

Supplementary Materials:

Table S1. Antiherpetic activity of the liposomal form of Σ -CRG/Ech

Compounds	CC ₅₀	Pretreatment of virus		Pretreatment of cells		Simultaneous treatment		Treatment of infected cells	
		IC ₅₀	SI	IC ₅₀	SI	IC ₅₀	SI	IC ₅₀	SI
		(μ /mL)		(μ /mL)		(μ /mL)		(μ /mL)	
Liposomes	>2000	9.5 \pm	211	57 \pm 7	35 \pm	47 \pm 6	42 \pm	68 \pm 9*	29
Σ-CRG/Ech		1,1*	\pm 23*		5*		6		\pm 4*
Σ-CRG/Ech	>1000	2.8 \pm 0.4	350 \pm 54	55 \pm 7	18 \pm 3	31 \pm 5	32 \pm 5	96 \pm 14	10 \pm 2
Empty liposomes	>2000	NA		NA		NA		NA	

Note: Values represent the means \pm standard deviations of three or more independent experiments; Σ -CRG/Ech - complex of carrageenan Σ -CRG with echinochrome A; Liposomes Σ -CRG/Ech - liposomal form of this complex; IC₅₀, concentration that inhibited 50% of viral plaque formation; SI, selectivity index (CC₅₀/IC₅₀); NA - no activity. * Significance of the differences between the parameters of complex of carrageenan Σ -CRG/Ech compared to the liposomal form of this complex ($p \leq 0.05$)

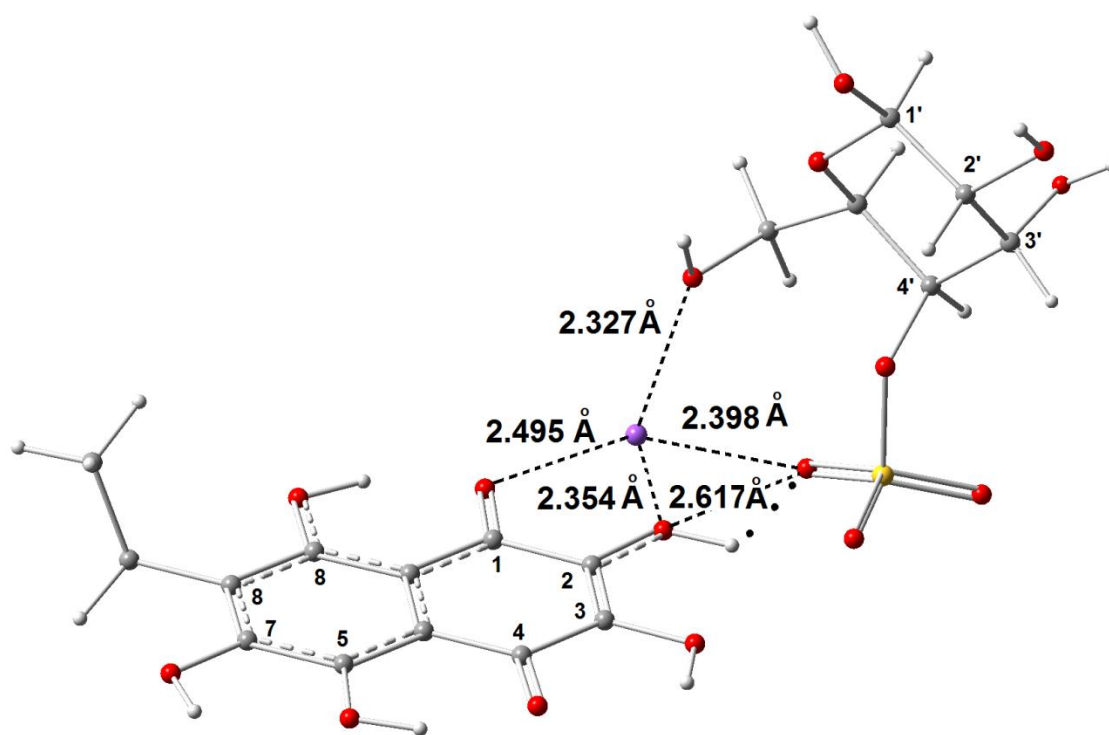


Figure S1. PCM-B3LYP/6-31+G(d,p) optimized geometries of the most stable complex κ -CRGS with Ech at positions C(1)=O and C(2)-OH («k-CRG-Na-1'2-Ech»)

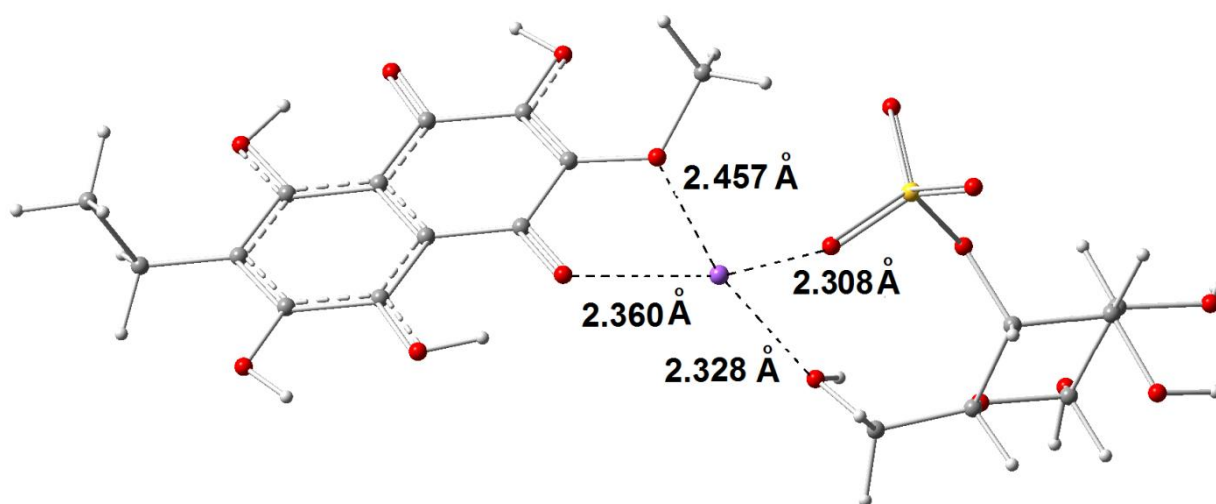


Figure S2. PCM-B3LYP/6-31+G(d,p) optimized geometries of the complex κ -CMS with 3-OMe-Ech at positions C(3)-O and C(4)=O («k-Car-Na-1'2-Ech»)
«k-Car-Na-3'4-3OMe-Ech»

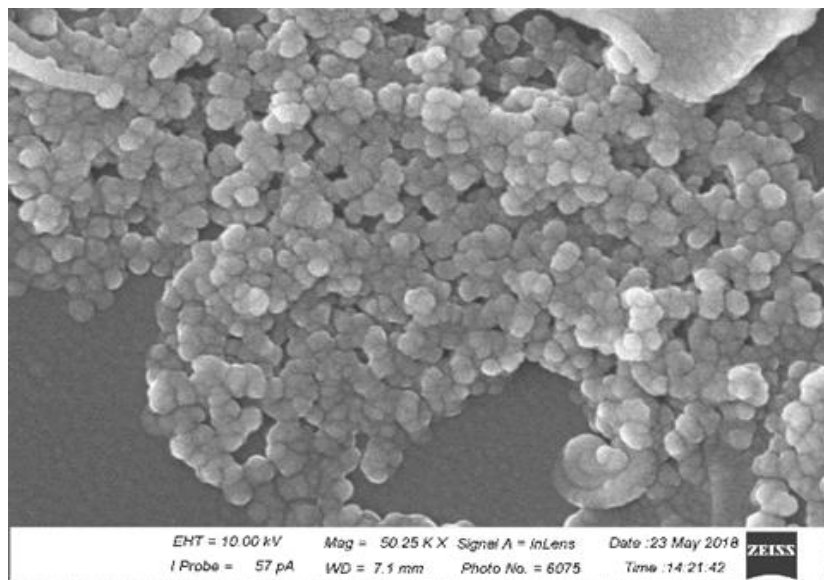


Figure S3. Scanning electron microscopy images of CRG/Ech-containing liposomes. Mag. = 50.87 KX, EHT = 10.00 kV