

Supplementary Material

Antioxidant and anticancer activities and protein interaction of the oxidovanadium(IV) naringin complex

Andrés Gonzalo Restrepo-Guerrero¹, Helen Goitia-Semenco¹, Luciana G. Naso¹, Marilin Rey², Pablo J. Gonzalez², Evelina G. Ferrer¹, Patricia A.M. Williams^{1,*}.

¹Centro de Química Inorgánica (CEQUINOR, UNLP, CONICET, asociado a CICPBA), Departamento de Química, Facultad de Ciencias Exactas, Universidad Nacional de La Plata, Bv. 120 N° 1465, CP 1900, La Plata, Buenos Aires, Argentina.

²Departamento de Física, Facultad de Bioquímica y Ciencias Biológicas, Universidad Nacional del Litoral and CONICET, S3000ZAA Santa Fe, Argentina.

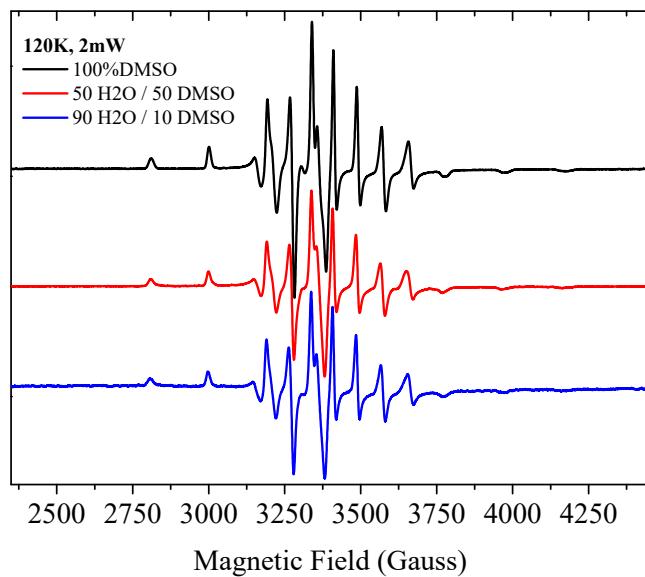


Figure S1. EPR spectrum of frozen solutions of VONarg at 120 K with different water:DMSO ratios.

Black: 100% DMSO, Red: 50% water pH 7, Blue: 90% water pH 7.

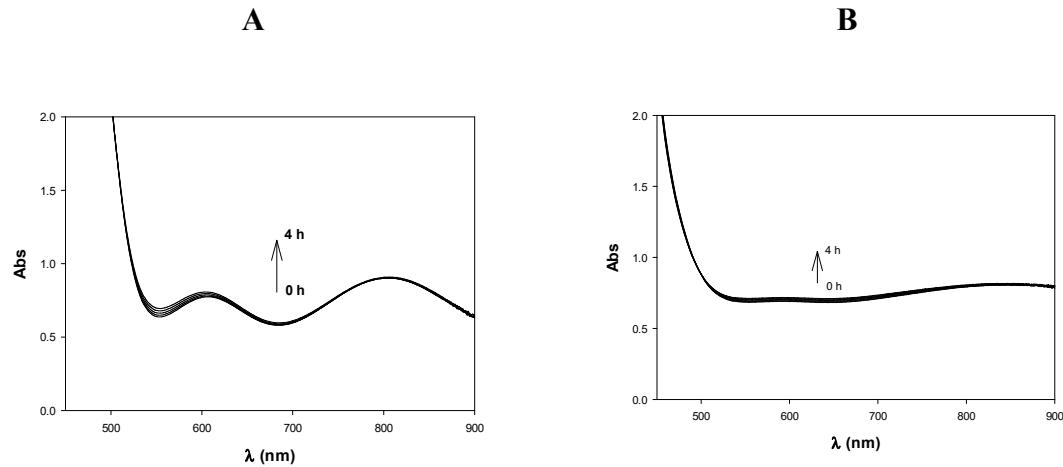


Figure S2. UV-Vis spectra recorded after 4h of 0.005 M solutions of $[VO(Narg)_2] \cdot 8H_2O$ in: A) DMSO;

B) DMSO/H₂O 1/99.

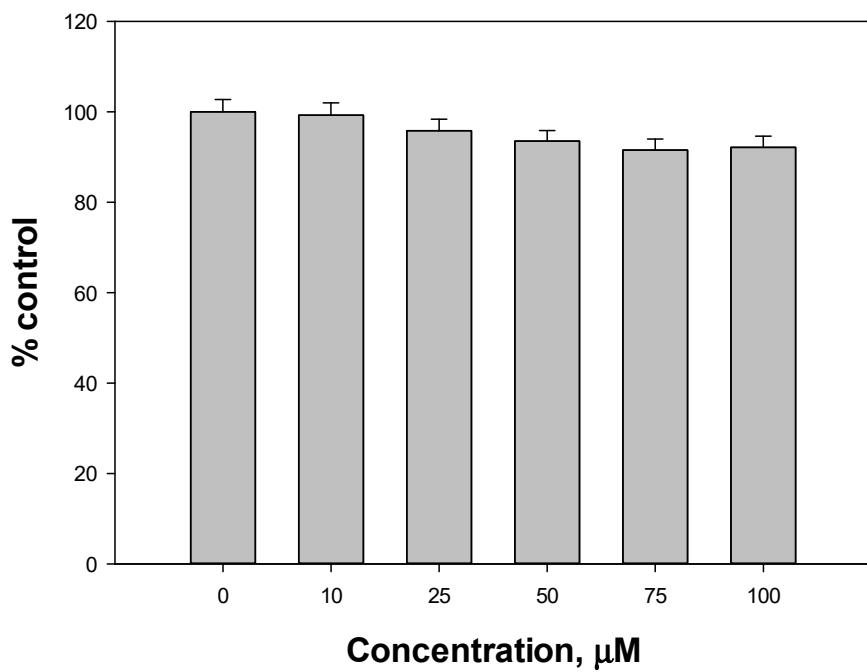


Figure S3. Cell viability assay at different VONarg concentrations after treatment for 24 h on HEK293 cells. The results are expressed as a percentage of the control level and represent the mean \pm the standard error of the mean (SEM) from three separate experiments.

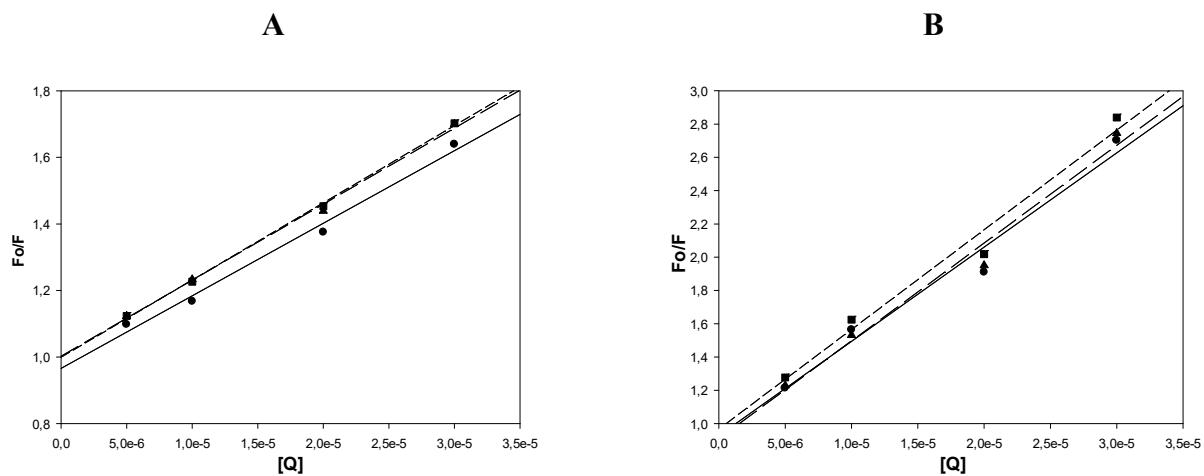


Figure S4. Stern-Volmer plot of the fluorescence quenching of BSA with different concentrations of naringin (A) and VONarg (B) systems, 5, 10, 20 and 30 μM : (●) 298 K; (▲) 303 K; (■) 310 K, [BSA] = 6 μM , $\lambda_{\text{ex}} = 280 \text{ nm}$.