Phyllodes tumors are fibroepithelial neoplasms that represents less than 1% of all breast tumors and are usually found in middle-aged women. The combination of this tumor along with hypoglycemia is very rare. The perioperative management of hypoglycemia in those cases is also quite difficult. Here, we are presenting a challenging case of phyllodes tumor of the breast associated with hypoglycemia, and perioperative management of hypoglycemia.

Keywords: Fibroepithelial neoplasm; Neuroendocrine tumor; Non-islet cell tumor-induced hypoglycaemia; Perioperative management

Introduction
Phyllodes tumor (from Greek: phullon leaf) was originally described by Chelius in 1827 [1] and first named by Johannes Muller in 1838. Phyllodes tumors are a fibroepithelial tumor composed of an epithelial and a cellular stromal component. They may be benign, borderline, or malignant depending on histologic features including stromal cellularity, infiltration at the tumor’s edge, and mitotic activity. All forms of phyllodes tumors are considered to have malignant potential. This is predominantly a tumor of adult women, with very few examples reported in adolescents (although there are case reports in men, especially if previously associated with gynaecomastia). Phyllodes tumors account for about 0.3%-0.5% of all female breast tumors [2]. Patients typically present with a firm, palpable mass. These tumors are very fast-growing, and can increase in size in just a few weeks. The median size of tumor is 4-7 cm, and it may cause fatigue, dyspnoea, and bone pain in metastatic disease, 20% tumours can grow larger than 10 cm. Tumours can also grow larger than 40 cm [3]. Occurrence is most common between the ages of 45 and 49 [4,5], prior to menopause. This is older than the typical age of patients with fibroadenoma, with which phyllodes tumors may be confused.

Non-islet cell tumor-induced hypoglycemia (NICTH), although more commonly associated with other mesenchymal tumors, is a rare paraneoplastic sequel of phyllodes tumor. Phyllodes tumors associated with non-islet cell tumor-induced hypoglycemia usually grows to greater than 10 cm. This tumor leading to non-islet cell tumor-induced hypoglycemia may develop over several years or even rapidly. The common treatment for phyllodes tumor is wide local excision [6]. Chemotherapy and radiation therapy are not effective for this tumor. The risk of developing local recurrence or metastases is related to the histologic grade. Despite wide excision, a very high percentage of surgeries yielded incomplete excision margins that required revision surgery [7]. Complete excision of the tumor with continued follow-up is key to management. Here we report a case of phyllodes tumor presenting with non-islet cell tumor-induced hypoglycemia posed for surgery, along with the management of hypoglycemia and perioperative management in intensive care.

Case Summary
A 38 year old non-diabetic, non-hypertensive female patient had an episode of drowsiness followed by brief unconsciousness on the morning of 9th august 2015. Patient was arousable after 10 minutes of unconsciousness and her relatives rushed her to a local nursing home. She was given intravenous fluids and she could move her limbs and her body normally. She had no slurring of speech, deviation of angle of mouth, loss of voluntary control of micturition and defecation, fever, sore throat, shortness of breath, palpitation or chest pain. While being shifted to intensive therapy unit (ITU) of Institute of Post Graduate Medical Education and Research & Seth Sukhlal Karnani Memorial hospital (IPGMER&SSKM) Kolkata, she became fully conscious and oriented. CT-Scan of brain was done immediately which was absolutely normal. Arterial blood gas analysis was unremarkable. Her capillary blood glucose (CBG) revealed a level of 57 mg/dl, but the patient seemed to have no symptoms of hypoglycemia. She was given 200 ml bottles of 25% dextrose over 15 minutes each. CBG after 1 hour revealed 113 mg/dl. Her routine blood tests were normal. She was put on normal diet and kept under strict CBG monitoring. Chest X-ray PA view was done next day which was normal. The patient continued to have repeated episodes of hypoglycemia over the next few day with CBG readings as low as 19 mg/dl, 27 mg/dl, 29 mg/dl especially during 3 am to 6 am in the mornings.

She was put under strict monitoring of heart rate, spo2, respiratory rate and blood pressure but these came out to be normal limit. Since she was having repeated hypoglycemic attacks in spite of adequate normal diet, she was put on infusion 25% dextrose 100 ml 4 hourly from 13th august, 2015. After that her blood glucose stabilised with levels >90 mg/dl. An endocrine referral was done and she was advised several tests like fasting cortisol, insulin antibody, insulin, serum c-peptide during hypoglycemic attacks, thyroid stimulating hormones(TSH), free thyroid hormones (T4, T3). Thyroid profile, fasting serum cortisol and insulin antibody level were within normal range. But Serum c-peptide value was 0.18 ng/ml (normal level 1.1-4.4 ng/ml) and serum insulin level was 0.68 miu/L (normal level
The patient had a history of morning sweating and drowsiness for the last 15 days before admission which got relieved with taking little food. On 2nd August she had a history of similar conditions followed by unconsciousness which was transient about 20 seconds and she was relieved by glucose drinks. The patient also had a past history of phyllodes tumor of right breast for the last 1 year under investigations & the tumor was increasing in size for last 6 months. All reports including bone scan, USG whole abdomen, CECT thorax, core biopsy and routine of last year were normal. A general surgery referral was done and she was advised CECT chest and abdomen which came out to be normal except for phyllodes tumor in CECT chest. The possibility of phyllodes tumor causing hypoglycemic attacks was thought of and she was planned for modified radical mastectomy (MRM) with stage 1 lymph node dissection on 19th August, 2015 (Figure 1).

**Figure 1:** Right breast mass impending to rupture.

For management of perioperative hypoglycemia, she was advised inj. octreotide 50 µg s/c 8 hourly upto 2 hours before OT, intravenous fluid 5% dextrose(500 ml) 8 hourly, 25% dextrose(100 ml) 6 hourly and sos, inj hydrocortisone 50 mg i.v. 8 hourly and glucagon challenge test was done which showed hypoglycemia was responsive to glucagon. Her operation was done with excision of a 4 kg mass from the right breast under general anesthesia with strict control of CBG. In operating room, premedication was given with inj. Glycopyrrolate (0.01 mg/kg), fentanyl (2 mcg/kg) and induction was done with inj. Thiopentone (5 mg/kg) and as muscle relaxant atracurium (loading dose-0.5 mg/kg) was used. Maintenance was done with propofol, nitrous oxide and isoflurane 1-2%. Fluid management intraoperatively was done with crystalloids and maintenance 10% dextrose given slowly 100 ml/hr to prevent hypoglycemia. Intraoperatively CBG monitoring was done half hourly. Intraoperative period was uneventful with stable hemodynamics and minor fluctuations in blood glucose level. She was extubated after the operation.

After operation her blood glucose started stabilising with most values>150 mg/dl. 25% dextrose infusion was gradually tapered off and ultimately stopped on postoperative third day. Now she could maintain her CBG with her normal diet only. Histopathological examination revealed large phyllodes tumor (23 cm × 19 cm × 14 cm) which was benign in origin & all nine level 1 axillary lymph nodes were reactive.

The routine laboratory investigations were unremarkable. Postoperative IGF1, IGF2 levels were measured and compared with preoperative value suggestive of non islets cell tumor induced hypoglycaemia (Table 1).

<table>
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<td>IGF2/IGF1</td>
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<td>11.03</td>
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**Table 1:** Comparison of intraoperative and postoperative Insulin antibody, IGF 1 & 2.

### Discussion

In rare cases large phyllodes tumors can present with hypoglycemia. This case highlights the diagnosis and management of non-islet cell tumour-induced hypoglycaemia caused by phyllodes tumor and perioperative anaesthetic management. Hypoglycaemia is a common medical problem in patients with diabetes mellitus. Hypoglycaemia is also associated with gastric bypass surgery [8]. Although rare, hypoglycaemia is occasionally encountered in non-diabetic, nongastric bypass patients where hypoglycaemia is usually a manifestation of pancreatic islet cell tumors producing insulin, primary or secondary adrenal insufficiency, pheochromocytoma, IGF1 secreting tumors, hypothyroidism, nonislet cell tumor induced hypoglycaemia (NICITH) or antibodies interfering with insulin receptors. In patients with recurrent hypoglycaemia and no history of diabetes or weight loss surgery, non-islet cell tumour-induced hypoglycaemia may be the underlying cause. Malignancies associated with insulin receptor antibodies, tumor necrosis factor (TNF) and tumors like pheochromocytoma, solitary fibrous tumors (independent of anatomical location), wilm’s tumors, metastatic hemangiopericytomas, mesotheliums, hepatocellular carcinomas, gastrointestinal stromal tumors (GIST), colorectal adenomas, osteosarcomas, rhabdomyosarcomas, leiomyosarcomas, paragangliomas, prostate cancers, breast cancers are known to cause non-islet cell tumor-induced hypoglycaemia [9,10].

Non-islet cell tumor-induced hypoglycaemia is a paraneoplastic process and tumors that cause hypoglycaemia are generally greater than 10 cm in diameter. Hypoglycaemia can occur in both fasting and non-fasting state by multiple mechanisms. Hypoglycaemia results from inhibited hepatic glycogenolysis, gluconeogenesis and diminished lipolysis in adipose tissue. Hypoglycaemia here indicates increased insulin-like activity in the body. Hypoglycaemia may also occur due to increased glucose utilization by the cells of large tumor. However the β-hydroxybutyrate level is usually low and the glucose response to glucagon stimulation is normal in these cases, showing that this is not a normal cause of hypoglycaemia responding to glucagon [11]. It is now proved that IGF is a key factor in the pathogenesis of non-islet cell tumor-induced hypoglycaemia [12-14].

The IGF-2 gene is situated on short arm of chromosome 11 (11p15) near the insulin (INS) gene. IGF-2 is normally a 7.5-kDa peptide, but in cases of NICITH, IGF-2 has high molecular weight (10- to 20-kDa) [15,16]. Mature IGF-2 is synthesized from a 180 amino acid preprohormone having a C-terminal extension of 89 amino acids known as the E-domain in the immature stage. During intracellular
processing the E-domain is cleaved, resulting in mature IGF-2, which if not cleaved from pro IGF-2 results in big IGF-2. Normally in plasma <10% of circulating IGF-2 is big IGF-2. Under physiologic conditions, IGF-2 is mainly produced by the liver. But several malignancies (like sarcomas) can produce large amounts of big IGF-2 [16] leading to a large amount of big IGF-2 at the level of target tissues. Mature IGF-2 has 47% sequence match with insulin, and also active biochemically, hence stimulates glucose metabolism pathways which leads to hypoglycaemia [15,17]. On the other hand, suppression of growth hormone occurs at pituitary gland by negative feedback mechanism which causes decrease in IGF-1 level. Therefore increased IGF-2/IGF-1 ratio points to the diagnosis of IGF2 producing NICTH.

Unlike IGF-1, IGF-2 regulation is independent of growth hormone influence [18]. IGF-1and IGF-2 both have hypoglycemic potential which is approximately 5% of insulin’s potency. Plasma concentrations of IGF-1and IGF-2 can be 1000 times greater than insulin in NICTH permitting them to cause hypoglycaemia [19]. More than 95% of IGF-2 in the circulation is bound to insulin like growth factor binding proteins (IGFBP) [20]. At the cellular level, IGF-2 binds with IGF-2 receptor which mediates functions like endocytosis, intracellular hormone transport and metabolism of circulating IGF-2 [21]. On the contrary, there are several reports of neoplasms having high levels of IGF-2 mRNA without exhibiting elevated hormone activity [22,23].

In this case, the pre-excision IGF-1 was 66.4 ng/ml which is low and IGF-2 level was 732.6 ng/ml which is towards higher level of normal range. But after surgery, both IGF-1 (174.6 ng/ml) and IGF-2 (390.2 ng/ml) level were within the normal range. The pre-operative IGF-2/ IGF-I ratio was 11.03 (normal<10) suggesting NICTH. The post excision IGF-2/IGF-1 ratio was 2.23 which further strengthened our diagnosis. This patient was suffering from recurrent hypoglycaemic attacks in spite of normal diet which was managed by continuous administration of dextrose containing fluids. Once the tumor is identified, its excision is the mainstay of treatment and complete resection leads to rapid resolution of hypoglycaemia. After removal of the tumor, the blood sugar level normalises and the IGF-2 level decreases, as seen in this case [6].

Endogenous hyperinsulinism and insulinomas come in close differential diagnosis with NICTH and needs to be identified. Endogenous hyperinsulinism can be caused by a primary beta-cell tumor, an antibody to insulin receptor or ectopic insulin secretion. This is diagnosed by measuring plasma insulin and c-peptide level during period of hypoglycemia. The findings in endogenous hyperinsulinism are plasma insulin>3 µu/ml, c-peptide>0.6 ng/ml with plasma glucose<55 mg/dl and symptoms of hypoglycemia. Insulinomas are insulin secreting pancreatic beta cell tumors. They are benign and a rare form of neuroendocrine tumor. They are a treatable cause of fatal hypoglycemia. The median age at presentation is in the third decade of life as part of multiple endocrinol neoplasia (MEN-1). CT or MRI detects approximately 70-80% of insulinomas, as well as metastasis in malignant insulinomas [24]. These were excluded easily in this patient by relevant laboratory investigations.

Conclusion

We reported a middle aged female patient with large right sided breast mass who underwent modified radical mastectomy with level 1 lymph node dissection. Histopathological examination suggested of a benign phyllodes tumor. The patient had symptomatic hypoglycemia secondary to her phyllodes tumor which suggested to non-islet cell tumor-induced hypoglycemia. This hypoglycemia was corrected after removal of the tumor.

References

