

Electronic Supplementary Information:

***In vivo* anti-tumor activity of the organometallic ruthenium(II)-arene complex [Ru(η^6 -*p*-cymene)Cl₂(pta)] (RAPTA-C) in human ovarian and colorectal carcinomas**

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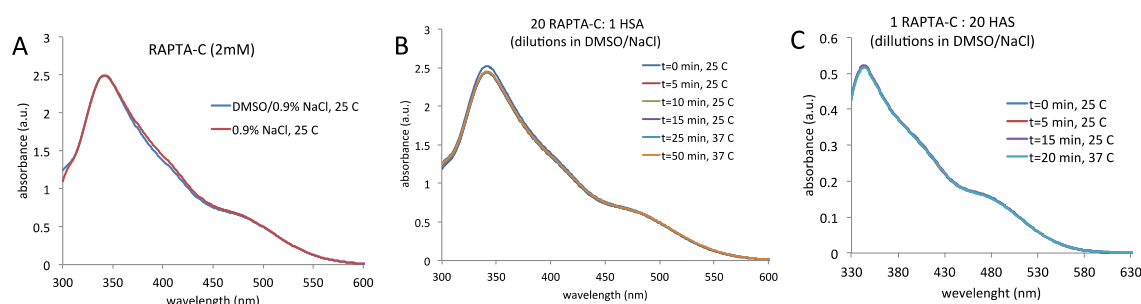


Figure S1. Spectral profiles for RAPTA-C solutions in various solvents recorded at 25°C or 37°C. HSA: human serum albumin. Absorption spectra of RAPTA-C solutions were recorded with a two-beam Varian Cary UV-Vis-NIR 500 Scan spectrophotometer in 1cm quartz cuvettes (Suprasil, Hellma, Müllheim, Germany) between 300 and 600 nm with an average scan speed of 600 nm/min at 25°C or 37°C. These spectra indicate that RAPTA-C is stable under the tested conditions.

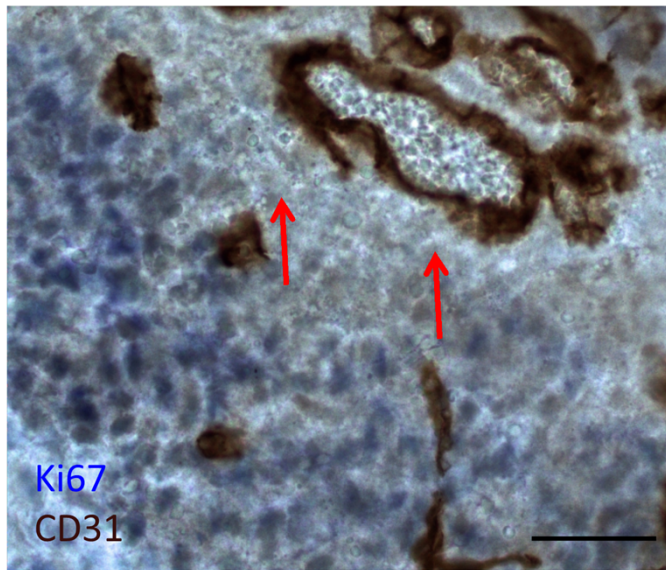


Figure S2. Image showing immunohistochemical staining of LS174T tumor treated with RAPTA-C at 100 mg kg^{-1} . The islands of tumor cells (red arrows) around the vasculature (CD31, in brown) were Ki-67 negative, meaning that tumor cells were in a quiescent state. On the tumor periphery Ki-67 (in blue) tumor cells were visible.

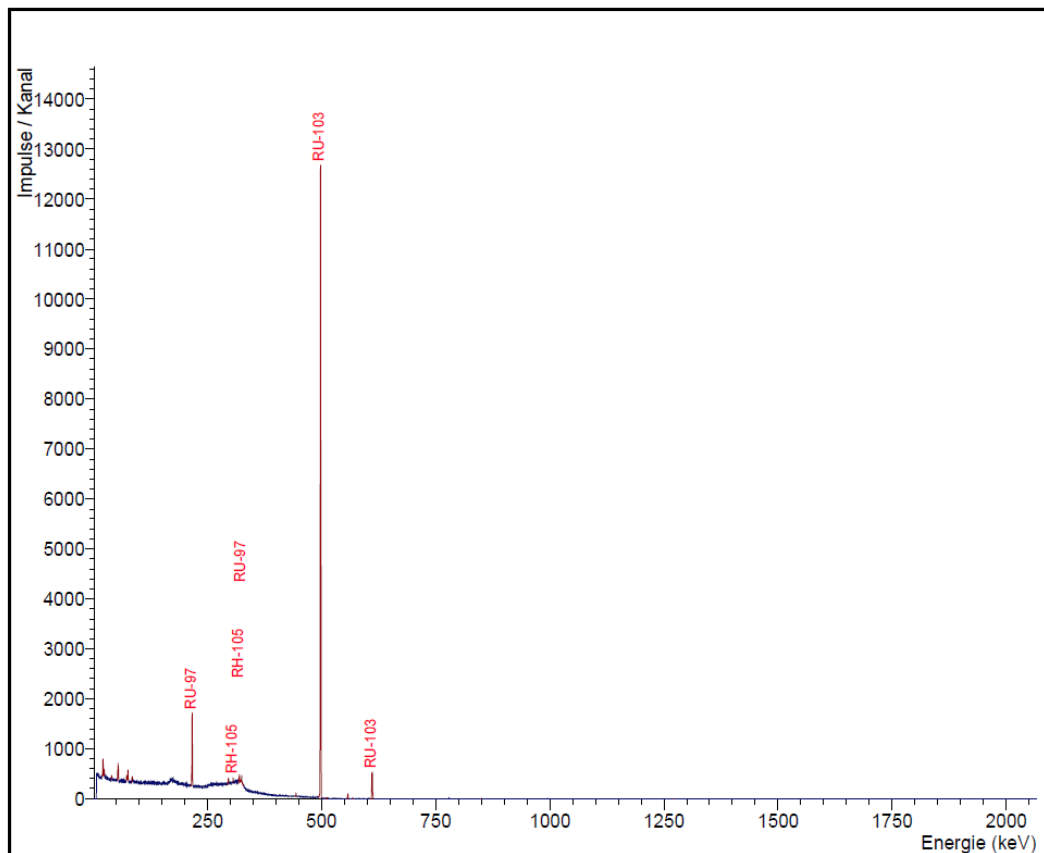


Figure S3. γ -ray spectrometry after work-up of the target material. The spectrum shows the γ -rays of ^{103}Ru (497 keV, 91% and 610 keV, 5.8%) and γ -rays of traces of ^{97}Ru (215, 85.6% and 324 keV, 10.8%) as well as ^{105}Ru (316 keV, 11.1%) are also visible.

Equation used to estimate ^{103}Ru activity

$$A_{^{103}\text{Ru}} = N_{^{102}\text{Ru}} \cdot \sigma \cdot \phi \cdot (1 - e^{-\lambda \cdot t})$$

A: activity [Bq]

N: number of ^{102}Ru atoms

σ : thermal cross section of ^{102}Ru [barn]

ϕ : neutron flux [$\text{n} \cdot \text{cm}^{-2} \cdot \text{s}^{-1}$]

λ : $\frac{\ln(2)}{T_{1/2}}$ [h^{-1}]

t: irradiation time [h]