This study re-examined the hyperactivity and disruption of *prepulse inhibition* induced by Nmethyl-D-aspartate stimulation ... of the rat ventral hippocampus

haloperidol or clozapine. While the hyperactivity is thought to depend on dopamine receptor activation in the nucleus accumbens, the dopamine D2-class receptor blocker haloperidol failed to antagonize the disruption of prepulse *inhibition* in previous studies. However, an ameliorative effect of the atypical

neuroleptic clozapine on disruption of prepulse inhibition was suggested by ...

We conclude that dopaminergic mechanisms are differentially involved in the hyperactivity and disruption of prepulse inhibition induced by Nmethyl-D-

and compared how both effects were affected by pretreatment with either

Biomedical Abstract

In the present study, bilateral infusion of Nmethyl-D-aspartate ... into the ventral hippocampus of Wistar rats increased ... disrupted *prepulse inhibition*. Both effects were observed immediately after infusion but disappeared 24h later. Injection of ..., 45min prior to Nmethyl-D-aspartate infusion, totally antagonized the hyperactivity but did not affect the disruption of *prepulse inhibition*.

aspartate stimulation of the ventral hippocampus. negative-activation

Inputs Supporting Sentences

Outputs

Mechanism Sentence

2. Relation: negative-activation

2. Regulated Entity: prepulse inhibition 3. Regulator Entity: Nmethyl-D-aspartate