

α 1-Adrenergic and 5-HT₂-serotonergic effects of some α -alkoxy- α -phenylethylamines on isolated rat thoracic aorta

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1. Ten racemic α -alkoxy- α -phenylethylamines were found to elicit concentration dependent contraction of the isolated rat thoracic aorta with apparent pot values in the 4.56-6.76 range. With one exception, which produces the same maximal contraction ($E(\max)$) as serotonin (5-HT), the E , values attained with these compounds are lower than those produced by either 5-HT or norepinephrine (NE). 2. Pretreatment with either prazosin or ketanserin (10^{-8} M) leads in most cases to decreased $E(\max)$ values and slopes in the dose-response curves. Apparent serotonergic ($pD_2(S)$) and adrenergic ($pD_2(A)$) pD_2 values going from 4.22 to 6.08 ($pD_2(S)$) and from 3.87 to 5.27 ($pD_2(A)$) were calculated from results obtained in the presence of prazosin or ketanserin, respectively. 3. In the 10^{-7} - 10^{-5} M range, and in contrast with the results obtained with the previous compounds BON [(±) 2-(2,5 dimethoxy-4-nitrophenyl)-2-methoxy-ethylamine] behaves as an antagonist to both 5-HT and NE (apparent $pA_2 = 7.08$ and 7.45