# Incidence of gestational trophoblastic disease in Tokat province, Turkey

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## Abstract

Objective: This study investigated the incidence of gestational trophoblastic disease (GTD) in Tokat province, Turkey.

Material and Methods: The medical records of patients who had been diagnosed and treated at one university hospital, six government hospitals, and one specialist hospital in Tokat province between January 2005 and December 2012 were evaluated retrospectively.

**Results:** During the study period, there were 59,754 births and GTD was diagnosed in 73 cases. The calculated GTD incidence was 1.22/1000. The mean age of the patients diagnosed with GTD was 28.6±7.3 (range 17-51) years. In GTD, complete moles occurred in 26%, partial moles in 74%, and no invasive moles, choriocarcinomas, or placental site trophoblastic tumours were found. Only two patients received chemotherapy (methotrexate). There was no mortality associated with the disease during follow-up.

**Conclusion:** The incidence of GTD in Tokat province was 1.2 per 1000 births. Early diagnosis, treatment, and follow-up play a critical role in preventing the morbidity and mortality associated with disease. The incidence of GTD, which has a high recovery rate with adequate treatment and follow-up, can be determined from regional and community-based research. (J Turk Ger Gynecol Assoc 2014; 15: 22-4)

Key words: Gestational trophoblastic disease, molar pregnancy, incidence Received: 11 October, 2013 Accepted: 09 November, 2013

## Introduction

Gestational trophoblastic disease (GTD) is classified histopathologically as complete or partial mole, invasive mole, placental site trophoblastic tumour, and gestational choriocarcinoma (1). Menarche age, parity, first pregnancy age, mole pregnancy history, time interval between previous pregnancies, genetic factors, malnutrition, viral infections, and socioeconomic level all predispose to GTD (2). The aetiology is not clear, but the disease is characterised by abnormal gametogenesis, fertilisation, and malignant transformation of trophoblastic tissue.

The incidence of GTD differs across geographic regions. The reported GTD incidence was 0.3-16 per 1000 pregnancies in Turkey, 0.6-1.2 in Europe and North America, 0.2-4.6 in Latin America, and 3.2-5.8 in Middle East countries (3, 4). This study determined the incidence of GTD in Tokat province, Turkey.

## Material and Methods

This study was performed between January 2005 and December 2012 and patients were collected from one university hospital, six government hospitals, and one private hospital in Tokat. The study protocol was approved by the Medical Ethics Committee and informed consent was taken. The patients diagnosed with GTD were confirmed histopathologically. Epidemiological data, laboratory and pathology results, and follow-up examination findings were obtained from the patients' records. Some information was obtained by telephoning patients when it was missing from the records. GTD was classified as complete and partial mole, invasive mole, placental site trophoblastic tumour, and gestational choriocarcinoma. Patients diagnosed with GTD were evaluated according to their age, pregnancy, birth, abortion number, and histopathology. All data were analysed using "PASW Statistics version 18.0" (PASW, Chicago, IL, USA). Data were given as mean±SD (standard deviation) and n (%).

#### Results

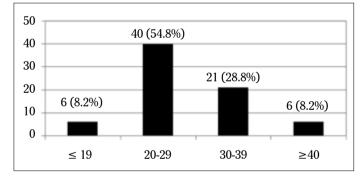
Between January 2005 and December 2012, there were 59,754 births in Tokat province. During this period, 73 cases were diagnosed as GTD histopathologically. The calculated GTD incidence was 1.22 per 1000 births. The mean age of the diagnosed patients was  $28.6 \pm 7.3$  (range 17-51) years. Many patients were aged 20-29 years (Figure 1). The mean numbers of pregnancies, births, and abortions were  $3.5 \pm 2.9$  (range 1-15),  $1.3 \pm 1.4$  (range 0-5), and  $1.1 \pm 2.3$  (range 0-13),

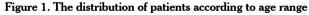


respectively. There was no history of molar pregnancy in any case. Comparing blood types, 45.9% were type A, 21.3% were type B, 31.1% were type O, and 1.6% were type AB. Vacuum curettage was performed in all patients for diagnosis and treatment. Histopathologically, 26% were complete moles, 74% were partial moles, and there were no invasive moles, choriocarcinomas, or placental site trophoblastic tumours (Figure 2). Two patients were treated for non-metastatic disease with single-agent chemotherapy (methotrexate) (Emthexate; Med Ilac, İstanbul, Turkey) and were cured. No hysterectomy was performed in any patient and there was no disease-dependent mortality during the follow-up period.

#### Discussion

Gestational trophoblastic disease, which arises from the abnormal proliferation of trophoblasts, is seen in 0.6-11.5 per 1000 pregnancies; however, this ratio differs among communities (2, 5, 6). The suggested GTD incidence is 0.3-16 per 1000 gestations (4). The incidence of GTD also differs among regions in Turkey. The incidence is 12.1 per 1000 births in Şanlıurfa, 8.1 in Van, 1.78 in Ankara, 3.35 in Konya and 5.9 in Istanbul (7-11). The reason for the increased GTD incidence in these cities might be the fact that these studies were performed in referral centres (10). Özalp et al. (12) conducted a society-based study, and found an incidence of 0.8 per 1000 live births and 0.6 per 1000 preg-





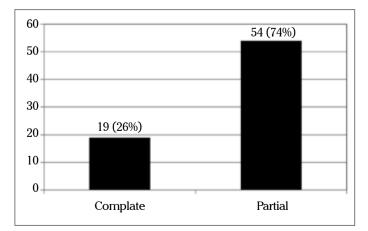


Figure 2. The distribution of patients according to histopathological diagnosis

nancies. Our study is the first conducted in Tokat and the GTD incidence was 1.22 per 1000 live births. It is apparent that the GTD incidence is lower when studies are based on large communities.

Çetin et al. (13) found that the incidence of GTD was 6.60 per 1000 births in Sivas province, so the incidence differs among regions. This rate is approximately six-fold higher than in our study. In comparison, Malatyalıoğlu et al. (14) reported a GTD incidence of 0.3 per 1000 pregnancies in Samsun. Unlike other studies, however, they calculated the incidence using the number of pregnancies, not births. Given the differences among regions, the incidence can differ between two cities, possibly because the calculation can be based on the number of pregnancies or births.

In our study, GTD was most common in the third decade of life and the mean patient age was  $28.6\pm7.3$  (range 17-51) years. In the literature, the disease is usually seen during the early and late fertility periods, *i.e.* younger than 20 and older than 35 years (15, 16). GTD in Turkey usually occurs at ages from 19-35 years (11, 17). We found that blood type A (45.9%) was the most frequent, and this is the most common maternal blood type A in the literature (8, 11).

Vacuum curettage, hysterectomy, and chemotherapy are usually used to treat GTD. In our study, vacuum curettage was performed in all patients, while no hysterectomies were done. In the literature, hysterectomy is usually used to treat old grand multiparity patients because malignant degeneration increases with age and the risk decreases from 20% to 3-10% with hysterectomy (18, 19).

Chemotherapy is given if the  $\beta$ -hCG level plateaus or increases or stays high for more than 6 months, choriocarcinoma is diagnosed histologically, or metastatic disease is present (20). Chemotherapy is based on the anatomical staging of the International Federation of Gynecology and Obstetrics (FIGO) and a prognostic scoring system devised by the World Health Organisation and modified by FIGO. Methotrexate or actinomycin is usually used as a single agent, while methotrexate, actinomycin-D, and cyclophosphamide (MAC) or etoposide, methotrexate, actinomycin-D, cyclophosphamide, and vincristine (EMA-CO) combinations are used as multiple treatment modalities (21). Cures are obtained in 100% of the low-risk group and 80-90% of the high-risk group (20, 22, 23). In our study, only two patients were treated with methotrexate because of persistent  $\beta$ -hCG. No chemotherapy-dependent morbidity occurred.

In conclusion, the incidence of GTD in Tokat province was 1.22 per 1000 births. Early diagnosis, treatment, and follow-up are important to prevent morbidity and mortality. We believe that sufficient treatment and follow-up will facilitate the cure of GTD and the incidence can be calculated more accurately by performing broad community-based studies.

*Ethics Committee Approval: Ethics committee approval was received for this study from the Medical Ethics Committee.* 

*Informed Consent:* Written informed consent was obtained from patient who participated in this study.

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