Organotin adducts of indomethacin: synthesis, crystal structures and spectral characterization of the first organotin complexes of indomethacin.

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The complexes $[Me_2(Indo)SnOSn(Indo)Me_2]_2$, (**1**) $[Bu_2(Indo)SnOSn(Indo)Bu_2]_2$, (**2**), where Hindo is indomethacin, have been prepared and structurally characterized by means of ^{119}Sn Mössbauer, vibrational, and NMR (^{1}H and ^{13}C) spectroscopy. The crystal structures of the complexes $[Me_2(Indo)SnOSn(Indo)Me_2]_2 \cdot 2C_4H_8O_2$, **3** and **2** have been determined by X-ray crystallography. Each structure is centro-symmetric and features a central rhombus Sn_2O_2 unit with two additional tin atoms linked at the O atoms. Pairs of tin atoms are bridged by bidentate carboxylate ligands and by a monoatomic bridging oxygen. C–H→π, π→π, stacking interactions, inter and intramolecular hydrogen bonds stabilize the structures **2** and **3**. Complexes **2** and **3** are self-assembled via π→π, C–H-π and stacking interactions.The complexes $[Me_2(Indo)SnOSn(Indo)Me_2]_2$, (**1**) $[Bu_2(Indo)SnOSn(Indo)Bu_2]_2$, (**2**), where Hindo is indomethacin, have been prepared and structurally characterized by means of ^{119}Sn Mössbauer, vibrational, and NMR spectroscopy. The crystal structures of the complexes $[Me_2(Indo)SnOSn(Indo)Me_2]_2 \cdot 2C_4H_8O_2$, **3** and **2** have been determined by X-ray crystallography. Complexes **2** and **3** are self-assembled via π→π, C–H-π and stacking interactions.

Słowa kluczowe

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