Adverse drug reaction of the month

Bipolar disorder associated with interferon-alpha treatment

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Interferon-alpha (Hoffman-La Roche Ltd, Basel, Switzerland) is a highly purified, bacterially synthesized agent with antiviral, antiproliferative and immunomodulatory properties.1 Interferon-alpha has been approved for the treatment of several types of cancer (eg, leukaemia, lymphoma, renal cell carcinoma and melanoma) and of chronic hepatitis.2,4 Usually causing flu-like symptoms, interferon-alpha’s most serious adverse effects are those related to the central nervous system, including psychiatric side-effects.1,5,6 Psychiatric adverse effects include anxiety, irritability, depression, psychosis and delirium1,6-9 and even suicidality.10

We report the precipitation of bipolar disorder after interferon-alpha therapy for chronic myeloid leukaemia in a previously healthy 47-year-old man. We believe that this constitutes the first report of development of bipolar disorder in association with interferon-alpha treatment.

Case report

A 47-year-old Jewish man of Eastern European descent was referred to our psychiatric emergency room by his haematologist due to insomnia. His parents had divorced when he was 15 years old. He was married and was the father of three children. He had always been introverted with a low self-image. He exhibited learning difficulties and bad social ties. After full-term military service, he worked at different jobs and for the last 10 years he had worked as a container contractor. There was no history of familial psychopathology; he had used flunitrazepam (2 mg/day) for sleep induction for the last year. Seven years ago he had developed Graves disease which was treated successfully with mercaptizol. No thyroid dysfunction was noted at follow-up.

One year previously, after a routine blood count revealed leukocytosis, he underwent a bone marrow biopsy and was diagnosed as having chronic myeloid leukaemia (Philadelphia positive). He was asymptomatic and there was no evidence of hepatosplenomegaly or lymph node enlargement. He began treatment with interferon-alpha 9 million units/day, which he had received for seven months until his admission.

While on this treatment, he reported flu-like symptoms and impotence, but continued to function. The haematologist decided to continue the interferon treatment due to partial improvement in the blood cell count. How-

ever, in the last three months the patient became gradually euphoric and hyperactive, and also exhibited tension, insomnia and talkativeness. He began to spend more money and bought a great deal of car devices. In the last month his condition worsened and finally he had not slept for a whole week and was very irritable. He felt very self-confident, thought that he was highly sociable, superior, and analytical and that he was going to become rich. He also showed increased interest in his wife’s sexual pleasures.

On admission he was agitated and had a fluent speech, which was difficult to stop. His affect was euphoric and labile. The thought process was circumstantial. He revealed megalomaniacal thoughts and complimented his doctor on being the greatest psychiatrist in the world! No hallucinations were revealed.

Physical examination was normal (including fundus). Laboratory tests revealed 18 000 leucocytes with 80% polymorphonucleocytes (without signs of chronic myeloid leukaemia); low thyroxine and free thyroxine index and high thyroid-stimulating hormone (another side effect of the interferon-alpha treatment). Brain computed tomography (CT) and electroencephalography (EEG) were normal. A Mini-Mental State Examination revealed normal cognitive functioning (score of 28).

A diagnosis of manic episode with psychotic features was made and the patient was started on perphenazine 32 mg/day and clonazepam 2 mg/day. Gradually, the hyperactivity and insomnia abated. His score on the Brief Psychiatric Rating Scale decreased from an initial score of 50 to 20 after a week (60% improvement). His irritability and delusions of superiority gradually abated and after two weeks he stopped the medication and discharged himself against medical advice. After a short period of euthymia, he developed a moderate depression which resolved slowly after six months with fluvoxamine (250 mg/day) and lithium (900 mg/day) therapy.

Discussion

At admission, the patient had no signs of chronic myeloid leukaemia and his CT and EEG were normal, as was neurological examination. Thus, we believe that the occurrence of the manic episode was not related to the chronic myeloid leukaemia. Also, he had not interrupted his benzodiazepine use, ruling out withdrawal as aetiology of the mania. The interruption of interferon-alpha did not lead to

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full recovery, however, so we assume that interferon-alpha precipitated the development of a latent bipolar disorder.

Jannsen et al 10 reported a short (four week) manic psychosis in a 49-year-old patient treated with interferon-alpha for hepatitis B. Van Thiel et al 11 reported manic exacerbation in two patients with bipolar disorder treated with interferon-alpha for hepatitis C. The interruption of interferon-alpha, together with a lithium dose increase, enabled a quick recovery in these patients (Van Thiel, personal communication, 1995).

The exact role of interferon-alpha in the origin of psychiatric side-effects is as yet unknown. Although its passage through the blood–brain barrier is limited due to its being a large hydrophilic protein, interferon-alpha has a direct toxic effect on vasculature and thus might diffuse into the brain. Psychiatric symptoms and EEG abnormalities during therapy have been described in detail. 6,10 Interferon-alpha or one of its metabolites could be directly neurotoxic, but may also induce other substances which interfere with neuronal excitation or neuroendocrine regulation. 12,13 Patients with brain metastases or other central nervous system abnormalities and those using narcotics may experience heightened neurotoxicity. Patients over 40 years old and those receiving doses higher than 6 million units are susceptible to psychiatric side-effects. 7 These side-effects usually clear within two to three weeks. 7

As the number of patients treated with interferon-alpha is increasing, its range of psychiatric side-effects should be well recognised. In most cases the management strategy will depend on the extent and severity of the problem, but will usually involve dosage reduction, intermittent breaks in treatment, or psychotropic treatment. 1,6–11

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