Case Reports

THYROTOXIC CRISIS AND TRANSIENT MYASTHENIA GRAVIS IN PREGNANCY

P. HORROCKS, M.B., Ch.B.
Senior House Officer

Withington Hospital, Manchester.

J. C. LEONARD, M.D., M.R.C.P.
Physician

Withington Hospital, Manchester.

With safer premedication for thyroidectomy, thyrotoxic crisis is now seldom seen postoperatively. It still occurs, however, when infections or surgical operations complicate uncontrolled thyrotoxicosis, and it is a rare complication of thyrotoxicosis during pregnancy and especially after delivery. We report here a patient who developed a severe thyrotoxic crisis in the ninth month of pregnancy, and whose illness was complicated by transient myasthenia gravis. So far as we can determine, there is no previous record of myasthenia occurring in a thyrotoxic crisis.

Case Report

The patient, a married woman aged 20 years, was seen early in her first pregnancy by her general practitioner on September 17th, 1964, complaining of nausea, vomiting and dizziness for which trifluc-perazine was prescribed. A tachycardia was noted then, and her doctor "wondered if she was thyrotoxic". No further action was taken and her pregnancy seemed to make good progress thereafter, although she had gained only 6.4 kg. (14 lb.) by the thirty-sixth week.

She was admitted to Withington Hospital as an emergency on March 24th, 1964. During the previous eight days, she had had severe bilateral throbbing frontal headaches, tiredness and a dry cough. Three days before admission, she developed profuse sweating and rigors; the headaches and fatigue became worse, she vomited twice and felt very ill. On the evening of admission, her temperature had risen to 106.4°F. and her pulse rate to 180/min.

On examination at the hospital, she was thin, flushed, apathetic; sweating and very ill; the hands were warm and the temperature was 103.4°F. There was a marked stare and von Graefe's sign was positive; there was no tremor. The thyroid gland was just palpable, and there was a thrill and a loud continuous murmur over it. The pulse rate was 144/min., B.P. 95/45 mm. Hg., and J.V.P. raised 10 cm. above the sternal angle; no oedema. The heart was hyperdynamic; there was a loud gallop rhythm and a basal ejection systolic murmur conducted to the neck. Examination of the chest was normal. The abdomen contained a gravid uterus consistent with a thirty-six weeks' pregnancy. The tendon reflexes were hyperactive.

Investigations. Hb 8.6 g./100 ml.; WBC 6,500/cu. mm., normal differential count. The blood urea, serum sodium, chloride and bicarbonate normal; serum potassium 2.95 mEq/l. Chest X-ray normal. ECG: sinus tachycardia. Lumbar puncture: normal CSF normal pressure. A twenty-four-hour specimen of urine contained 250 μg. of catecholamines. The serological tests for autoimmune thyroiditis were negative. Unfortunately, treatment with iodide was begun before a specimen of blood for estimation of protein-bound iodine was obtained.

On admission, at 1 a.m. on March 24th, she was gravely ill and a clinical diagnosis of thyrotoxic crisis was made. She was given an intravenous infusion of sodium iodide 1 g. six-hourly in 5 per cent dextrose; she also received phenobarbital 200 mg. and chlorpromazine 50 mg. eight-hourly, Lugol's iodine 10 min. t.d.s., prednisolone 10 mg. t.d.s., reserpine 1 mg. t.d.s., carbimazole 20 mg. t.d.s. and digoxin. Her condition rapidly improved (see chart), and ten hours later her temperature was 95.6°F., and the pulse rate 104 per minute. After twenty-four hours, sodium iodide was withdrawn for six hours but her condition immediately relapsed. After a further six hours the temperature was 104.2°F., and the pulse rate 180 per minute. The original dosage of sodium iodide was restored and she again responded; by mid-day on March 25th, the temperature was normal and the pulse rate 100 per minute. This treatment was continued and after her condition had remained satisfactory for 24 hours a further attempt was made to reduce the dosage of sodium iodide because of the theoretical risk of acute iodism. At the same time, guanethidine was added in an attempt to suppress the manifestations of thyrotoxicosis. A further deterioration with weakness, cough, fever and tachycardia again ensued within a few hours of reducing the dosage of sodium iodide and her condition rapidly became alarming. She became semicomatose and a left sixth nerve palsy developed. Within four hours, she developed ptosis, divergent squint, impairment of upward gaze of the eyes, a bilateral lower motor neurone facial palsy, dysphagia, dysarthria and a "myasthenic snarl". 10 mg. of edrophonium chloride (Tensilon) was given intravenously and caused a dramatic temporary remission of the myasthenia.

Large doses of sodium iodide were re-introduced intravenously, steroids were gradually withdrawn for fear of exacerbating the myasthenia, neostigmine was given by mouth, and a course of chloramphenicol was given in case an underlying infection was responsible for precipitating and perpetuating the thyrotoxic crisis. The latter again came under control within a few hours, but after 24 hours the myasthenia was unchanged. Neostigmine was then added to the intravenous infusion in a dosage of 7.5 mg. (later 9 mg.) six-hourly; after a further 24 hours there was slow improvement in the neurological signs. Her general condition then began to improve and after two days the thyrotoxicosis was under control and the myasthenia had become completely suppressed. On March 30th an alarming episode occurred. Whist talking to a nurse, the patient suddenly lost consciousness and had a generalised tonic convulsion. This was thought to be due to overdosage of neostig-
mine, as the patient was still receiving this drug by slow intravenous infusion; the drug was immediately withdrawn and the patient given 0.5 mg. of atropine. She gradually recovered consciousness and was apparently no worse for this attack. From then onwards, her progress was rapid: the myasthenia never relapsed and no further neostigmine was required. Intravenous iodide was also withdrawn after this attack, but had to be given two days later in the face of returning fever and tachycardia. It was only after a further four days that intravenous iodide could be finally withdrawn—thirteen days after admission. Throughout this period, the patient had also received carbimazole 20 mg. t.d.s., Lugol's iodine 10 min. t.d.s. and digoxin. Reserpine, guanethidine, phenobarbitone and chlorpromazine...
were gradually discontinued. A course of intramuscular iron was given for her anaemia. The thyrotoxicosis was well-controlled and she began to regain some of the weight lost during her illness. By the time she went into labour on April 26th, the patient was taking only carbimazole, Lugol's iodine and digoxin. Labour was uneventful, and the patient was delivered of a healthy girl weighing 6 lbs. 3 oz. (2.9 kg.). The infant was clinically normal and made excellent progress. No trouble ensued in the peripartum and Lugol's iodine and digoxin were withdrawn. She was discharged home taking carbimazole 10 mg. t.d.s. on May 11th.

She was seen at frequent intervals in the outpatients' clinic and remained well. The dose of carbimazole was reduced to 10 mg. b.d. on August 8th. On September 12th, carbimazole was discontinued and Lugol's iodine substituted. She was re-admitted to hospital on September 29th., and subtotal thyroidectomy was performed by Mr. Alan Nicholson on October 5th. On the next day she developed tetany, and the serum calcium was 7.9 mg./100 ml. She was given seven injections of 10 ml. of 10 per cent calcium gluconate during the next two days, and was given calciferol by mouth in gradually decreasing doses. The tetany rapidly subsided and all treatment was discontinued on October 24th. When she was last seen, on May 19th, 1965, she was quite well. The histological report of the thyroid gland was as follows: "There is mild epithelial hyperactivity throughout the gland, but colloid production is abundant; no lymphoid infiltration present. Extra sections have been taken from the posterior aspect of the thyroid lobes but no parathyroid glands have been encountered." (Dr. James Davson.)

Discussion

In his excellent description of thyrotoxic crisis, Lahey (1928) emphasises that most patients are extremely agitated and nervous, with a marked tremor and often delirium. He recognised, however, that some were very apathetic (as in our case) and similar patients have been reported culminating in thyrotoxic coma (Weaver, Jones and Smith, 1956).

Myasthenia gravis and thyrotoxicosis are known to co-exist more commonly than would be expected by chance, but the nature of the association is very obscure. Sometimes a "seesaw" effect has been described, the myasthenia appearing as the thyrotoxicosis comes under control (Robbins and Burkle, 1960; Monro, 1963). Weickhardt and Redmond (1960) throw doubt on this and point out that sometimes myasthenia may appear when the patient is severely thyrotoxic, as in our own case. It is clear from published cases only that there is no constant pattern. One factor which may have operated in our own patient is that the myasthenia followed large doses of chlorpromazine, which is thought sometimes to precipitate weakness in a myasthenic patient. (McQuillan, Gross and Johns, 1963).

Our patient was given so many different agents during the height of her illness that it may be thought difficult to identify those that were therapeutically most useful. The management of thyrotoxic crisis in recent years has been improved by the introduction of chlorpromazine, and possibly also by corticosteroids, guanethidine and reserpine; the latter two drugs are useful in suppressing the peripheral manifestations of thyrotoxicosis whilst not curing the disease. Corticosteroids are advised because of the possibility of adrenocortical exhaustion in thyrotoxic crisis. In spite of the use of all these drugs, however, it was our distinct impression that intravenous sodium iodide was by far the most effective agent in the management of our patient. We repeatedly observed that increasing the concentration of iodide in the infusion quickly led to reduction in fever and tachycardia, whilst both relapsed whenever iodide was discontinued. A daily dose of 4 g. of iodide was tolerated without evidence of ill effect (see chart).

It is therefore suggested that in the management of thyrotoxic crisis, intravenous sodium iodide should be given in a dosage of at least 0.5 g. six-hourly. Reductions in the dose should be made cautiously, and the dose increased if fever and tachycardia return or are uncontrolled. The patient should be nursed in an oxygen tent; skilled nursing care and observation are essential. Hyperpyrexia and restlessness should be controlled by chlorpromazine, tepid sponging and pheno- barbitone. Intravenous hydrocortisone should also be given. Guanethidine and reserpine may be used for their symptomatic value. A careful search must be made for any precipitating infection, and appropriate treatment instituted. Heart failure must be controlled by digoxin; the value of propranolol in severe thyrotoxicosis is currently being assessed. Carbimazole should be started immediately, but will not have much therapeutic effect for two weeks.

Summary

A woman who developed a thyrotoxic crisis in the ninth month of pregnancy is described. At the height of her illness, she developed neostigmine-responsive myasthenia lasting four days. The management of thyrotoxic crisis is discussed: although chlorpromazine, phenobarbitone, corticosteroids, guanethidine and reserpine probably played a useful part, it is emphasised that the use of these drugs does not render superfluous the use of intravenous iodide; in our patient this appeared to be the most effective agent.

We are very grateful to Dr. L. A. Liversedge, Dr. Donald Longson, Mr. R. H. Martin and Mr. Alan Nicholson for invaluable help in the management of this patient. We also thank Sister K. Murphy and her nursing staff for their devoted and efficient care of this patient, without which she would not have survived.

REFERENCES


APLASTIC ANAEMIA DUE TO PHENYL BUTAZONE

E. A. CAMERON, M.B., Ch.B., M.R.C.P.E.,
Consultant Physician.

A. A. EISEN, M.R.C.S., L.R.C.P.
Registrar.

L. M. NIRANJAN, M.B., B.S.
House Physician.

From Ashington General Hospital, Northumberland.

It has been estimated (Annotation, 1962) that in any given week in Great Britain alone, 100,000 people are taking phenylbutazone. There are several well known side effects from this drug, and a considerable number of cases in which marrow depression has occurred have already been reported (Leonard, 1953; Kersley and Mandell, 1963; Kaplin, 1955; Venning, 1957; Hale and DeGruchy, 1960; Rankin, 1961; Lander and Bonnin, 1962; McCarthy and Chalmers, 1964; Humble, 1964; Woodliff and Dougan, 1964). Our search of the literature has revealed 15 cases of aplastic anaemia in patients taking phenylbutazone; in 7 of these cases the marrow failure proved fatal.

We record the following case of fatal aplastic anaemia because it re-emphasises the possible dangers of this drug and the need for extreme wariness when it is being used.

Case Report

A 79-year-old widow was admitted to hospital on the 6th of November, 1964. Pernicious anaemia had been diagnosed in 1954 and had been well controlled with cytamen given every three weeks. About the end of 1963 she had complained of “arthritic pains” in the hands and legs, and her own doctor prescribed phenylbutazone, 100 mg. tablets, one thrice daily. In fact she took the tablets rather erratically, so that the total amount taken over a period of 11 months was approximately 10 g. (about 100 tablets). Three weeks prior to admission spontaneous bruises appeared on the right arm, and shortly after she noticed what transpired to be a typical purpuric rash on both lower limbs. Phenylbutazone was stopped on October 14th, 1964.

On admission she was well nourished, rather obese and obviously anaemic, but not jaundiced or cyanosed. The tongue was covered with small purpuric lesions, also noted in other parts of the buccal cavity. There was a one centimetre bruise on the lower lip, which was superficially infected. There was no lymphadenopathy. Pulse regular, 100/min., normal volume; the blood pressure was 130/80 mm. Hg. JVP not raised; no sacral or ankle oedema. The heart was not clinically enlarged and heart sounds were normal. The liver and spleen were impalpable. The central nervous system was normal. In the right fundus there were haemorrhages around the disc margin; the left fundus was normal. There was no evidence of peripheral neuropathy or of lateral column involvement as might have occurred in association with her known pernicious anaemia.

Initial Investigations: Hb. 5.6 g./100 ml., RBC 2,000,000/cu. mm. (retics 0.2%), PCV 19%, MCHC 31%, ESR 136 mm./1 hour, WBC 2,100/cu. mm. (Polys. 8%, Lymph. 90%, Lymphoblasts 1%, Monos. 1%). Platelets 45,000/cu. mm. Bleeding Time (McFarlane) 24 min., Clotting Time (Capillary) 4 min., Prothrombin Activity (Thrombo Test) 100% of average normal, Direct Coombs Test negative, Serum Iron 157 µg./100 ml. Blood Group ‘A’ positive.

Blood film: The blood film showed normal erythrocytes with occasional late normoblasts; an occasional primitive leucocyte was present and agranulocytosis was obvious. Platelets were very sparse.

Sternal Marrow: Sections of the marrow showed 95% fat with only an occasional island of haemopoietic tissue (Fig. 1a & b). These islands were mainly lymphocytes and showed no polymorphonuclear cells or megakaryocytes. Differential count showed a leuco-erythroid ratio of 5 to 1. The leucopoietic tissue was 75% lymphocytes with a few lymphoblasts. Primitive cells were not frequent. Erythropoiesis was normoblastic with no arrest of maturation. The overall picture was that of aplasia, affecting most severely thrombocytes and granulocytes; to a slightly less degree the erythrocytes.
Thyrotoxic crisis and transient myasthenia gravis in pregnancy.
P. Horrocks and J. C. Leonard

Postgrad Med J 1966 42: 46-49
doi: 10.1136/pgmj.42.483.46

Updated information and services can be found at:
http://pmj.bmj.com/content/42/483/46.citation

These include:

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/