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THE REACTION OF POTASSIUM GERMYL WITH ORGANIC HALIDES

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Robert Michael Dreyfuss
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THE REACTION OF POTASSIUM GERMYL WITH ORGANIC HALIDES

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ABSTRACT

Many organic halides react with potassium germyl in diglyme to give the corresponding organogermanes. However, halides for which nucleophilic substitutions are slow react with potassium germyl to give principally the corresponding hydrocarbons. Data relevant to the mechanism of the latter type of reaction are presented. A method for the preparation and standardization of stock solutions of potassium germyl is described. A new compound, digermylmethane (1,3-digermapropane), is reported, and its nmr, mass and infrared spectra are discussed.
Introduction

The germyl anion, $\text{GeH}_3^-$, was first prepared by Kraus and Carney by the reaction of germane, $\text{GeH}_4$, with sodium-ammonia solutions.\textsuperscript{1} Teal and Kraus found the germyl anion to be an excellent nucleophile, capable of converting primary alkyl halides to the corresponding alkylgermanes.\textsuperscript{2} However, simple displacement did not occur with all organic halides. Sodium germyl and bromobenzene did not yield the expected phenylgermane, but rather benzene and $\text{GeH}_2$.\textsuperscript{3}

\[
\text{NaGeH}_3 + \text{C}_6\text{H}_5\text{Br} \rightarrow \text{NaBr} + \text{C}_6\text{H}_6 + \text{GeH}_2
\]

Methylene bromide also did not react to give simple germyl anion substitution. The reported products were methylgermane and aminogerma.\textsuperscript{2}

\[
2\text{NaGeH}_3 + \text{CH}_2\text{Br}_2 + \text{NH}_3 \rightarrow \text{CH}_3\text{GeH}_3 + \text{GeH}_3\text{NH}_2 + 2\text{NaBr}
\]

In each of the latter reactions, a hydrogen atom replaced a halogen atom. It was the aim of this study to determine the types of halides for which this type of reaction occurs, and to investigate the reaction mechanism. Reactions were carried out in diglyme (bis(methoxyethyl)ether) because of the ease of preparation of potassium germyl in this solvent and because of the hope that diglyme might prove to be a more favorable medium than liquid ammonia for germyl anion substitutions and that it might enable us to prepare the elusive compound digermymethane.
Experimental

Reagents.- Germane was obtained from the Matheson Company; its purity was checked by infrared and mass spectrometry and by vapor pressure measurements at -111.6° (180 torr as compared to the literature value, 181 torr). Diglyme (Ansul Ether 141) was purified by distillation from potassium hydroxide followed by vacuum distillation from lithium aluminum hydride at approximately 40°. Organic halides of reagent grade were used without purification; lower grade materials were purified by distillation from phosphorus pentoxide or by gas chromatography. The identity and purity of volatile substances were determined by infrared spectrometry with Perkin-Elmer Infracord spectrometers (Models 137 and 137B), by mass spectrometry with a Consolidated Engineering Corp. mass spectrometer (Model 21-103C) or a Varian M-66 mass spectrometer, and by gas chromatography with a Varian Aerograph, Model A-90-P.

Preparation and Standardization of Potassium Germyl.- Potassium germyl was prepared in the apparatus shown in Figure 1. A magnetic stir bar and twenty grams of powdered reagent-grade potassium hydroxide were placed in flask A. Sixty or seventy milliliters of diglyme was then pipetted into the flask. The diglyme was degassed at -78° by pumping through stopcock B; stopcock B was then closed. The diglyme slurry was stirred while holding it at 0°, and stopcock B was opened to a section of the vacuum line containing 30 mmols of germane. Stirring was continued for about one hour, during which essentially all the germane was absorbed by the slurry; the remainder was condensed into a -196° trap.
The apparatus was then inverted, a tube leading to a tank of dry nitrogen was attached at stopcock B, and an atmosphere of dry nitrogen was introduced. (The tube was alternately flushed with nitrogen and evacuated several times to ensure the absence of air.) The solution was filtered by gravity flow through the medium-porosity filter C into the long-necked Erlenmeyer flask E. When the filtration was complete, dry nitrogen at atmospheric pressure was introduced through stopcock D, using the same precautions noted above. A and C were removed from E, and E was stoppered with a ground glass sleeve joint attached to an open stopcock. After a few minutes of flushing, the latter stopcock and stopcock D were almost simultaneously closed.

This procedure yielded a clear, colorless solution of potassium germyl which could be stored at 0° without apparent decomposition. The solution was standardized as follows: Using the precautions described above, dry nitrogen at atmospheric pressure was again introduced at stopcock D. When the pressure in the flask slightly exceeded one atmosphere, the flask was unstoppered and seated into a sleeve joint sheath with a sidearm as shown in Figure 2. Dry nitrogen was flushed through the sidearm; this nitrogen and the nitrogen simultaneously flushing the flask vented through the narrow top of the sheath. A long glass pipet was clamped over the solution as shown in Figure 2 and was flushed by drawing in nitrogen with a greased syringe, followed by expulsion of the nitrogen through the side arm of the three-way stopcock. When the system was completely purged of air, the pipet was lowered and potassium germyl solution was sucked up to one of several calibrated marks on the
pipet. The pipet was then redrawn into the sheath, and the stock solution flask was removed and restoppered as before. An identical flask, previously evacuated, was then opened with nitrogen flushing and was fitted to the sheath. After the system had been purged, the potassium gemyl in the pipet was transferred to the new flask. The flask was restoppered as before, cooled to -196°, and evacuated.

An excess of degassed water was then distilled onto the frozen potassium gemyl solution; the mixture was warmed to room temperature, and the evolved germane was distilled, with pumping, through a -111.6° trap into a -196° trap. The germane was quantitatively measured by PVT measurements, and its purity was checked by vapor pressure and infrared measurements. A known excess of 0.1 M HCl was added to the residue, and the solution was titrated with 0.1 M NaOH to the phenolphthalein endpoint. In accordance with the following hydrolysis reaction,

\[
\text{GeH}_3^- + \text{H}_2\text{O} \rightarrow \text{GeH}_4 + \text{OH}^-
\]

the germane produced equalled the titrated hydroxide within ±1 percent.

**Qualitative Aryl Halide Experiments.** An aliquot of 0.2 M KGeH₃ was transferred to a long-necked Erlenmeyer flask as described in the standardization procedure. Excess bromobenzene was condensed onto the frozen solution at -196°, and the mixture was allowed to warm to room temperature. Immediately a white precipitate formed which soon developed a yellow color; after one half hour, the reaction mixture was orange. The solution was then pumped to dryness through traps at -45°, -160°, and -196°. The first trap contained diglyme and benzene, the second,
benzene and digermane, and the third, germane, as indicated by infrared and mass spectra. No trace of any compound containing a carbon-germanium bond was detected in these samples. A similar experiment using iodo-benzene also showed benzene, germane and digermane to be the only volatile products.

Adding the potassium germyl solution dropwise to bromobenzene or iodobenzene in other experiments gave identical results. In an attempt to detect biphenyl formation, the contents of the -45° trap were analyzed by gas chromatography. The retention time of biphenyl was determined using a 4-ft by 1/4-in. 20% silicone GE SF-96 on firebrick column at 230°. Under these conditions, an aliquot of the unknown showed no evidence of biphenyl.

Quantitative Aryl Halide Experiments.- Excess chlorobenzene was condensed onto a diglyme solution containing 1.33 mmols of potassium germyl. After two hours at room temperature, a faint yellow cloudiness had appeared. The mixture was then distilled, with pumping, for fifteen minutes; the volatile products were collected in traps at -95° and -196°. A total of 0.132 mmol of germanium (as germane and digermane) was found in the -196° trap. (The digermane was separated by fractional condensation in a -160° trap.) Water was then distilled into the solution, and germane and digermane (1.10 mmols of germanium) were produced. To the contents of the -95° trap was added 10 ml of a solution of toluene in diglyme which contained 1.0 ml of toluene (9.41 mmol) per 100 ml of solution. The mixture was analyzed gas-chromatographically for benzene using the GE SF-96 column at 65°; less than 0.01 mmol was found.
The results of experiments in which excesses of bromobenzene, 4-bromo-m-xylene and 2-bromo-m-xylene were condensed onto potassium germyl solutions are presented in Table I. After the indicated reaction times, the volatile materials were collected in traps at -95° and -196°. The germane and digermane in the -196° traps were separated, identified, and the respective amounts measured. The -95° traps were analyzed for benzene or xylene as described above. The solid residues were dissolved by boiling in a slightly basic 3% H₂O₂ solution. Excess peroxide was destroyed by acidification and further boiling, and the resulting solution then analyzed volumetrically for germanium by the method of Tchakirian.⁶ We have no explanation for the low total germanium analyses in the iodobenzene experiments.

Reaction of Bromobenzene with Potassium Germyl-d₄.- Germane-d₄ was prepared from GeCl₄ and LiAlD₄ according to a standard method.⁷ A small portion was pyrolyzed, and the hydrogen was analyzed by mass spectrometry; 92 atom percent of the hydrogen in the sample was deuterium.

Reagent grade potassium was distilled from a sidearm into a reaction vessel equipped with a break-off seal and an outlet to the vacuum line. About 0.5 ml of diglyme was distilled from lithium aluminum hydride onto the potassium mirror. Germane-d₄ (0.366 mmol) was condensed in, the vessel was sealed off, and the mixture was allowed to react for three days at -20°. At the end of this time, the solution was warmed to 0°, the vessel was attached to the vacuum line, and the break-off seal was opened. The solution was pumped on through two liquid
Table I

Reactions of Potassium Germyl with Aryl Halides

<table>
<thead>
<tr>
<th>Halide</th>
<th>Rxn. time, min.</th>
<th>KGeH₃ mmols</th>
<th>Germane + 2xDigermane formed, mmols</th>
<th>Nonvolatile Ge, mmols</th>
<th>Hydrocarbon formed, mmols</th>
<th>Hydrocarbon KGeH₃</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bromobenzene</td>
<td>5</td>
<td>1.33</td>
<td>.597</td>
<td>.72</td>
<td>.85</td>
<td>.64</td>
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<td></td>
<td>90</td>
<td>1.33</td>
<td>.589</td>
<td>.74</td>
<td>.81</td>
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<td></td>
<td>1200</td>
<td>1.33</td>
<td>.602</td>
<td>.66</td>
<td>.85</td>
<td>.64</td>
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<td>Iodobenzene</td>
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<td>1.36</td>
<td>.364</td>
<td>.60</td>
<td>.90</td>
<td>.66</td>
</tr>
<tr>
<td></td>
<td>90</td>
<td>1.36</td>
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<td>.63</td>
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<td></td>
<td>90</td>
<td>1.36</td>
<td>.334</td>
<td>.65</td>
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<td>.67</td>
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<td>90</td>
<td>1.57</td>
<td>.491</td>
<td>.65</td>
<td>1.01</td>
<td>.64</td>
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<td>4-Br-m-xylene</td>
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<td>1.57</td>
<td>.555</td>
<td>.90</td>
<td>.82</td>
<td>.52</td>
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<tr>
<td>2-Br-m-xylene</td>
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<td>1.57</td>
<td>.517</td>
<td>.88</td>
<td>.50</td>
<td>.38</td>
</tr>
</tbody>
</table>

* Benzene formed in the reactions of the halobenzenes; m-xylene formed in the runs with the bromo-m-xylene.
nitrogen traps with a Toepler pump; 0.98 mmol of hydrogen (of which 86 atom per cent was deuterium) was collected.

An excess of bromobenzene was condensed onto the potassium germyl, and the solution was allowed to react one half hour at room temperature. The volatiles were then passed through a -95° trap and a -196° trap. The fraction in the -95° trap was analyzed by mass spectrometry and was found to have peaks at m/e values of 78, 79, and 80, with relative intensities 30.2:100:5.7, respectively. A sample of ordinary benzene was shown to have 77, 78, and 79 peaks with relative intensities 19.8:100:7.1; from these data we calculate that 87% of the benzene produced in the above reaction contained one deuterium atom.

**Reaction of Phenylsodium with Germyle Bromide in Diglyme.** - A dispersion of sodium (2.42 mmols) was converted to phenylsodium by reaction with chlorobenzene in p-xylene. (100% yield corresponds to 1.21 mmol of NaC₆H₅.) The stirred slurry of phenylsodium was degassed by pumping, after which a solution of 1.29 mmol of GeH₃Br in 10 ml of diglyme was added quickly. Effervescence began at once and continued for one-half hour. The mixture was then cooled to -196° and Toepler-pumped through traps at -22°, -95°, and -196°; 0.082 mmol of hydrogen was collected. Water was then condensed onto the reaction mixture and the mixture warmed to room temperature and Toepler-pumped through the cold traps; this time 0.256 mmol of hydrogen was collected. The other volatile products of the reaction were germane, digermane, trigermane, benzene and methyl vinyl ether as determined by infrared and mass
spectroscopy. No phenylgermane was detected. The hydrogen produced in the reaction presumably was a result of unreacted sodium metal in the slurry.

Neopentyl and Neopentyl Halides. - About 1.5 mmols of neopentyl chloride (1-chloro-2,2-dimethylpropane), obtained from Matheson, Coleman and Bell and purified by gas chromatography, was condensed onto an equal number of millimoles of potassium germyl solution. A slight cloudiness appeared after several hours at room temperature, but analysis of the volatiles showed that essentially no reaction had taken place.

About 1.5 mmols of neopentyl bromide, obtained from K and K Laboratories and purified by gas chromatography, was condensed onto an equal number of millimoles of potassium germyl solution. A white cloudiness formed within a few minutes at room temperature; this soon became slightly yellow. After two hours, the volatile products were analyzed and were found to consist of germane, digermane and an approximately 1:1 mixture of neopentane and an alkylgermane, probably neopentylgermane. An attempt was made to separate the latter two compounds using the silicone GE SF-96 chromatograph column at room temperature. The neopentane was isolated as the pure compound (identified by comparison of its ir spectrum with that of a known sample of the material); the alkylgermane had an ir spectrum with the most intense bands at 845 and 2100 cm\(^{-1}\), typical of alkylgermanes.\(^{10}\) However, the bands found in neopentane were also present; therefore there is some doubt as to the purity of the isolated alkylgermane. The mass spectrum of this material
contained an envelope of peaks at m/e 141-147 and a much more intense envelope from m/e 120-135. The first band could be due to ions of formula $C_5\text{GeH}_x^+$, while the second could be due to $C_4\text{GeH}_x^+$, the most stable carbonium ion formed by fragmentation of neopentylgermane.\footnote{Neophil chloride (1-chloro-2-methyl-2-phenylpropane) was obtained from Fluka A. G. It did not react with potassium germyl. Neophil bromide was prepared by bromination of the chloride Grignard reagent. In separate experiments, potassium germyl solutions (0.2 and 0.01 M) were added dropwise to equivalent amounts of neophil bromide. The reaction was allowed to proceed two hours, during which time a yellow-white precipitate formed. The volatiles were analyzed by gas chromatography using a 20-ft by 3/8-in. silicone SE-30 on chromosorb W column at 110°. Some tert-butylbenzene, but no iso-butylbenzene could be detected.}

Neophil chloride (1-chloro-2-methyl-2-phenylpropane) was obtained from Fluka A. G. It did not react with potassium germyl. Neophil bromide was prepared by bromination of the chloride Grignard reagent. In separate experiments, potassium germyl solutions (0.2 and 0.01 M) were added dropwise to equivalent amounts of neophil bromide. The reaction was allowed to proceed two hours, during which time a yellow-white precipitate formed. The volatiles were analyzed by gas chromatography using a 20-ft by 3/8-in. silicone SE-30 on chromosorb W column at 110°. Some tert-butylbenzene, but no iso-butylbenzene could be detected.

**Haloalkane Experiments.** Bromomethane (1.62 mmols) was condensed onto a frozen solution of 1.12 mmols of potassium germyl. The reaction vessel was warmed, with shaking, to room temperature. After five minutes a white precipitate had formed. The volatile materials were Toepler-pumped for two hours through a -78° trap and two -196° traps. No non-condensable gas was found. The contents of the -196° traps were pumped through a -160° trap; 0.034 mmol of germyl was isolated. The contents of the -160° trap were passed through a -111.6° trap, and 1.07 mmols of methylgermyl was obtained (v.p. 47 mm at -83°, literature value 46.3 torr). A trace of dimethylgermyl and 0.40 mmol of bromomethane were found in the residue.
The same experiment was carried out using iodomethane. Again, an essentially quantitative formation of methylgermane was found (1.12 mmols KGeH₃ yielded 1.10 mmols CH₃GeH₃). However, about 0.01 mmol of methane was found.

The results of several experiments with di- and trihalomethanes are summarized in Table II. In most cases the halide was distilled from phosphorus pentoxide onto the frozen potassium germyl solution at -196°, and the reactants were allowed to warm to room temperature. In other cases, potassium germyl solution was added dropwise through a dropping funnel to the pure halide, and in one case, a 1% solution of the dihalide in diglyme was added dropwise to the germyl solution. After reaction had proceeded for the stated period of time, the volatile materials were fractionally condensed using -45°, -111.6°, -160°, and -196° traps. Infrared spectra of the contents of the -45° trap indicated solvent and traces of organogermanes which were then discarded. The -196° trap contained germane and the -160° trap methylgermane; these were identified by their ir spectra and vapor pressures, and were quantitatively measured. The -111.6° trap contained digermylmethane, dihalomethane, digermane, digermylethane, and digermylpropane, as indicated by mass and ir spectrometry. The latter mixture was treated in either of two ways to effect isolation of pure digermylmethane. Digermane and dibromomethane could be removed by hydrolysis overnight with a saturated aqueous solution of potassium hydroxide (the water being subsequently removed by passing through magnesium perchlorate) or by gas chromatography. Either method is effective; however hydrolysis
<table>
<thead>
<tr>
<th>Organic halide</th>
<th>Time, hr</th>
<th>KGeH₃ / Halide</th>
<th>GeH₄ / KGeH₃</th>
<th>CH₂GeH₃ / KGeH₃</th>
<th>CH₂(GeH₃)₂ / KGeH₃</th>
</tr>
</thead>
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<tr>
<td>CH₂Br₂</td>
<td>2</td>
<td>.25</td>
<td>.034</td>
<td>.20</td>
<td>.025</td>
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<tr>
<td>CH₂Br₂</td>
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<td>.45</td>
<td>.060</td>
<td>.25</td>
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<td>CH₂Br₂</td>
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<td>.038</td>
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<td>.06</td>
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<td>CH₂Br₂</td>
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<td>.05</td>
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<tr>
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<td>2.0</td>
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<td>.061</td>
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<td>CH₂Cl₂</td>
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<td>CHCl₃</td>
<td>2</td>
<td>1.0</td>
<td>.41</td>
<td>.028</td>
<td>.07</td>
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</table>

**a** Halide condensed onto potassium germyl;  
**b** Germyl added dropwise to halide;  
**c** Halide solution added dropwise to potassium germyl;  
**d** These values are only approximate; they are estimates of yield of 90% pure compound;  
**e** A trace of methyl bromide could be detected by ir as a product of this reaction;  
**f** A significant amount of dichloromethane was formed in this reaction.
is easier for greater than 0.2 mmol of product. Dichloromethane proved impossible to separate from digermylmethane; using the 20-ft by 3/8-in. silicone SE-30 column a one-minute difference in retention times was obtained after twenty minutes. Even then the peaks overlapped to a small extent. The other germylalkanes were not detected chromatographically, yet they were detected by mass spectrometry.

**Digermylmethane Mass Spectrum.** - The mass spectrum, obtained with a Varian M-66 mass spectrometer, is recorded in Table III. Sample pressure was 5 x 10^{-7} torr; sample temperature was 90° at the inlet and 125° at the analyzer. The electron energy was 70 eV, and the electron current was 30 µA. The peaks may be grouped into envelopes corresponding to the ions GeH_x^+ (m/e 70-80), GeCH_x^+ (m/e 82-96), Ge_2H_x^+ (140-156), and Ge_2CH_x^+ (152-172). The reported spectrum of disilylmethane shows analogous envelopes, including one corresponding to the anomalous ion Si_2H_x^+.\textsuperscript{13}

**Digermylmethane nmr Spectrum.** - The spectrum of a neat sample prepared from dichloromethane was recorded on a Varian A60 spectrometer and consisted of a 1:2:1 triplet and a 1:6:15:20:15:6:1 septet, 3.84 ppm and 0.33 ppm downfield from external tetramethylsilane, respectively; \( J_{C-H-Ge-H} \) was 4.00 Hz. Three tiny peaks could be detected under the expanded septet; these may have been part of a quartet, of which one peak was hidden by the digermylmethane peaks. Such a feature would be expected for contamination by digermylethane. The spectrum of digermylmethane is entirely analogous to that reported for disilylmethane.\textsuperscript{14}
Table III
Mass Spectrum of Digermylmethane

<table>
<thead>
<tr>
<th>m/e</th>
<th>Intensity</th>
<th>m/e</th>
<th>Intensity</th>
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<td>169</td>
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<td>93</td>
<td>2.8</td>
<td>73</td>
<td>5.6</td>
</tr>
<tr>
<td>154</td>
<td>2.8</td>
<td>92</td>
<td>2.1</td>
<td>72</td>
<td>9.9</td>
</tr>
<tr>
<td>153</td>
<td>3.5</td>
<td>91</td>
<td>31</td>
<td>71</td>
<td>2.8</td>
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<tr>
<td></td>
<td></td>
<td>90</td>
<td>31</td>
<td>70</td>
<td>5.6</td>
</tr>
</tbody>
</table>
Digermylmethane Infrared Spectrum. - The infrared spectrum of digermylmethane was recorded on Perkin-Elmer Infracord spectrometers (Models 137 and 137B). A 10-cm NaCl cell with a sample pressure of 10 mm was used, and the spectrum was recorded from 4000 to 650 cm\(^{-1}\). The sample was then transferred to a KBr cell, and the spectrum was recorded from 800 to 400 cm\(^{-1}\). The observed bands, with tentative vibrational assignments, are listed in Table IV.

The assignments were made by analogy with previous assignments for related compounds, notably disilylmethane\(^{15}\) and methylgermane\(^{16}\). In digermylmethane, the Ge-C asymmetric stretch has a high frequency, 680 cm\(^{-1}\), compared to Ge-C stretches in methylgermane, 601.6 cm\(^{-1}\), and in ethylgermane,\(^{17}\) 610 cm\(^{-1}\). However, this shift is consistent with that observed for the silyl analogs. In disilylmethane the C-Si asymmetric stretch frequency is 773 cm\(^{-1}\) whereas in methylsilane,\(^{18}\) it is 701 cm\(^{-1}\).
Table IV
Infrared Spectrum of Digermylmethane

<table>
<thead>
<tr>
<th>Band cm(^{-1})</th>
<th>Intensity</th>
<th>Assignment</th>
</tr>
</thead>
<tbody>
<tr>
<td>3000</td>
<td>w</td>
<td>CH(_2) stretch</td>
</tr>
<tr>
<td>2160</td>
<td>sh</td>
<td></td>
</tr>
<tr>
<td>2100</td>
<td>vs</td>
<td>GeH(_3) stretch</td>
</tr>
<tr>
<td>2030</td>
<td>sh</td>
<td></td>
</tr>
<tr>
<td>1380</td>
<td>w, br</td>
<td>CH(_2) scissor</td>
</tr>
<tr>
<td>1140</td>
<td>w</td>
<td>CH(_2) twist</td>
</tr>
<tr>
<td>1060</td>
<td>mw</td>
<td>CH(_2) wag</td>
</tr>
<tr>
<td>1050</td>
<td></td>
<td></td>
</tr>
<tr>
<td>960</td>
<td>br, sh</td>
<td>GeH(_3) asym. deformations</td>
</tr>
<tr>
<td>880</td>
<td>br, m</td>
<td></td>
</tr>
<tr>
<td>840</td>
<td>vs</td>
<td>GeH(_3) sym. deformation</td>
</tr>
<tr>
<td>755</td>
<td>br, m</td>
<td>CH(_2) rock</td>
</tr>
<tr>
<td>680</td>
<td>s</td>
<td>Ge-C asym. stretch</td>
</tr>
</tbody>
</table>
Results and Discussion

Potassium germyl reacts with organic halides in two different ways. The reaction which produces an organogermane undoubtedly involves nucleophilic attack by a germyl anion on a carbon atom, with simultaneous halide ion displacement. The reaction which produces a hydrocarbon and GeH₂ polymer probably involves nucleophilic attack by a germyl anion on a halogen atom, with simultaneous or subsequent formation of a C-H bond and cleavage of a Ge-H bond. A possible mechanism for the latter, hydrogen-substitution, reaction is indicated below.

\[
RX + GeH_3^- \rightarrow R^- + XGeH_3 \rightarrow RH + X^- + \frac{1}{n}(GeH_2)_n
\]

Halogen-atom-attack mechanisms have been proposed for many reactions of organophosphines\textsuperscript{19,20} and for the reaction of organic halides with potassium triphenylsilyl\textsuperscript{21} and sodium triphenylgermyl.\textsuperscript{22}

The hydrogen substitution reaction predominates in cases where one would expect halide ion displacement by the germyl anion to be slow. Consequently high yields of the hydrogen-substituted products are obtained in the reactions with aromatic halides,\textsuperscript{23} neopentyl and neophil halides,\textsuperscript{24} and dihalomethanes.\textsuperscript{25} However, even in the case of iodo-methane, a trace of the hydrogen-substituted product, methane, is obtained; it appears that the two types of reaction always occur simultaneously.

In the reaction of potassium germyl-\textsuperscript{d₃} with bromobenzene, it was observed that the hydrogen atom that replaced the bromine atom was
originally bonded to germanium. This means that a phenyl anion, if it is formed, deprotonates the nascent bromogerme in preference to the solvent, diglyme. However, we have observed that when sodium phenyl is introduced to diglyme containing excess bromogerme, the principal reaction is the cleavage of the diglyme. 26 Hence we conclude that no free phenyl anion is formed in the reaction of the germyl anion with bromobenzene. For a carbanion-type mechanism to be valid, a concerted reaction, in which germyl anion deprotonation occurs almost simultaneously with bromine cation removal, must be assumed. Perhaps a four-center activated complex 27 is formed; such a mechanism is almost equivalent to a cage reaction 28 in which the phenyl anion reacts with bromogerme before it can be dispersed and react with the solvent. This type of mechanism is indicated below.

\[
RX + \text{GeH}_3^- \rightarrow \left[ X^- \right. \begin{array}{c} \text{H} \end{array} \left. \begin{array}{c} \text{GeH}_2 \end{array} \right] \xrightarrow{\text{+}} \text{RH} + X^- + \frac{1}{n}(\text{GeH}_2)_n
\]

Both the hydrogen-substitution reaction and the germyl-substitution reaction accelerate as one goes from a chloride to a bromide to an iodide but the acceleration is more pronounced in the former than in the latter reaction. Thus much more benzene is obtained from bromobenzene than from chlorobenzene, more methylgermane is obtained from dibromomethane than from dichloromethane, and a detectable amount of methane is obtained from iodomethane, but not from bromomethane. The trends in rates may be explained by considering the activated complexes in the two types of reaction. Nucleophilic attack by germyl anion at either
carbon or halogen results in an activated complex in which the carbon-
halogen bond is partially broken. The carbon-iodine bond is weaker
than the carbon-chlorine bond; therefore both types of reaction are en-
hanced when a carbon-iodine, rather than a carbon-chlorine, bond is
involved. However, there are further effects (which do not apply to
the nucleophilic attack on carbon) that enhance nucleophilic attack on
halogen when one goes from a chloride to an iodide. Because iodine is
less electronegative than chlorine, a bonded iodine atom is more sus-
ceptible to attack by an anion that is a bonded chlorine atom. Further-
more, iodine is "softer" than chlorine and therefore might be expected
to react more readily with the soft germyl anion. Thus one would ex-
pect that the overall accelerative effect of going from a chloride to
a bromide to an iodide would be more pronounced for the hydrogen-substi-
tution reaction than for the germyl-substitution reaction.

It should be pointed out that a free-radical mechanism, analogous
to that proposed for the reaction of organic halides with organotin
hydrides,\textsuperscript{29} will account for most of the experimental results. Trace
amounts of peroxo compounds in the diglyme solvent could act as initi-
ators. However, in the reaction of neophil bromide, the intermediate
neophil radical would be expected to rearrange to yield iso-butylben-
zene;\textsuperscript{30,31} only tert-butylbenzene was formed. Also, one might expect
phenyl radicals to cleave the diglyme in the reaction of bromobenzene
with potassium germyl-\textsuperscript{d}{\textsubscript{3}}; however, no such cleavage products were found.
These two negative observations are consistent with a radical mechanism
if one assumes that abstraction of a germanium-bonded hydrogen by the
intermediate radical is so fast that all other reactions, including rearrangement, are precluded.

Certain mechanisms for the hydrogen substitution reaction can be ruled out. The aromatic halide substitutions cannot be explained by a benzene intermediate because of the observed reactivity of 2-bromo-m-xylene toward potassium germyl. Organogermane formation and subsequent decomposition to GeH₂ and hydrocarbon is ruled out because phenylgermane and digermylmethane are known to be stable compounds, even in basic solution. Finally, nucleophilic attack on carbon by a hydrogen atom of the germyl anion would be expected to show the same trends as nucleophilic attack by the germanium atom of the germyl anion; one would expect the same relative yields of hydrocarbon and organogermane regardless of the organic halide involved. This result was not found.

In addition to undergoing the germyl and hydrogen substitutions, dihalo- and trihalomethanes react in yet another way with potassium germyl, i.e., to form digermyl ethane and digermyl propane. The latter products can be explained by the deprotonation of the halomethanes by potassium germyl. Indeed, the yield of GeH₄ increases on going from dibromomethane to dichloromethane to trichloromethane, in accord with the expected acid strengths of the halomethanes. The carbanions formed can go on to attack unreacted halomethane, forming haloethanes, halo-propanes, etc., which then react with germyl anion for form the corresponding digermylalkanes.

\[ \text{CH}_2\text{X}_2 + \text{GeH}_3^- \rightarrow \text{GeH}_4 + \text{CHX}_2^- \]
-21-

\[ \text{CHX}_2^- + \text{CH}_2X_2 \rightarrow \text{CH}_2XCHX_2 + X^- \]

\[ \text{CH}_2XCHX_2 + 3\text{GeH}_3^- \rightarrow \text{GeH}_2\text{CH}_2\text{CH}_2\text{GeH}_3 + \frac{1}{n}(\text{GeH}_2) + 3X^- \]

etc.

Acknowledgement. - This research was supported by the U.S. Atomic Energy Commission.
References

(2) G. K. Teal and C. A. Kraus, ibid., 72, 4706 (1950).
(3) S. N. Glarum and C. A. Kraus, ibid., 72, 5398 (1950).
(5) Materials available from Varian Aerograph, Walnut Creek, California.
(22) a. C. A. Kraus and H. S. Nutting, ibid., 54, 1623 (1932).
    See p. 293.
    31, 2027 (1966).
(27) Ref. 24, p. 79.
(28) Ref. 24, p. 475.
    (1964).
(34) Unreported observations
Figure Captions

Figure 1.- Apparatus for the preparation and filtration of KGeH$_3$ solutions.

Figure 2.- Apparatus for pipetting KGeH$_3$ solutions.
APPENDIX

An Attempt to Prepare Potassium Germylsulfonate

Abstract

An attempt was made to prepare $\text{KGeH}_3\text{SO}_3$ by the reaction of potassium germyl with sulfur trioxide or with sulfur trioxide-amine adducts. Adducts of $\text{SO}_3$ with dimethylformamide, trimethylamine, and pyridine were tried; solvents used were diglyme, dimethylformamide and dimethyl sulfoxide. In all cases reaction occurred but indications are that if $\text{KGeH}_3\text{SO}_3$ can be made, it decomposes at temperatures well below 0°.
Potassium germyl has been shown to react with Lewis acids. The compounds potassium germyltrihydroborate (KGeH₃BH₃)¹ and potassium 2-germaacetate (KGeH₃CO₂)² can be prepared by the reactions of potassium germyl with diborane and carbon dioxide respectively. However, the reactions of potassium germyl with the Lewis acids CS₂, BH₃CO, BF₃,³ and SO₂⁴ are more complicated and yield decomposition products. An attempt to react SO₃ with KGeH₃ solutions in monoglyme and diglyme yielded a white solid which rapidly yellowed at 0°, giving off germane.⁵ Making the hypothesis that the desired product KGeH₃SO₃ was incompatible with unreacted SO₃, we investigated the reactions of KGeH₃ with amine adducts of SO₃ in the hope that these less reactive species would not decompose the product. We found that even under these milder conditions, we were unable to prepare potassium germylsulfonate.
Reagents. - Standardized diglyme solutions of potassium germyl were prepared as described in the first part of this thesis. It was found that the same techniques could be used to prepared standardized solutions in dimethylformamide (DMF).

Spectroquality dimethylformamide and reagent grade dimethyl sulfoxide were obtained from Matheson, Coleman and Bell and used without further purification. Trimethylamine was obtained from the Matheson Company and passed through a -78° trap prior to use. Its purity was checked by vapor pressure measurements at 0° (v.p. 670 torr; literature value 680 torr). Technical grade sulfur trioxide-pyridine was obtained from Aldrich Chemical Company and used without purification.

A standardized, anhydrous solution of SO$_3$:DMF in DMF was prepared by first adding reagent grade sulfan (obtained from Baker and Adamson) to DMF at 0° in a dry nitrogen atmosphere. DMF was then vacuum-distilled from this solution into a flask identical to those used for storing potassium germyl solutions. SO$_3$ was then vacuum-distilled out of sulfan and condensed onto the DMF at -78°. The solution was warmed to 0° with shaking; the SO$_3$ dissolved leaving a clear solution with a small amount of undissolved SO$_3$:DMF. The solution was stored at 0° and aliquots of the solution were taken using the same apparatus used for pipetting KGeH$_3$ solutions. By distilling water into an aliquot of the solution, the titer was found to be 1.30 M SO$_3$:DMF. This did not change over a period of one month.
Sulfur trioxide:trimethylamine (SO₃:TMA) was prepared by the reaction of excess trimethylamine with a stirred solution of SO₃:DMF. A white precipitate formed which was filtered and dried under vacuum. The material melted with decomposition at 238°-240° (lit. 240°). Volatile products were identified by their infrared spectra and vapor pressures.

Reaction of SO₃ with KGeH₃ in DMF. - An unstandardized solution of KGeH₃ in DMF was prepared from 2.5 mols GeH₄ in 15 ml solvent. The solution was stirred for one hour at 0° in the presence of excess SO₃ vapor. The solution gradually yellowed and no precipitation occurred. The solution was then distilled with pumping, through -130° and -196° traps. The -130° trap contained SO₃ and DMF, the -196° trap GeH₄. Monoglyme was then condensed onto the frozen solution; on warming to room temperature a small amount of yellow-white precipitate formed.

Reaction of SO₃:DMF with KGeH₃. - To 4.12 mmol KGeH₃ in diglyme at 0° was added dropwise with stirring 4.00 mmol SO₃:DMF in DMF. A white precipitate formed immediately; after standing 5 minutes, it and the solution had yellowed. The reaction mixture was distilled, with pumping, for 15 minutes through -130° and -196° traps. The -130° trap contained DMF and a trace of Ge₂H₆; the -196° trap contained GeH₄ (0.82 mmol). The reaction mixture was filtered under vacuum. The precipitate was dried under vacuum and a portion dissolved in basic H₂O₂ and analyzed volumetrically for germanium; it was found to be 11% Ge by weight (calculated for KGeH₃SO₃, 37.0 %). Another portion of solid was weighed and sealed in an evacuated ampoule equipped with a break-off seal. The
Ampoule was kept in a 400° muffle furnace for one hour, after which the vessel was opened on the vacuum line and the volatiles analyzed. A trace of hydrogen and \( \text{SO}_2 \) were found along with diglyme pyrolysis products. The pyrolyzed solid was black and was insoluble in basic \( \text{H}_2\text{O}_2 \).

The same amounts of \( \text{KGeH}_3 \) and \( \text{SO}_3\cdot\text{DMF} \) were allowed to react as above, the only difference being the temperature, -20°; the reaction mixture was distilled for 30 minutes at -22° and 0.40 mmol GeH\(_4\) were measured. On warming to 0° and letting the mixture decompose for 60 minutes, another 0.72 mmol GeH\(_4\) were measured. The dried precipitate was found to have a Ge content of 8\% by weight.

To \( \text{KGeH}_3 \) (1.56 mmol) in DMF at 0° was added \( \text{SO}_3\cdot\text{DMF} \) (2.30 mmol) in DMF. No apparent change occurred. The colorless solution was distilled, with pumping, for 15 minutes through -78°, -130° and -196° traps. The -78° trap contained solvent only, the -130° trap Ge\(_2\)H\(_6\) (0.11 mmol) and the -196° trap GeH\(_4\) (0.27 mmol). Pumping was continued to remove most of the solvent; the solution became slightly yellow. Monoglyme was condensed onto the solution at -196° and the solution allowed to warm to room temperature; a yellow-white precipitate had formed.

The experiment was repeated adding the reagents in reverse order. The mixture was distilled, with pumping, for 3 hours through -78°, -130° and -196° traps. The -78° trap contained solvent only, the -130° trap Ge\(_2\)H\(_6\) (0.34 mmol) and the -196° trap GeH\(_4\) (0.56 mmol). The solution did not yellow. Treatment of the solution with monoglyme yielded a
white precipitate which was filtered, dried and analyzed for germanium; it contained 5% Ge by weight.

Reaction of $\text{SO}_3$-pyridine with KGeH$_3$. To 1.66 mmol $\text{SO}_3$-pyridine at 0° was added with stirring an unstandardized solution of KGeH$_3$ in DMF prepared from 2.3 mmol GeH$_4$. The $\text{SO}_3$-pyridine dissolved immediately forming a pink solution and, over a period of 15 minutes, GeH$_4$ (0.40 mmol). Monoglyme was condensed onto the solution at -196° and on warming, a white precipitate formed.

Reaction of $\text{SO}_3$-TMA with KGeH$_3$. To 0.806 mmol $\text{SO}_3$-TMA at -45° was added 1.18 mmol KGeH$_3$ in DMF. The slurry was stirred for 2 hours at -45° after which it was Toepler-pumped through -78°, -130° and -196° traps. No non-condensable gas was detected. The -78° trap and the -130° trap contained a trace of TMA while the -196° trap contained GeH$_4$ (0.09 mmol). The solution was allowed to warm to 0° and after 3 hours pumping was resumed; by this time the solution had yellowed. No non-condensable gas was detected but the -130° trap contained TMA (0.269 mmol) and the -196° trap GeH$_4$ (0.362 mmol). The solution was warmed to room temperature and allowed to decompose overnight. When pumping was resumed, H$_2$ (0.140 mmol), TMA (0.369 mmol), and GeH$_4$ (0.10 mmol) were found. To the mixture was added 5 ml 1M H$_3$PO$_4$; the mixture was kept at 50° for 30 minutes after which it was pumped on again. The only volatile products were traces of GeH$_4$, Ge$_2$H$_6$, and Ge$_3$H$_8$. There was no trace of SO$_2$. The reaction mixture was filtered and a 0.2 M BaCl$_2$ solution added to the solution; a white precipitate formed indicating the presence of
sulfate anion. This solution was centrifuged and a drop of 3% H₂O₂ added to the supernate. No precipitate formed, indicating the absence of sulfite anion.

When SO₃:TMA was stirred at -45° with potassium germyl in diglyme a small amount of germane was again produced; on warming to 0° again decomposition occurred. In dimethylsulfoxide at room temperature, KGeH₃ and SO₃:TMA yielded a solution which darkens from yellow to blood red after 15 minutes. Large amounts of GeH₄ and TMA were produced.

**Evidence for Sulfide Anion Formation.** - To approximately 1 mmol SO₃:TMA at 0° was added approximately 1 mmol KGeH₃ in DMF. The solution yellowed immediately and was warmed to room temperature where it was allowed to react further for four hours; a small amount of yellow-white precipitate was formed. Most of the DMF was pumped off overnight along with the other volatiles after which 10 ml monoglyme was added to the mixture; a heavier yellow-white precipitate was formed. The mixture was filtered under vacuum; to a portion of the precipitate was added 1 ml 6 M HCl and a small piece of zinc. A piece of lead acetate-treated filter paper held over the solution turned slightly brown, indicating the presence of sulfide anion. The presence of sulfide anion was also observed in precipitates formed by the reaction of SO₃:DMF with KGeH₃ in diglyme.
Results and Discussion

We have been unable to synthesize KGeH$_3$SO$_3$ from the reaction of KGeH$_3$ with SO$_3$ adducts. The formation of trimethylamine from SO$_3$:trimethylamine indicates the desired reaction may have occurred, followed by decomposition of the anticipated product. Decomposition yields products that do not form SO$_2$ when treated with aqueous acid; therefore formation of HSO$_3^-$, S$_2$O$_6^{2-}$, and SO$_2$:amine complexes must be ruled out. The presence of S$^-$ and SO$_4^{2-}$ in reaction mixtures suggests possible decomposition reactions:

\[
2\text{KGeH}_3\text{SO}_3 \rightarrow \text{GeS} + \frac{1}{n}(\text{GeH}_2)_n + \text{K}_2\text{SO}_4 + 2\text{H}_2\text{O}
\]

\[
\text{KGeH}_3\text{SO}_3 + \text{KGeH}_3 \rightarrow \text{GeS} + \frac{1}{n}(\text{GeH}_2)_n + 2\text{KOH} + \text{H}_2\text{O}
\]

The possibility also exists that KGeH$_3$SO$_3$ is unstable in the presence of amines. This condition might be avoided by reacting KGeH$_3$ with salts of chlorosulfonate or dithionate anions.
References

(2) P. M. Kuznesof and W. L. Jolly, ibid., 7, 2574 (1968).
(4) Unreported observations
(5) Unreported observations of M. Goodrow
Acknowledgments

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