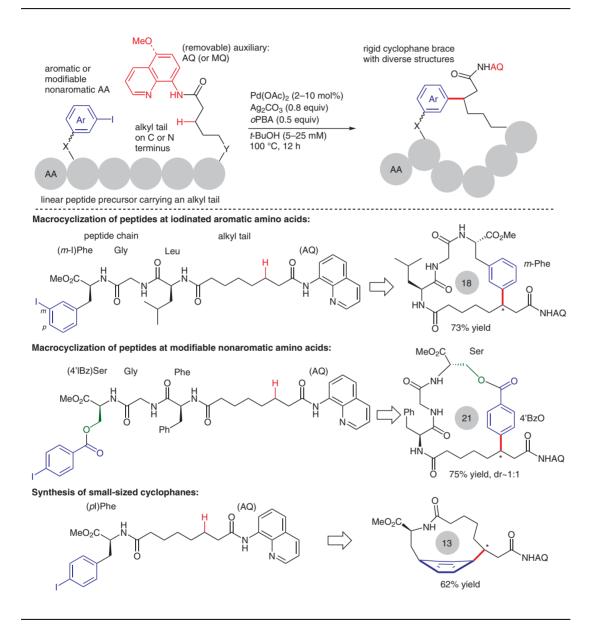
X. ZHANG, G. LU, M. SUN, M. MAHANKALI, Y. MA, M. ZHANG, W. HUA, Y. HU, Q. WANG, J. CHEN, G. HE*, X. QI*, W. SHEN*, P. LIU*, G. CHEN* (NANKAI UNIVERSITY, TIANJIN AND NATIONAL INSTITUTE OF BIOLOGICAL SCIENCES, BEIJING, P. R. OF CHINA; UNIVERSITY OF PITTSBURGH, CALIFORNIA INSTITUTE FOR BIOMEDICAL RESEARCH, LA JOLLA, AND THE PENNSYLVANIA STATE UNIVERSITY, UNIVERSITY PARK, USA) A General Strategy for Synthesis of Cyclophane-Braced Peptide Macrocycles via Palladium-Catalysed Intramolecular sp³ C-H Arylation

Nat. Chem. 2018, 10, 540-548.

Synthesis of Cyclophane-Braced Peptide Macrocycles



Significance: New efficient methods for the intramolecular cyclization of peptides are important in terms of the development of drugs based on cyclic peptides. The authors report a powerful method for constructing new types of peptide macrocycles through palladium-catalyzed, aminoquinoline-directed, intramolecular C(sp³)–H arylation reactions.

SYNFACTS Contributors: Hisashi Yamamoto, Takahiro Sawano Synfacts 2019, 15(03), 0317 Published online: 15.02.2019 DOI: 10.1055/s-0037-1611466; Reg-No.: H00419SF

Comment: The cyclization of readily accessible linear peptide precursors selectively proceeds at side chains of either aromatic or modified non-aromatic amino acids units to provide a variety of cyclophane-braced peptide macrocycles containing small-sized cyclophanes.

Category

Peptide Chemistry

Key words

palladium catalysis
C-H arylation
cyclophanes
macrocycles
peptide macrocycles

