

AN IN VITRO STUDY ON THE ANTIMICROBIAL EFFECT OF SELECT HETEROCYCLIC NITROGEN COMPOUNDS

Y.S.R. REDDY, SAMA VENKATESH, UMA SHANKAR MISRA, B. SURESH, Md
AFZAL AZAM and M.SETHURAMAN

Department of Pharmaceutical chemistry, J.S.S College of Pharmacy,
Rocklands, Ooty – 643 001.

Received: 3 August, 1995

Accepted: 9 August, 1995

ABSTRACT: *In vitro* antimicrobial activity of seventeen heterocyclic nitrogen compounds {2substituted pyrido [1,2-a] pyrimidin -4-oxo-1 [4]-3-Carbonitriles and ethyl 5-amino-3(substituted) – pyrazole-3-Carboxylates} was tested against *Escherichia coli*, *Bacillus subtilis* and *candida albicans* strains, Antimicrobial activity was measured using standard two –fold serial dilution method and the minimum inhibitory concentration (MIC) values are determined, The MIC of pyrazoles and pyrido[1,2 –a] pyrimidines are found to be 25 µg/ml to 50 µg/ml against *E. Coli* and 50 µg/ml against *B.subtilis* respectively. Both pyrazole and pyrido [1,2-a] Pyrimidines are found to be totally ineffective against *C. albicans*.

INTRODUCTION

The roles played by many pyrazoles, pyridine and pyrimidine compounds are well known for their therapeutic properties and are widely used to treat various diseases. 1-8 The antibacterial, analgesic and anti-inflammatory effects of some pyrido [1,2-a] pyrimidines and pyrazole compounds are reported earlier elsewhere.⁹ The present work was undertaken to assess the antimicrobial activity of select heterocyclic nitrogen compounds.

MATERIALS & METHODS

The authentic bacterial and fungal cultures were procured from central drug research Institute, Lucknow and also from National collection of Industrial micro organisms, Pune. We selected three strains for use in the present study. They are *Coli* (2345 NCIM), *B.subtilis* (2439, NCIM) and *C albicans* (27, CDRI). The above strains were maintained in our laboratory by periodic subculturing

technique, the fungal culture was maintained on Sabouraud Dextrose Agar (SDA) and bacterial strains in nutrient broth, the inhibition studies were carried out by two – fold serial dilution method and the MIC values were calculated.¹⁰

RESULTS AND DISCUSSIONS

Results of antimicrobial effect of seven heterocyclic nitrogen compounds are presented in Table 1 and 2. It is evident from the data that pyrazole compounds are found to show inhibition against *E.coli* strain (MIC:-25 µg/ml -50 µg/ml). The pyrido [1,2 – a] pyrimidin compounds are also found to be effective in inhibiting *B. subtilis* strains (MIC : 50 µg/ml).

However, both pyrazole and pyrido [1,2-a] pyrimidine compounds are totally devoid of inhibitory action against *C. albicans*. The low anti-microbial effect exhibited by all the test compounds was reflected by their high MIC values. The overall antimicrobial

activity observed in the present study on heterocyclic nitrogen compounds are not **ACKNOWLEDGEMENT** comparable with than of standard drugs such The authors wish to place on regard their as ampicillin and clotrimazole. However, the heartfelt thanks to His Holiness Jagadguru present findings of this study are subject to Sri shivarathree Deshkendra Maha confirmation through future studies Swamigalavaru of Sri Suttur Mutt for information through future studies providing necessary facilities indifferent models with more number of strains.

Table – 1
Antimicrobial effect of Pyrido [1,2-a] pyrimidines

Substituent	MIC values (µg/ml)			
“R”	E.coli	B.subtilis		C. albicans
Dicyclohexylamino Diethylamino N-isopropylamino Ethylamino Ampicillin Clotrimazole	No activity “ “ “ 12.5 -	50 “ “ “ 12.5 -		No activity “ “ “ -12.5
Substituent “R”	R1	MIC values (µg/ml)		
		E.coli	B.subtilis	C. albicans
p-Bromo phenyl	Hydrogen	50	NA *	NA *

Table – 2
Antimicrobial effect of Pyrazoles

Substituent	MIC values (µg/ml)			
“R”	E.coli	B.subtilis	C. albicans	
Dicyclohexylamino Diethylamino N-isopropylamino Ethylamino Ampicillin Clotrimazole	No activity “ “ “ 12.5 -	50 “ “ “ 12.5 -	No activity “ “ “ -12.5	
Substituent “R”	R1	MIC values (µg/ml)		
		E.coli	B.subtilis	C. albicans
p-Bromo phenyl	Hydrogen	50	NA*	NA*
p-Chloro phenyl	“	“	“	“
p-Nitro phenyl	“	“	25	“
p-Hydroxy phenyl	“	“	“	“
Thiozolidine	“	“	NA*	“
o-Methoxy phenyl	“	25	50	“
p-Bromo phenyl	2,4 – Dinitrophenyl	“	NA*	“
p-Chloro phenyl	“	“	25	“
NA* indicates no activity	“	“	50	“
p-Nitro phenyl	“	“	“	“

REFERENCES

- 1 Ahluwalia, N.K Dutta Uttara, Sharma, H.R., J Indian Chem Soc, 64(4), 221-223, 1987.
- 2 Mogaliah, K., and Reddy Vijayender,K,. J Indian chem.. Soc., 62(3), 256-261,1985.
- 3 Sunil Desai, B., M.Pharm Dissertation, Karmataka University, Dharwad, Unpublished (1990)
- 4 Pathak, U.S., Devani, M.B., Shishoo, C.J {ate; H.H Gandhi T.P and Patel R.B Indian J. chem., 27 () 602-604, 1988.
- 5 Betzing Hans, graf Erich Leyer sigurd, chem. Abstr., 99,70572f, 1983.
- 6 Pfizer Inc., Chem Abstr, 99,212418m, 1981
- 7 Suto, Y., Shimoji, Y., Fujita H., et al, J Med chem., 23,927, 1989.
- 8 Ohno sachio Mizokushi, Kioshi Kamastursmur, chem., abstr, 23,927, 1983.
- 9 Nargund, L.V.G Reddy, Y.S.R and Robin Jose, Indian drugs., 29(1), 45-46, 1991.
- 10 Indian pharmacopoeia, 3rd edition, Vol II, A 90, 1985.