IBDA/LiBr-MEDIATED FUNCTIONALISATION OF ent-TRACHYLOBAN-19-OIC ACID

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ent-Trachyloban-19-oic acid (1) was isolated from sunflower *Helianthus annuus* L. [1-4]. It exhibited weak cytotoxic activity against gastric carcinoma [5] and displayed *in vivo* anti-inflammatory activity [4].

In order to obtain new polyfunctionalized tetracyclic terpenoids, the transformation of *ent*-trachyloban-19-oic acid (1) was carried out by studying a PhI(OAc)₂/LiBr reaction [6].

As a result, a mixture of polyfunctionalized tetracyclic diterpenoids (2) and (3) was obtained, which was purified by column chromatography. The major compound (2) (61%) was identified as the functionalized at C-12 and C-16 ent-kaurenic derivative. The minor compound (3) (11%) had an *ent*-atisanic structure and it was functionalized at C-13, C-15 and C-17 atoms. The structure and stereochemistry of the newly synthesized compounds (2) and (3) were established on the basis of their IR, NMR spectral and GS-MS data.



In conclusion, the oxidative conversion of *ent*-trachyloban-19-oic acid (1) under treatment with IBDA/LiBr led basically to *ent*-kaurenic carbonic framework derivatives.

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