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A: You can use the following regex to match the pattern with 2 subpatterns (starts with and ends with) and a space in between: `^\s*(.*?)\s*$` OR `^\s*(.*?)\s*(.*)\s*$` This should capture the string you are looking for. Demo

Amino acids have important roles in nitrogen metabolism and in regulating normal physiological functions in both the central nervous system and the peripheral nervous system (Kominami and Hikosaka, 1997; Spira et al., 2001). The amino acids glutamate and aspartate are particularly important in the normal function of the central nervous system in the mammalian brain (Dudley and Will, 1998; Norenberg et al., 2001). Glutamate is the predominant excitatory neurotransmitter in the brain, and the aspartate is a co-agonist with glutamate. It is well known that the administration of glutamate causes both acute and chronic neuronal damage (Olney et al., 1992; Kominami and Hikosaka, 1997; Norenberg et al., 2001). Therefore, it is important to develop methods of relieving the symptoms of neuronal damage caused by excess exogenous glutamate. Glutamate in the brain is highly susceptible to oxidation by free radicals, and this leads to neurotoxicity (Fenizia et al., 2004; MacDonald, 2004). It has been reported that naturally occurring sulfur-containing amino acids such as cysteine and glutathione and a derivative of glutamate, N-acetylcysteine (NAC), are effective in the treatment of neuronal damage induced by glutamate

excess (Kirin et al., 1996; MacDonald, 2004). NAC is a drug that has been approved for the treatment of acetaminophen overdose and the treatment of acute liver failure (Kisin et al., 2000). However, the use of NAC is restricted due to its high cost. Therefore, it is desirable to develop a substance which can be used as an effective and inexpensive treatment of neuronal damage caused by glutamate excess. Many studies have been conducted in recent years to develop novel cysteine compounds which are effective as a glutamate antagonist. One such compound is cysteamine, which has been used as an anti-ulcer drug. Although, the administration of cysteamine does not cause neuronal damage or any other adverse effects, cysteamine has little affinity for NMDA receptors and this limits its 6d1f23a050

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