

Primary Fallopian Tube Cancer: An Unusual Case of Inguinal Lymphadenopathy

Zeinab Nazari (MD)¹, Jila Torabizadeh (MD)², Taraneh Geran (MD)¹, Hamed Jafarpour (MD)^{3*}, Amir Shamshirian (BSc)⁴

¹ Assistant Professor, Department of Gynecology, Imam Khomeini Hospital, Mazandaran University of Medical Sciences, Sari, Iran

² Assistant Professor, Department of Pathology, Imam Khomeini Hospital, Mazandaran University of Medical Sciences, Sari, Iran

³ Medical Student, Student Research Committee, Mazandaran University of Medical Science, Sari, Iran

⁴ Graduate, Student Research Committee, Mazandaran University of Medical Sciences, Sari, Iran.

ARTICLE INFO

Article type:
Case report

Article History:
Received: 06-Dec-2018
Accepted: 21-Apr-2019

Key words:
Fallopian tube cancer
Lymphadenopathy
Lymphatic metastases

ABSTRACT

Background: Primary fallopian tube cancers (PFTCs) are rare gynecological malignancies by the prevalence of 0.3-1%. The PFTCs occur in individuals within the age range of 18-88 years, more specifically at the age range of 40-65 years with the mean age of 55 years. The PFTC usually is observed with the chronic inflammation of the fallopian tube, infertility, tuberculous salpingitis, and tubal endometriosis. This study aimed to discuss the diagnosis of PFTC and the follow-up procedures of such patients. Moreover, it investigated inguinal lymph node metastasis as an uncommon occurrence and also reported a review of the literature about PFTC.

Case report: The case in this study was a 57-year-old woman (G5P5) referring to the clinic of Imam Hospital of Sari, Iran, in May 2016 after right inguinal lymphadenopathy. Total bilateral salpingo-oophorectomy with omentectomy, abdominal-hysterectomy, exploratory-laparotomy, as well as pelvic and para-aortic lymphadenectomy were performed for the patient. The pathogenic report indicated metastatic adenocarcinoma of a small tumor at the end of the right fallopian tube.

Conclusion: Since it is difficult to conduct initial diagnosis after primary surgery and definitive diagnosis of the disease, the surgeons complete the previous surgical procedure by performing an additional surgery. However, this issue seems to increase mortality among patients. As a result, it is essential to conduct more comprehensive studies to find the effective methods of diagnosis and apply the best medical management protocols for a better treatment of the disease, and therefore reduce mortality.

► Please cite this paper as:

Nazari Z, Torabizadeh J, Geran T, Jafarpour H, Shamshirian A. Primary Fallopian Tube Cancer: An Unusual Case of Inguinal Lymphadenopathy. Journal of Midwifery and Reproductive Health. 2019; 7(3): 1851-1855. DOI: 10.22038/jmrh.2019.36724.1403

Introduction

Primary fallopian tube cancers (PFTCs) are rare gynecological malignancies with the prevalence of 0.3-1%. The PFTCs occur in individuals within the age range of 18-88 years, specifically 40-65 years, with the mean age of 55 years (1-3). The PFTC usually is observed with the chronic inflammation of the fallopian tube, infertility, tuberculous salpingitis, and tubal endometriosis (4, 5). Most of PFTC metastasis occur via hematogenous, lymphatic, and peritoneal routes, which is usually asymptomatic (6). About 0-10% of patients are diagnosed with this disease before the surgery and most of

the diagnoses occur during the surgery and postoperative pathology (2, 7, 8). This disease appears with symptoms, such as vaginal discharge and bleeding, the colicky abdominal pain of tubal peristaltic, and abdominal/pelvic mass. Patients are diagnosed with no specific clinical signs and symptoms, leading to extensive pelvic metastases in advanced stages (1, 2, 9). Regarding the treatment, abdominal hysterectomy, bilateral salpingo-oophorectomy with omentectomy, appendectomy, peritoneal washings are the early therapeutic procedures for this disease (10, 11). In previous studies, a

* *Corresponding author:* Hamed Jafarpour, Medical Student, Student Research Committee, Mazandaran University of Medical Science, Sari, Iran. Tel: +989371143879; Email: hamed.jafarpour7@gmail.com

few number of patients were reported with inguinal lymph node metastasis caused by fallopian tube cancer. This case study aimed to discuss the diagnosis and the follow-up procedure of patients diagnosed with PFTC and inguinal lymph node metastasis.

Case report

A 57-year-old G₅P₅ postmenopausal woman with a history of right inguinal lymphadenectomy referred to Gynecology and Obstetrics Clinic of Imam Hospital of Sari, Iran, in May 2016. Written informed consent was obtained from the patient for the publication of this case report as well as accompanying images. Metastatic adenocarcinoma was indicated in the pathologic report (Figure 1). Her endoscopy and colonoscopy, as well as pelvic and abdominal CT scans were normal. Mammography showed fibrocystic changes. No abnormalities were found in the gynecological evaluation and Pap smear test showed atrophy. The endometrial line was 10 mm in transvaginal sonography. Endometrial biopsy showed a few endometrial glands with tubal metaplasia. Moreover, endocervical curettage showed

squamous cell metaplasia. All laboratory data were normal and tumor markers, including carcinoembryonic antigen, cancer antigen 25, 19-9, 15-3, were negative. Histological evaluation was performed and the result showed the serous papillary carcinoma of the ovary as the origin of the tumor; therefore, an exploratory laparotomy was also performed for her.

In the performed surgery, abdominal and pelvic organs were normal, except the small tumor at the end of the right fallopian tube (Figure 2). The surgical procedures included total abdominal hysterectomy, bilateral salpingo-oophorectomy with omentectomy, followed by pelvic and para-aortic lymphadenectomy. The final pathologic report indicated senile atrophy in endometrium adenomyosis in the myometrium, a benign simple cyst was observed in the left ovary; however, nothing remarkable was detected in the left fallopian tube. There was 2 cm tumor in the right fallopian tube without perineural or intralymphatic invasion. On the other hand, right ovary was free of tumor. All lymph nodes, omentum, and peritoneal washing cytology were negative for malignancies.

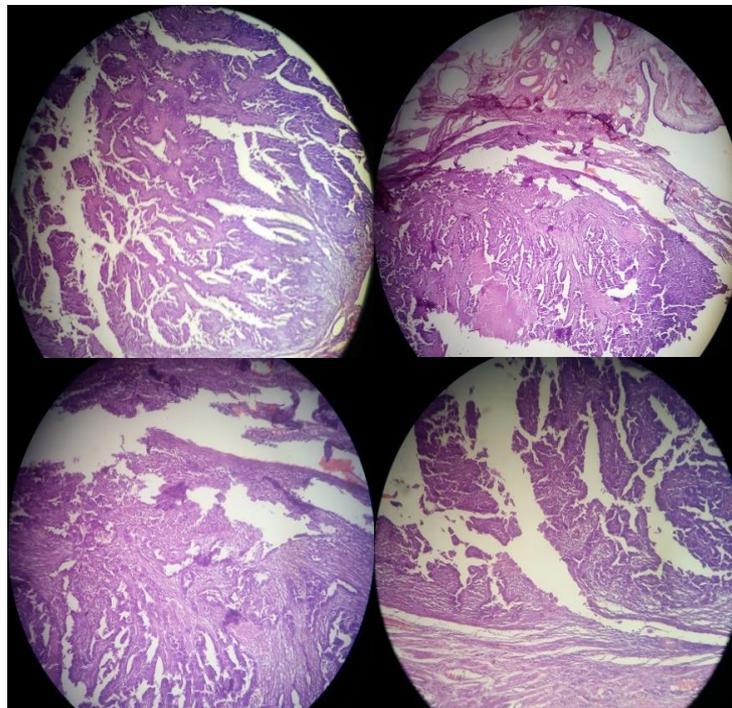


Figure 1. Metastatic adenocarcinoma of the right fallopian tube in the inguinal lymph node (hematoxylin and eosin, 100×)



Figure 2. Small tumor at the end of the right fallopian tube

Discussion

The primary carcinoma of the fallopian tube, known as rare aggressive gynecological malignancy, was first identified by Reynaud in 1847, which involves less than 1% of genital tract cancers. This disease mostly occurs among menopausal women (11-13). According to the previous studies, the incidence of this carcinoma can be much higher due to a large number of the misdiagnosis of PFTCs as ovarian cancers (12, 14-16). However, the etiology of this malignancy remains unclear. Although there are no specific symptoms for its diagnosis, some studies reported that this malignancy is mostly associated with some symptoms, such as abdominal pain due to tubal distention, pelvic inflammation, abnormal vaginal bleeding and discharge, followed by nulliparity, infertility, and Latzko's triad (1, 14, 17). Moreover, it is important to distinguish PFTC from epithelial ovarian cancer that are similar both clinically and histologically (18). Nevertheless, PFTC is not usually diagnosed before the surgical specimen due to its nonspecific symptoms (19).

The importance of lymphatic expansion was first introduced by Tamimi and Figge as the main cause of the spread of fallopian tube cancer (20). The lymphatic drainage of the fallopian tube may indicate its metastatic pattern and it follows the fundus of the uterus and ovaries. For example, the proximal part of the tube that is near the uterus drains into the para-aortic nodes, fimbriae drains to the pelvic nodes from the distal portion, and the round

ligament may drain to the inguinal lymph nodes through the lymphatic ducts (19, 21-23).

According to previous studies, it is clear that lymphadenopathy is a rare sign of primary fallopian tube cancer and only a few number of patients have been reported so far. Several studies reported that lymphatic node metastases frequently occurs in women who are the victim of emergency obstetric care and have indicated the involvement of abdominal and pelvic, supradiaphragmatic lymph nodes and inguinal lymph nodes (24-27). Furthermore, it is documented that the frequency of inguinal lymph nodes involvement is less than lymphadenopathy of other sites (e.g., pelvic and supradiaphragmatic) and it has been observed in only a few cases (28-30).

The most expressed antigen associated with PFTC is CA-125 that is not enough to detect the tumor progression; however, it can be a useful marker during the patient's follow-up (31). According to the obtained results of the current study, several possible tumor markers, such as CEA, CA125, CA19-9, and CA15-3, were negative.

According to previous studies, Winter-Roach et al. reported the case of inguinal lymph node metastasis as an unusual presentation of fallopian tube carcinoma in 2000. Total abdominal hysterectomy, bilateral salpingo-oophorectomy, omentectomy, pelvic, and para-aortic lymphadenectomy were performed for this patient and it became clear that in addition to the right fallopian tube cancer, implants were present on the ovarian serosa. Evidently, it has been the first case of PFTC that reported in this way (22).

Moreover, Euscher et al. presented a case of right inguinal lymph node caused by fallopian tube cancer similar to our case in a study named "Serous Carcinoma of the Ovary, Fallopian Tube, or Peritoneum Presenting as Lymphadenopathy" during a 20-year period (1982-2002) (6). Moreover, Jennifer Cho et al. reported a 72-year-old G3P3 postmenopausal woman diagnosed with PFTC with left inguinal lymph node metastasis in 2006 (19). Furthermore, Hviid et al. described a case with metastases to both ovaries and one inguinal lymph node in a study named "atypical debut of symptoms of fallopian tube cancer" in 2013. (32)

Conclusion

As mentioned, PFTC is an extremely rare problem that makes a challenge for surgeons and its causes are still unknown. Since it is difficult to conduct initial diagnosis after primary surgery and definitive diagnosis of the disease, the surgeons complete the previous surgical procedure by performing an additional surgery. However, this issue seems to increase mortality among patients. As a result, it is essential to conduct more comprehensive studies to find the effective methods of diagnosis and apply the best medical management protocols for a better treatment of the disease, and therefore reduce mortality.

Acknowledgments

The authors would like to express their thanks to the Imam Khomeini Hospital of Mazandaran University of Medical Sciences, Sari, Iran, and also the cooperation of staff working at this hospital.

Conflicts of Interest

Authors declare no conflicts of interest.

References

1. Ajithkumar TV, Minimole AL, John MM, Ashokkumar OS. Primary fallopian tube carcinoma. *Obstetrical & Gynecological Survey*. 2005; 60(4):247-252.
2. Kalampokas E, Kalampokas T, Tourountous I. Primary fallopian tube carcinoma. *European Journal of Obstetrics and Gynecology and Reproductive Biology*. 2013; 169(2):155-161.
3. Nazari Z, Torabizadeh Z, Rokhghireh S. Fallopian tube adenocarcinoma presenting with vaginal bleeding: a case report. *Journal of Mazandaran University of Medical Sciences*. 2012; 22(90):127-130.
4. Gadducci A, Landoni F, Sartori E, Maggino T, Zola P, Gabriele A, et al. Analysis of treatment failures and survival of patients with fallopian tube carcinoma: a cooperation task force (CTF) study. *Gynecologic Oncology*. 2001; 81(2):150-159.
5. Demopoulos RI, Aronov R, Mesia A. Clues to the pathogenesis of fallopian tube carcinoma: a morphological and immunohistochemical case control study. *International Journal of Gynecological Pathology*. 2001; 20(2):128-132.
6. Euscher ED, Silva EG, Deavers MT, Elishaev E, Gershenson DM, Malpica A. Serous carcinoma of the ovary, fallopian tube, or peritoneum presenting as lymphadenopathy. *The American Journal of Surgical Pathology*. 2004; 28(9):1217-1223.
7. Eddy GL, Copeland LJ, Gershenson DM, Atkinson EN, Wharton JT, Rutledge FN. Fallopian tube carcinoma. *Obstetrics and Gynecology*. 1984; 64(4):546-552.
8. Mei-Liu M, Chou BQ, Ziang J. Diagnosis of primary adenocarcinoma of the fallopian tube. *Journal of Cancer Research and Clinical Oncology*. 1985; 110(2):136-140.
9. Alvarado-Cabrero I, Navani SS, Young RH, Scully RE. Tumors of the fimbriated end of the fallopian tube: a clinicopathologic analysis of 20 cases, including nine carcinomas. *International Journal of Gynecological Pathology*. 1997; 16(3):189-196.
10. Takeshima N, Hasumi K. Treatment of fallopian tube cancer. *Archives of Gynecology and Obstetrics*. 2000; 264(1):13-19.
11. Riska A, Leminen A. Updating on primary fallopian tube carcinoma. *Acta Obstetrica et Gynecologica Scandinavica*. 2007; 86(12):1419-1426.
12. Hariprasad P, Hariprasad S, Srinivas T. Primary bilateral fallopian tube carcinoma the report of a single case with review of the literature. *Journal of Clinical and Diagnostic Research*. 2013; 7(5):930.
13. Lau HY, Chen YJ, Yen MS, Chen RF, Yeh SO, Twu NF. Primary fallopian tube carcinoma: a clinicopathologic analysis and literature review. *Journal of the Chinese Medical Association*. 2013; 76(10):583-587.
14. Veliscu A, Marinescu B, Costoiu L, Damian C, Stanescu C, Chiutu L, et al. Bilateral primary fallopian tube carcinoma: a case report. *Romanian Journal of Morphology and Embryology*. 2013; 54(4):1183-1187.
15. Kindelberger DW, Lee Y, Miron A, Hirsch MS, Feltmate C, Medeiros F, et al. Intraepithelial carcinoma of the fimbria and pelvic serous carcinoma: evidence for a causal relationship. *The American Journal of Surgical Pathology*. 2007; 31(2):161-169.
16. Callahan MJ, Crum CP, Medeiros F, Kindelberger DW, Elvin JA, Garber JE, et al. Primary fallopian tube malignancies in brca-positive women undergoing surgery for ovarian cancer risk reduction. *Journal of Clinical Oncology*. 2007; 25(25):3985-3990.
17. Chaudhry S, Hussain R, Zuberi MM, Zaidi Z. Rare primary fallopian tube carcinoma; a gynaecologist's dilemma. *The Journal of the Pakistan Medical Association*. 2016; 66(1):107-110.
18. Hu CY, Taymor ML, Hertig AT. Primary carcinoma of the Fallopian tube. *American Journal of Obstetrics and Gynecology*. 1950; 59(1):58-67.
19. Cho J, Grumbine FC, Díaz-Montes TP. Inguinal node metastasis as the initial presentation of

- primary fallopian tube cancer. *Gynecologic Oncology*. 2006; 103(2):759-762.
20. Tamimi HK, Figge DC. Adenocarcinoma of the uterine tube: potential for lymph node metastases. *American Journal of Obstetrics and Gynecology*. 1981; 141(2):132-137.
 21. Deffieux X, Morice P, Thoury A, Camatte S, Duvillard P, Castaigne D. Anatomy of pelvic and para-aortic nodal spread in patients with primary fallopian tube carcinoma. *Journal of the American College of Surgeons*. 2005; 200(1):45-48.
 22. Winter-Roach BA, Tjalma WA, Nordin AJ, Naik R, de Barros Lopes A, Monaghan JM. Inguinal lymph node metastasis: an unusual presentation of fallopian tube carcinoma. *Gynecologic Oncology*. 2001; 81(2):324-325.
 23. Plentl AA, Friedman EA. Lymphatic system of the female genitalia. The morphologic basis of oncologic diagnosis and therapy. *Major Problems in Obstetrics and Gynecology*. 1971; 2:223.
 24. Euscher ED, Silva EG, Deavers MT, Elishaev E, Gershenson DM, Malpica A. Serous carcinoma of the ovary, fallopian tube, or peritoneum presenting as lymphadenopathy. *The American Journal of Surgical Pathology*. 2004; 28(9):1217-1223.
 25. Miyaishi O, Iida KI, Saga S, Sato T. An autopsy case of serous papillary carcinoma of peritoneum with distant metastases but no peritoneal dissemination. *Gynecologic Oncology*. 1994; 55(3):448-452.
 26. Scholz HS, Lax S, Tamussino KF, Petru E. Inguinal lymph node metastasis as the only manifestation of lymphatic spread in ovarian cancer: a case report. *Gynecologic Oncology*. 1999; 75(3): 517-518.
 27. Wu PC, Qu JY, Lang JH, Huang RL, Tang MY, Lian LJ. Lymph node metastasis of ovarian cancer: a preliminary survey of 74 cases of lymphadenectomy. *American Journal of Obstetrics and Gynecology*. 1986; 155(5):1103-1108.
 28. Maaßen nV, Hiller K. Glandular inclusions in lymph nodes: pattern of distribution and metaplastic transformation. *Archives of Gynecology and Obstetrics*. 1994; 255(1):1-8.
 29. Prade M, Spatz A, Bentley R, Duvillard P, Bognel C, Robboy SJ. Borderline and malignant serous tumor arising in pelvic lymph nodes: evidence of origin in benign glandular inclusions. *International Journal of Gynecological Pathology*. 1995; 14(1):87-91.
 30. Shiraki M, Otis CN, Donovan JT, Powell JL. Ovarian serous borderline epithelial tumors with multiple retroperitoneal nodal involvement: metastasis or malignant transformation of epithelial glandular inclusions? *Gynecologic Oncology*. 1992; 46(2): 255-258.
 31. Niloff JM, Knapp RC, Schaetzl E, Reynolds C, Bast JR. CA125 antigen levels in obstetric and gynecologic patients. *Obstetrics & Gynecology*. 1984; 64(5):703-707.
 32. Hviid MM, Teklay B, Jensen PT. Atypical debut of symptoms of fallopian tube cancer. *Ugeskr Laeger*. 2013; 175(37):2114-2115.