

# **Synthesis and in vitro evaluation of new ethyl and methyl quinoxaline-7-carboxylate 1,4-di-N-oxide against *Entamoeba histolytica***

**Blanca Estela Duque-Montaño a, Lilia Citlalli Gómez-Caro b, Mario Sanchez-Sanchez b, Antonio Monge c, Efrén Hernández-Baltazar a, Gildardo Rivera b, Oscar Torres-Angeles**

a Laboratorio de Microbiología, Facultad de Farmacia, Universidad Autónoma del Estado de Morelos, Av. Universidad 1001, 62209 Cuernavaca, México

b Centro de Biotecnología Genómica, Instituto Politécnico Nacional, Blvd. del Maestro, s/n, Esq. Elías Piña, 88710 Reynosa, México

c Unidad de Investigación y Desarrollo de Medicamentos, Centro de Investigación en Farmacobiología Aplicada, Universidad de Navarra, 31008 Pamplona, Spain

***Bioorganic & Medicinal Chemistry, Volume 21 , 2013***

## **Abstract**

In our search for new antiamoebic agents, a new series of ethyl and methyl quinoxaline-7-carboxylate 1,4-di-N-oxide derivatives have been synthesized using the Beirut reaction. All compounds were characterized by spectroscopic techniques and elemental analysis. Antiamoebic activity was evaluated in vitro against *Entamoeba histolytica* strain HM1:IMSS by the microdilution method, and the structure–activity relationship was analyzed. We found that eleven quinoxaline derivatives showed greater activity than metronidazole and nitazoxanide with IC<sub>50</sub> values in the range 1.99–0.35 μM. Compounds T-001 and T-016 shows IC<sub>50</sub> values of 1.41 and 1.47 μM, respectively, with a value of selectivity index >60.