

**Supporting Information to:****Antitrypanosomal Activity of a Diterpene and Lignans Isolated from *Aristolochia cymbifera***

Patrícia Sartorelli<sup>1</sup>, Camila Salomone Carvalho<sup>1</sup>, Juliana Quero Reimão<sup>2</sup>, Harri Lorenzi<sup>3</sup>,  
André Gustavo Tempone<sup>2</sup>

**Affiliation**

<sup>1</sup> Departamento de Ciências Exatas e da Terra, Universidade Federal de São Paulo  
Diadema/São Paulo, Brazil

<sup>2</sup> Laboratório de Toxinologia Aplicada, Departamento de Parasitologia, Instituto Adolfo Lutz,  
São Paulo, Brazil

<sup>3</sup> Instituto Plantarum de Estudos da Flora, Nova Odessa/ São Paulo, Brazil

**Correspondence*****Patrícia Sartorelli***

Departamento de Ciências Exatas e da Terra

Universidade Federal de São Paulo

R. Prof. Artur Ridel

275, CEP 09972-270

Diadema/São Paulo

Brazil

Tel.: +55-11-4059-3618

Fax: +55-11-4043-6428

psartorelli@unifesp.br

**Extraction and isolation**

Air-dried powdered leaves of *A. cymbifera* (100 g) were extracted with MeOH (3 × 250 mL) at room temperature. After evaporation *in vacuo*, the MeOH residue (2.34 g) was submitted to column chromatography over 30 g silica gel 60 (0.063–0.200 mm; Merck) (20 × 5 cm) by vacuum liquid chromatography, eluted with hexane and mixtures of hexane–EtOAc of increasing polarity. This procedure resulted in 12 fractions of 100 mL each (AC-1: hexane;

AC-2: hexane–EtOAc 95:5; AC-3: hexane–EtOAc 9:1; AC-4: hexane–EtOAc 85:15; AC-5: hexane–EtOAc 8:2; AC-6: hexane–EtOAc 75:25; AC-7: hexane–EtOAc 7:3; AC-8: hexane–EtOAc 65:35; AC-9: hexane–EtOAc 6:4; AC-10: hexane–EtOAc 55:45; AC-11: hexane–EtOAc 5:5; AC-12: hexane–EtOAc 4:6). Fractions AC-5 (175 mg), AC-7 (67 mg), AC-8 (51 mg), AC-9 (118 mg), and AC-10 (29 mg) exhibited significant activity against trypomastigote forms of *T. cruzi*, and the <sup>1</sup>H-NMR data of these fractions revealed the presence of diterpene (AC-5) and a mixture of lignans (AC-7–AC-10). Thus, these active fractions were further purified by thin-layer chromatography to afford the six active compounds. TLC analyses were performed with precoated TLC sheets of silica gel eluting with different mixtures of MeOH in CH<sub>2</sub>Cl<sub>2</sub> and EtOAc in hexane. Plates were observed under a UV lamp (254 nm and 365 nm). The purities of tested compounds were inferred by analysis of <sup>1</sup>H-NMR spectra.