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ic addition on the product of (8-13) using methanesulfonyl chloride or p-toluenesulfonyl chloride in an appropriate solvent such as dichloromethane in the presence of a base such as triethylamine or pyridine. Alternatively, the compound of formula (8-13) may be synthesized by reacting an amine of formula (8-4) with (14-8) in the presence of a dehydrating agent such as phosphorus trichloride or P(S)t-butyl chloride in an appropriate solvent such as dimethylformamide or the like in the presence of a base such as triethylamine or pyridine. Each of the intermediates of formula (3-1) to (3-16) can be prepared by known processes or analogous known processes. The compounds of formula (1-1), (1-2), (1-3), (1-4), (2-1), (3-3), (3-4), (4-2), and (4-3) represented by the following chemical formulas [1-1a], [1-2a], [1-3a], [1-4a], [2-1a], [3-3a], [3-4a], [4-2a], and [4-3a], respectively, can be prepared by the following scheme. Said compounds may be prepared by known processes or analogous known processes, such as the processes described in the scheme, or analogous known processes. wherein, X1 is halogen, hydroxy, nitro, or cyano; R13 and R14 are each independently an optionally substituted aryl group; each of R15 and R16 is independently an optionally substituted

alkyl group, cycloalkyl group, heterocyclic group, aryl group, aralkyl group, or alkenyl group; and R17 is a protecting group. Said compounds (3-1a), (3-2a), (3-3a), (4-1a), (4-2a), (4-3a), (4-4a), (5-1a), (5-2a), (5-3a), (5-4a), and (5-5a) represented by the following chemical formulas [3-1 f988f36e3a

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