Title	BrF3-KHF2: An air-stable fluorinating reagent
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Citation	Journal of fluorine chemistry, 168, 55-60 https://doi.org/10.1016/j.jfluchem.2014.08.019
Issue Date	2014-12
Doc URL	http://hdl.handle.net/2115/57927
Type	article (author version)
File Information	BrF3.pdf

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BrF₃-KHF₂: An air-stable fluorinating reagent

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Keywords: stable fluorinating reagent, KHF₂, BrF₃, desulfurizing fluorination

Abstract

BrF₃-KHF₂, an air-stable solid prepared from BrF₃ and KHF₂, was used in the various fluorination reactions, including desulfurizing fluorination reactions of benzylic sulfides, ketone and aldehyde dithioacetals, (phenylthio)glycosides, and trimethyl trithioorthocarboxylates. As the results, one to three fluorine atoms were selectively introduced to the substrates.

1. Introduction

Organofluorine compounds are widely used, as medicines, pesticides, functional materials, and so on [1]. They are generally prepared artificially using fluorinating reagents because organofluorine compounds are rare in nature. Therefore, the role of the fluorinating reagent is important for synthesizing desired organofluorine compounds, and many fluorinating reagents have been produced and used [2]. However, most of them are sensitive to moisture, and special skills and equipments are required for their use. Therefore, stable fluorinating reagent is desirable [3]. Recently, we reported the

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preparation of a new stable fluorinating reagent, IF₅-pyridine-HF, and its use in various fluorination reactions [4]. BrF₃ has been also used as a fluorinating reagent, and is more reactive than IF₅ [5]. Therefore, we attempted to synthesize a new stable fluorinating reagent from BrF₃.

2. Results and discussion

BrF₃ is known to make a complex of MBF₄ where M is Cs, Rb, K [6], or Me₄N [7]. However, their ability as a fluorinating reagent has not yet been studied. We attempted to synthesize a stable complex from BrF₃. Addition of BrF₃ to KF in CH₂Cl₂ was performed at -78 °C, and the cooling bath was removed. When the temperature reached room temperature, a violent exothermal reaction took place. As BrF₃ violently reacts with CH₂Cl₂ at room temperature, this result shows that free BrF₃ remains in the mixture and causes the violent reaction. On the other hand, when BrF₃ was added to an excess amount of KHF₂ in CH₂Cl₂, the exothermal reaction did not occur even after reaching room temperature. A slightly reddish supernatant was removed by decantation and the remaining solid was washed with CH₂Cl₂ several times. The remaining solvent was removed by blowing a nitrogen gas to the solid. The resulting pale yellow solid is air-stable and can be stored in a Teflon™ bottle in the refrigerator [8]. We applied this BrF₃-KHF₂ complex in various fluorination reactions.

2.1. Desulfurizing difluorination of benzylic sulfide **1** using BrF₃-KHF₂

Initially, BrF₃-KHF₂ was used in a desulfurizing difluorination reaction of benzylic sulfide. When 2-[4-chlorophenyl]thio]-1,2-diphenylethanone (**1a**) was added to a

suspension of BrF₃-KHF₂ in CH₂Cl₂ at room temperature, the solution color became dark red, and 2,2-difluoro-1,2-diphenylethanone (**2a**) was formed in 88% yield. Although the yield of **2a** was comparable to that obtained by using IF₅-pyridine-HF [4a], the reaction was completed in a shorter time (15 min versus 5 h) (Table 1).

Table 1

Comparison of reactivity of BrF₃-KHF₂ and IF₅-pyridine-HF in desulfurizing difluorination of **1a**^a

Entry	Fluorinating reagent	Reaction time	Yield of 2a (%) ^b
1	IF ₅ -pyridine-HF	5 h	88
2	BrF ₃ -KHF ₂	15 min	89

^a 2eq of fluorinating reagent to **1a** was used.

^b Isolated yield based on **1a** used.

In the reaction of BrF₃-KHF₂ with benzylic sulfides containing an electron-withdrawing group (**1b-d**), the corresponding desulfurizing difluorination products (**2a-c**) were obtained in high yields as shown in Table 2.

Table 2Desulfurizing difluorination of **1** with BrF₃-KHF₂^a

Entry	Substrate 1	Reaction time	Product 2	Yield (%) ^b
1	<p>1b</p>	3 h	<p>2b</p>	(84)
2	<p>1c</p>	15 min	<p>2c</p>	76 (85) ^c
3	<p>1d</p>	15 min	<p>2a</p>	91 (99) ^c

^aIf otherwise not mentioned, the reaction was carried out in CH₂Cl₂ at room temperature using 2 eq of BrF₃-KHF₂ to **1**.

^bIsolated yield based on **1** used. In parentheses, ¹⁹F NMR yield.

^c3eq of BrF₃-KHF₂ to **1** was used.

2.2. Reaction of aldehyde and ketone dithioacetal **3** with BrF₃-KHF₂

BrF₃-KHF₂ was also applied to the reaction with the ketone and aldehyde dithioacetals [3c, 5b, 9]. Reactions with diphenyl dithioacetals of aldehydes (**3a-c**) and ketones (**3d,e**) were completed in 1 h, and the corresponding *gem*-difluorides (**4a-e**) were obtained in

good yields, as shown in Table 3.

Table 3.

The reaction of aldehyde and ketone thioacetals with $\text{BrF}_3\text{-KHF}_2^{\text{a}}$

Entry	Substrate 3	Reaction time (min)	Product 4	Yield (%) ^b
1	3a	45	4a	(91)
2	3b	45	4b	81(90)
3	3c	15	4c	91(99)
4	3d	45	4d	(84)

5	3e	60	4e	(81) ^c
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^aIf otherwise not mentioned, the reaction was carried out at room temperature using 2.2 eq of BrF₃-KHF₂ to **3**.

^bIsolated yield based on **3**. In parentheses, ¹⁹F NMR yield.

^cThe reaction was carried out at 0 °C.

2.3. Synthesis of glycosyl fluorides **6** by the reaction of (phenylthio)glycosides **5** with BrF₃-KHF₂

Glycosyl fluorides have been widely used as glycosyl donors in glycosidation reactions [10]. They are generally prepared from the corresponding thioglycosides using a fluorination reagent with or without an oxidizing agent [11]. We applied BrF₃-KHF₂ for the synthesis of glycosyl fluorides (**6**) from the corresponding (phenylthio)glycosides (**5**). Both pyranosyl fluoride (**6a**) and furanosyl fluorides (**6b-d**) were prepared in good yield by the reaction of the corresponding (phenylthio)glycosides (**5a-d**) with BrF₃-KHF₂ in CH₂Cl₂ (Table 4). In the reaction with furanosyl derivatives, only one isomer was selectively formed (Entries 2-4).

Table 4.

The reaction of (phenylthio)glycosides with BrF₃-KHF₂ ^a

Entry	Substrate 5	Reaction	Product 6	Yield
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		time		(%) ^b
1	5a	4 h	6a	83(86) ^c ($\alpha : \beta = 63:37$)
2	5b	15 min	6b	89(94) ^d α only
3	5c	15 min	6c	87 β only
4	5d	15 min	6d	(66) β only

^aIf otherwise not mentioned, the reaction was carried out at room temperature using 1.1 eq of BrF₃-KHF₂ to **5**.

^bIsolated yield based on **5**. In parentheses, ¹⁹F NMR yield.

^c 1.5 eq of BrF₃-KHF₂ to **5** was used.

^d The reaction was carried out at 0 °C.

2.4. Reaction of trimethyl trithioorthocarboxylates **7** with BrF₃-KHF₂

A tris(methylthio)methyl group can be introduced to the electron rich aromatic ring and α -position of the ester group by a reaction with tris(methylthio)methyl cation species generated from dimethyl trithiocarbonate [12]. The reaction of *N,N*-dimethyl-4-[tris(methylthio)methyl]aniline (**7a**) with BrF₃-KHF₂ was completed in 15 min at 0 °C, and the tris(methylthio)methyl group was converted to the trifluoromethyl group. However, bromination at the aromatic ring took place concurrently and 2-bromo-*N,N*-dimethyl-4-(trifluoromethyl)aniline (**8a**) was formed in moderate yield (Entry 1 in Table 5). Similarly, in the reaction of *N*-methyl-3-tris(methylthio)methylindole (**7b**), 5-bromo-1-methyl-3-(trifluoromethyl)-1*H*-indole (**8b**) was obtained selectively (Entry 2). On the other hand, in the reaction of ethyl 2,2-dimethyl-3,3,3-tris(methylthio)propanoate (**7c**) with BrF₃-KHF₂, only two fluorine atoms were introduced and one methylthio group remained (Entry 3).

Table 5.

Reaction of trimethyl trithioorthocarboxylates **7** with BrF₃-KHF₂^a

Entry	Substrate 7	Reaction time	Product 8	Yield (%) ^b
1	<p>7a</p>	15 min	<p>8a</p>	52 (62)
2	<p>7b</p>	15 min	<p>8b</p>	76 (83)

3	<p>7c</p>	1 h	<p>8c</p>	81
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^aThe reaction was carried out in CH₂Cl₂ at 0 °C using 3.2 eq of BrF₃-KHF₂ to **7**.

^bIsolated yield based on **7**. In parentheses, ¹⁹F NMR yield.

3. Conclusion

A new air-stable fluorinating reagent, BrF₃-KHF₂, was prepared by the reaction of BrF₃ with KHF₂. The reagent was shown to be more reactive than the previously reported IF₅-pyridine-HF in desulfurizing difluorination reaction of the benzylic sulfide. The reagent was successively applied to desulfurizing fluorination reactions of dithioacetals, (phenylthio)glycosides, and trimethyl trithioorthocarboxylates.

4. Experimental

4.1. General

The melting points were measured with a Yanagimoto micro melting-point apparatus. The IR spectra were recorded using a JASCO FT/IR-410. The ¹H NMR (400 MHz) spectra, ¹⁹F NMR (376 MHz) spectra, and ¹³C NMR (100 MHz) were recorded in CDCl₃ on a JEOL JNM-A400II FT NMR and the chemical shift, δ, is referred to TMS (¹H, ¹³C) and CFC₃ (¹⁹F), respectively. BrF₃ in a cylinder was purchased from Galaxy Chemicals, LLC and used without purification. BrF₃ was transferred from cylinder to a Teflon™ bottle through a Teflon™ tube using nitrogen pressure. BrF₃ decomposes in air

by humidity under emitting HF fume, and should be handled in a bench hood with rubber-gloved hands under nitrogen atmosphere. BrF_3 reacts violently with most of organic solvents at room temperature and a special care is required for its use.

4.2. Preparation of $\text{BrF}_3\text{-KHF}_2$

To a suspension of KHF_2 (3.4 g, 44 mmol) in CH_2Cl_2 (10 mL) in a Teflon™ bottle, BrF_3 (3.0g, 22 mmol) was slowly added through a Teflon™ tube at $-78\text{ }^\circ\text{C}$. The resulting mixture was stirred at $-78\text{ }^\circ\text{C}$ for 30 min, and the cooling bath was removed and temperature was allowed to reach room temperature. A slightly reddish supernatant was removed using a Teflon™ pipette, and the remaining solid was washed with CH_2Cl_2 (10 mL) several times, until CH_2Cl_2 became almost colorless. The remaining solvent was removed by stirring under nitrogen stream for a few hours. The resulting pale yellow solid (5.4 g) was stored in a Teflon™ bottle in the refrigerator. It is slightly hygroscopic, and therefore, it should be used as quickly as possible to minimize contact with moisture.

4.2. Desulfurizing difluorination of benzylic sulfides **1** with $\text{BrF}_3\text{-KHF}_2$

4.2.1. 2,2-Difluoro-1,2-diphenylethanone (**2a**)

To a suspension of $\text{BrF}_3\text{-KHF}_2$ (129 mg) in CH_2Cl_2 (2.4 mL) in Teflon™ bottle, **1a** (101 mg, 0.3 mmol) in CH_2Cl_2 (1.0 mL) was added at room temperature, and the mixture was stirred at room temperature for 15 min. Then, H_2O (5 mL) was added to the reaction mixture and the resulting product was extracted with CH_2Cl_2 (5 mL X 3). The combined organic layer was washed with saturated aqueous NaHCO_3 (5 mL) and

saturated aqueous $\text{Na}_2\text{S}_2\text{O}_3$ (5 mL), and dried over MgSO_4 . After concentration under reduced pressure, **2a** was isolated by column chromatography (silica gel, hexane-ether) in 89% yield. IR (neat) 1703 (C=O), 1450, 1256, 1135 cm^{-1} ; ^1H NMR (400MHz, CDCl_3) δ 7.43-7.61 (m, 8H), 8.02-8.04 (m, 2H); ^{19}F NMR (376MHz, CDCl_3) δ -98.12 (s, 2F); ^{13}C NMR (100MHz, CDCl_3) δ 188.9 (t, $^2J_{\text{C-F}} = 30.7$ Hz), 134.2, 133.1 (t, $^2J_{\text{C-F}} = 24.9$ Hz), 132.1, 130.9, 130.3 (t, $^4J_{\text{C-F}} = 2.9$ Hz, 2C), 128.8 (2C), 128.6 (2C), 125.6 (t, $^3J_{\text{C-F}} = 5.8$ Hz, 2C), 116.9 (t, $^1J_{\text{C-F}} = 253.9$ Hz); HRMS (EI) calcd for $\text{C}_{14}\text{H}_{10}\text{F}_2\text{O}$ 232.0700, found 232.0683.

4.2.2. Butyl 2,2-difluoro-2-phenylacetate (**2b**)

IR (neat) 2963, 1764 (C=O), 1265, 1105 cm^{-1} ; ^1H NMR (400 MHz, CDCl_3) δ 7.62-7.60 (m, 2H), 7.49-7.45 (m, 3H), 4.24 (t, $J = 6.6$ Hz, 2H), 1.68-1.60 (m, 2H), 1.37-1.28 (m, 2H), 0.90 (t, $J = 7.4$ Hz, 3H); ^{19}F NMR (376 MHz, CDCl_3) δ -104.65 (s, 2F); ^{13}C NMR (100 MHz) δ 164.3 (t, $^2J_{\text{C-F}} = 35.7$ Hz), 132.8 (t, $^2J_{\text{C-F}} = 25.8$ Hz), 130.9, 128.6 (2C), 125.4 (t, $^3J_{\text{C-F}} = 6.2$ Hz, 2C), 113.4 (t, $^1J_{\text{C-F}} = 251.9$ Hz), 66.8, 30.2, 18.9, 13.5; HRMS (EI) calcd for $\text{C}_{12}\text{H}_{14}\text{F}_2\text{O}_2$ 228.0962, found 228.0956.

4.2.3. *N,N*-Diethyl-2,2-difluoro-2-phenylacetamide (**2c**)

IR (neat) 2979, 1669 (C=O), 1452, 1364, 1260, 1093 cm^{-1} ; ^1H NMR (400MHz, CDCl_3) δ 7.56 (d, $J = 7.0$ Hz, 2H), 7.44-7.49 (m, 3H), 3.42 (q, $J = 7.2$ Hz, 2H), 3.25 (q, $J = 7.2$ Hz, 2H), 1.17 (t, $J = 7.2$ Hz, 3H), 1.03 (t, $J = 7.0$ Hz, 3H); ^{19}F NMR (376MHz, CDCl_3) δ -95.41 (s, 2F); ^{13}C NMR (100MHz, CDCl_3) δ 162.7 (t, $^2J_{\text{C-F}} = 29.7$ Hz), 133.9 (t, $^2J_{\text{C-F}} = 23.6$ Hz), 130.7 (t, $^4J_{\text{C-F}} = 1.9$ Hz, 2C), 128.7, 125.1 (t, $^3J_{\text{C-F}} = 5.8$ Hz, 2C),

115.5 (t, $^1J_{C-F} = 251.5$ Hz), 42.0 (t, $^4J_{C-F} = 3.8$ Hz), 41.4, 13.7, 12.2; HRMS(EI) calcd for $C_{12}H_{15}F_2NO$ 227.1122, found 227.1128.

4.3. The reaction of aldehyde and ketone thioacetals **3** with BrF_3 - KHF_2

4.3.1. 1-(Difluoromethyl)naphthalene (**4a**)

The reaction was carried out as in the case of **2a** using 2.2 eq of BrF_3 - KHF_2 to **3a**, and yield of **4a** was determined to be 91 % by ^{19}F NMR using fluorobenzene as an internal standard. IR (neat) 1514, 1349, 1242 cm^{-1} ; 1H NMR δ 8.19-7.49 (m, 7H), 7.14 (t, $J = 55.8$ Hz, 1H); ^{19}F NMR δ -111.48 (d, $J = 56.0$ Hz, 2F) (lit.[13] -111.38 (d, $J = 55.2$ Hz)); ^{13}C NMR δ 133.7, 131.5, 129.7, 129.5 (t, $^2J_{C-F} = 21.1$ Hz), 128.7, 127.1, 126.4, 124.8 (t, $^3J_{C-F} = 8.6$ Hz), 124.6, 123.5, 115.4 (t, $^1J_{C-F} = 239.5$ Hz).

4.3.2. 4-(Difluoromethyl)-1,1'-biphenyl (**4b**)

White solid. mp 71-72 $^{\circ}C$ (lit.[14] 77.0-77.5 $^{\circ}C$); IR (KBr) 1414, 1380, 1226, 1077, 1024, 767 cm^{-1} ; 1H NMR δ 7.69-7.39 (m, 9H), 6.70 (t, $J = 56.5$ Hz, 1H); ^{19}F NMR δ -110.98 (d, $J = 57.3$ Hz, 2F); ^{13}C NMR δ 143.7 (t, $^5J_{C-F} = 1.9$ Hz), 140.2, 133.2 (t, $^2J_{C-F} = 22.1$ Hz), 128.9 (2C), 127.9, 127.4 (2C), 127.2 (2C), 126.0 (t, $^3J_{C-F} = 6.2$ Hz, 2C), 114.7 (t, $^1J_{C-F} = 238.5$ Hz).

4.3.3. Methyl 4-(difluoromethyl)benzoate (**4c**)

White solid. mp 38 $^{\circ}C$ (lit.[15] 36.5-37.0 $^{\circ}C$); IR (KBr) 1724 (C=O), 1442, 1281 cm^{-1} ; 1H NMR δ 8.13 (d, $J = 8.0$ Hz, 2H), 7.59 (d, $J = 8.1$ Hz, 2H), 6.70 (t, $J = 56.7$ Hz,

1H), 3.95 (s, 3H); ^{19}F NMR δ -112.86 (d, J = 57.9 Hz, 2F); ^{13}C NMR δ 166.2, 138.4 (t, $^2J_{\text{C-F}}$ = 22.5 Hz), 132.3, 129.9 (2C), 125.6 (t, $^3J_{\text{C-F}}$ = 6.3 Hz, 2C), 114.0 (t, $^1J_{\text{C-F}}$ = 240.9 Hz), 52.3.

4.3.4. 9,9-Difluoro-9H-fluorene (**4d**)

White solid. mp 46-48 °C (lit.[16] 47-48 °C). IR (KBr) 1918, 1454, 1261 cm^{-1} ; ^1H NMR δ 7.62 (d, J = 7.0 Hz, 2H), 7.56 (d, J = 7.3 Hz, 2H), 7.45 (dd, J = 7.5, 7.5 Hz, 2H), 7.33 (dd, J = 7.6, 7.6 Hz, 2H); ^{19}F NMR δ -112.12 (s, 2F); ^{13}C NMR δ 139.4 (t, $^3J_{\text{C-F}}$ = 5.3 Hz, 2C), 137.9 (t, $^2J_{\text{C-F}}$ = 25.1 Hz, 2C), 132.0 (2C), 128.7 (2C), 123.7 (2C), 123.2 (t, $^1J_{\text{C-F}}$ = 244.0 Hz), 120.3 (2C).

4.3.5. 2,2-Difluoroadamantane (**4e**)

White solid. mp 102-103 °C (lit.[17] 104-105 °C); IR (KBr) 2938, 2917, 1389, 1121 cm^{-1} ; ^1H NMR δ 2.18 (brs, 2H), 1.97 (brs, 2H), 1.94 (brs, 2H), 1.86 (brs, 2H), 1.78-1.72 (m, 6H); ^{19}F NMR δ -100.41 (s, 2F); ^{13}C NMR δ 125.5 (t, $^1J_{\text{C-F}}$ = 248.2 Hz), 36.6 (2C), 35.8 (t, $^2J_{\text{C-F}}$ = 4.0 Hz, 2C), 34.0 (t, $^3J_{\text{C-F}}$ = 4.0 Hz, 4C), 26.4.

4.4. The reaction of phenylthioglycosides **5** with $\text{BrF}_3\text{-KHF}_2$

4.4.1. 2,3,4,5-Tetra-O-acetyl-D-glucopyranosyl fluoride (**6a**)

The reaction was carried out as in the case of **2a** using 1.5 eq of $\text{BrF}_3\text{-KHF}_2$ to **5a**, and **6a** was isolated in 83% yield. The ratio of α -isomer : β -isomer was determined to be 63:37 from ^1H NMR spectra. (**6a- α**) mp 104-106 °C. IR (neat) 2958, 1748 (C=O),

1379, 1230, 1038 cm^{-1} . ^1H NMR δ = 5.76 (dd, J = 53.4, 2.76 Hz, 1H), 5.50 (dd, J = 9.9, 9.9 Hz, 1H), 5.16 (dd, J = 9.9, 9.9 Hz, 1H), 4.96 (ddd, J = 24.6, 10.4, 2.8 Hz, 1H), 4.29 (dd, J = 12.2, 3.8 Hz, 1H), 4.21-4.13 (m, 2H), 2.14 (s, 3H), 2.11 (s, 3H), 2.05 (s, 3H), 2.03 (s, 3H); ^{19}F NMR δ -150.34 (dd, J = 52.5, 25.3 Hz, 1F); ^{13}C NMR δ = 170.5, 169.9, 169.8, 169.4, 103.7 (d, $^1J_{\text{C-F}}$ = 228.8 Hz), 70.1 (d, $^2J_{\text{C-F}}$ = 24.8 Hz), 69.7 (d, $^3J_{\text{C-F}}$ = 4.1 Hz), 69.3, 67.2, 61.1, 20.6, 20.5, 20.4 (2C). (**6a- β**) mp 77-78 $^{\circ}\text{C}$. IR (neat) 2942, 1761, 1439, 1378, 1227, 1109, 1042 cm^{-1} ; ^1H NMR δ 5.37 (dd, J = 52.0, 6.1 Hz, 1H), 5.22-5.20 (m, 2H), 5.18-5.08 (brs, 1H), 4.29-4.20 (m, 2H), 3.93-3.88 (s, 1H), 2.11 (s, 6H), 2.05 (s, 6H); ^{19}F NMR δ -137.83 (1F, dd, J = 51.9, 10.4 Hz); ^{13}C NMR δ 170.5, 170.0, 169.2, 169.1, 106.1 (d, $^1J_{\text{C-F}}$ = 219.2 Hz), 72.0 (d, $^3J_{\text{C-F}}$ = 4.1 Hz), 71.7 (d, $^3J_{\text{C-F}}$ = 8.3 Hz), 71.1 (d, $^2J_{\text{C-F}}$ = 28.9 Hz), 67.4, 61.7, 20.6-20.5 (4C); HRMS (EI) calcd for $\text{C}_{14}\text{H}_{20}\text{O}_9\text{F}$ ($\text{M}^+ + \text{H}$) 351.1091, found 351.1115.

*2,3;5,6-di-O-Isopropylidene- α -D-mannofuranosyl fluoride (**6b- α**)*

IR (neat) 2989, 1374, 1212, 1130, 1070, 972, 849 cm^{-1} ; ^1H NMR δ 5.69 (d, J = 59.5 Hz, 1H), 4.77-4.43 (m, 2H), 4.43-4.38 (m, 1H), 4.18-4.05 (m, 3H), 1.46 (s, 6H), 1.39 (s, 3H), 1.35 (s, 3H); ^{19}F NMR δ -129.25 (dd, J = 59.5, 6.7 Hz, 1F); ^{13}C NMR δ 113.6 (d, $^1J_{\text{C-F}}$ = 221.6 Hz), 113.2, 109.4, 84.7, (d, $^2J_{\text{C-F}}$ = 42.2 Hz), 82.6, 78.6, 72.7, 66.6, 26.9, 25.8, 25.1, 24.5; HRMS (EI) calcd for $\text{C}_{12}\text{H}_{19}\text{O}_5\text{F}$ ($\text{M}^+ + \text{H}$) 263.1295, found 263.1317.

*4.4.4. 2,3-O-Isopropylidene-5-O-benzoyl- β -D-ribofuranosyl fluoride (**6c**)*

IR (neat) 2990, 1725, 1273, 1094, 977, 714 cm^{-1} ; ^1H NMR δ 8.07 (d, J = 8.2 Hz, 2H), 7.61-7.56 (m, 1H), 7.48-7.42 (m, 2H), 5.83 (d, J = 61.8 Hz, 1H), 4.88-4.85 (m, 2H), 4.71-4.67 (m, 1H), 4.45-4.37 (m, 2H), 1.50 (s, 3H), 1.35 (s, 3H); ^{19}F NMR δ -116.44 (d,

$J = 60.9$ Hz, 1F){ lit.[18] -115.85 (dq, $J = 61.6, 4.0$ Hz, 1F)}; ^{13}C NMR $\delta = 166.1, 133.4, 129.9$ (2C), $129.6, 128.5$ (2C), 115.4 (d, $^1J_{\text{C-F}} = 223.1$ Hz), $113.3, 86.5$ (d, $^3J_{\text{C-F}} = 3.2$ Hz), 85.1 (d, $^2J_{\text{C-F}} = 40.8$ Hz), $81.0, 64.7, 26.4, 25.0$.

4.4.5. 2,3,5-Tri-*O*-benzyl- β -D-arabinofuranosyl fluoride (**6d**)

White solid. mp $78-79$ °C (lit.[19] $77-78$ °C); IR (KBr) $3062, 3030, 2865, 1454, 1115, 1028, 738, 698$ cm^{-1} ; ^1H NMR δ $7.30-7.17$ (m, 15H), 5.79 (d, $J = 61.5$ Hz, 1H), $4.73-4.45$ (m, 7H), 4.17 (dd, $J = 9.3, 2.2$ Hz, 1H), 3.96 (dd, $J = 5.1, 2.0$ Hz, 1H), $3.64-3.57$ (m, 2H); ^{19}F NMR δ -121.23 (dd, $J = 61.6, 9.2$ Hz, 1F); ^{13}C NMR δ $137.9, 137.7, 137.2, 127.7-128.5$ (15C), 108.3 (d, $^1J_{\text{C-F}} = 229.9$ Hz), 84.5 (d, $^2J_{\text{C-F}} = 21.5$ Hz), $82.4, 81.5, 73.5, 72.6, 72.4, 71.5$.

4.5. Reaction of trimethyl trithioorthocarboxylates **7** with $\text{BrF}_3\text{-KHF}_2$

4.5.1. 2-Bromo-*N,N*-dimethyl-4-(trifluoromethyl)aniline (**8a**)

The reaction was carried out as in the case of **2a** at 0 °C using 3.2 eq of $\text{BrF}_3\text{-KHF}_2$ to **7a**, and **8a** was isolated in 52% yield. IR (neat) $2952, 2874, 2842, 2791, 1608, 1324, 1123$ cm^{-1} ; ^1H NMR δ 7.79 (s, 1H), 7.49 (d, $J = 7.5$ Hz, 1H), 7.09 (d, $J = 7.5$ Hz, 1H), 2.87 (s, 6H); ^{19}F NMR δ -62.52 (s, 3F); ^{13}C NMR δ $154.9, 131.3$, (q, $^3J_{\text{C-F}} = 3.8$ Hz), 125.3 (q, $^3J_{\text{C-F}} = 3.8$ Hz), 125.2 (q, $^2J_{\text{C-F}} = 34.3$ Hz), 123.7 (t, $^1J_{\text{C-F}} = 276.0$ Hz), $120.3, 117.9, 43.8$ (2C); HRMS (EI) calcd for $\text{C}_9\text{H}_8\text{BrF}_3\text{N}$ ($\text{M}^+ - 1$) 265.9791 , found 265.9792 .

4.5.2. 5-Bromo-1-methyl-3-(trifluoromethyl)-1*H*-indole (**8b**)

White solid. mp 60 °C (lit.[20] $58-60$ °C); IR (KBr) $1558, 1473, 1235, 1095$ cm^{-1} ; ^1H

NMR (DMSO- d_6) δ 8.08 (s, 1H), 7.72 (s, 1H), 7.60 (d, $J = 8.9$ Hz, 1H), 7.46 (dd, $J = 8.9, 1.9$ Hz, 1H), 3.86 (s, 3H); ^{19}F NMR (DMSO- d_6) δ -54.8 (s, 3F); ^{13}C NMR (DMSO- d_6) δ 135.4, 131.8 (q, $^3J_{\text{C-F}} = 4.9$ Hz), 125.4, 124.8 (q, $^3J_{\text{C-F}} = 2.2$ Hz), 124.3 (q, $^1J_{\text{C-F}} = 270.0$ Hz), 120.4, 113.9, 113.3, 120.6 (q, $^2J_{\text{C-F}} = 37.2$ Hz), 33.2.

4.5.3. Ethyl 3,3-difluoro-2,2-dimethyl-3-(methylthio)propanoate (**8c**)

IR (neat) 2988, 2938, 1737, 1274, 1175, 1034 cm^{-1} ; ^1H NMR δ 4.20 (q, $J = 7.3$ Hz, 2H), 2.29 (s, 3H), 1.40 (s, 6H), 1.28 (t, $J = 7.3$ Hz, 3H); ^{19}F NMR δ -84.67 (s, 2F); ^{13}C NMR δ 171.7 (t, $^3J_{\text{C-F}} = 2.8$ Hz), 131.5 (t, $^1J_{\text{C-F}} = 289$ Hz), 61.5, 51.6 (t, $^3J_{\text{C-F}} = 22.0$ Hz), 20.7 (t, $^3J_{\text{C-F}} = 3.1$ Hz, 2C), 13.9, 9.9 (t, $^2J_{\text{C-F}} = 5.3$ Hz); HRMS (EI) calcd for $\text{C}_8\text{H}_{14}\text{F}_2\text{O}_2\text{S}$ 212.0683, found 262.0682.

Acknowledgment

We are grateful to prof. Hermann-Josef Frohn (University of Duisburg-Essen, Germany) for his kind advice on the handling of BrF_3 .

References

- [1] (a) P. Kirsh, in Modern Fluoroorganic Chemistry, Wiley-VCH; Weinheim, 2004, pp. 203-277.
- (b) T. Hiyama, in : H. Yamamoto, (Ed.), Organofluorine Compounds, Springer-Verlag Heidelberg, 2000, pp. 183-233.
- (c) R. F. Anderson, J. O. Punderson, in: R. E. Banks, (Ed.), Organofluorine Chemicals and Their Industrial Applications, Ellis Horwood LTD., Chichester, 1979, pp. 123-247.
- [2] Recent reviews and books on fluorination reagent, see: (a) R. P. Singh, J. M.

Shreeve, *Synthesis* (2002) 2561-2578.

(b) K. L. Kirk, *Org. Process Res. Dev.* 12 (2008) 305-321.

(c) K. Uneyama, in *Organofluorine Chemistry*, Blackwell Publishing, Oxford, 2006.

[3] Recent reviews and books on air stable fluorination reagent, see: (a) R. P. Singh, J. M. Shreeve, *Acc. Chem. Res.* 37 (2004) 31-44.

(b) P. T. Nyffeler, S. G. Durón, M. D. Burkart, S. P. Vincent, C.-H. Wong, *Angew. Chem. Int. Ed.* 44 (2005) 192-212.

(c) S. Hara, in: K. K. Laali, (Ed.), *Advances in Organic Synthesis*, Bentham Science Publishers LTD., Hilversum, 2006; p 49-60.

(d) N. Al-Maharik, D. O'Hagan, *Aldrichimica Acta* 44 (2011) 65-75.

[4] (a) S. Hara, M. Monoi, R. Umemura, C. Fuse, *Tetrahedron* 68 (2012) 10145-10150.

(b) M. Kunigami, S. Hara, *J. Fluorine Chem.* (2014) in press.

[5] As for the reviews of the fluorination using BrF_3 , see: (a) S. Rozen, *Acc. Chem. Res.* 38 (2005) 803-812. (b) S. Rozen, *Adv. Synth. Catal.* 352 (2010) 2691-2707.

[6] (a) S. Siegel, *Acta Cryst.* 9 (1956) 493-495.

(b) K. O. Christe, C. J. Schack, *Inorg. Chem.* 9 (1970) 1852-1858.

[7] W. W. Wilson, K. O. Christe, *Inorg. Chem.* 28 (1989) 4172-4175.

[8] We didn't have any information about the structure of this solid. But it was conveniently used as $\text{BrF}_3 \cdot 2(\text{KHF}_2)$ (MW 215) because, two equivalent of KHF_2 to BrF_3 was used to make it. This solid is insoluble in most of organic solvents, and a slightly hygroscopic.

[9] As for the recent review articles of *gem*-difluoride synthesis from thioacetals, see:

(a) V. Hugenberg, G. Haufe, *J. Fluorine Chem.* 143 (2012) 238-262.

(b) T. Fuchigami, S. Inagi, *Chem. Commun.* 47 (2011) 10211-10223.

- (c) M. Shimizu, T. Hiyama, *Angew. Chem. Int. Ed.* 44 (2005) 214-231.
- (d) M. Kuroboshi, K. Kanie, T. Hiyama, *Adv. Synth. Catal.* 343 (2001) 235-250.
- (e) V. P. Reddy, G. K. S. Prakash, G. A. Olah, in: K. K. Laali (Ed.), *Advances in Organic Synthesis*, Bentham Science Publishers LTD., Hilversum, 2006, pp.183-211.
- (f) V. P. Reddy, M. Perambuduru, R. Alleti, in: K. K. Laali (Ed.), *Advances in Organic Synthesis*, Bentham Science Publishers LTD., Hilversum, 2006, pp.327-351.
- [10] (a) M. Shimizu, H. Togo, M. Yokoyama, *Synthesis* (1998) 799-822.
- (b) K. Toshima, *Carbohydr. Res.* 327 (2000) 15-26.
- (c) T. Mukaiyama, *Angew. Chem. Int. Ed.* 43 (2004) 5590-5614.
- [11] (a) K. C. Nicolau, R. E. Dolle, D. P. Papahatjis, J. L. Randall, *J. Am. Chem. Soc.* 106 (1984) 4189-4192.
- (b) J. C. López, P. B. Albert, C. Uriel, S. Valverde, A. M. Gómez, *J. Org. Chem.* 72 (2007) 10268-10271.
- (c) T. Sawamura, S. Kuribayashi, S. Inagi, T. Fuchigami, *Adv. Synth. Catal.* 352 (2010) 2757-2760.
- (d) S. Tsegay, R. J. Williams, S. J. Williams, *Carbohydr. Res.* 357 (2012) 16-22.
- (e) K. Suzuki, Y. Ito, O. Kanie, *Carbohydr. Res.* 359 (2012) 81-91.
- (f) G. Mugunthan, K. P. R. Kartha, *Tetrahedron Lett.* 53 (2012) 5631-5634.
- [12] M. Barbero, S. Cadamuro, I. Degani, R. Fochi, A. Gatti, V. Regondi, *Synthesis* (1988) 22-25.
- [13] G. K. S. Prakash, S. K. Ganesh, J.-P. Jones, A. Kulkarni, K. Masood, J. K. Swabeck, G. A. Olah, *Angew. Chem. Int. Ed.* 51 (2012) 12090-12094.
- [14] K. Fujikawa, Y. Fujioka, A. Kobayashi, H. Amii, *Org. Lett.* 13 (2011) 5560-5563.
- [15] T. Furuya, T. Fukuhara, S. Hara, *J. Fluorine Chem.* 126 (2005) 721-725.

- [16] F. E. Ray, C. E. Albertson, J. Am. Chem. Soc. 70 (1948) 1954-1955.
- [17] G. A. Olah, M. Nojima, I. Kerekes, J. Am. Chem. Soc. 96 (1974) 925-927.
- [18] M. Rapp, X. Cai, W. Xu, W. R. Dolbier Jr., S. F. Wnuk, J. Fluorine Chem. 130 (2009) 321-328.
- [19] W. A. Szarek, G. Gryniewicz, Chem. Lett. (1984) 1751-1754.
- [20] M. M. Bastos, L. M. U. Mayer, E. C. S. Figueira, M. Soares, W. B. Kover, N. Boechat, J. Heterocycl. Chem. 45 (2008) 969-973.