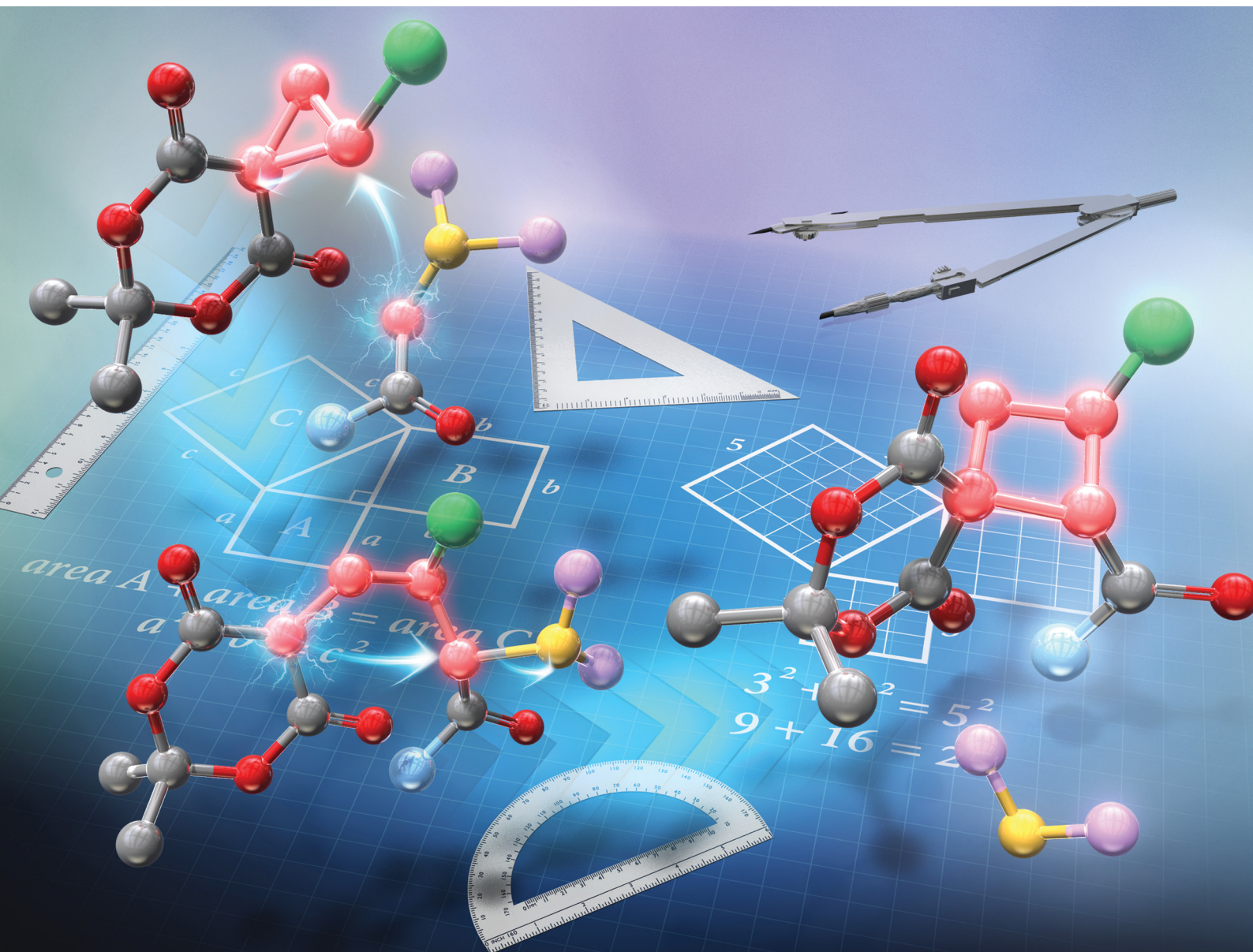


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ISSN 1359-7345

COMMUNICATION

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Cite this: *Chem. Commun.*, 2024, 60, 4537

Received 11th December 2023,  
Accepted 7th March 2024

DOI: 10.1039/d3cc06033k

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# Ring expansion of spirocyclopropanes with stabilized sulfonium ylides: highly diastereoselective synthesis of cyclobutanes†

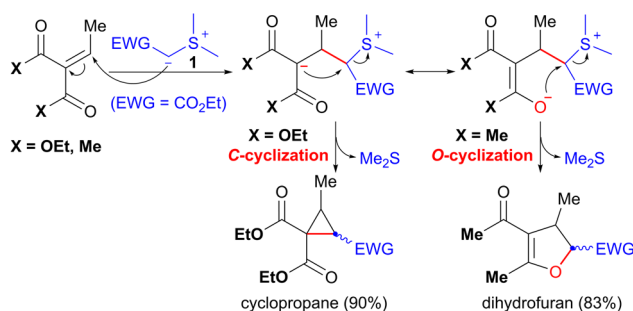
Hisanori Nambu,<sup>a</sup> Yuta Onuki,<sup>a</sup> Kana Aso,<sup>a</sup> Momoka Kanamori,<sup>a</sup> Keisuke Tomohara,<sup>b</sup> Kiyoshi Tsuge<sup>c</sup> and Takayuki Yakura<sup>\*a</sup>

**A novel method was devised for regioselective ring expansion of Meldrum's acid-derived spirocyclopropanes to spirocyclobutanes with stabilized sulfonium ylides, affording 1,2-trans-disubstituted 6,8-dioxaspiro[3.5]nonane-5,9-diones in up to 87% yields without the formation of any isomers. The aforementioned reaction was also applied to the barbituric acid-derived spirocyclopropane, resulting in the formation of the corresponding cyclobutanes.**

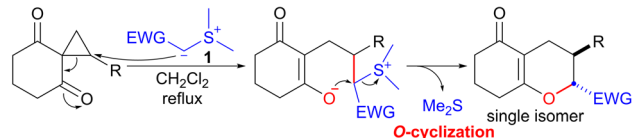
Sulfonium ylides stabilized by electron-withdrawing groups (EWG) have been used as a versatile methylene synthon in the synthesis of a variety of carbo- and heterocyclic compounds.<sup>1,2</sup> As a pioneering work, Payne reported that the reaction of  $\alpha,\beta$ -unsaturated diethylmalonate with EWG-stabilized sulfonium ylide **1** (EWG = CO<sub>2</sub>Et) afforded cyclopropane in 90% yield (Scheme 1A).<sup>3</sup> In this reaction, the Michael addition of **1** followed by S<sub>N</sub>2-type cyclization of the carbanion (C-cyclization) proceeded with the concomitant release of the sulfide. In contrast, the reaction of the corresponding 1,3-diketone with stabilized sulfonium ylide **1** unexpectedly produced dihydrofuran in 83% yield through enolate cyclization (O-cyclization, Scheme 1A).<sup>3</sup> The regioselectivity of these reactions may be attributed to the inherent difference between esters and ketones. Recently, we reported the ring-opening cyclization of spirocyclopropanes<sup>4</sup> with EWG-stabilized sulfonium ylides **1** to afford hexahydrobenzopyranone as a single isomer *via* the regioselective ring-opening of cyclopropane with sulfonium ylide **1** and subsequent S<sub>N</sub>2-type O-cyclization

(Scheme 1B, eqn (1)).<sup>4f</sup> Considering the similar reactivity of cyclopropane and carbon-carbon double bonds, we expected that the reaction of ester-derived spirocyclopropane **2** with stabilized sulfonium ylide **1** would provide spirocyclobutane **3** through C-cyclization (Scheme 1C). Because cyclobutane is a useful scaffold found in several biologically active natural products and pharmaceutically active compounds,<sup>5</sup> the development of a synthetic method for cyclobutane is currently the subject of intense research.<sup>6</sup> Although several instances of

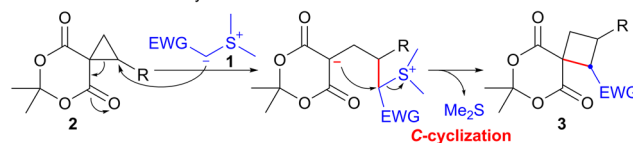
**A. Background:** Regioselectivity in reactions of  $\alpha,\beta$ -unsaturated carbonyl compounds with stabilized sulfonium ylide **1**



**B. Previous work:** Ring-opening cyclization of cyclohexane-1,3-dione-2-spirocyclopropanes with stabilized sulfonium ylides **1**



**C. This work:** Ring expansion of Meldrum's acid-derived spirocyclopropanes **2** with stabilized sulfonium ylides **1**



**Scheme 1** Reactions of various carbonyl compounds with stabilized sulfonium ylides **1** as nucleophiles.

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† Electronic supplementary information (ESI) available: General procedures, analytical data, and NMR spectra (PDF). CCDC 2312076 (3a). For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3cc06033k>

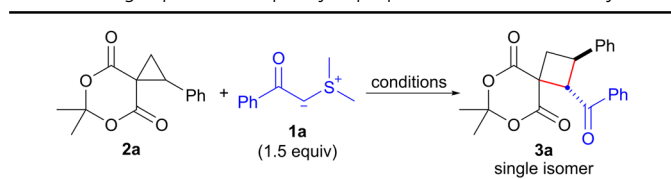


cyclopropane to cyclobutane ring expansion have been documented thus far,<sup>7,8</sup> to the best of our knowledge, there have been no examples of sulfonium ylide-mediated ring expansion (*C*-cyclization).<sup>9</sup> Herein, we describe the ring expansion of Meldrum's acid-derived spirocyclopropanes **2** to spirocyclobutanes **3** using EWG-stabilized sulfonium ylides **1** (Scheme 1C).

Initially, we examined the reaction of 6,6-dimethyl-1-phenyl-5,7-dioxaspiro[2.5]octane-4,8-dione (**2a**)<sup>10</sup> with dimethylsulfonium benzoylmethylide (**1a**) as an EWG-stabilized sulfonium ylide (Table 1). The ring expansion of **1a** proceeded under the reaction conditions previously reported by our group (1.5 equiv. of **1a** in refluxing CH<sub>2</sub>Cl<sub>2</sub>),<sup>4f</sup> affording 1-benzoyl-7,7-dimethyl-2-phenyl-6,8-dioxaspiro[3.5]nonane-5,9-dione (**3a**) after 24 h in 75% yield (entry 1). Notably, no isomer formation was observed during this process. The structure of **3a** including its stereochemistry was confirmed by a single-crystal X-ray diffraction analysis. This analysis revealed that the structure corresponds to that of cyclobutane with a 1,2-*trans* configuration (see ESI† for details). Screening of the solvents at reflux revealed that benzene and halogenated solvents, such as dichloromethane and 1,2-dichloroethane, were suitable for this reaction (entry 1 *vs.* entries 2–5). Finally, we found that chlorobenzene at 80 °C was the most effective and afforded **3a** in 86% yield after 6 h (entry 6).

After determining the optimal conditions, we investigated the reaction of spirocyclopropane **2a** using a range of sulfonium ylides **1** that are stabilized by carbonyl functional groups (Table 2). The reaction with 1.5 equiv. of *p*-methoxybenzoyl sulfonium ylide **1b** in chlorobenzene at 80 °C afforded the corresponding spirocyclobutane **3b** as the sole product after 6 h in 74% yield (entry 2). The use of *m*- and *o*-methoxybenzoyl sulfonium ylides **1c**<sup>11</sup> and **1d**<sup>12</sup> provided the corresponding products **3c** and **3d** in 86% and 87% yields, respectively (entries 3 and 4). The reaction with sulfonium ylide **1e** bearing a *p*-nitro group as a strong EWG decreased the product yield, and a significantly longer reaction time was required to achieve full conversion (61% yield, 24 h, entry 5 *vs.* entry 1). In contrast, the reaction with *p*-chlorobenzoyl sulfonium ylide **1f** under the optimized conditions proceeded smoothly to completion within 5 h, furnishing **3f** in 83% yield (entry 6). We also

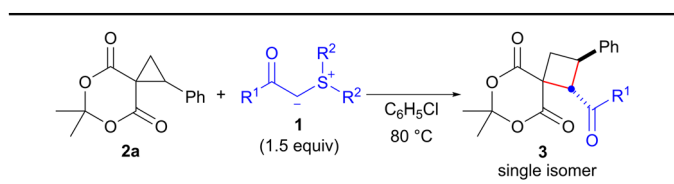
Table 1 Ring expansion of spirocyclopropane **2a** with sulfonium ylide **1a**



Entry	Solvent	Temp.	Time (h)	Yield <sup>a</sup> (%)
1	CH <sub>2</sub> Cl <sub>2</sub>	Reflux	24	75
2	EtOAc	Reflux	4.5	60
3	Benzene	Reflux	5	74
4	CH <sub>3</sub> CN	Reflux	24	59
5	ClCH <sub>2</sub> CH <sub>2</sub> Cl	Reflux	7	75
6	C <sub>6</sub> H <sub>5</sub> Cl	80 °C	6	86

<sup>a</sup> Isolated yield.

Table 2 Ring expansion of spirocyclopropane **2a** with sulfonium ylides **1a–h**



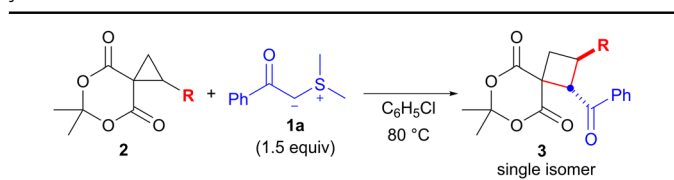
Entry	Sulfonium ylide		Time (h)	Product	
	R <sup>1</sup>	R <sup>2</sup>			Yield <sup>a</sup> (%)
1	<b>1a</b>	C <sub>6</sub> H <sub>5</sub>	Me	6	<b>3a</b> 86
2	<b>1b</b>	<i>p</i> -MeOC <sub>6</sub> H <sub>4</sub>	Me	6	<b>3b</b> 74
3	<b>1c</b>	<i>m</i> -MeOC <sub>6</sub> H <sub>4</sub>	Me	6	<b>3c</b> 86
4	<b>1d</b>	<i>o</i> -MeOC <sub>6</sub> H <sub>4</sub>	Me	6	<b>3d</b> 87
5	<b>1e</b>	<i>p</i> -NO <sub>2</sub> C <sub>6</sub> H <sub>4</sub>	Me	24	<b>3e</b> 61
6	<b>1f</b>	<i>p</i> -ClC <sub>6</sub> H <sub>4</sub>	Me	5	<b>3f</b> 83
7	<b>1g</b>	Me	–(CH <sub>2</sub> ) <sub>4</sub> –	24	<b>3g</b> 36
8	<b>1h</b>	EtO	Me	23	<b>3h</b> 53

<sup>a</sup> Isolated yield.

investigated the suitability of an acetyl sulfonium ylide for this reaction. To this end, we used tetrahydrothiophenium acetylmethylide (**1g**) because of the difficulty in preparing dimethylsulfonium acetylmethylide. The reaction of **2a** with **1g** afforded the desired product **3g** as a single isomer, albeit with a prolonged reaction time and lower yield (24 h, 36% yield, entry 7 *vs.* entry 1). Moreover, ethoxycarbonyl group-substituted sulfonium ylide **1h** was used in the present protocol, and the corresponding cyclobutane **3h** was obtained in 53% yield after 23 h (entry 8).

Next, we examined the scope of the reaction with the spirocyclopropane substrates **2** using benzoyl-substituted sulfonium ylide **1a** (Table 3). Treatment of spirocyclopropanes **2b**, **2c** and **2d**, which possess *p*-acetoxy-, *p*-methyl-, and *p*-bromophenyl groups on the cyclopropane, respectively, with **1a** under the optimized conditions (chlorobenzene at 80 °C), afforded the corresponding products **3i**, **3j**, and **3k** in 64%–80% yields with perfect diastereoselectivities (entries 1–3). The

Table 3 Ring expansion of spirocyclopropanes **2b–h** with sulfonium ylides **1a**



Entry	Spirocyclopropane		Time (h)	Product	
	R				Yield <sup>a</sup> (%)
1	<b>2b</b>	<i>p</i> -AcOC <sub>6</sub> H <sub>4</sub>	3	<b>3i</b>	64
2	<b>2c</b>	<i>p</i> -MeC <sub>6</sub> H <sub>4</sub>	6	<b>3j</b>	74
3	<b>2d</b>	<i>p</i> -BrC <sub>6</sub> H <sub>4</sub>	6	<b>3k</b>	80
4	<b>2e</b>	<i>m</i> -MeOC <sub>6</sub> H <sub>4</sub>	12	<b>3l</b>	69
5	<b>2f</b>	2-naphthyl	48	<b>3m</b>	80
6	<b>2g</b>	Vinyl	24	<b>3n</b>	68
7	<b>2h</b>	H	24	<b>3o</b>	26

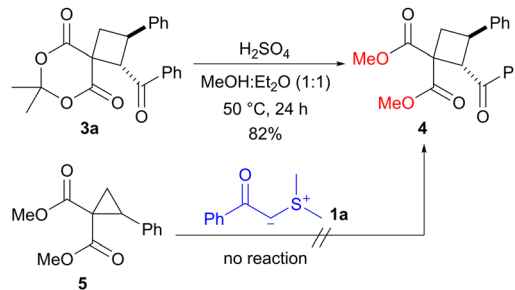
<sup>a</sup> Isolated yield.



reaction of *m*-methoxyphenyl-substituted spirocyclopropane **2e** for 12 h provided cyclobutane **3l** in 69% yield (entry 4). Although spirocyclopropane **2f**, which possesses a 2-naphthyl group, required a relatively long reaction time (48 h), **3m** was obtained in a good yield (80%, entry 5). There was a concern that the use of vinyl-substituted spirocyclopropane **2g** would compete with the conjugate addition, but the reaction of **2g** proceeded uneventfully and afforded the desired product **3n** in 68% yield (entry 6). Finally, the reaction of the simple spirocyclopropane **2h** ( $R = H$ )<sup>13</sup> was investigated (entry 7). A 2',3'-nonsubstituted spirocyclopropane was found to be less reactive than an aryl-substituted one,<sup>4e</sup> which resulted in a lower yield of product **3o** (26% yield).

A plausible mechanism for the ring expansion of spirocyclopropane **2** with sulfonium ylide **1**, stabilized by an acyl group, is shown in Scheme 2. The ring opening of spirocyclopropane **2** would proceed through the nucleophilic attack of the carbanion in **1** on the electrophilic cyclopropane carbon possessing an  $R^1$  substituent in **A**. This reaction would lead to the formation of betaine intermediates **B** and **C**.  $S_N2$ -type *C*-cyclization of the carbanion in **B** would occur smoothly to afford *trans*-product **3** with the concomitant release of dimethyl sulfide. In contrast, the *C*-cyclization of **C** would hardly proceed owing to the severe steric repulsion between the acyl group ( $R^2CO$ ) and substituent  $R^1$  in **C**. Consequently, intermediate **C** could be converted into cyclization precursor **B** through reversible intramolecular proton transfer *via* the stabilized sulfonium ylide **D**,<sup>14,15</sup> finally providing *trans*-isomer **3**.

To demonstrate the utility of the present protocol, we examined the conversion of spirocyclobutane **3a** into highly substituted non-spiro cyclobutane **4** (Scheme 3). The treatment of **3a** with sulfuric acid in methanol/diethyl ether (1 : 1) at 50 °C led to a transesterification process, resulting in the formation of dimethyl ester. The reaction yielded the corresponding cyclobutane **4** in 82% yield. Since the reaction of dimethyl 2-phenylcyclopropane-1,1-dicarboxylate (**5**) with sulfonium

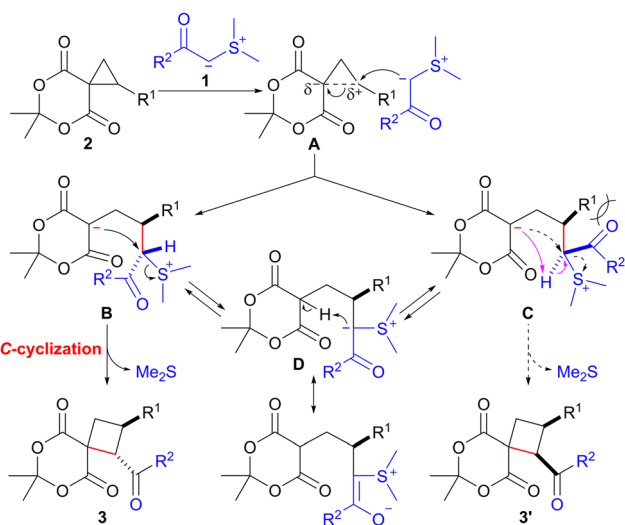


Scheme 3 Conversion of spirocyclobutane **3a** into cyclobutane **4**.

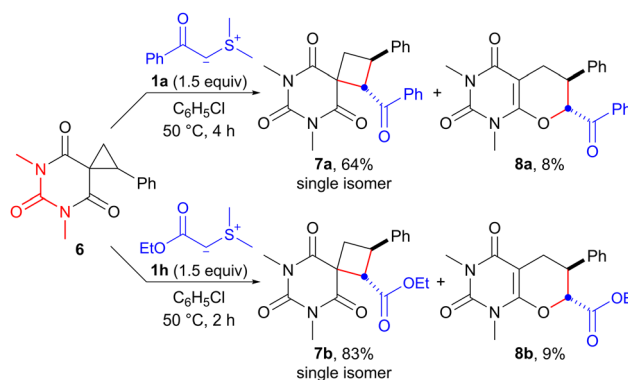
ylide **1a** did not proceed,<sup>4f,16</sup> spiro form **3a** was required for the synthesis of diester **4**. This ring-expansion reaction of spirocyclopropanes could be a useful method for the preparation of substituted cyclobutanes.

Having achieved ring expansion of ester-derived spirocyclopropanes, we further investigated the reaction of an amide-derived spirocyclopropane with an EWG-stabilized sulfonium ylide. The reaction of spirocyclopropane **6**,<sup>17</sup> derived from barbituric acid, with sulfonium ylides **1a** and **1h** in chlorobenzene proceeded smoothly at 50 °C to provide the corresponding spirocyclobutanes **7a** and **7b** in 64% and 83% yields, respectively (Scheme 4). Interestingly, unexpected products **8a** and **8b**, which indicated that  $S_N2$ -type *O*-cyclization of the enolate ion instead of the carbanion would occur, were also obtained in 8% and 9% yields, respectively. Although the results are still preliminary, the reaction of barbituric acid-derived spirocyclopropane with sulfonium ylide exhibits promise as a synthetic method of spirobarbiturate cyclobutane analogs. These compounds have potential as pharmaceutical agents.<sup>18</sup>

In conclusion, we devised a novel method for regioselective ring expansion of cyclopropanes to cyclobutanes using stabilized sulfonium ylides. Meldrum's acid-derived spirocyclobutanes with EWG-stabilized sulfonium ylides afforded the corresponding spirocyclobutanes as single diastereomers in yields of up to 87%. The present reaction provides an efficient route to highly substituted cyclobutanes. To the best of our knowledge, this is the first example of a ring expansion of cyclopropanes with sulfonium ylides. This reaction may be envisaged as a



Scheme 2 Plausible reaction mechanism.



Scheme 4 Ring expansion of barbituric acid-derived spirocyclopropane **6** with sulfonium ylides **1a** and **1h**.



formal [3+1] cycloaddition, facilitating the construction of the four-membered ring system.<sup>19</sup> The expansion reaction could be applied to the transformation of barbituric acid-derived spirocyclopropane into the corresponding spirocyclobutane. Ongoing efforts are being made to apply the present method to the synthesis of a variety of cyclobutane derivatives.

This research was supported, in part, by a Grant-in-Aid for Scientific Research (C) (Grant No. 21K05048) from the Ministry of Education, Culture, Sports, Science and Technology, Japan and by a Grant from Hoansha Foundation (Osaka, Japan).

## Conflicts of interest

There are no conflicts to declare.

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