Cytotoxic Activity and Chemical Composition of the Root Extracts from the Mexican Species *Linum scabrellum:* Mechanism of Action of the Active Compound 6-methoxypodophyllotoxin

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Supplementary Material



Figure 1. ¹H NMR and ¹³C NMR spectra (400 and 100 MHz, $CDCl_3$) of (-)- pinoresinol (5)



Figure 2. ¹H NMR and ¹³C NMR spectra (400 and 100 MHz, CDCl₃) of 6-methoxypodophyllotoxin (6)



Figure 3 . HSQC and COSY spectra (400 MHz, CDCl3) of 6-methoxypodophyllotoxin (6)



Figure 4. Positive FABMS and HPLC of 6-methoxypodophyllotoxin (6). 6-methoxypodophyllotoxin (6) was analyzed by high performance liquid chromatography, using a Waters 600 pump and Waters 2996 photodiode array detector. The analytical analysis was carried out with an isocratic solvent system (60% CH₃CN, 40% H₂O with 0.0125% TFA) through a Shiseido Capcell pak C18 column (4.6 mm I.D. x 250 mm, 5 μ m), and a flow rate of 1 mL/min. An injection volume of 20 μ L, three injections were performed for each sample.



Figure 5. ¹H NMR, ¹³C and HSQC NMR spectra (400 and 100 MHz, acetone-d₆) of coniferin (**7**)



Figure 6. ¹H NMR and HSQC NMR spectra (400 and 100 MHz, acetone-d6) of 6methoxypodophyllotoxin 7-*O*-β-D-glucopyranoside (**8**)