Adding metoclopramide to paroxetine induced extrapyramidal symptoms and hyperprolactinemia in a depressed woman: a case report

Ryohei Igata
Hikaru Hori
Kiyokazu Atake
Asuka Katsuki
Jun Nakamura
Department of Psychiatry, University of Occupational and Environmental Health, Kitakyushu, Japan

Abstract: A 54-year-old Japanese woman was diagnosed with major depressive disorder and prescribed paroxetine 20 mg/day. In around May 2013, the patient experienced gastric discomfort, so metoclopramide was prescribed. Beginning on June 4, 2013, the patient was given metoclopramide, 10 mg intravenously, twice per week. On the seventh day after beginning metoclopramide, facial hot flushes, increased sweating, muscle rigidity, and galactorrhea were noted. Extrapyramidal symptoms (EPS) rapidly subsided in response to an intramuscular injection of biperiden. Blood biochemical tests revealed an elevated serum prolactin level of 44 ng/mL. After stopping metoclopramide, EPS disappeared. Serum prolactin level decreased to 15 ng/mL after 4 weeks. In our case, although no adverse reactions had previously occurred following the administration of metoclopramide, the patient developed EPS and hyperprolactinemia following the administration of this antiemetic in combination with paroxetine. Paroxetine and metoclopramide are mainly metabolized by CYP2D6, and they are inhibitors for CYP2D6. We report a case with EPS and hyperprolactinemia whose plasma paroxetine and metoclopramide level rapidly increased after the addition of metoclopramide. Our experience warrants the issuing of a precaution that adverse reactions may arise following the coadministration of metoclopramide and paroxetine even at their respective standard dose levels.

Keywords: metoclopramide, paroxetine, extrapyramidal symptoms, SSRI, hyperprolactinemia, depression

Introduction
Selective serotonin reuptake inhibitors (SSRIs) are the first-line antidepressants used in primary care and psychiatric practices. Paroxetine, one of the most potent SSRIs, is widely used in the treatment of depression and is a strong selective CYP2D6 inhibitor. Metoclopramide is a drug with a highly potent antiemetic effect and is considered to cause relatively few adverse reactions. In this report, the extrapyramidal symptoms and hyperprolactinemia occurred in a patient receiving a standard dose of metoclopramide concomitantly with an SSRI for the relief of gastrointestinal symptoms, which were due to the exacerbation of depression.

Case report
Written informed consent was obtained from the patient to publish this paper. A 54-year-old Japanese woman presented with depressed mood, psychomotor retardation, and loss of interest. She was diagnosed with major depressive disorder according to the Diagnostic and Statistical Manual of Mental Disorders, 4th Edition (DSM-IV-TR) in 2006 and was subsequently prescribed paroxetine 20 mg/day. In around May 2013,
that is often prescribed for the management of depression, prolactinemia following the administration of this antiemetic. 

In our case, the extrapyramidal symptoms and hyperprolactinemia occurred in a patient receiving a standard dose of metoclopramide concomitantly with an SSRI for the relief of gastrointestinal symptoms, which were due to the exacerbation of depression. Our experience with this patient warrants the issuing of a precaution that adverse reactions may arise following the coadministration of metoclopramide and paroxetine even at their respective standard dose levels.

Discussion
Metoclopramide is a drug with a highly potent antiemetic effect and is considered to cause relatively few adverse reactions. Several reports described that metoclopramide caused extrapyramidal symptoms via its inhibitory effect on dopaminergic neurons. In most of these papers, it was speculated that the extrapyramidal symptoms were evoked when the drug was administered in high doses. In our case, however, although no adverse reactions had previously occurred following the administration of metoclopramide, the patient developed extrapyramidal symptoms and hyperprolactinemia following the administration of this antiemetic in combination with paroxetine. Paroxetine is an SSRI that is often prescribed for the management of depression, panic disorder, and obsessive–compulsive disorder. There have been several reports indicating the occurrence of extrapyramidal symptoms due to SSRI administration. The underlying mechanism of these symptoms is thought to be due to excessive serotonin, which exerts an inhibitory effect on nigrostriatal dopaminergic neurons.

Moreover, paroxetine and metoclopramide are mainly metabolized by cytochrome P450 (CYP) 2D6. Paroxetine is a potent inhibitor for CYP2D6, and metoclopramide is a moderate inhibitor for CYP2D6. We report a case with extrapyramidal symptoms and hyperprolactinemia whose plasma paroxetine and metoclopramide level rapidly increased after the addition of metoclopramide.

Disclosure
The authors report no conflicts of interest in this work.

References