Live birth after ovarian tissue autotransplantation following overnight transportation before cryopreservation

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Objective: To describe the first live birth after transplantation of ovarian tissue following overnight transportation of the tissue before freezing.

Design: Technical note.

Setting: University department of obstetrics and gynecology.

Patient(s): A 25-year-old cancer survivor with previous Hodgkin disease and relapse.

Intervention(s): The ovarian tissue was kept cool for >20 hours in a special transport medium and a special cooling device before it was cryopreserved. After premature ovarian failure due to preconditioning chemotherapy for bone marrow transplantation, the cryopreserved ovarian tissue was transplanted orthotopically.

Main Outcome Measure(s): Resumption of ovarian function after transplantation, recovery of fertility, and pregnancy.

Result(s): Ovarian function returned in the patient 3 months after transplantation, as shown by follicle development and estrogen production. During the fifth menstrual cycle, mild stimulation with FSH was initiated in accordance with a low-dose protocol. When ultrasonography revealed a follicle 18–20 mm in size in the ovarian graft, hCG was added and the patient had sexual intercourse at the optimal time point. On day 14 of the luteal phase, hCG concentration and vaginal echography confirmed a viable intrauterine pregnancy, which resulted in a healthy live birth.

Conclusion(s): Overnight transportation of ovarian tissue appears to be possible in combination with appropriate transportation logistics. However, further investigations are needed before this procedure can be offered as a chance for women to preserve fertility independently of direct access to a tissue-processing bank. (Fertil Steril® 2012;97:387–90. ©2012 by American Society for Reproductive Medicine.)

Key Words: Fertility preservation, ovarian tissue cryopreservation, ovarian tissue transplantation, orthotopic, pregnancy

s a result of improvements in oncologic treatment, most young cancer patients are now achieving prolonged survival. The 5-year survival rate for all cancers combined is currently >64% in women (1). However, loss of ovarian function and therefore fertility is one of the most common long-term adverse effects affecting premenopausal patients

treated with chemotherapy and/or radiation therapy. Conditioning therapy for bone marrow transplantation (BMT) is considered to be the most gonadotoxic form of treatment, leading to premature ovarian failure in >80% of cases even when performed during childhood (2).

For these young patients, ensuring their reproductive capacity after oncologic treatment has become a major concern, because it is directly related to their quality of life. A number of strategies have therefore been developed in recent years to enable these patients to have children using their own gametes. When radiotherapy alone is administered, it is possible to place the ovaries outside of the radiation field. When chemotherapy can be postponed, it is possible to use ovarian stimulation to obtain oocytes, which can be frozen in either a fertilized or an unfertilized state (3, 4).

Cryopreservation of ovarian tissue is another, very promising, alternative (5). Ovarian tissue can be extracted laparoscopically without significantly delaying gonadotoxic therapy. The tissue can be cryopreserved at centers specializing in reproductive medicine and can be transplanted into the pelvic cavity (or to a heterotopic site for oocyte

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retrieval and in vitro fertilization) if the women experience premature ovarian failure. Donnez et al. (6) reported the first live birth after autotransplantation of cryopreserved ovarian tissue in humans. Orthotopic reimplantation has so far led to the birth of 17 healthy babies (7–9).

Herein, we report a live birth after ovarian transplantation in a survivor of Hodgkin lymphoma. This is the first live birth after ovarian tissue transplantation in which the tissue was transported chilled on ice overnight before freezing.

CASE REPORT

A 25-year-old woman was diagnosed with nodular sclerosing Hodgkin lymphoma, clinical stage IIA. The patient received six courses of ABVD chemotherapy—adriamycin (doxorubicin), bleomycin, vincristine, and dacarbazine—and mediastinal radiotherapy to a dose of 30 Gy. Because the chemotherapy-associated amenorrhea rate with the ABVD regimen is very low, fertility preservation was not carried out up to this point (10). Remission was achieved, and the patient reported regular menstrual cycles of ~28 days.

However, the disease recurred after 2 years. Intensive chemotherapy consisting of two courses of Dexa-BEAM—dexamethasone, BCNU (carmustine), etoposide, Ara-C (cytarabine), and melphalan—was administered before autologous BMT was performed. Before the start of this treatment, about two-thirds of the ovarian cortex was removed laparoscopically from both ovaries. A histologic reference sample showed numerous primordial and primary follicles.

After BMT, the patient was considered to be disease free, but she became amenorrheic and developed vasomotor symptoms. Cyclical hormone replacement therapy was administered with $\rm E_2$ and dydrogesterone (Femoston) for a 5-year period. During this time, she had regular menstrual cycles. However, amenorrhea returned when the hormone replacement therapy was stopped, and her hormonal status confirmed premature ovarian failure (antimullerian hormone 0.1 $\rm ng/mL$).

Transportation and Cryopreservation Procedure

After laparoscopic removal at Dresden University Hospital, the ovarian tissue was rinsed for blood and cells with 0.9% saline solution in sterile conditions and placed in 30-mL plastic tubes containing a special medium for transporting ovarian tissue, precooled to 4°C (Brahma I; CryoBioSystem). The plastic tubes were placed in a special isolated transportation box (DeltaT) with cooling elements to ensure optimal conditions during transport and to preserve the viability of the tissue by maintaining a stable temperature of 5–8°C for 36 hours. The package was sent by an express overnight transportation service and reached the Reproductive Medicine Laboratory in Bonn within 20 hours.

Immediately after arrival in Bonn, the biopsies were dissected in medium for ovarian tissue manipulation (Brahma II; CryoBioSystem) (11). The biopsy specimens were then cut into pieces measuring $\sim\!1\times2\times1$ mm and equilibrated in a freezing solution containing 1.5 mol/L dimethylsulfoxide (DMSO) in Leibovitz medium. The pieces of ovarian tissue were then frozen in standard cryopreservation containers (1.8 mL

Nunc cryovials) using a slow-cooling protocol with a closed freezing system (Icecube 14S; Sylab) with autoseeding (12).

Thawing and Retransplantation

Five years after complete remission, the patient requested autotransplantation of the cryopreserved ovarian tissue. The tissue was sent in a shipping container cooled with liquid nitrogen at -196°C to the Reproductive Medicine Laboratory in Erlangen.

Thawing was fast in a water bath with warm water. The tissue fragments were released from the protective cryopreservation medium in reverse order with the addition of 0.25 mol/L sucrose. The thawed tissue was transplanted into a 1.5-cm deep pouch of peritoneum in the region of the broad ligament, below the right tube. Six fragments of ovarian tissue 1-2 mm in size were introduced into this pouch, and the pocket was closed with a Vicryl suture (Fig. 1).

At the same time, chromopertubation was performed, which showed bilateral tubal patency. Andrologic examination of the partner exhibited normal sperm parameters.

RESULTS

The preoperative FSH and LH levels were high and $\rm E_2$ levels low. Three months after transplantation, a fall in gonadotropins (LH to 3.9 U/L, FSH to 11.1 U/L) and a rise in $\rm E_2$ (to 114 pg/mL) was detected. Transvaginal ultrasonography showed an antral follicle in the ovarian graft that had been transplanted into the pelvic wall. The patient afterwards reported her first postoperative menstrual period.

As she urgently desired to have children during the following 4 months, cycle monitoring was performed, and when follicles in the transplanted graft reached 17–18 mm in size, ovulation was triggered with 5,000 U hCG (Predalon). The patient was encouraged to have regular sexual intercourse, but a pregnancy was not achieved. During the fifth menstrual cycle, therefore, mild stimulation with 25 U FSH

FIGURE 1



View of the transplantation site. The frozen/thawed tissue was transplanted into a deep pouch of peritoneum in the region of the broad ligament, below the right tube.

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