Obesity and Breast Cancer: Adipose Tissue, Adipocytokines, Chronic Inflammation, and Hypoxia

Obesity and Meme Kanseri: Yağ Dokusu, Adipositokinler, Kronik İnflamasyon ve Hipoksi

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Abstract
Obesity that has represented defense and warfare in the stone age and beauty, success, authority, and welfare in the middle ages, has now turned into a chronic and undesirable low-grade inflammatory disease that adversely affects the lifespan and quality of life. It results in serious complications such as Type 2 diabetes, cardiovascular disease, and cancer and is considered a pandemic all over the world. Chronic inflammation in obesity is attributed to an increase in the release of pro-inflammatory factors. This makes it a risk factor for many types of cancers, such as colon, gastric, breast, and prostate carcinomas, suggesting a cause and effect relationship between obesity and cancer. These pro-inflammatory factors act as signal transducers for tumor growth and progression. In recent years, animal studies have shown that adipocyte hypoxia can result in tumor growth in adipocytes and stromal vascular tissue. However, the mechanisms associated with obesity and cancer are still not completely understood. In the present study, we discussed the relationship between inflamed adipose tissue and cancer in obese patients, together with the contribution of interaction between hormones, cytokines, growth factors, and angiogenesis.

Keywords: Obesity; breast cancer; inflammation; cytokines; adipose tissue

Introduction
Obesity, an increasing pandemic in the world, is defined as an abnormal or excessive accumulation of fat to the extent that it impairs human health. According to the 2014 report of World Health Organization (WHO), more than 1.9 billion adults are overweight (39%) and over 600 million adults are obese (13%) worldwide (1). In the United States alone, this rate has been reported to be 69% and 35%, respectively (2). Obesity leads to increased health problems, thereby reducing life expectancy (3). According to a recent study, association of

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Taş devrinde savunma, mücade gücü, kuvveti, orta çağda güzellik, başarı, iktidar ve zenginliği simgeleyen obezite; günümüzde yaşam süresini ve yaşam kalitesini olumsuz etkileyen kronik ve istenmeyen düşük dereceli inflamasyon bir hastalığı dönüştürüyor. Tip 2 diabetes, kardiyovasküler hastalıklar ve kanser gibi ciddi kompleksiyonlara neden olmakta ve tüm dünyada bir pandemi olarak kabul edilmektedir. Obezitede kronik inflamasyon, proinflamatur faktörlerin salınımındaki artış ve obezite ile kanser arasında neden ve sonuç ilişkisini oluşturduğu düşünülmektedir. Bu proinflamatur faktörler, tümör büyümesi ve ilerlemesi için sinyal transduktörleri görevi görür. Bununla birlikte, obezite ve kanser ile ilişkili mekanizmalar hâlâ tam olarak anlaşılamamıştır. Bu çalışmada, obez hastalarda inflame adipoz doku ve kanser arasındaki ili̇ki̇nin; hormonlar, sitokinler, büyümeye faktörleri ve anjiyogenez arası etkileşimin katkısıyla birlikte tari̇ş tilahi̇ma amaçlanmıştır.

Anahtar kelimeler: Obezite; meme kanseri; inflamasyon; sitokinler, adipoz doku

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Body mass index (BMI) with death due to all causes has increased from 23.7 kg/m² in the cohort that was evaluated from 1976 to 1978 to 27.0 kg/m² in the cohort that was evaluated from 2003 to 2013 (4).

Obesity, which is the second leading cause of preventable deaths after cigarette smoking, is a multifactorial disease (5). Genetic factors have a penetrance of 25 to 40% in obesity, whereas environmental factors, such as low the level of physical activity due to sedentary lifestyle and consumption of foods rich in simple carbohydrates and oil, are also involved. Earlier studies report that obesity increases the susceptibility to cardiovascular disease, type 2 diabetes mellitus (DM), obstructive sleep apnea, osteoarthritis, and many types of cancers (3, 6-10). According to “The Turkish Diabetes Epidemiology Study (TURDEP-II),” obesity increased by 44% in Turkey in 12 years. Interestingly, the average age of the adult population has also increased by four years in the 12-year period since TURDEP-I. The average male and female height increased by 1 cm each, whereas the body weight increased by 8 kg in men and 6 kg in women, waist circumference by 7 cm in men and 6 cm in women, the hip circumference by 2 cm in men and 7 cm in women. According to the TURDEP-II study, obesity prevalence was 32% in Turkey (11, 12). These results indicate that obesity and its consequences are important social health problems all over the world and more research is required in this regard.

Obesity involves an excessive accumulation of fat in adipocytes and is accompanied by a low-grade chronic inflammation. Recent studies indicate that fat tissue functions as an endocrine organ (13) and has a heterogeneous structure, consisting of preadipocytes, fibroblasts, endothelial cells, nerve cells, and macrophages around mature adipocytes (7). Majority of cells found in fat tissue are brown, white, and beige adipocytes (14). White fat tissue acts as the primary endocrine organ with metabolically active secretions. For this reason, adipocytes behave differently with different physiologic reactions in lipid and glucose metabolism, and during blood pressure regulation, angiogenesis, and homeostasis. Accumulation of fat beyond the threshold limit leads to hypoxia and necrosis, which, in turn, initiate chronic subclinical inflammation and metabolic disorders, as well as changes in carcinogenesis, tumor progression, and formation of metastases (9).

Fat tissue is known as the largest endocrine organ of the body owing to the secretion of many cytokines and hormones. An increase in adipose tissue leads to an increase in the synthesis of leptin, also known as the starvation hormone. Chronic inflammation in enlarged adipose tissue augments the secretion of IL-6 and TNF. Since increased levels of these cytokines, along with leptin, are also reported in cancer patients, obesity is considered to be a cause of cancer development, growth, progression, and metastasis (15).

Obesity is characterized by fat tissue hypoxia associated with chronic type 1 inflammation, decrease in adiponectin, increase in leptin, death of adipocytes, mitochondrial dysfunction, and cancer development as a result of insulin resistance, increased proinflammatory cytokines (IL-1, IL-6, TNF-α, and PAI-1), and infiltration of macrophages. Hypoxia-induced factor-1α (HIF-1α), released during hypoxia, regulates the transcription of the genes associated with carcinogenesis and is associated with increased formation of metastases (10, 16-19). The relationship between inflammation and cancer was first described by Virchow more than 100 years ago, who observed that leukocytes were increased in the neoplastic tissue (20-22). Together with the relationship between chronic inflammation and cancer, these findings reveal many common mechanisms for the association between obesity and cancer.

**Relationship Between Breast Cancer and Obesity**

Although breast cancer has not yet acquired a pandemic aspect as obesity, it is the most common type of cancer in women worldwide, and the leading cause of death by cancer (23). The development of breast cancer in the premenopausal or postmenopausal period is important in terms of assessing the risk factors. Various factors, including genetic, hormonal, and environmental factors, influence the development of the malignant tumor. Weight gain and obesity pose a po-
Several studies exist that have investigated the relationship between obesity and breast cancer. However, the results of these studies are not consistent. Some researchers advocate that the risk of breast cancer increases in both premenopausal and postmenopausal periods if BMI is greater than 30 kg/m² (24-27). While some researchers indicate that obesity decreases breast cancer risk in the premenopausal period but increases the risk in the postmenopausal period (28-32). In "Nurses’ Health Study," which included 87,143 women in menopause period followed up for a long time, it was reported that women who gained 10 kg or more weight after menopause had a higher risk of breast cancer than those who maintained their weight (33). In studies reporting premenopausal obesity to be protective against breast cancer, pathogenesis has not yet been completely elucidated. However, breast cancer in the premenopausal period was positively associated with estrogen and androgen levels in the circulatory system (34). Estrogen, estrogen receptor (ER), leptin, and adiponectin play a pivotal role in this mechanism. It has been shown that estrogen, a potent mitogen for breast cells, is involved in the development of breast tumors. Estrogen is synthesized primarily in the ovary and placenta with the aromatase activity in the premenopausal period. Its aromatase activity in the fat tissue and epidermis is more effective after menopause that increases the conversion of androgens to estrogen. Therefore, fat tissue in the breast is an important source of local estrogen production (18). Breast is a modified apocrine gland that produces milk after delivery. After menopause, adipose tissue is the only source of estrogen production through the aromatization of androstenedione, which is a 19-carbon steroid. Increased aromatase activity and androstenedione production in obesity lead to a larger total estrogen pool in obese women (35). Thus, anti-estrogenic compounds and aromatase inhibitors are the methods used to treat breast cancer (36).

Obesity is also considered a risk factor in relapse of breast cancer and poor survival (37-39). A meta-analysis published in 2011 reported that the lifestyle changes are the foremost step in the prevention and treatment of obesity. These lifestyle changes can decrease breast cancer progression and increase overall survival (40).

Earlier studies have indicated that high levels of leptin in blood have a mitogenic effect, which might be attributed to leptin resistance associated with obesity (41, 42). Leptin, when binds to its receptor, can result in breast cancer cell proliferation and angiogenesis; it also stimulates the estrogen receptor pathway by increasing aromatization (43). Adiponectin, with antagonistic effects to leptin, has anti-proliferative, pro-apoptotic, and anti-estrogen effects and its low levels play a role in the progression of breast cancer (44, 45). Recently, different studies have reported the relationship between adiponectin and leptin and breast cancer (46-48).

In addition, hyperinsulinemia and hyperglycemia that develop with obesity affect breast cancer development and progression (49). Insulin directly stimulates cell proliferation in breast cancer cell lines and reduces sex hormone-binding globulin (SHBG) levels, thereby increasing free estradiol. Some researchers suggest that insulin-like growth factor-1 (IGF-1) levels, which rise in premenopausal women, are associated with the risk of developing breast cancer. Increased level of insulin is one of the reasons for breast cancer recurrence and death in breast cancer survivors. Therefore, targeting insulin as a therapeutic modality in breast cancer could be an option in the adjuvant treatment of breast cancer. It seems that insulin may trigger the activation of a cascade of proliferative and anti-apoptotic events in the cancer cells. Metformin, an oral anti-diabetic known for 50 years, may also have direct effects on cancer cells. Metformin causes Her-2 suppression through inhibition of mTOR in breast cancer cells. Yurekli et al. reported the use of metformin as an adjuvant therapy in breast cancer patients to target insulin reduction and alter Her-2 oncogene-based molecular pathogenetic steps in breast cancer (50).

A nutritious, energy balanced diet with macronutrients and physical activity are the main determinants of IGF-1 levels. The changes in IGF-1 axis due to nutritional and lifestyle changes can mediate the develop-
ment and progression of cancer (18, 51). Metabolic changes due to insulin resistance and cytokines produced from adipose tissue play a significant role in making the tumors more aggressive. Insulin acts through its receptors found in normal cells and cancer cells and stimulates cell growth by allowing glucose and amino acids to enter the cells. Insulin and IGF-1 stimulate estrogen synthesis and the tyrosine kinase signaling pathway. Binding of insulin to IGF-1 receptor leads to mitogenicity and transformation, reduction in the synthesis of IGF-binding proteins (IGFBP-1 and IGFBP-2), and increased free IGF-1 levels. The PI3K-Akt signaling pathway, the main pathway of insulin and IGF-1 effect, upon activation by high levels of insulin and IGF-1 transduces the signal to the nucleus and inhibits p27 function by phosphorylation, thereby inhibiting cell growth.

HER-2/neu (c-erbB-2) is an oncogene that encodes the tyrosine kinase receptor involved in the pathogenesis of breast cancer. Insulin, IGF-1, and estrogen stimulate proliferation of breast cancer cells via a signaling pathway activated by HER-2/neu. Increased IGF-1 receptor stimulation has been shown to play an important role in the development of resistance against monoclonal antibody called trastuzumab, developed against HER-2/neu (52). The anti-estrogenic drug, tamoxifen, is one of the strongest medications used in the treatment of breast cancer and has been shown to reduce IGF-1 levels. The main reason for its ineffectiveness is its insufficiency to block estrogen receptor due to high levels of insulin. This explains why patients with postmenopausal diabetic breast cancer are desensitized to its effects. Metformin that has shown to break insulin resistance in cell culture studies reduces the growth of breast cancer cells that are resistant to tamoxifen (53-55). It is also used in the treatment of type 2 diabetes and has been shown by multiple studies to prolong the survival in breast cancer by decreasing AMP kinase pathway and insulin levels. Again, metformin reduces the cardiotoxic effect of trastuzumab in HER-2-positive patients. This effect is seen not only in patients who have type 2 diabetes but also in patients without type 2 diabetes (55).

Breast cancer, as a result of direct effect of insulin or indirectly by increasing IGF-1 receptor levels, may develop resistance to treatment by increasing the aggressiveness of tumor cells. Cell proliferation and angiogenesis are stimulated by proinflammatory cytokines, such as IL-6, IL-1β, tumor necrosis factor-α (TNF-α), C reactive protein (CRP), and PAI-1, that are released during chronic low-grade inflammation in insulin resistance.

One of the problems encountered in the treatment of patients with breast cancer is weight gain during or after treatment. It affects the treatment negatively through the mechanisms described below. Insulin resistance and high levels of insulin or IGF-1 can be controlled by simple, easily applicable, and harmless approaches, such as exercise and nutritional regulation. Metabolic changes caused by reduced activity and high caloric diet are involved both in the pathogenesis of breast cancer and negative prognosis of breast cancer. Since the mechanisms involved in pathogenesis could be improved with exercise and medical nutrition treatments, it is important that patients diagnosed with breast cancer do not gain weight. Additionally, those who are overweight during the diagnosis must lose weight in a controlled manner. In this way, protection against breast cancer becomes possible and the chances of successful treatment could be increased.

**Obesity-Related Comorbid Conditions Affect Breast Cancer Development and Surveillance**

**Dyslipidemia**

Hyperlipidemia, which often accompanies type 2 diabetes and obesity, is an important metabolic problem. In addition, dyslipidemia and hypercholesterolemia are found to be associated with aggressive breast cancer growth. In particular, it affects the ER ligand, thus contributing to the growth of ER-positive tumors. Many studies have demonstrated the role of statins in the treatment of hyperlipidemia to prolong the survival in breast cancer patients. One of the most recent studies in this regard is the observational BIG 1-98 study (56), in which statin added to the adjuvant endocrine therapy, as
a part of the treatment of ER-positive breast cancer, reduced total cholesterol, thus decreasing the cholesterol metabolite 27-hydroxycholesterol (27HC). 27HC is a molecule that acts as an estrogen receptor ligand and its elevated levels in blood accelerate the tumor growth (57). In the BIG 1-98 study, 318 ER-positive patients in the post-menopausal period were evaluated after treatment with letrozole, 106 patients with tamoxifen, 189 patients initially with tamoxifen followed by letrozole, and 176 patients initially with letrozole followed by tamoxifen. The hazard ratio (HR) for only endocrine therapy before statin was added to the treatment was 0.82 for disease-free survival, 0.83 for disease relapse, and 0.81 for disease survival without distant. The addition of statin to treatment protocol improved the parameters; HR ratios were 0.79, 0.76, and 0.74, respectively (56). However, apart from this observational study, there is an absolute need for randomized, controlled studies to reach a final and foolproof conclusion.

**Nonsteroidal Anti-Inflammatory Drugs**

The recent surge in the incidence of osteoarthritis is attributed to mechanical load on joints in obese patients and increases the frequency of the use of nonsteroidal anti-inflammatory drugs (NSAIDs). A study has reported NSAIDs to have a protective effect against breast cancer and may even reduce mortality in patients with breast cancer (58). Potential mechanisms include the ability of NSAIDs to inhibit prostaglandin synthesis, while prostaglandins increase the estrogen level by increasing aromatase gene expression. In breast cancer, studies have reported an increased expression of COX-2 and prostaglandin levels, with COX-2 gene responsible for angiogenesis and apoptosis (59-62). The animal studies have shown elevated nitric oxide levels after aspirin use to have an inhibitory effect on tumor cell growth (63). A meta-analysis involving 6 cohorts and 8 case-control studies published in 2001 reported that NSAIDs could slightly reduce the risk of breast cancer (64). In another meta-analysis of 3,133 patients published in the same year, a 24% reduction in the risk was observed with NSAID use for more than two months, especially when used more than eight years (65).

However, a cohort study comprising 28,695 patients, published in 2008, where patients were observed for 7.5 years, 847 breast cancer cases were found to have a higher NSAID use; the relation was not causal but coincidental. There was no relationship between NSAID use frequency and dose and breast cancer. The decrease in breast cancer incidence with NSAID use, detected in previous studies, could not be demonstrated in this study (66).

The Vitamin and Lifestyle (VITAL) study consisting of 35,323 postmenopausal patients reported that the use of NSAIDs increases breast cancer risk, and the HR was 1.26 (67). To conclude, even if NSAIDs have been considered protective against breast cancer, the dose and duration of the use of these drugs remain to be resolved.

**Can Weight Loss Reduce the Incidence of Breast Cancer?**

The risk of cancer increases with the increase in body weight. However, the question remains whether a loss in body weight is associated with a reduction in cancer incidence. Many diseases can be prevented with a reduction in weight; however, whether this is also true for breast cancer is still not known. In a study of 67,000 postmenopausal women, no reduction in breast cancer incidence was observed after weight loss. In this study, a linear relationship was found between BMI and breast cancer. Patients with BMI 35 kg/m² and above, when compared to patients with BMI less than 25 kg/m², had a 60% increased risk of invasive breast cancer (HR: 1.58). These patients also had cancer detected in more advanced stages (tumor size HR: 2.12, lymph node positivity HR: 1.89, local or distant metastasis HR: 1.89), and had a higher level of mortality in the obese patients (HR: 2.25, BMI: 30-35 kg/m²; HR:1.37, BMI: 35 kg/m²; and above HR: 2.11). The same study also demonstrated that the incidence of ER and progesterone receptor (PR)-negative breast cancer was not increased in patients who developed breast cancer. The patients, who were followed up throughout the study and had BMI of 25 kg/m² at the beginning of the study, reported an increased breast cancer risk with a 5% weight gain (HR: 1.36). However, patients with an
initial high BMI did not report a significant reduction in breast cancer after weight loss. From these studies, the following can be deduced: obese patients should be advised to lose weight; even after weight loss, these patients should be followed up owing to the increased risk of developing malignancy. From the above study, it appeared that increased white fat tissue in the postmenopausal period increases estrogen receptors in the breast tissue, leading to an increase in the breast cancer incidence (68). On the other hand, Schauer et al. showed that in patients with severe obesity, bariatric surgery was associated with a lower risk of cancer, particularly obesity-associated cancers, such as postmenopausal breast cancer, endometrial cancer, and colon cancer (69). One of the key points in the pathogenesis of the relationship between obesity and breast cancer is an interaction between the stromal microenvironment and normal epithelial cells. The stromal microenvironment includes adipose stromal cells and adipose stem cells (ASC). Increased accumulation of ASCs in obesity leads to increased adipogenesis and angiogenesis, which, in turn, accelerates cancer growth. Studies have demonstrated that ASCs obtained from patients with BMI above 30 kg/m² enhanced tumorigenesis and led to a change in the genetic profile of breast cancer cells. Another study reported increased secretion of leptin by ASCs in patients with BMI over 30 kg/m² as compared to those with BMI under 25 kg/m² (70).

Leptin is the most important weight controlling adipokine in white fat tissue and thus has a key role in the relationship between obesity and cancer. It is reported to affect survival, migration, and growth of cancer cells by increasing the expression of anti-apoptotic proteins; estrogen; pro-inflammatory cytokines, such as TNF-α and IL-6; and angiogenic factors, such as VEGF and hypoxia-induced factors (HIF-1α) (18, 19, 71-74). In a cell culture study that investigated the effect of leptin on breast cancer, leptin (obASC) from obese patients led to increased proliferation of ER-positive breast cancer cells, whereas no such effect was observed in case of leptin (InASC) from patients with BMI less than 25 kg/m². Increased epithelial-mesenchymal transfor-

mation and enhanced expression of metastatic genes (SERPINE1, MMP-2, and IL-6) were detected in the breast cancer cell culture performed with obASC. An absence of leptin decreased the tumor size and number of metastatic lesions (70). Previous studies have shown that leptin enhanced the tumor survival by altering various signaling pathways in tumor cells, including MAP-kinase, JAK2-STAT3, and PI3K-AKT pathways (75). The relationship between leptin and breast cancer is also related to menopause with an inverse relationship between increased leptin levels and cancer development in premenopausal patients. However, the level of leptin in postmenopausal patients increases the risk of breast cancer (76).

In a meta-analysis performed by Niu et al., patients with different breast diseases were evaluated. In 23 studies, 2,058 patients with breast cancer, 2,078 healthy controls, and 285 controls with light breast problem were included. Leptin levels in circulation were the lowest in healthy controls. The ranking for all groups was as follows: healthy control, control with light breast disease, patients with breast cancer, and patients with lymph node metastasis. Serum leptin levels in postmenopausal ER-positive breast cancer patients were found to be the lowest in clinical and pathologic classifications (72, 73, 76). Increased ectopic lipoaccumulation in obesity leads to tissue hypoxia, inflammation, and reactive angiogenesis, leading to changes in tumorigenesis and tumor characteristics (metastasis, invasion) (77). Adiponectin is one of the major adipokines secreted by the adipose tissue, with a potent anti-inflammatory activity. Adiponectin increases insulin sensitivity, allows regulation of many growth factors at the receptor level, and inhibits angiogenesis. Its decreased levels in obese people lead to increased activity of growth factors at the receptor level, thereby causing tumorigenesis (78, 79). The effect of adiponectin on breast cancer is closely related to the condition of menopause with different results. One study found a close association between low adiponectin and premenopausal breast cancer (80), whereas another study, which included 4,249 patients, no such as-
sociation was reported (81). In post-menopausal patients, however, the relationship between low adiponectin and breast cancer incidence has been clearly demonstrated (79, 82). In postmenopausal patients, the relationship between obesity and breast cancer can be simply explained as a consequence of the relationship between many metabolic factors, endocrines, receptors, inflammation, stromal cells, and tumor microenvironment, which are triggered by increased adipose tissue. Furthermore, in vitro studies have clearly demonstrated the antitumor and apoptosis-inducing effects of adiponectin; these effects are independent of ER and PR positivity (83).

Conclusion
In conclusion, increased adipose tissue mass and adipose inflammation lead to increased risk of tumor growth and progression through the involvement of many complex and intertwined pathways. In this regard, healthy lifestyle and nutrition are the first steps in prevention.

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