3,4-Methylenedioxymethamphetamine (MDMA) Induced Hyperthermia-The Role of Pro-Inflammatory Cytokines

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**INTRODUCTION**

The main 3,4-methylenedioxymethamphetamine (MDMA)-induced adverse effect is disruption of normal thermoregulation leading to life threatening hyperthermia which is exacerbated by high ambient temperature and linked to chronic neurotoxicity [1-3]. Although the focus of the majority of research on MDMA-induced loss of thermoregulation has been on brain serotonergic and dopaminergic systems, results obtained from recent studies suggest that microglial activation and the subsequent release of the pyrogen interleukin-1β could also contribute to these effects [4, 5]. The aim of this study is to extend our understanding of the underlying mechanisms in the brain leading to the disruption of normal thermoregulation. We have examined the effect of interleukin 1 receptor antagonist (IL-1RA) on MDMA-induced hyperthermia and behavioural effects.

**METHODS**

Male Sprague Dawley rats maintained at normal ambient temperature (22 ± 1°C) were administered 10 mg/kg (i.p.) of MDMA and core body temperature and behavioural measurements were taken every 30 min for 4 hours. The interleukin 1 receptor antagonist (IL-1RA, 40 mg/kg i.p.) was administered 30 min prior to MDMA administration. Core body temperature measurements were analysed using a two-way ANOVA with Bonferroni’s post-hoc test. Behavioural data were compared using the Kruskal-Wallis test. P < 0.05 was taken as significant for all analyses.

**RESULTS**

Fig. (1) illustrates the core body temperature response for each treatment group. No significant changes in core body temperature were recorded for Saline+Saline or IL-1RA+Saline treated rats (P > 0.05, n=5-9). After Saline+MDMA treatment, core body temperature increased by 2.4 °C and significant hyperthermic response was maintained from 30 min up to 240 min post MDMA compared to control group (P < 0.05, n=11). Although IL-1RA+MDMA treated animals showed similar hyperthermic response as MDMA+Saline group up to 120 min, all animals pre-treated with IL-1RA showed a significant decrease in body temperature at 180, 210 and 240 min. MDMA-treated animals had significantly increased behaviour scores over saline-only treated controls (P < 0.05, n=12). IL-1RA pre-treatment did not produce any significant effect on MDMA-induced behavioural effect.

**DISCUSSION**

IL-1RA pre-treatment significantly attenuated the magnitude and duration of hyperthermia induced by MDMA administration. IL-1RA pre-treatment had no significant impact on MDMA-induced behavioural stimulation. The results of this study indicate that the mechanisms underlying MDMA induced hyperthermia may include the release of the endogenous pyrogen interleukin-1β in addition to serotonin.
and dopamine mediated hyperthermic response and behavioural stimulation.

**CONCLUSION**

The present findings are consistent with the hypothesis that inflammatory processes play a role in MDMA induced adverse effects [3,5] and increase in expression of pro-inflammatory cytokines further exacerbate and sustain the hyperthermia caused by MDMA.

**REFERENCES**


