HETEROCYCLES, Vol.12, No.1 , 1979

BIOMINETIC SYNTHESIS OF β -AMINO-4-AMINO-6-CARBOXY-5-METHYL-2-PYRIMIDINEPROPIONIC ACID DERIVATIVES

K<u>azuo</u> A<u>chiwa</u> and S<u>igeki</u> S<u>asaki</u> Faculty of Pharmaceutical Sciences, University of Tokyo, Bunkyo-ku, Tokyo 113, Japan

Biominetic synthesis of β -amino-4-amino-6-carboxy-5-methyl-2-pyrimidinepropionic acid derivative (1), a biologically important part of Bleomycins, one of effective antitumor antibiotics, was described.

From the consideration on the speculation that the β -amino acid (1) may be derived from N- α -aspartyl- β -methylasparagine, a dipeptide, by dehydration, amination, and dehydrogenation. We selected 4-amino-3-t-butoxycarbonyl-4-methylisoxazole (2) as the starting matrial.

Reaction of 2 with N-benzyloxycarbonylaspartic acid β -methyl ester α -chloride gave the corresponding 4-acylamino-3-t-butoxycarbonyl-4-methylisoxazole (3) in the optically active form. Succesive hydrogenation of 3 in ethyl acetate-acetic acid (5%) afforded the β -N-t-benzyloxycarbonyl-6-t-butoxycarbonyl-4-hydroxy-5-methyl-2pyrimidinepropionic acid methyl erter (4). Further treatment of 4 with SOCl₂-DMF and lig. NH₃ gave β -N-benzyloxycarbonylamino-6-t-butoxycarbonyl-4-amino-5-methyl-2-pyrimidinepropionic acid amide (5). Unfortunately, the optical activity of 4 and 5 was lost perhaps in the stage of the formation of pyrimidine ring.

Further investigations along this line are actively under way.