**INTRODUCTION**

Phyllode tumour is a breast neoplasm that usually occurs in patients aged 30–70 [1, 2]. It affects about 0.3–1% of women with diagnosed breast neoplasms [3, 4, 5, 6]. The most frequently noted size of phyllode tumour is 2–7 cm, yet there have often been cases of lesions reaching even 20–30 cm. In some patients the lesion fills the whole breast leading to the destruction of the gland and skin [7, 8].

Histologically, the tumour is formed from stroma of connective tissue and epithelial elements [7, 4]. In rare cases it can be a component of ductal cancer [8]. According to the WHO classification there are 3 types of phyllode tumours: low, medium and high malignant potential [9]. Low malignancy tumours are characterised by fast local growth without tendency to form distant metastases. However, even in such cases there is a significant risk of recurrence. Local recurrence affecting all types of phyllode tumour is estimated at 8–36% [4, 9]. For medium malignant potential lesions the risk of recurrence is 25% and 5% for distant metastases [10]. In the case of high malignant potential tumours the risk of recurrence and distant metastases is 25% in each case [10]. It is estimated that local recurrence does not worsen the prognosis [10].

On ultrasound examination, phyllode tumour is a polycyclic lesion of mixed echogenicity in which fluid spaces are present. Its structure bears resemblance to the cross-section of a cabbage. On mammography examination, the neoplasm is usually observed as shadowing with blurred contours and polycyclic structures. Most frequently it is not a spiculic lesion. Microscopic verification of the neoplasm is conducted with the use of fine-needle aspiration biopsy (BACC), oligobiopsy, mammotome biopsy or vacuum-assisted biopsy (VAB). However, these methods have their diagnostic limitations due to the fact that tumours are often histologically non-homogenous [11]. Phyllode tumour can be composed of adenofibroma or ductal cancer. Sometimes, various parts of the tumour have different grades of malignant potential [2]. In such cases there is no possibility to obtain diagnostic material with any of the above-mentioned methods, microscopic verification is performed with the use of so-called excisional surgical biopsy [12].

The treatment of choice in the case of phyllode tumour is excision of the tumour together with a 1–2 cm margin of healthy tissue. However, standardised treatment in the case of tumour of high malignant potential or lesions affecting the whole breast is mastectomy. In patients with axillary lymph nodes metastases, modified radical mastectomy is recommended [13, 14]. Fortunately, in the majority of patients, the course of the disease and prognosis are good [13].

**CASE DESCRIPTION**

A female Caucasian patient aged 13 was admitted to the Department of Children’s Surgery in order to diagnose a significant swelling and enlargement of her right breast. She was a schoolgirl in secondary school and came from a middle income family living in a small Polish village. On the basis of the obtained medical history it was known that the patient herself had found an insignificant thickening in her right breast five months prior to hospitalisation. Within the next two months she had observed a significant growth of the tumour which resulted in striae of the skin. This persuaded the girl’s mother to bring her daughter to the family doctor who qualified the tumour present in the breast as being changes typical for the period of puberty. The patient was not directed to any additional examinations or to consultation with a specialist.

Three months later, the right breast was even more enlarged, inappropriately in comparison to her left breast, and the disproportion and asymmetry between breasts was extremely significant. The girl’s right breast was more sensitive to touch, significantly enlarged, and of denser consistency in comparison to her left breast. On her right breast there were increasingly more visible striae of the skin.
Two months later, the patient, after having had her next appointment with a family doctor, was directed to a surgical outpatient clinic. On the basis of physical examination she was instructed to have antibiotic therapy and observation. She then again had a medical consultation in the surgical outpatient clinic. The girl’s constantly enlarging breast encouraged the doctors to hospitalise her and the patient was directed to the Department of Children’s Surgery.

In the department, she had an ultrasound examination of the breast (USG) and fine-needle aspiration biopsy (BACC) supervised by USG. Cytologic examination found numerous instances of protein mass, the presence of ductal epithelial cells, cells characterised by pleomorphism of the nucleus and presence of tiny intranucleus vacuoles. Microscopic diagnosis indicated mild breast cell proliferation, so-called juvenile cell proliferation. The pathologist suggested further diagnostics and treatment in an endocrinological outpatient clinic. However, the endocrinologist had no suggestions for treatment.

After the patient’s stay at the Department of Children’s Surgery, she was admitted to an oncologic surgery outpatient clinic, where on the basis of clinical and ultrasound examination of the breast she was diagnosed with phyllodes tumour. Repeated fine-needle aspiration biopsy performed in the outpatient clinic confirmed the clinical diagnosis and was the basis for directing the patient to the Breast Surgery Department.

A chest radiograph and abdominal ultrasonography showed no pathological changes. The results of various laboratory tests revealed no abnormalities. Ultrasound examination of the breasts found a well-separated pathological non-homogenous mass, low-echogenic, with scarce signs of vascular flow within the right breast. The lesion affected the breast between 3 – 9 o’clock, occupying about half of the breast. The size of the lesion was estimated at 120 × 70 × 35 mm. No pathological changes were seen in the left breast.

The patient was qualified for surgical treatment – quadrantectomy of the right breast.

Under general anaesthesia, the tumour was reached through an incision in the submammary fold. The lesion was well-separated, filling half of the breast (Fig. 1). The tumour was removed completely with a 2-cm margin of healthy tissues. A reconstructive operation of the nipple and the gland was performed.

There were no complications in the postoperative course. The patient was released home in good condition six days after the operation. On the basis of pathological examination, a final diagnosis was established – phyllodes tumour with low malignant potential. The patient remains under constant oncologic observation. There has been no recurrence of the disease to-date. Cosmetic results after the procedure are very good – the shape of the breast is natural and the scar is barely visible (Fig. 2, Fig. 3).

DISCUSSION

The diagnostics of breast diseases in children and teenagers is often conducted inappropriately. The reason for that can be little clinical experience of general practitioners (GPs), pathologists, paediatricians and gynaecologists in that field, as well as the rare incidence of this type of lesion in children. An additional factor making the establishment of diagnosis more difficult is the patient’s place of residence. Small towns and villages have no specialist diagnostic facilities to treat breast diseases [15]. Centres for the early detection of cancer and outpatient clinics of breast diseases
are located in bigger cities. There is no plan for training GPs in this matter. No organisational plan has been devised which would list guidelines where patients suffering from breast diseases should be directed. Neither has it been stated whether it should lie within the competence of oncologists or paediatricians. Maslach et al. [15] showed, that people who live in the rural areas of Poland usually have worse access to health care services and therefore worse access to effective oncological treatment. The authors underlined that especially the situation of young women suffering from breast cancer is the most severe in the group living in rural areas. Moreover, such a group is also characterized by the lowest 5-year survival rate [15].

However, the problem is not insignificant. The incidence of breast cancer in Poland has been on the increase. The disease affects younger and younger women. Genetic investigations conducted on an increasing scale enabled determination of a family thread with mutations which are inherited and which predispose to the development of breast cancer. In such cases, even teenagers can suffer from breast cancer. However, such people have some preventive methods at their disposal. Furthermore, it is also known that some benign lesions diagnosed in teenage years can increase the risk of the incidence of breast cancer in the future. According to Garcia-Closas et al. [10], the detection of some benign lesions during adolescence is an additional risk factor for developing breast cancer in subsequent years [10].

Delayed diagnostics and treatment of phyllodestumour always pose a risk to the patient, whereas delayed introduction of appropriate treatment can lead to an increased cancerous-transformation risk of the lesion and the need to undergo breast amputation. Such a procedure is performed in teenage patients and it can permanently change their psyche and influence their entire lives. That is why it is extremely important to formulate a diagnostic algorithm concerning breast diseases in children [16, 17]. The problem is especially difficult in smaller towns. The diagnostics which took far too long and was mismatched in the clinical case described in the presented study of a girl living in a village with only 305 inhabitants confirms that it is vital to take the steps.

REFERENCES