Thyrotoxicosis followed by hypothyroidism in a patient on lithium

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Abstract

While hypothyroidism is common in lithium-treated patients, thyrotoxicosis is rarely reported. We present a female patient on lithium for maintenance therapy of bipolar affective disorder, who developed thyrotoxicosis for few months which was followed by hypothyroidism which continued. There was no further thyrotoxicosis episode during a five year follow up period. While she was treated for thyroid dysfunction, lithium was continued. There was no clinical impact on the maintenance of the bipolar affective disorder during the follow up period; she was maintained well in the community.

Case Report

A 61-year-old female patient was suffering bipolar affective disorder for 23 years and was maintained on lithium along with various psychotropic medications over the years. In the past she had comorbid diagnoses of alcohol dependence syndrome, obsessive compulsive disorder, benzodiazepine (temazepam) and nicotine dependence. She also experienced paranoid symptoms secondary to alcohol dependence. She was successfully treated for these conditions. There was history of self-harm attempts by overdose of prescribed medications a few times.

She developed thyrotoxicosis after being on lithium for around 18 years. There was no history of thyroid abnormality in the past and she was maintaining euthyroid state during the course of lithium therapy until she developed thyrotoxicosis. She was compliant to lithium. During thyrotoxicosis, she had many instances of confusion suggestive of delirium.

She was admitted to medicine ward and commenced on carbimazole. Gradually the thyrotoxicosis symptoms became stable. The free T4 levels remained high for about three months at 57, 72, 40 pmol/L (normal range: 12.0-22.0 pmol/L); and the TSH values were 0.01 mU/L (normal range: 0.27-4.20 mU/L). Lithium was continued during the thyrotoxicosis phase. Close monitoring of thyroid status was maintained. After around 5 months of treatment for thyrotoxicosis, the thyroid-stimulating hormone (TSH) level increased beyond the normal range. Consequently to that, the dose of carbimazole was decreased. However TSH level remained high and it was considered that she had developed acquired hypothyroidism possibly due to carbimazole and lithium. Thyroxine was started. The TSH remained high (up to 9.5 mU/L) for around 7 months before becoming normal; although there were a few fluctuations.

The lithium level was within therapeutic range at the onset of thyrotoxicosis. The lithium levels of the patient had fluctuated and she had one recording of lithium level higher than 1.2 mmol/L around 5 months before the onset of thyrotoxicosis; and a few similar recordings around one year following the thyrotoxicosis episode. These were dealt by reducing the lithium dose and repeating the tests. TSH and lithium levels remained within usual range during the follow-up period over 5 years after the described phase of thyroid dysfunction. Besides lithium, she also had olanzapine as another psychotropic medication which was continued during thyrotoxicosis and at follow up. There were no specific episodes of bipolar affective disorder during or following the active phase of thyroid abnormality. Mood symptoms remained under control without any exacerbations. There were occasional obsessive doubts and anxiety symptoms. Her psychiatric treatment continued as usual; and she did not require any additional psychiatric input like crisis intervention, support from home treatment team or admission.

Gradually the thyroid abnormality was contained. The thyroid function tests became normal. During a five year follow up period, she did not have any further episode of thyrotoxicosis. Her thyroid function tests remained mostly within normal range with an occasional brief phase of dysregulation, needing adjustment in thyroxine dose.
Discussion

The thyroid abnormality in this patient occurred after 18 years of being on lithium at an age of 56. Thyroid abnormality associated with lithium has been reported to be more common in women and increased with age, which fits in to the description of the case described here. It may be highlighted that lithium could be continued while the thyroid dysfunction was treated. Considering the mood symptoms she maintained well in community. In this case the thyrotoxicosis was short lasting; which was followed by a prolonged hypothyroidism. Opposite presentation i.e. thyrotoxicosis following hypothyroidism has also been reported in a lithium-treated patient.

The relationship of lithium and hyperthyroidism which occurs rarely seems not to be casual, since hyperthyroidism in these patients occurs three times that of the normal population. There is clearly an increased risk of thyrotoxicosis associated with long-term lithium therapy. Various reasons have been considered for this association e.g. diffuse toxic goiter, multinodular toxic goiter, silent/painless thyroiditis have been reported in these subjects. Autoimmune mechanisms for the thyrotoxicosis have also been suggested. The toxic and immunomodulatory roles of lithium and perhaps genetic and dietary factors too have been implicated. According to these reports, it appears that lithium associated thyrotoxicosis is a heterogeneous condition with different kinds of underlying thyroid pathologies and the mechanisms remain uncertain.

The initial thyrotoxicosis in this case for few months could possibly due to thyroiditis; while the later acquired hypothyroidism could be due to either carbimazole or lithium, or both. Transient thyrotoxicosis has been reported in lithium-treated patients. It has been postulated that lithium might directly damage thyroid follicular cells and that subsequent release of thyroglobulin into the circulation might be a cause of transient thyrotoxicosis. There is a report where thyroiditis improved after withdrawal of lithium, which is consistent with the theory of lithium’s direct toxic effect on thyrocytes. As the patient was euthyroid before the onset of thyrotoxicosis and was compliant to lithium when the thyrotoxicosis symptoms appeared, probability of latent hyperthyroidism was unlikely.

There are reports of unmasking of latent hyperthyroidism when lithium is being completely or partially discontinued. As lithium suppresses thyroid function, so the early stages of thyrotoxicosis may be clinically undetectable. The symptoms and signs of thyrotoxicosis may be confused with the side effects of lithium and if lithium is withdrawn severe rebound thyrotoxicity may occur.

Conclusions

In conclusion, the findings suggest that although rare, thyrotoxicosis appears to be associated with long-term lithium therapy and reemphasize periodic clinical evaluation for thyroid dysfunction and thyroid function tests in lithium-treated patients, irrespective of the record of euthyroid state while on lithium. In the presented case, lithium remained effective in its maintenance role for bipolar affective disorder, following the onset of thyroid dysfunction, its course and treatment. There appears no need to discontinue lithium in the presence of thyroid abnormality in well maintained bipolar patients, especially if the thyroid abnormality is controlled.

References