rightarrow \pi^*$ transition of their benzenoid ring. Their fluorescence gives a relatively sharp line characteristic of the metal ion. This phenomenon based on the intramolecular energy transfer was tried to apply for the specific and sensitive determination of europium in the present paper.

Kononenko et al. investigated the fluorometry of europium and samarium in o-phenanthroline thenoyltrifluoroacetone system, Ballard and Edwards that of europium in TTA-trioctylphosphate system and Stanley et al. that of europium, samarium, terbium and gadolinium using TTA, benzoylaceton and dibenzoylmethane by computer spectrum stripping techniques.

The fluorescence intensity of β-diketone chelates in the solvent extraction

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method decreases rapidly as the ligand concentration increases because of the emission based on the intramolecular energy transfer. In the present work, the authors examined the fluorometric determination of europium as benzoyl trifluoroacetone-(BFA)-triocetylphosphine oxide (TOPO) complex by using the synergistic effect in solvent extraction.13

II. EXPERIMENTALS

Reagents.

A 2 × 10⁻³ M stock solution of europium was prepared by dissolving 0.0352 g europium oxide (Shin-Etsu Chem. Ind. Co., Ltd. 99.9 per cent Eu₂O₃) with concentrated hydrochloric acid, by evaporating to dryness and by diluting to 100 ml with redistilled water. The other rare earth solutions were prepared in a manner similar to the europium solution from their oxides of 99.99 per cent in purity.

Benzoyl trifluoroacetone and triocetylphosphine oxide (Dojindo & Co., Ltd Research Laboratory) were used without further purification. n-Hexane, purified by the distillation, was used as the solvent for the extraction of the chelates.

Rhodamine B was used as an internal standard. The other chemicals were reagent grade materials.

Apparatus.

A Hitachi spectrophotometer, Model EPU-2 equipped with an attachment for fluorescence measurement was used for obtaining spectra and intensity. A mercury arc was used as light source and the fused silica cell of 10 mm in light path as sample container. The pH measurements were performed with a Hitachi-Horiba Model M-4 pH meter.

Procedure.

An aliquot of a solution containing over 0.03 µg of europium and 1 ml of buffered solution (1.0 M NaAc-HAc) were taken into a 20 ml beaker. The pH of the solution was then adjusted to the desired value by the addition of 1 M hydrochloric acid or 1 M sodium hydroxide. The aqueous phase was transferred into a 25 ml separating funnel and finally diluted to 10 ml with water. After 10 ml of n-hexane solution consisting of 5 × 10⁻⁴ M BFA and 2 × 10⁻² M TOPO was added, the funnel was shaken for 30 min., and allowed to settle. The organic phase was carefully separated through a filter paper. The fluorescence intensity was measured at 612 mµ against a 0.3 µg per ml of rhodamine B as an internal standard, by using an excitation wavelength of 365 mµ.

III. RESULTS AND DISCUSSION

Fluorescence Spectrum.

The fluorescence spectrum for the n-hexane solution of europium-BFA-TOPO system containing 1.0 × 10⁻⁶ M of europium, 5 × 10⁻⁴ M of BFA and 2 × 10⁻² M of TOPO, respectively, is shown in Fig. 1. The spectrum has a maximum peak at 612 mµ, which is characteristic of europium ion. Three tenth microgramme per ml of rhodamine B was used as an internal standard, and n-hexane was chosen as an extractant solvent, because it has no absorbance in near ultraviolet region. The fluorescence intensity was measured throughout the experiment at 612 mµ.

Effect of pH.

To establish the optimum pH value of the aqueous phase for the fluorometry
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of europium, the dependence of the fluorescence intensity on the pH value was investigated. In Fig. 2 the fluorescence intensity is shown as a function of pH for the three systems:

System I: Eu $5 \times 10^{-7} \text{ M}$, BFA $5 \times 10^{-4} \text{ M}$, TOPO $2 \times 10^{-2} \text{ M}$
System II: Eu $1 \times 10^{-5} \text{ M}$, BFA $1 \times 10^{-2} \text{ M}$, TOPO $1 \times 10^{-2} \text{ M}$
System III: Eu $1 \times 10^{-6} \text{ M}$, BFA $1 \times 10^{-3} \text{ M}$, TOPO $1 \times 10^{-2} \text{ M}$

It is seen from the figure that nearly constant intensities are obtained through a considerably wide range of pH (pH value from 3 to 5.5) for all the systems. Taking this result into account, the pH value was kept at 4.50 throughout the following experiments.

Fig. 2. Effect of pH on fluorescence intensity.
I: Eu $5 \times 10^{-7} \text{ M}$, BFA $5 \times 10^{-4} \text{ M}$, TOPO $2 \times 10^{-2} \text{ M}$
II: Eu $1 \times 10^{-5} \text{ M}$, BFA $1 \times 10^{-2} \text{ M}$, TOPO $1 \times 10^{-2} \text{ M}$
III: Eu $1 \times 10^{-6} \text{ M}$, BFA $1 \times 10^{-3} \text{ M}$, TOPO $1 \times 10^{-2} \text{ M}$
Effect of TOPO Concentration.

In the solvent extraction, some metal chelates can be often extracted at a lower pH region by adding such Lewis bases as trioctylphosphine oxide. If this phenomenon, generally called synergistic effect, can be applied to the present system, the chelates could be extracted quantitatively even at the considerably low concentration of the chelating agent. This would be useful for the sensitive fluorometry of the element. Then, the effect of TOPO concentration on the fluorescence intensity was examined. As shown in Fig. 3, a nearly constant intensity was obtained when the concentration of TOPO was over $2 \times 10^{-4} \text{M}$. The authors used $2 \times 10^{-2} \text{M}$ of TOPO was used in all other experiments.

Fig. 3. Effect of TOPO concentration.
Eu: $5 \times 10^{-7} \text{M}$, BFA: $5 \times 10^{-4} \text{M}$

Fig. 4. Effect of the concentration of the chelating agent.
Eu: $5 \times 10^{-7} \text{M}$, TOPO: $2 \times 10^{-2} \text{M}$
Effect of the Concentration of the Chelating Agent.

After the europium chelate (5 × 10⁻⁷ M) was extracted at pH 4.5 with the chelating agent of various concentrations, the fluorescence intensity was measured at 612 μm. The intensity decreased rapidly in inverse proportion to the reagent concentration in the range between 10⁻² and 10⁻³ M. This is because of the fluorescence caused by the intramolecular energy transfer as described above. As the reagent concentration became further low, the reagent effect on the intensity slightly softened, as shown in Fig. 4. In consideration of sensitivity and practical use, 5 × 10⁻⁴ M of BFA was used throughout the other experiments.

Effect of Temperature.

The fluorescence intensity was measured at various temperatures. As shown in Fig. 5, the intensity decreased gradually as the temperature was raised. The decrease in the intensity was 0.55 per cent degree.

Effect of Standing and Shaking Time.

The intensity of the n-hexane solution of europium-BFA-TOPO chelate was measured at 612 μm at appropriate intervals of time to examine the effect of the standing and shaking time. As shown in Fig. 6, little influence of the standing and shaking time was observed upon the intensity of the organic phase for 10 min. and over.

Calibration Curves.

A calibration curve was made by extracting a different amount of europium into 10 ml portion of n-hexane solution containing 5 × 10⁻⁴ M of BFA and 2 × 10⁻² M of TOPO from 10 ml portion of aqueous solution at pH 4.50. The fluorescence intensity was measured at 612 μm against 0.3 μg per ml of rhodamine B as an internal standard. A good linear relationship was obtained by plotting the intensity of the organic phase against the concentration of europium added in the aqueous solution, as shown by curve I in Fig. 7. The other curve in the figure (curve II) was made by extracting the aqueous solution into n-hexane solution containing 5 × 10⁻⁵ M of BFA and 2 × 10⁻² M of TOPO and by measuring the intensity under the most sensitive condition.
of the present instrument. The curve shows that $2 \times 10^{-8} M$ of europium ($3.04 \times 10^{-3}$ µg per 10 ml) was detectable in the present method.

**Effect of Diverse Ions.**

The effect of diverse ions was examined with the solution containing 10.76 µg of europium and various diverse ions. The pH value of the aqueous phase was adjusted at pH 4.50. Of the other rare earth ions tested, only samarium gave a
positive error, while common di- and tervalent cations had little effect on the fluorescence intensity except iron (III) which caused a negative error, as shown in Table I.

Table 1. Effect of co-existing ions
Europium taken, 0.76 µg.

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<th>Ion</th>
<th>Added, µg</th>
<th>Eu found, µg</th>
<th>Ion</th>
<th>Added, µg</th>
<th>Eu found, µg</th>
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(4) R. M. Dagnall, R. Smith and T. S. West, Analyst, 92, 358 (1967)