

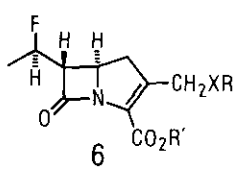
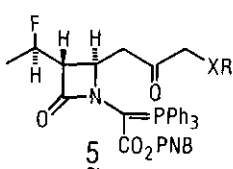
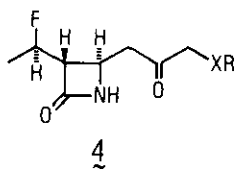
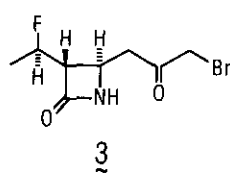
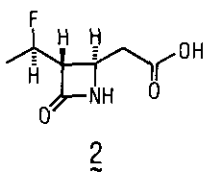
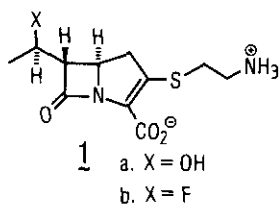
SYNTHESIS OF CARBAPENEMS WITH A  $RXCH_2$ -SUBSTITUENT IN THE 3-POSITIONJohannes G. de Vries\*<sup>1</sup>, Gerhard Sigmund, and Georg Vorisek

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**Abstract** - Carbapenems **6** were prepared by reacting mercaptans with bromoketone **3** followed by a Wittig sequence, or better through advanced intermediate **9**. The carbapenem carboxylates were chemically unstable; stability was not improved by changing the oxidation state of sulfur.

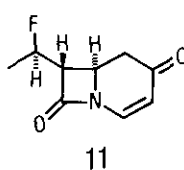
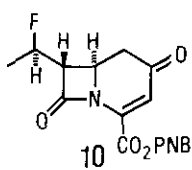
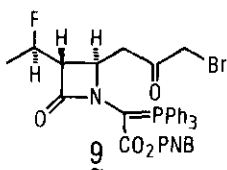
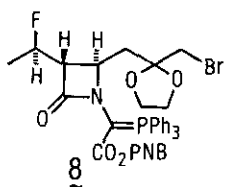
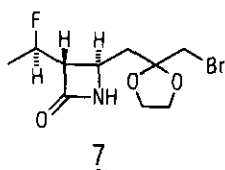
Ever since the discovery of thienamycin (**1a**) a large number of analogues have been synthesized in an attempt to improve the chemical and metabolic instability of thienamycin while retaining its potent antibacterial activity<sup>2</sup>. Our efforts in this area have been concentrated on carbapenems with an  $R$ -1-fluoroethyl side chain in the 6-position<sup>3</sup>. In this paper we report our synthetic work on carbapenems bearing a  $CH_2XR$  side chain in the 3-position<sup>4</sup>: we hoped that insertion of a methylene group in between nucleus and sulfur side chain would provide additional chemical and metabolic stability. From the viewpoint of synthetic efficiency introduction of the variable  $RX$  group should occur as late as possible in the synthetic sequence. We therefore chose 3-acetoxymethylcarbapenem ester **6a** as intermediate, based upon the knowledge that 3-acetoxymethylcephalosporins react with mercaptans to give the thio substituted analogues<sup>5</sup>.

Starting with acid **2**<sup>3a,b</sup> we were able to synthesize the bromoketone **3**<sup>6</sup> (41%<sup>7</sup>, mp 71-73°C) by a Nierenstein sequence<sup>8</sup> (see Reaction Conditions). Reaction with NaOAc in DMF gave the acetoxy compound **4a** in 88% yield which was transformed into the desired 3-acetoxymethylcarbapenem **6a** by the Wittig procedure developed by Woodward<sup>9</sup> (44% over 4 steps). Unfortunately, reaction of **6a** with lithium benzyl sulfide (THF, -30°C) led to unidentifiable products and reaction with  $PhCH_2SH/Et_3N$  ( $CH_2Cl_2$ , RT) gave only recovered **6a**. Addition of  $Pd(Ph_3P)_4$  to the latter reaction had no beneficial effect. Before investigating other strategies we decided to synthesize one representative of **6** by an alternative route. Reaction of **3** with  $PNBO_2CNHCH_2CH_2SH$ <sup>10</sup> gave **4b** (76%), which was converted to **6b** in the usual way (20% yield over 4 steps). Hydrogenation of **6b** gave **6c** in 19% yield (purity 80% by <sup>1</sup>H NMR)



- 4,5: a. RX = OAc  
b. RX = SCH<sub>2</sub>CH<sub>2</sub>NHCO<sub>2</sub>PNB  
c. RX = SCH<sub>2</sub>-3-Pyr.

- 6: a. RX = OAc R' = PNB  
b. RX = SCH<sub>2</sub>CH<sub>2</sub>NHCO<sub>2</sub>PNB R' = PNB  
c. RX = SCH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub> R' = H  
d. RX = SCH<sub>2</sub>-3-Pyr R' = PNB  
e. RX = SCH<sub>2</sub>-3-Pyr R' = H  
f. RX = SOCH<sub>2</sub>CH<sub>2</sub>NHCO<sub>2</sub>PNB R' = PNB  
g. RX = SOCH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub> R' = H  
h. RX = SOCH<sub>2</sub>-3-Pyr R' = PNB  
i. RX = SOCH<sub>2</sub>-3-Pyr R' = H  
j. RX = SO<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>NHCO<sub>2</sub>PNB R' = PNB  
k. RX = H R' = H  
l. RX = SO<sub>2</sub>CH<sub>2</sub>-3-Pyr R' = PNB  
m. RX = SO<sub>2</sub>CH<sub>2</sub>-3-Pyr R' = H



after RP-18 column chromatography and lyophilization. Though  $\tilde{6c}$  was very unstable<sup>11</sup>, we were able to establish that the compound had a satisfactory antibacterial spectrum and had improved renal dehydrogenase stability compared to the fluorinated thienamycin  $\tilde{1b}^{3b}$ . Encouraged by this finding we set out to synthesize more analogues, soon to find out that the sequence  $\tilde{4} \rightarrow \tilde{5}$  is of limited value because of the incompatibility of many functional groups with  $\text{SOCl}_2$ . So, clearly the synthetic sequence had to be adapted in such a way as to allow introduction of the variable side chain at a later stage. Thus we protected bromoketone  $\tilde{3}$  as the ketal  $\tilde{7}$  (77%) which was transformed into phosphorane  $\tilde{8}$  (74%). The ketal function in  $\tilde{8}$  was very stable and deprotection turned out to be a serious problem. After many unsuccessful attempts we found that  $\tilde{8}$  could be hydrolyzed in  $\text{CH}_2\text{Cl}_2$  solution by brief treatment with a large excess of 70% aqueous  $\text{HClO}_4$  (99%)<sup>12</sup>. The resulting bromoketone was very unstable and was best immediately reacted with mercaptan and base. A solution of  $\tilde{9}$  in  $\text{CH}_3\text{CN}$  left overnight gave  $\tilde{10}$  and  $\tilde{11}$  as main products. Reaction of  $\tilde{9}$  with 3-pyridylmethyl mercaptan and  $\text{Et}_3\text{N}$  gave  $\tilde{5c}$  (100%) which was cyclized to  $\tilde{6d}$  (64%), which we had not been able to obtain by the original sequence:  $\tilde{4c}$  had perished in the second step of the Woodward sequence. Catalytic hydrogenation of  $\tilde{6d}$  did produce the expected carbaopenem potassium salt  $\tilde{6e}$  (3%), but this compound was so unstable that no biological data could be obtained. We then sought to find out if the stability of these compounds is influenced by the oxidation state of the sulfur in the side chain. Oxidation of  $\tilde{6b}$  and  $\tilde{6d}$  with 1 eq. of MCPBA gave the sulfoxides  $\tilde{6f}$  (51%) and  $\tilde{6h}$  (61%); with 2 eq. MCPBA the sulfones  $\tilde{6j}$  (41%) and  $\tilde{6l}$  (47%) were obtained. Deprotection of the

$\tilde{2} \rightarrow \tilde{3}$ :	1. $(\text{CH}_3)_2\text{C}(\text{CH}_2)_2\text{O}_2\text{CCl}$ , $\text{Et}_3\text{N}$ , $\text{CH}_2\text{Cl}_2$ , -78°C, 11 $\text{CH}_2\text{N}_2$ , -30°C $\rightarrow$ -10°C, 11. HBr, -40°C.
$\tilde{3} \rightarrow \tilde{4b}$ :	$\text{PNBO}_2\text{CNHCH}_2\text{CH}_2\text{SH}$ , $\text{t-Pr}_2\text{NEt}$ , $\text{CH}_2\text{Cl}_2$ , 0°C.
$\tilde{3} \rightarrow \tilde{4c}$ :	3-pyridyl $\text{CH}_2\text{SH}$ , $\text{Et}_3\text{N}$ , $\text{CH}_2\text{Cl}_2$ , 0°C.
$\tilde{4} \rightarrow \tilde{5}$ :	1. $\text{PNBO}_2\text{CCH}(\text{OH})_2$ , benzene, azeotropic reflux, 11 $\text{SOCl}_2$ , $\text{Et}_3\text{N}$ , THF, -20°C, 11 $\text{Ph}_3\text{P}$ , THF, RT.
$\tilde{7} \rightarrow \tilde{8}$ :	Toluene, 1% hydroquinone, reflux.
$\tilde{6b}, \tilde{6d}, \tilde{6f}, \tilde{6h} \rightarrow \tilde{6j}, \tilde{6l}$ :	1 eq. MCPBA, $\text{CH}_2\text{Cl}_2$ , -30°C.
$\tilde{6b}, \tilde{6d}, \tilde{6f}, \tilde{6h} \rightarrow \tilde{6j}, \tilde{6l}$ :	2 eq. MCPBA, $\text{CH}_2\text{Cl}_2$ , 0°C.
$\tilde{6c}, \tilde{6e}, \tilde{6g}, \tilde{6i}, \tilde{6k}, \tilde{6m}$ :	$(\text{CH}_2\text{OH})_2$ , PPTS, benzene, azeotropic reflux.
$\tilde{3} \rightarrow \tilde{7}$ :	buffer pH 7.
$\tilde{8} \rightarrow \tilde{9}$ :	70% aq. $\text{HClO}_4$ , $\text{CH}_2\text{Cl}_2$ , 8 min.
$\tilde{9} \rightarrow \tilde{10} + \tilde{11}$ :	$\text{CH}_3\text{CN}$ , RT, 16 hrs.

REACTION CONDITIONS

sulfoxides 6f and 6h and the sulfone 6l proceeded as usual to give 6g (15%), 6j (46%) and 6m (47%), respectively; but the stability of these compounds was very similar to those of the parent sulfides. Deprotection of 6j was accompanied by reduction of the allylic sulfone, giving 3-methylcarbapenem carboxylate 6k<sup>13</sup>.

Since it is possible that the instability of these carbapenems is due to the presence of a potential leaving group on methyl we have now shifted our attention to aminoalkyl substituted carbapenems and we will report on this in due course.

#### ACKNOWLEDGEMENTS

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6. Selected spectral data (NMR in CDCl<sub>3</sub> or D<sub>2</sub>O, UV, in H<sub>2</sub>O): 3j: <sup>1</sup>H NMR: 6.28 (1H, br s, NH), 4.96 (1H, ddq, J = 48.6, 7.5, 6.3 Hz, HCF), 4.00 (1H, ddd, J = 9.5, 3.5, 2.5 Hz, H-4), 3.91 (s, 2H, CH<sub>2</sub>Br), 3.25 and 2.96 (2H, ABX, J<sub>AB</sub> = 18.5 Hz, J = 7.5, 3.5 Hz, CH<sub>2</sub>CO), 3.02 (1H, ddd, J = 18.0, 7.5, 2.5 Hz, H-3), 1.47 (3H, dd, J = 24.5, 6.3 Hz, CH<sub>3</sub>). 4a: <sup>1</sup>H NMR: 4.66 (2H, s, CH<sub>2</sub>O), 2.17 (3H, s, Ac). 4b: <sup>1</sup>H NMR: 5.24 (2H, s, OCH<sub>2</sub>), 3.29 (2H, s, CH<sub>2</sub>S). 4c: <sup>1</sup>H NMR: 3.70 (2H, s, CH<sub>2</sub>Py), 3.12 (2H, s, CH<sub>2</sub>S). 6a: <sup>1</sup>H NMR: 5.29 and 5.05 (2H, AB, J = 15.5 Hz, CH<sub>2</sub>OAc), 2.10 (3H, s, Ac), IR (KBr): 1783, 1744, 1710 cm<sup>-1</sup>. 6b: <sup>1</sup>H NMR: 3.80 and 3.66 (2H, AB, J = 13.5 Hz, CH<sub>2</sub>S). IR (CH<sub>2</sub>Cl<sub>2</sub>): 1783, 1724 cm<sup>-1</sup>. 6c: IR (KBr): 1761 cm<sup>-1</sup>, UV:  $\lambda_{\max}$  = 272 ( $\epsilon$ =5900) nm. 6d: <sup>1</sup>H NMR: 3.70 (2H, s, CH<sub>2</sub>Py), 3.82 and 3.69 (2H, AB, J = 14.0 Hz, CH<sub>2</sub>S), IR (CH<sub>2</sub>Cl<sub>2</sub>): 1781, 1721 cm<sup>-1</sup>. 6e: <sup>1</sup>H NMR: 3.32 (1H, ddd, J = 27.4, 5.1, 2.9 Hz, H-6), 1.36 (3H, dd, J = 25.0, 6.3 Hz, CH<sub>3</sub>). 6f: IR (CH<sub>2</sub>Cl<sub>2</sub>): 1789, 1716, 1700,

1048  $\text{cm}^{-1}$ .  $\underline{6g}$ :  $^1\text{H NMR}$ : 4.32 and 4.19 (2H, AB,  $J = 13.5$  Hz,  $\text{CH}_2\text{SO}$ ).  $\underline{6h}$ : IR ( $\text{CH}_2\text{Cl}_2$ ): 1786, 1718, 1055  $\text{cm}^{-1}$ .  $\underline{6i}$ : IR (KBr): 1762, 1595, 1387, 1029  $\text{cm}^{-1}$ . UV:  $\lambda_{\text{max}} = 263$  (6.400), 269 (5900) nm.  $\underline{6j}$ :  $^1\text{H NMR}$ : 4.59 and 4.27 (2H, AB,  $J_{\text{AB}} = 14.0$  Hz,  $\text{CH}_2\text{SO}_2$ ). IR ( $\text{CH}_2\text{Cl}_2$ ): 1789, 1726, 1320, 1114  $\text{cm}^{-1}$ .  $\underline{6k}$ :  $^1\text{H NMR}$ : 2.01 (3H, s,  $=\text{C}-\text{CH}_3$ ).  $\underline{6l}$ :  $^1\text{H NMR}$ : 4.60 and 4.32 (2H, AB,  $J_{\text{AB}} = 14.2$  Hz,  $\text{CH}_2\text{SO}_2$ ), 4.30 (2H, s,  $\text{CH}_2\text{Py}$ ), IR ( $\text{CH}_2\text{Cl}_2$ ): 1787, 1719, 1331, 1120  $\text{cm}^{-1}$ .  $\underline{6m}$ :  $^1\text{H NMR}$ : 4.22 (1H, ddd,  $J = 9.5, 9.0, 3.0$  Hz, H-5), 3.61 (1H, ddd,  $J = 28.1, 4.5, 3.0$  Hz, H-6).  $\underline{7}$ :  $^1\text{H NMR}$ : 4.06 (4H, M,  $\text{CH}_2\text{CH}_2$ ), 3.36 (2H, s,  $\text{CH}_2\text{Br}$ ), 2.42 and 2.13 (2H, ABX,  $J_{\text{AB}} = 14.6$  Hz,  $J = 9.7, 3.2$  Hz,  $\text{CH}_2$ ).  $\underline{10}$ :  $^1\text{H NMR}$ : 6.25 (1H, s,  $=\text{CH}$ ), 2.95 and 2.66 (2H, ABX,  $J_{\text{AB}} = 16.2$  Hz,  $J = 5.4$  and  $13.7$  Hz, H-5).  $\underline{11}$ :  $^1\text{H NMR}$ : 7.44 (1H, d,  $J = 7.6$  Hz, H-2), 5.54 (1H, d,  $J = 7.6$  Hz, H-3), 2.67 and 2.84 (2H, ABX,  $J_{\text{AB}} = 16.0$  Hz,  $J = 13.0, 6.2$  Hz, H-5). IR ( $\text{CH}_2\text{Cl}_2$ ): 1775, 1672, 1585, 1380  $\text{cm}^{-1}$ . MS (calcd. for  $\text{C}_9\text{H}_{10}\text{FNO}_3$ : 183):  $m/e$  183 (M, 18), 136 (M- $\text{CH}_3\text{CHF}$ , 16), 96 (100).

7. The only side product we have been able to isolate was the isobutyl ester of  $\underline{2}$ , but use of other activated derivatives such as the acid chloride or imidazolide did not lead to improved yields of  $\underline{3}$ .
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11. We were unsuccessful in our attempts to convert  $\underline{6c}$  to the *N*-acetimidoyl or *N*-formimidoyl derivatives, a strategy which was successfully applied by Merck chemists to stabilize thienamycin: W.J. Leanza, K.J. Wildonger, T.W. Miller, and B.G. Christensen, *J. Med. Chem.* 1979, **22**, 1435.
12. We were not able to hydrolyze  $\underline{8}$  using milder methods.
13. A similar allylic reduction was observed by Sankyo chemists: Ref. 4c.

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