



Synthesis, Cytotoxic and Antimicrobial activities of 5-benzylidene-2-[(pyridine-2-ylmethylene)hydrazono]-thiazolidin-4-one Derivatives

Eraldo A.G. NETO ¹, Janete M. ARAÚJO ², Maria D. RODRIGUES ², Teresinha G. SILVA ²,
Silene C. NASCIMENTO ², Gardênia C.G. MILITÃO ², Alexandre J.S. GÓES ² & José G. LIMA ^{1*}

¹ Departamento de Ciências Farmacêuticas, Universidade Federal de Pernambuco,
Recife 50670-901, Brazil

² Departamento de Antibióticos, Universidade Federal de Pernambuco, Recife 50670-901, Brazil

SUMMARY. A novel series of 5-benzylidene-2-[(pyridine-2-ylmethylene)hydrazono]-thiazolidin-4-ones **3a-i** has been synthesized. 2-[(Pyridine-2-ylmethylene)hydrazono]-thiazolidin-4-ones **2a-c** were also obtained and used as intermediates to give the target compounds. The *in vitro* cytotoxic activity was evaluated for both series. The findings obtained showed that the compounds **2a**, **2b**, **3b** and **3c** were effective against the HEP-2 cell lines with IC₅₀ in the 1.6 - 0.5 µg/mL range, whereas the compounds **2a** (IC₅₀= 3.6 µg/mL), **2b** (IC₅₀= 2.4 µg/mL) and **3f** (IC₅₀= 3.5 µg/mL) showed good inhibitory effects against HT-29 cell lines. As complementary biological test, all 4-thiazolidinones were evaluated for antimicrobial activity against various bacterial and fungal species.

KEY WORDS: Antimicrobial activity, Antiproliferative activity, Cytotoxicity, 4-Thiazolidinone.

* Author to whom correspondence should be addressed. E-mail: jgildolima@gmail.com