Toxicity of Fucoxanthin on Balb/c Mice Splenocytes and Thymocytes

Alexander Lykov, Lubov Rachkovskaya, Olga Poveshchenko, Maria Surovtseva, Irina Kim, Edmund Rachkovsky, Maxim Korolev, Anastasiya Kotlyarova, Andrey Letyagin

Research Institute of Clinical and Experimental Lymphology- Branch of the Institute of Cytology and Genetics SB RAS

Novosibirsk, Russia

Ruslan Gevorgiz, Svetlana Zheleznova

A.O. Kovalevsky Institute of Biology of the Southern Seas

RAS

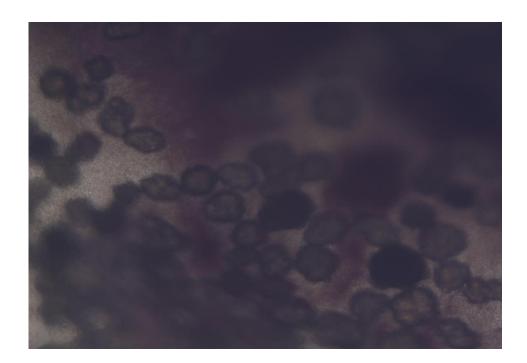
Sevastopol, Russia

Fucoxanthin (Fx) is a naturally marine carotenoid in chloroplasts of brown seaweeds (brown algae) and diatoms possess various health benefits, such as antioxidant and anticancer properties, anti-obesity, and anti-diabetic activities.

Fucoxanthin (Fx) was extracted from *Cylindrotheca closterium* (EhrenbERG) using a series of steps. Briefly, fucoxanthin isolation with ethanol, then concentrated, and purified by precipitation, and dried.

Modification of γ -Al₂O₃/PDMS composition was carried out by immobilizing the Fx in the aqueous phase (Fx@Al/PDMS) by physical adsorption, then the obtained matrices was subjected to drying and short-term low-temperature heat treatment up to 120 $^{\circ}$ C, the obtained matrices were loose powder materials.

Splenocyte and thymocytes were extracted from Balb/c male mice, then crushed by homogenization, and twice washed in phosphate-buffer saline, and transplanted in RPMI-1640 (Biolot, Russia) medium with the addition of 80 µg/mL of gentamicin, 2 mmol L-glutamine, 5 mmol HEPES-buffer and 10% FCS at a concentration 10⁶ cells/mL.



Microphotograph of modified porous material based on γ - AL_2O_3 and polydimethylsiloxane by fucoxanthin in splenocytes suspension (x10).

EFFECT OF FUCOXANTHIN ALONE OR IN COMBINATION WITH POROUS MATERIAL BASE ON γ -AL $_2$ O $_3$ ON LYMPHOCYTES PROPERTIES IN VITRO (M \pm SD)

Splenocytes (24 h)				
Basal	100	1.37±0.23	NA	8.39±0.6
96% Ethanol	92±20	1.26±0.28	NA	7.87±0.53
Fx (ethanol extract)	110±24	1.5±0.33	NA	10.64±0.66*
AI/PDMS	120±30	1.64±0.41	NA	10.57±1.06 *
Fx@AI/PDMS	130±24*	1.79±0.33*	NA	9.36±1.03
Splenocytes (120 h)				
Basal	100	0.9±0.19	1.51±0.31	6.97±0.79
96% Ethanol	75±10 *	0.68±0.09 *	1.71±0.34	7.69±1.04
Fx (ethanol extract)	78±15 *	0.7±0.13 *	1.62±0.31	9.41±0.78*
AI/PDMS	116±33	1.05±0.29	1.58±0.36	8.58±0.74*
Fx@AI/PDMS	89±11	0.81±0.1	1.7±0.33	8.46±0.74*
Thymocytes (24 h)				
Basal	100	0.69±0.21	NA	8.62±0.59
96% Ethanol	67±12 *	0.46±0.08 *	NA	7.72±0.75*
Fx (ethanol extract)	94±40	0.65±0.27	NA	12.1±0.53*
AI/PDMS	98±25	0.68±0.17	NA	10.17±1.01 *
Fx@AI/PDMS	115±48	0.8±0.33	NA	9.24±1.07
Thymocytes (120 h)				
Basal	100	0.55±0.07	1.07±0.11	6.95±0.92
96% Ethanol	119±22	0.66±0.12	1.11±0.03	6.75±1.0
Fx (ethanol extract)	109±32	0.6±0.17	0.96±0.08	9.28±0.78*
AI/PDMS	116±24	0.64±0.13	1.18±0.11	8.21±0.68*
Fx@AI/PDMS	153±48 *	0.84±0.32*	1.15±0.13	8.83±0.9*

Note. MPO, myeloperoxidase activity; NO, nitric oxide; Fx, ethanol extract of fucoxanthin; Al/PDMS, porous aluminum oxide with polydimethylsiloxane particles; Fx@Al/PDMS, fucoxanthin immobilized on porous aluminum oxide with polydimethylsiloxane particles; NA, not analyzed; *p<0.05 compared with control.