Thymoma with Myasthenia Gravis in Pregnancy: A Case Report

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ABSTRACT

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Keywords

thymoma, thymic carcinoma, mediastinal mass, myasthenia gravis, pregnancy

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INTRODUCTION

Thymoma is the commonest mediastinal tumour but a rare type of thymic tumour of unknown aetiology. The overall incidence of thymoma is 0.13 per 100,000 persons per year which is even rarer in pregnancy. Until this date, the specific link between thymoma and pregnancy has not been identified. Presentations are also variable and include chest pain, respiratory insufficiency, superior vena cava syndrome or non-specific symptoms like fever and weight loss. An Autoimmune paraneoplastic disease which is often linked with thymoma includes myasthenia gravis (MG), hypogammaglobinaemia, aplastic anaemia and systemic lupus erythematosus (SLE). Here we describe a case of thymoma in pregnancy who had successfully delivered and underwent surgical removal postpartum.

Thymoma and thymic carcinoma, also known as thymic year. However, thymectomy was only scheduled in 2019 as epithelial tumours (TETs) are two types of rare cancers of she declined for surgery after the chemotherapy. thymus that can form in the cells that cover the outside Throughout 2015 till 2019, she had multiple admissions to surface of the thymus. The overall incidence of thymoma intensive care unit (ICU) for MG crisis. Despite being is 0.13 per 100,000 person years. Thymoma incidence is advised to undergo thymectomy, she persistently refused noted to be higher among Asians and Pacific Islanders surgery due to worry of surgical risks. During a followcompared to Whites and Hispanics for unknown reasons.¹ up in 2019 prior to her scheduled surgery, she was found Thymoma patients may present with chest pain, to be pregnant and thus referred to us. She denied respiratory insufficiency, superior vena cava syndrome or difficulty in breathing or swallowing or muscle weakness. even with non-specific symptoms like fever and weight Examination revealed a normal built lady, with bilateral eye loss. Autoimmune paraneoplastic disease is often linked ptosis. There was a palpable neck mass. The latest with thymoma which includes myasthenia gravis (MG), computerized tomography (CT) scan prior to her hypogammaglobinaemia, aplastic anaemia and systemic pregnancy showed a mediastinal mass measuring 3x9x8.6 lupus erythematosus (SLE).² Despite being the most cm, with suspicious metastatic lesion to the liver. common mediastinal tumour, the presentation during pregnancy is rare.3 Here we describe a case of thymoma in A multidisciplinary team (MDT) meeting involving the pregnancy who had successfully delivered and underwent cardiothoracic surgical removal postpartum.

CASE REPORT

A 37-year-old lady, Gravida 5 Para 4 was referred to us (specify clinic and institution) at 12 weeks period of gestation. The patient was diagnosed with a thymic cancer in 2015 and had completed chemotherapy in the same thorax performed at 30 weeks showed markedly enlarged

surgeon, neuromedical team and obstetrician was arranged. Decision was made for the patient to continue with her pregnancy with possibility of early delivery should her condition worsened. Nuchal translucency scan at 12 weeks and detail scan at 20 weeks were both normal. Throughout the pregnancy, she remained well without any MG crisis. Fetal growth was appropriate to age. Magnetic resonance imaging (MRI)

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thymoma. Hence, she was planned for delivery at 32 symptoms and the drop in postpartum period making the weeks. Following the delivery, she was to be scheduled for thymectomy. However, the couple refused caesarean section at 32 weeks. She was managed as an inpatient to monitor for signs and symptoms of MG crisis. An elective caesarean section was performed at 35 weeks under combined spinal epidural anaesthesia . She was given intravenous immunoglobulin (IVIG) 5 days before the delivery. The caesarean section was uneventful with blood loss of 200 mls. She delivered a female baby weighing 2.15 kg. Debulking thymectomy and lung resection were performed on Day 10 postpartum. Intraoperatively, there was a thymic mass which had invaded into the right and left upper lobe of the lung, and pericardium invading into the adventitia layer just above the aorta. The procedure went well and she was discharged 4 days later.

DISCUSSION

Thymoma is a rare type of thymic tumour of unknown actiology. It is even rarer in pregnancy. The earliest case reported was in 1959 and since then there are about 20 other cases reported. Until now, a specific link between thymoma and pregnancy has not been identified.4,5 About 50-70% of thymoma is associated with paraneoplastic syndrome with MG being the commonest of 30-50%.6 The course of MG in pregnancy and its outcome is unpredictable. It is shown that 40% of the patients experienced exacerbation of symptoms, 31% of patients remained stable throughout pregnancy while 29% have disease improvement.7 Disease exacerbation usually occurs in the first trimester and in the puerperium while improvement of disease happens in second and third trimester as mentioned in our case.

Symptoms worsening is resulted from imbalance in the immune system from the sex hormone exposure as there is evidence that oestrogen enhances cytokine and immunoglobulin production in MG patients. Another factor that contributes to changes of disease course in pregnancy is alpha-fetoprotein which is very effective in inhibiting the binding of AChR antibodies to acetylcholine receptors. The alpha-fetoprotein levels are high in late pregnancy resulting in improvement of MG

disease worsen.8

The standard management of patients with thymoma and MG is thymectomy. In women who had not undergone thymectomy, it is found that they have higher incidence of exacerbation during pregnancy as compared to those who had undergone the surgery. It is also seen that infants of thymectomised mothers had less risk of developing neonatal MG.9 Therefore, women with thymoma and MG should undergo thymectomy during pre-pregnancy. However, in our case, the patient was found to be pregnant before the operation was scheduled. Therefore, the surgery was postponed after delivery as thymectomy is a major surgical procedure with adverse implications if performed during pregnancy.10

In MG, vaginal delivery is the preferred mode as the uterus which does not consist of striated muscle is not affected by AChR antibodies. However, in second stage of labour, where striated muscle is involved, the delivery may require assistance either with forceps or vacuum.7 Caesarean section should only be reserved for obstetrics indication as the stress of surgery can cause worsening of MG symptoms.¹¹ Unfortunately for our patient, the size of her thymoma was noted to be increasing, thus a premature caesarean section was scheduled. As general anaesthesia and narcotics should be avoided due to potential synergistic effect on AChR antibodies, the caesarean section for our patient was done under combined spinal epidural as per recommendation.12

During postpartum period, patients are encouraged to breastfeed as long as they are not on contraindicated medications and the disease is well controlled. They should be informed that the course of previous pregnancies cannot predict outcome of the pregnancies.¹¹ Continuous subsequent hormonal contraception without hormone free period, barrier or intrauterine device are good contraception options for women with MG.12 In conclusion, management of thymoma in pregnancy is challenging. However, with good discussion and management involving team, it can result in better patient multidisplinary prognosis and good outcome.

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DISCLOSURE STATEMENT

All of the authors have no conflicts of interests to declare.

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