Supporting Information

Development of a Concise Synthetic Approach to Access Oroxin A

Haijun Chena,*, Guihua Hea, Cailong Li², Longrong Donga, Xiaobo Xiea, Jianlei Wua,

Yu Gaoa, Jia Zhoub,*

aCollege of Chemistry, Fuzhou University, Fuzhou, Fujian 350108, China

bChemical Biology Program, Department of Pharmacology and Toxicology,
University of Texas Medical Branch, Galveston, Texas 77555, United States

Corresponding authors:
*Haijun Chen, PhD
College of Chemistry
Fuzhou University
Fuzhou, Fujian 350108, China
Email: chenhaij@gmail.com

*Jia Zhou, PhD
Chemical Biology Program
Department of Pharmacology and Toxicology
University of Texas Medical Branch
Galveston, Texas 77555, United States
Email: jizhou@utmb.edu

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1. Experimental section

**General:** All commercially available starting materials and solvents were reagent grade, and used without further purification. Reactions were performed under a nitrogen atmosphere in dry glassware with magnetic stirring. Preparative column chromatography was performed using silica gel 60, particle size 0.063-0.200 mm (70-230 mesh, flash). Analytical TLC was carried out employing silica gel 60 F254 plates (Merck, Darmstadt). Visualization of the developed chromatograms was performed with detection by UV (254 nm). NMR spectra were recorded on a Bruker-400 (1H, 400 MHz; 13C, 100 MHz) spectrometer. 1H and 13C NMR spectra were recorded with TMS as an internal reference. Chemical shifts were expressed in ppm, and J values were given in Hz. Melting point was determined using the X-4 melting point apparatus (Beijing Taike Co., Ltd.) and uncorrected. High-resolution mass spectra (HRMS) were obtained from Thermo Fisher Scientific Exactive Plus mass spectrometer.
2. Copies of $^1$H and $^{13}$C NMR spectra of Oroxin A
$^1$H NMR spectra of Synthetic Oroxin A

$^1$H NMR spectra from *Process Biochem.*, **2013**, 48, 1744-1748.
1H NMR spectra of Synthetic Oroxin A (Part 1)

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1H NMR spectra of Synthetic Oroxin A (Part 2, contains the signals of OH/glucoside)

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$^1$H NMR spectra of Synthetic Oroxin A (Part 3)
$^{13}$C NMR spectra of Synthetic Oroxin A

$^{13}$C NMR spectra from *Process Biochem.*, 2013, 48, 1744-1748.