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Biomimetic Total Synthesis of (±)-Garcibracteatone *Org. Lett.* **2012**, *14*, 5162–5164.

## Total Synthesis (±)-Garcibracteatone

**Significance:** Garcibracteatone (**K**) is the structurally most complex polycyclic polyprenylated acylphlorogucinol natural product that has so far been isolated. The four-step total synthesis presented makes use of a biomimetic radical cascade reaction to build up four rings in one transformation. Additionally, the previously unknown relative stereochemistry at C-5 was assigned.

Comment: Precursor **F** for the key transformation is synthesized from phloroglucinol **A** in three steps by Friedel–Crafts acylation followed by subsequent diprenylation and alkylation with (±)-lavandulyl iodide (**E**). Oxidation of **F** by using Mn(OAc)<sub>3</sub>–Cu(OAc)<sub>2</sub> initiates a radical cascade, which ultimately leads to the formation of the natural product garcibracteatone **K** (14% yield) along with its C5-epimer **L** (8% yield). This key transformation constructs four rings and five stereocenters.

SYNFACTS Contributors: Erick M. Carreira, Stefan Diethelm Synfacts 2013, 9(1), 0001 Published online: 17.12.2012 DOI: 10.1055/s-0032-1317856; Reg-No.: C02812SF