

Activation of protein tyrosine kinase: A possible requirement for fixed-bacteria and lipopolysaccharide-induced increase in human natural killer cell activity

Puente, J.

Salas, M. A.

Canon, C.

Miranda, D.

Wolf, M. E.

Mosnaim, A. D.

Preincubation of peripheral blood lymphocytes (PBL) from drug-free, healthy volunteers with either the protein tyrosine kinase inhibitor genistein (GNT, n = 10, final concentration 200 μ M) or the protein kinase A activator dibutyryl-cyclic-AMP (cAMP, n = 11, final concentration 10 μ M), resulted in a significant inhibition of natural killer cell activity (NKCA, expressed as percentage of specific chromium release). With the exception of 4 out of the 11 cAMP-treated samples, individual values for NKCA in the drug preincubated specimens were at least 20% below the same subject baseline activity, furthermore, NK lytic function was non-detectable in 4 out of the 10 and in 1 out of the 11 samples presented with either GNT or cAMP, respectively. PBL preincubation with glutaraldehyde-fixed Gram-negative bacteria (GNB, n = 13, final GNB-to-effector cell ratio of 50: 1) resulted in a statistically significant increase in NKCA (baseline ($\bar{x} \pm SD$) of 21.6 ± 16.4 and bacteria treated samples of 41.