LA-UR -83-1548

Ş

CONF-830565--2

Los Alamos National Laboratory is operated by the University of California for the United States De, artment of Energy under contract W-7405-ENG-36.



TITLE: A HIGH-Z ORGANIC-SCINTILLATION SOLUTION

LA-UR--83-1548

DE83 012774

AUTHOR(S): I. B. Berlman, EG&G/Santa Barbara J. M. Fluornoy, EG&G/Santa Barbara C. B. Ashford, EG&C/Santa Barbara Peter B. Lyons, P-14

SUBMITTED TO: ADVANCES IN SCINTILLATOR COUNTING CONFERENCE Banf?, Alberta, Canada May 15-18, 1993

DISCLAIMER

This report was prepared as an account of work sponsored by an agency of the United States Government. Neither the United States Government nor any agency thereof, nor any of their employees, makes any warranty, express or implied, or assumes any legal liability or responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference horein to any specific commercial product, process, or service by trade name, trademark, manufacturer, or otherwise does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof.

By acceptance or this article, the public cognizes that the U.S. Government retains a nonexclusive, royalty-tree license to publish or reproduce the published form of this contribution, or to allow others to do so, for U.S. Government purposes.

The Los Alamos National Laboratory requests that the publisher identify this article as work performed under the auspices of the U.S. Department of Energy.



DISTRIBUTION OF THIS DOCUMENT IS UNLIMITED-1

A HIGH-Z ORGANIC-SCINTILLATION SOLUTION*

by

I.B. Berlman,** L.A. Franks, S.S. Lutz, J.M. Flournoy, and C.B. Ashford EG&G, Santa Barbara Operations Goleta, California 93117

and

P.B. Lyons University of California Los Alamos National Laboratory Los Alamos, New Mexico 87545

INTRODUCTION

It is now believed that the properties needed to make a high-Z liquid scintillator are known. In an earlier study¹ it was demonstrated that a close encounter is a prerequisite for a heavy atom to quench an excited molecule. More recently² it was shown that a very fast scintillator may be obtained by selectively quenching the solute by strategically substituting a relatively heavy atom on the solute chromophore.

In the present experiment, an attempt is made to raise the average Z of a scintillation solution with as little attendant quenching as possible. Since high-Z atoms quench by means of a close encounter, as stated above, such encounters are minimized by the use of alkyl groups substituted on the solvent, solute, and heavy atoms. The aromatic compound 1,2,4-trimethylbenzene (pseudocumene) is used as the solvent; 4,4"-ci(5-tridecyl)-p-terphenyl (SG-180) as the solute; and tetrabutyltin as the high-Z material.

******Presently at Hebrew University, Jerusalem, Israel.

^{*}This work was performed under the auspices of the U.S. Department of Energy under Contract No. DE-AC08-83NV10282. NOTE: By acceptance of this article, the publisher and/or recipient acknowledges the U.S. Government's right to retain a nonexclusive royalty-free license in and to any copyright covering this paper. Reference to a company or product name does not imply approval or recommendation of the product by the U.S. Department of Energy to the exclusion of others that may be suitable.

To establish the validity of our ideas, various experiments have been performed with less protected solvents, and heavy atoms. These include benzene, toluene, p-terphenyl, bromobutane, and bromobenzene.

PROCEDURE

Various solutions of p-terphenyl or 4,4"-di(5-tridecyl)-pterphenyl (SG-180) in benzene, toluene, or pseudocumene, were tested. To these solutions various amounts of a quencher such as tetrabutyltin, bromobenzene, or bromobutane, or of a diluent such as 2,2,4-trimethylpentane (isooctane) were added.

When large quantities of tetrabutyltin were added, there was a diluting of both the solvent and solute. An attempt was made to separate the quenching effect from the dilution effects by substituting isooctane for tetrabutyltin. In addition the decreases in pulse height are measured with a straight dilution of just the solute by addition of pure solvent.

Before the solutions were tested in a Suprasil spectrometer cell they were placed in an ultrasonic bath first to dissolve the material and then to outgas the solution.

Each sample solution was evaluated by exposing it to 50-ps electron pulses from the DOE/EG&G electron linear accelerator (linac) and measuring a number of parameters of the resultant fluorescence pulse. These included the 10-90% pulse response rise time (PRT), the pulse full width at half maximum (FWHM), the decay time from 0.7 maximum to (1.7e) the integral rise time (IRT) (i.e., the 10-90% rise time of the integral of the pulse), and the integral of the fluorescence pulse (yield). Figure 1 shows a schematic diagram of the experimental arrangement.

Standard solutions were tested each day to assess variations in detection efficiency of the system.

RESULTS

In one set of experiments our standard was a solution of pseudocumene, as solvent, and SG-180 as solute, at a high concentration of 0.14 M. To this solution various amounts of tetrabutyltin were added, and the resultant fluorescence characteristics were measured. The data are assembled in Table 1.

When tetrabutyltin in a concentration of 75% by volume is added to the standard solution, the average pulse height is decreased

٦



Figure 1. Schematic diagram of test configuration used for scintillator evaluation. Linac was operated at 6 MeV with 50 ps (FWHM) electron pulses at 360 Hz.

Table 1. Quenching by tetrabutyltin in a high-concentration solution of SG-180 in pseudocumene (see Glossary for abbreviations).

SG-180 (moles/liter)*	TBSN (volume %)	PRT (ns)	FWHI1 (ns)	Decay Time (ns)	IRT (ns)	Yield**
0.14 0.07 0.04	50 75	0.51 0.55 0.51	2.0 1.9 2.3	1.7 1.4 1.6	5.9 4.1 3.8	0.8 0.3 0.2

*Takes into account dilution by quencher. Effect of dilution with isooctane is shown in last three lin_s of Table 4. **Yield values from different data sets are not directly comparable due to slight day-to-day variations in detection efficiency.

only by a factor of three. This is just a little more than what would be expected from a pure dilution effect where the diluent dues not transfer excitation energy to the basic solution. It is of interest to note that the decay time remains essentially

unchanged at about 1.6 ns, a result that would indicate that almost no quenching of the solute takes place. Part of the success of this experiment is due to the large solubility of the solute since it does not precipitate out on addition of such large quéntities of quencher.

In a second set of experiments the effectiveness of alkyl chains on the solvent in reducing the quenching by heavy atoms was tested on less concentrated samples than those mentioned above. The data are assembled in Table 2. Most of the pulse parameters show little dependence on TBSN concentration or on the degree of methylation of the solvent. However, the decay time and integral rise time increase somewhat as tetrabutyltin is added. This is probably due to the increased contribution

Solute	Concentration (moles/liter)*	Solvenc	TBSN (volume ")	PRT (ns)	FLIHIT (HS)	Decay Time (ns)	IRT (ns)	Yield**
ТР Тр Тр	0.025 0.018 0.013	8 8 8	28.6 50.0	0.73	2.7 2.8 3.0	1.6 1.7 2.0	4.7 4.6 5.0	0.52 0.19 0.10
TP TP TP	0.025 0.018 0.013	T T T	28.6 50.0	0.69 0.71 0.65	2.4 2.6 2.9	1.6 1.7 2.0	4.5 4.4 5.5	0.37 0.15 0.06
ТР ТР ТР	0.025 0.010 0.013	PC PC PC	28.6 50.0	0.73 0.75 	2.6 2.8 3.3	1.7 1.0 2 2	4.8 4.6 5.5	0.63 0.29 0.16
SG-180 SG-180 SG-180	0.025 0.010 0.013	T T T	28.6 50.0	0.81 0.81 0.77	2.5 2.5 2.9	1.6 1.9 2.2	4.5 4.5 4.9	0.49 0.22 0.11
SG-130 SG-120 SG-180	0.025 0 018 0.013	PC PC PC	2C.6 50.0	0.75 0.83 0.71	2.4 2.6 2.9	1.6 1.7 2.2	4.7 4.6 4.9	0.4£ 0.30 0.17
4-0r-TP 4-8r-TP 4-0r-TP	0.025 0.018 0.013	PC PC PC	28.6 50.0	0.51 0.55 0.41	1.5 1.6 1.5	1.0 1.3 1.9	2.8 3.3 4.8	0.12 0.06 0.04
3-0r-TP 3-8r-TP 3-Gr-TP	0.025 0.018 0.013	PC PC PC	28.6 50.0	0.53	1.6 1.6 1.6	1.2 1.4 1.5	3.6 3.5 4.4	0.17 0.09 0.05

Table 2. Fluorescence characteristics of some scintillation solutions with low solute concentrations quenched by TBSN (see Glossary for abbreviations).

*Takes into account dilution by quencher.

*Yield values from different data sets are not directly comparable due to day-to-day variations in detection efficiency.

Solute	Concentration* (Holes/liter)	Solvent	Quencher	Concentration (volume <)	PRT (ns)	F\.1H11 (ns)	Decay Time (ns)	IRT (ns)	Yield**
TP TP TP	0.025 0.025 0.025	8 8 8	8r8t Br8t	 1 2	0.73 0.73 0.71	2.7 2.6 2.6	1.7 1.6 1.6	4.8 4.8 4.6	0.47 0.40 0.37
TP TP TP	0.025 0.025 0.025	8 8 8	BrBz BrDz	1 2	0.73 0.57 0.49	2.7 1.8 1.6	1.7 1.4 1.2	4.8 3.6 3.1	0.47 0.16 0.09
TP TP TP	0.025 0.025 0.925	T T T	BrBt BrDt	1 2	0.73 0.71 0.71	2.5 2.5 2.5	1.6 1.6 1.6	4.6 4.7 3.8	0.57 0.48 0.43
TP TP TP	0.025 0.025 0.025	T T T	BrBz BrBz	1 2	0.71 0.57 0.57	2.5 2.0 1.8	1.6 1.4 1.5	4.5 3.3 3.9	0.56 0.23 0.16
TP TP TP	0.025 0.025 0.025	PC PC PC	BrBt BrBt	1 2	0.73 0.73 0.71	2.5 2.4 2.4	1.7 1.6 1.6	4.9 4.4 4.5	0.40 0.35 0.29
TP TP TP	0.025 0.025 0.025	PC PC PC	Br8z Br9z	1 2	0.73 0.63 0.61	2.5 2.2 2.0	1.7 1.5 1.5	4.9 3.9 3.8	0.40 0.22 0.16
SG-100 SG-180 SG-180	0.025 0.025 0.025	9 B 8	Br8t Br8t	1 2	0.77 0.77 0.75	2.6 2.5 2.5	1.6 1.6 1.6	4.6 4.5 4.5	0.40 0.35 0.31
SG-180 SG-180 SG-180	0.025 0.025 0.025	8 8 8	BrBz BrBz	1 2	0.77 0.61 0.55	2.6 2.0 1.7	1.6 1.5 1.2	4.6 3.6 3.8	0.40 0.17 0.09
SG-180 SG-120 SG-120	0.025 C.025 0.025	PC PC PC	Brßt BrDt	1 2	0.71 0.69 0.71	2.4 2.3 2.3	1.6 1.5 1.5	4.6 4.1 4.0	0.49 0.41 0.35
SG-180 5G-180 SG-180	0.025 0.025 0.025	PC PC PC	BrBz BrBz	1 2	0.71 0.67 0.61	2 4 2.1 1.9	1.6 1.4 1.4	4.6 3.8 3.4	0.49 0.28 0.20

Table 3. Fluorescence quenching by bromobenzene and bromobutane (see Glossary for abbreviations).

.

*Takes into account dilution by quencher. **Yiald values from different data sets are not directly comparable due to day-to-day variations in detection efficiency.

yield if both the solute and solvent are diluted by an <u>inert</u> solvent. Curves above the dashed line represent varying degrees of energy transfer; curves below the dashed line indicate quenching.

Isooctane is seen to be an active diluent for both p-terphenyl and SG-180 in all solvents. The effect is greatest in the case of SG-180 in pseudocumene and is least with p-terphenyl

Solute	Concentration (moles/liter)*	Solvent	lsooctane (volume 3)	PRT (ns)	FWHH (ns)	Decay Time (ns)	IRT (ns)	Yield**
TP TP TP	0.025 0.018 0.013	8 8 8	28.6 50.0	0.73 0.81 0.81	2.7 2.9 3.3	1.7 1.8 2.2	4.8 4.8 5.5	0.54 0.40 0.29
TP TP TP	0.025 0.018 0.013	• T T T .	28.6 50.0	0.71 0.79 0.91	2.4 2.7 3.1	1.6 1.7 2.0	4.8 4.7 5.2	0.41 0.33 0.26
TP TP TP	0.025 0.013 0.013	PC PC PC	28.6 50.0	0.73 0.79 0.91	2.7· 2.9 3.4	1.7 1.9 2.4	5.0 5.0 5.6	0.61 0.52 0.42
SG-180 SG-130 SG-180	0.025 0.013 0.013	8 8 8	28.6 50.0	0.77 0.77 0.71	2.8 :3.3 3.9	1.6 2.2 3.4	4.5 5.3 6.7	0.72 0.46 0.26
SG-180 SG-180 SG-180	0.025 0.018 0.013	T T T	28.6 50.0	0.73 0.83 0.85	2.5 3.2 3.9	1.5 2.0 3.2	4.3 5.2 6.4	0.87 0.60 0.36
SG-180 SG-180 SG-180	0.025 0.018 0.013	PC PC PC	28.6 50.0	0.69 0.81 0.89	2.5 3.1 3.8	1.6 1.9 3.2	4.5 4.0 6.7	1.04 0.75 0.50
SG-180 SG-180 SG-180	0.140 0.079 0.040	PC PC PC	50.0 75.0	0.51 0.57 0.65	2.0 2.2 2.5	1.7 1.6 1.6	5.9 5.1 4.7	0.84 0.72 0.52

Table 4. Modification in fluorescence characteristics induced by dilution with isooctane (see Glossary for abbreviations).

*Takes into account dilution by isooctane.

*Yield values from different data sets are not directly comparable due to day-today variations in detection efficiency.

in benzene, where the diluent is nearly inert. These results suggest a selective interaction between the isooctane and methyl groups on the solvent. In fact, addition of only 2% pseudocumene to an isooctane solution of SG-180 increased the yield by 50%. If the yield had varied linearly with the composition between pure pseudocumene and pure isooctane, only a 12% increase should have occurred. It is also noted that the tetrabutyltin quenches the benzene and toluene solutions to a greater degree than those with pseudocumene. The SG-180 solution in pseudocumene was quenched the least. While 50% added tetrabutyltin caused about 80% reduction in the yield of p-terphenyl in benzene, only 1% added bromobenzene produced the same effect. Thus the combined shielding of the butyl groups on the tin and the tridecyl groups on SG-180 essentially eliminate quenching by the tin.



Figure 2. Dilution and quenching effects on light yield of initially 0.025 M solutions of p-terphenyl in benzene (2a) and pseudocumene (2b). Solutions were diluted with benzene (2a-circles), or pseudocumene (2b-circles), isooctane (squares), and tetrabutlytin (triangles) and irradiated with 50-ps burst of 6-MeV electrons. Dashed curve represents effect expected from diluting both solvent and solute with an inert solvent.



Figure 3. Dilution and quenching effects on the light yield of initially 0.025 M solutions of SG-180 in toluene (3a) and pseudocumene (3b). Solutions were diluted in toluene (3a-circles) or pseudocumene (3b-circles), isooctane (squares), and tetrabutyltin (triangles) and irradiated with 50-ps burst of 6-MeV electrons. Dashed curve represents effect expected from dilution both solvent and solute with an inert solvent. Figure 4 shows a pulse height spectrum (and associated photopeak) that was obtained when a solution consisting of 0.035 M of SG-180 in 25% by volume of pseudocumene and 75% by volume of tetrabutyltin was exposed to a ⁵⁷Co source. For comparison, the pulse height spectrum from a 0.14 M SG-180 solution in pseudocumene without added tin is also shown. The enhanced photoelectric interaction resulting from the tin is evident and suggests immediate applications to x-ray diagnostics.



Figure 4. Pulse height spectra from irradiation with ⁵⁷Co (122-keV gamma-ray) of (a) 0.035 m SG-180 in a solvent of 25% pseudocumene and 75% tetrabutyltin (19% tin by weight) and (b) 0.14 M SG-180 in pseudocumene without added tin. Abscissa is not energy-calibrated.

ACKNOWLEDGEMENT

We wish to thank Dr. Shlomo Gershuni for providing us with the SG-180.

GLOSSARY

.

.

SG-180	- 4,4"-di(5-tridecyl)-p-terphenyl
PC	- pseudocumene
TBSN	- tetrabutyltin
T	- toluene
TP	- p-terphenyl
3-Br-TP	- 3-bromo-p-terphenyl
4-Br-TP	- 4-bromo-p-terphenyl
В	- benzene
BrBt	- bromobutane
BrBz	- bromobenzene
	•

.

REFERENCES/BIBLIOGRAPHY

- 1. I.B. Berlman, J. Phys. Chem., 77, pp 562-567 (1973).
- 2. In press.
- 3. F. Hirayama and S. Lipsky, J. Chem. Phys., 51, p 3616 (1969).
- 4. F. Hirayama, W. Rothman, and S. Lipsky, Chem. Phys. Lett., 5, pp 296-298 (1970).