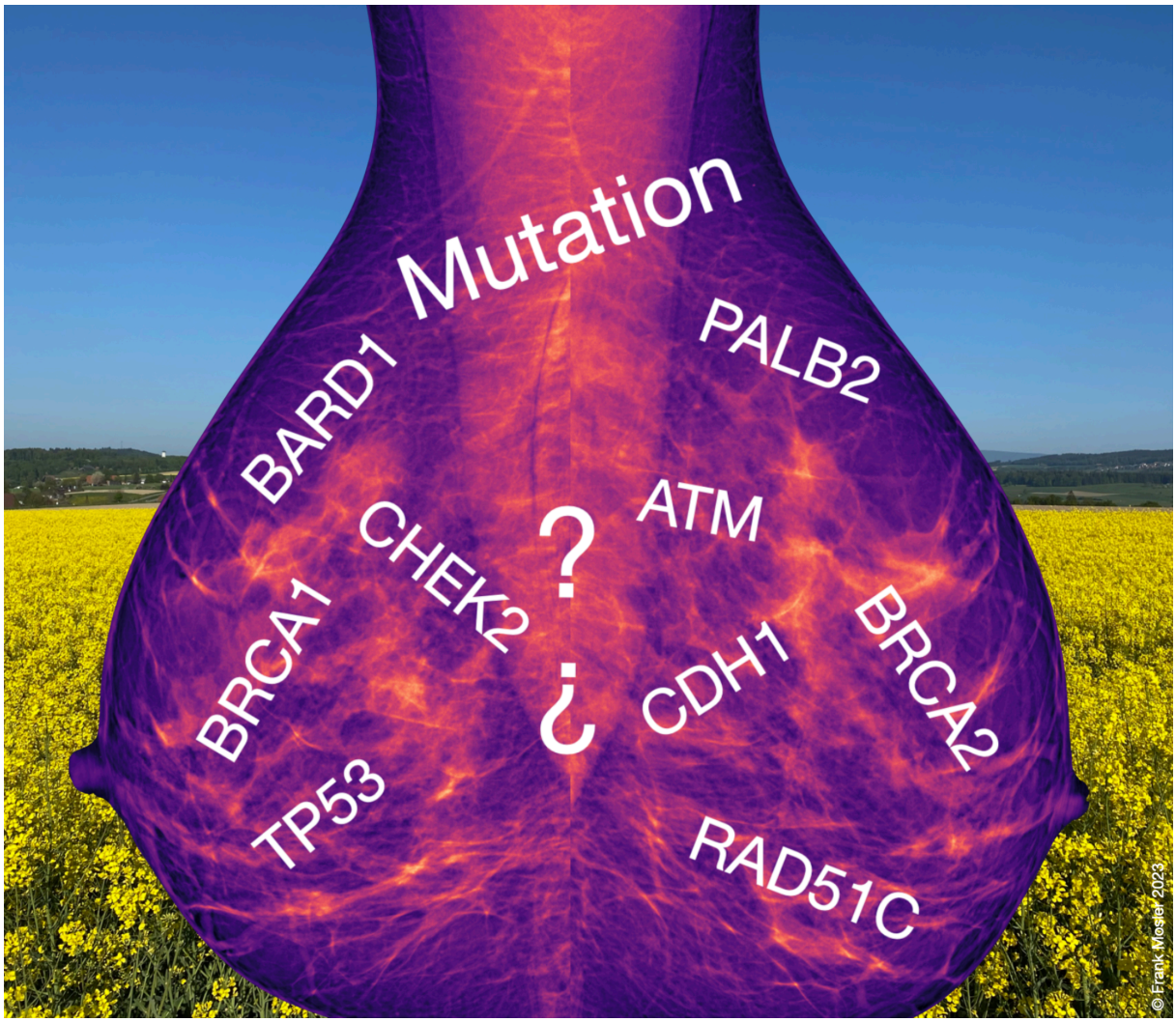
Original article - Swiss J. Rad. Nucl. Med. (2023) 2; DOI: 10.59667/sjoranm.v2i1

Early detection in cases of familial breast cancer predisposition: What is appropriate and beneficial for the individual seeking advice?



Florian Dammann[1], Susanne Ditz[2]

[1]Department of Diagnostic, Interventional and Pediatric Radiology, Inselspital Bern, University of Bern, Bern Switzerland
[2]Psycho-oncology and Psychosomatics at the University Women's Hospital, University Hospital, Heidelberg Germany

Swiss Journal of Radiology and Nuclear Medicine - SJORANM GmbH - CH-3072 Ostermundigen bei Bern - Switzerland

Asymptomatic women with a family history of breast cancer should first receive comprehensive and detailed counselling with a risk-benefit assessment of diagnostic or preventive measures. The medical counselling should enable individuals seeking advice to make a participatory decision.

[1]Corresponding author: f.dammann@gmx.com - received: 15.07.2023, Published: 25.08.2023

Abstract

A "high-risk situation" is present when the lifetime risk of developing breast cancer is ≥ 30%. Currently, the most accurate risk assessment is provided by the Tyrer-Cuzick model. This takes into account several factors including the presence of certain risk genes, age, family history of breast and ovarian cancer, as well as mammographic breast density. In addition to BRCA1 and BRCA2 several other risk genes are known that can be tested using gene panels. However specific familial risk constellations are prerequisites for indicating a genetic test. Prior to conducting a genetic test comprehensive counselling should take place and the individual seeking advice should be given time to consider. The individual seeking advice faces a series of questions regarding the potential implications of a genetic test which not only affect herself but also her environment.

In high-risk situations prophylactic mastectomy is an established surgical measure and intensified surveillance is a conservative approach. The latter includes semi-annual clinical breast examination with ultrasound starting from age 25 as well as an annual MRI mammography which exhibits the highest reliability compared to other imaging methods.

Medical consultation aims to enable a participatory decision-making process for the individual seeking advice. This requires comprehensive information for the individual seeking advice and incorporation of her preferences. Evidence-based decision aids from professional societies can improve the decisions of the individual seeking advice.

Keywords: *breast cancer – high risk – intensified surveillance – genetic testing - psychological aspects of patient management - participatory decision-making*

Despite now having good chances of cure and significantly improved survival rates a diagnosis of breast cancer is perceived as potentially life-threatening and affects the entire family and friends. Fears of disease-specific and psychosocial limitations take center stage along with the fear of a much too early end of life. This is particularly true when there is a suspected hereditary predisposition which substantially increases the risk of developing breast cancer especially at a young age.

This article explains how individual risk can be determined and what to consider in risk counselling. In particular it presents the option of intensified surveillance and possible alternatives.

Assessment of the Risk Situation

To assess the risk situation it is first necessary to examine the environment of the individual seeking advice. Concrete methodological guidelines for this are provided by the S3 Guideline for Breast Cancer from the Association of Scientific Medical Societies in Germany (AWMF) [1], the guidelines/recommendations of the Gynecological Oncology Working Group (AGO) of the German Society of Gynecology and Obstetrics (DGGG) and the German Cancer Society (DKG) [2], as well as the German Consortium for Hereditary Breast and Ovarian Cancer [3]. The information and recommendations derived from these three organizations largely coincide for practical use.

What Risk Groups exist?

The risk of developing breast cancer over the course of one's life, referred to as "lifetime risk" can be categorized into three classes: based on the epidemiological understanding that approximately one in eight women will develop breast cancer in their lifetime [4], the normal lifetime risk is 10-13%. High risk is classified at a lifetime risk of

30% for breast cancer or at a 20% heterozygous risk for the presence of a breast cancer gene based on pedigree analysis [1;3]. An intermediate risk is present when the lifetime risk falls between the two aforementioned risk groups.

When is there a "Family History" of Breast Cancer?

Approximately 30% of all women with breast cancer have a family history of the disease. A family history is considered to be present when multiple women (and possibly men) in the biological relatives of the individual seeking advice have been diagnosed with breast cancer and/or ovarian cancer in specific constellations (Tab. 1).

she truly ready to handle the knowledge of the presence of a breast cancer gene and thus a potentially existential threat along with the associated consequences?

Genetic Factors

The currently most well-known risk genes are the BReast CAncer (BRCA) genes, BRCA1 and BRCA2. These were the first breast cancer genes discovered in the 1990s. Since that time, it has been understood that carriers of these genes develop breast cancer approximately 20 years earlier, have a lifetime risk of 50-80% for breast cancer, a contralateral breast cancer risk of 60% and a lifetime risk of 10-40% for ovarian cancer [5-7].

Recommendations/Statements	
Genetic testing should be offered when there is a familial or individual history that is associated with at least a 10% probability of detecting a mutation. This is true if in one line of the family: • At least three women have been diagnosed with breast cancer. • At least two women have been diagnosed with breast cancer, one of them before the age of 51. • At least one woman has breast cancer and one woman has ovarian cancer. • At least two women have ovarian cancer. • At least one woman has breast and ovarian cancer. • At least one woman aged 35 or younger has been diagnosed with breast cancer. • At least one woman aged 50 or younger has bilateral breast cancer. • At least one man has breast cancer and one woman has breast and ovarian cancer. An appropriate time to think things over should be observed before carrying out the diagnostic procedure.	**Level of recommendation: Grade B** **Level of Evidence 5/2a (LoE2a = for mutation probability)**

Tab. 1: Inclusion criteria for a genetic investigation due to "family history" of breast cancer.

In these constellations a detection rate of at least 10% for genetic mutations can be expected. Thus the inclusion criteria for a genetic investigation would be met. Therefore before undergoing genetic testing it is essential to have a risk counselling session to determine the individual's lifetime risk.

Furthermore it is important to ensure that the individual seeking advice is given sufficient time for consideration before undergoing genetic testing: is

In recent years numerous variants of other gene regions that could be associated with breast cancer risk have been identified [8]. To more accurately assess the significance of individual gene defects two research groups from the USA and Europe have compiled data from case-control studies. The EU-funded Breast Cancer Association Consortium examined the impact of a total of 34 gene variants on breast cancer risk. The analysis involved 44 studies with 113’927 women (60’466 with

breast cancer and 53'461 healthy women as a control group) from 25 countries [9].

The US-based CARRIERS Consortium focused on results from 12 cohorts that described the effect of 28 gene variants in 64'791 women (32'247 with breast cancer and 32'544 without breast cancer) [10].

Both studies found a significant association between breast cancer risk and genetic alterations in the following eight genes: BRCA1, BRCA2, PALB2, BARD1, RAD51C, RAD51D, ATM, and CHEK2 [9]. The European study also identified a significant association with changes in the TP53 gene while the US study associated pathogenic variants of the CDH1 gene. For alterations in most of the other tested genes a significant correlation with an elevated breast cancer risk could not be described [9].

BRCA1 & BRCA2: Tip of the Iceberg

The BRCA1 and BRCA2 genes stood out in both studies: In the US study an increase in breast cancer risk by a factor of 7.62 for BRCA1 variants and a factor of 5.23 for BRCA2 variants was determined [10]. In the European study the results were even more pronounced: the odds ratios were 10.57 (BRCA1) and 5.85 (BRCA2) [9]. PALB2 is also considered a high-risk gene [2]. Moreover pathogenic variants in the ATM and CHEK2 genes were associated with a lifetime risk of over 20% [9]. Other moderately penetrant risk genes identified include BARD1, RAD51C, and RAD51D [11]. Additionally, it was observed that different breast cancer subtypes are influenced by these gene variants in distinct ways: While alterations in BRCA1, BRCA2, and PALB2 were more frequently discovered in triple-negative breast cancer, women with mutations in the ATM or CHEK2 genes were more likely to have estrogen receptor-positive tumors [9]

Increased Risk in Other Cancer Types

Pathogenic variants of breast cancer genes not only elevate the risk of developing breast cancer. Alterations in BRCA1 and BRCA2 also raise the risk of ovarian cancer in women and prostate cancer in men. Carriers of BRCA1 mutations have an elevated risk of colorectal cancer while carriers of BRCA2 mutations are more susceptible to other cancer types than the average population. Mutations in the CHEK2 gene increase the risk of colorectal cancer. ATM mutations are possibly linked to an increased risk of pancreatic cancer. Other associations such as between hereditary non-polyposis colo-rectal cancer and breast cancer could not be confirmed [9]. However several other cancer syndromes are recognized to come with an elevated risk of breast cancer such as Li-Fraumeni syndrome [2]. Since some years now, commercially available gene panels have the capability to identify the presence of risk genes [3]. Some of these provide a polygenetic risk score (PRS) as a single value to estimate an individual's genetic susceptibility to developing cancer.

Non-Genetic Risks

A specific risk situation arises following radiation therapy in the thoracic region. A typical scenario is radiation exposure to the thorax during childhood or adolescence such as in cases of Hodgkin's lymphoma leading to an increased risk of developing breast cancer [12-14].

How Lifetime Risk is determined

The individual's lifetime risk of developing breast cancer is influenced not only by genetic factors but also by a variety of other variables. Over the past years several models have been introduced aiming to achieve both high pre-

dictive accuracy and user-friendly applicability.

Currently the Tyrer-Cuzick model [15] achieves the highest predictive accuracy. It takes into account multiple genes and their penetrance family history endo-genous hormone status potential hormone replacement therapy (HRT), risk factors like age and BMI, and almost uniquely mammographic breast density. The model can be accessed free of charge via the web interface of the In-ternational Breast Cancer Intervention Study (IBIS) [16]

substantially reduces the risk of breast cancer [19]. Bilateral prophylactic mastectomy gained public attention a few years ago when Angelina Jolie a prominent individual facing a high-risk situation openly discussed her experience. However such a procedure can have significant psychological and psycho-sexual impacts on the individuals, an area that remains insufficiently explored [20; 21]. For carriers of BRCA1 or BRCA2 mutations, due to the increased risk of ovarian cancer consideration should also be given to risk-reducing salpingo-oophorectomy [2].

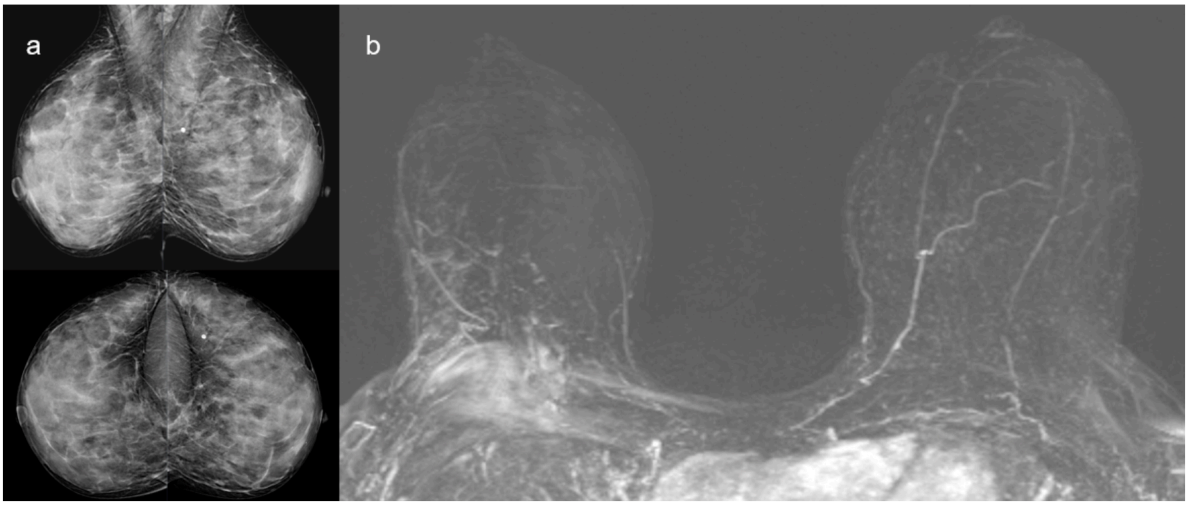


Fig. 1: 38-year-old woman whose mother was diagnosed with breast cancer at the age of 37.
(a) The mammography, due to the age-related low tissue transparency (ACR Type D), has limited diagnostic value.
(b) However, the MR mammography (here the maximum intensity projection of the latest image from the contrast-enhanced dynamic series) does not show any pathological contrast enhancement and effectively rules out the presence of breast cancer with high certainty.

What Preventive Measures for which Level of Risk?

For individuals with a "normal" lifetime risk of breast cancer participation in a systematic breast screening program is recommended. The basis of this approach is a mammography at 2-year intervals between the ages of 50 and 70 or 75 [17].

For a "moderate" risk situation there are no standardized recommendations. In cases of a "high-risk" constellation conservative methods of care are summarized under the term "intensified surveillance" [18]. As a surgical measure mastectomy is an option which involves removing the mammary gland and

What is Intensified Surveillance?

The goal of intensified surveillance is twofold: to detect newly occurring breast cancer as early as possible while minimizing the potential harmful effects of diagnostic procedures. These harmful effects include not only physical damage from ionizing radiation exposure or biopsy-related effects but also emotional distress reactions. The key aspects of intensified surveillance are therefore the choice of intervals and diagnostic methods. Regarding diagnostic methods all known breast examination techniques are applicable for intensified surveillance: clinical examination involving physical inspection, palpation, ultra-sound, mammo-

graphy, and magnetic resonance imaging (MRI) with potential image-guided biopsies. Breast self-examination does not play a role within intensified surveillance.

Compared to systematic breast cancer screening intensified surveillance faces distinct challenges. These challenges include higher proliferation rates and faster growth of breast cancers specific tumor biology as seen in BRCA1-associated malignancies often presenting with a "benign" morphology and the relatively younger age at first manifestation resulting in reduced breast tissue transparency in young women and consequently reduced mammography sensitivity for early detection. These characteristics are not accounted for in systematic breast cancer screening programs.

of detecting breast cancers at an early stage. The impact of new genetic diagnostics on the other hand is still uncertain [1].

Therefore the AWMF recommends intensified surveillance with MR mammography within a quality-assured program for women with pathogenic BRCA1/2 mutations and for women with a lifetime risk of ≥ 30% due to other reasons [1]. This recommendation is supported by recent data showing high negative predictive value and a sensitivity of nearly 90% for MR mammography (Fig. 1) [22]. In a recent study comparing MR mammography, ultra-sound, and mammography in a high-risk group 21-36% of breast cancers were detectable solely through MR mammography (Fig. 2) whereas only 1% to 3% were detected solely through

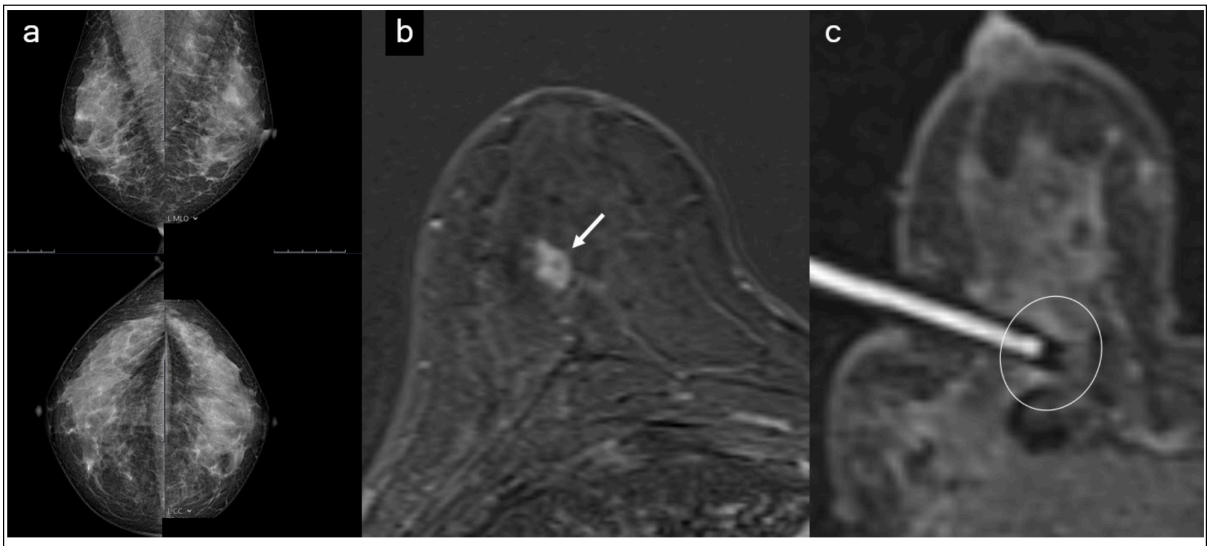


Fig. 2: Figure 2: 43-year-old woman with BRCA2 mutation.
(a) Mammography does not reveal any significant findings, but due to low tissue transparency (ACR Type C), it has limited diagnostic value. An ultrasound-guided core biopsy of an complex echo area in the upper outer quadrant of the right breast did not show evidence of malignancy.
(b) MR mammography exhibited a non-mass enhancement in the right breast (arrow).
(c) MR-guided vacuum biopsy of the area (circle) confirmed the diagnosis of ductal cancer in situ (DCIS).

For these reasons recommendations from medical societies are based on the specific benefit-risk ratio of available methods in high-risk situations. The guideline from the AWMF notes that the diagnostic accuracy of mammography and MRI is published but a reduction in mortality through these methods in high-risk situations has not yet been proven. However it is clear that intensified surveillance with MRI is capable

ultrasound or mammography, respectively [22].

The AGO recommends a multimodal approach to intensified surveillance starting at the age of 25, including semi-annual clinical breast examination, and ultrasound, as well as annual MR mammography [2].

Case Presentation
Julia, a 36-year-old woman and mother of two daughters has a 3 year younger sister who recently discovered a lump in her breast. Subsequent diagnosis confirmed breast cancer. Since then Julia has been gripped by a strong fear that she might also be afflicted with breast cancer.

Though she hasn't detected any lumps in her own breasts Julia has been experiencing a sensation of pressure and occasional discomfort in both breasts lately. She is deeply concerned about whether these sensations could potentially be indicative of breast cancer. Seeking guidance, she schedules an appointment with her gynecologist and asks for advice on what steps she should take next.

Fear
Experiencing fear is principally normal and healthy. Fear arises when we find ourselves in situations we are not yet equipped to handle or understand. The fear response has evolved over time to ensure human survival. Therefore fear is not an illness in itself; it triggers actions for personal protection and self-care such as seeking advice undergoing genetic testing, and participating in an intensified surveillance program when facing a suspected or confirmed high-risk situation for breast cancer.

The trigger for fear is not a specific perception but rather how we evaluate that perception. When fear takes up increasing space and becomes an energy drain it can reach a pathological level. Particularly when fear significantly impacts quality of life or daily functioning, psychological support might be necessary.

What do the guidelines say?
As a physician in the consultation room we acknowledge and address Julia's fears and concerns. In the first step it is important to recognize that we are not facing a patient, but an individual seeking advice. Therefore our response is guided by the principle of "primum nil nocere" which means "first do no harm" [2]. This principle encompasses both potential physical and psychological harm. Therefore before offering any preventive measures a comprehensive and thorough consultation with a balanced consideration of benefits and risks should take place [2]. The evidence-based decision aids provided by professional societies can enhance the decisions made by those seeking advice [1].

How can we advise the individual seeking guidance?
It is important to conduct the consultation in a way that enables participatory decision-making for the individual seeking advice. This requires comprehensive information provision, addressing their questions, and taking their preferences into account. This applies not only to the decision regarding the extent of risk assessment such as whether to explore "only" familial or individual burdens but also extends to the consideration of predictive diagnostics, including genetic testing. It is crucial that the individuals have sufficient time to absorb the information and make their own decisions.

During the consultation it is essential that the options presented are not seen as obligations by the individual seeking advice. Particularly the decision-making process regarding genetic testing is open-ended. Several questions need to be clarified for the individual seeking advice (Tab. 2).

In addition to the responsibility for oneself family responsibilities also play a role in the decision-making process. For instance, mothers might contemplate

their children's future, women in partnerships may consider the impact on their relationship, and young women might ponder the implications for their desire to have children.

analyses not only assess predispositions, but also have psychological and social consequences for those being tested and their relatives directly influencing their life and family planning.

Questions to be answered
What is the aim of the examination for me?
What risks are associated with the examination?
What does the disease under investigation mean for me and my family?
What are the consequences of an abnormal test result for us?
What options are available when a disease/genetic condition is identified ? Are they suitable for me?
What are the alternatives if I decide not to undergo the examination?
Who will support my decision?

Tab. 2: Questions that the individuals seeking advice should be able to answer for themselves after a consultation on genetic diagnostics [23].

Engaging with the options is a process of becoming that orients itself toward the future. Experiences of the family and others who have faced or are facing the same life challenge are taken into consideration. Satisfaction with the decision can arise when the individual seeking advice can be certain that they are only asked or required to undertake what they believe they are capable of handling.

Auxiliary Communication

The collaboration between the physician and the individual seeking advice within the context of genetic counselling in high-risk situations is a significant challenge for both communication partners. The diversity of diagnostic, therapeutic, and preventive options requires a high degree of orientation not only from medical professionals but also from those affected and their families. Genetic

Experience shows that inadequate communication and misunderstanding between the physician and the individual seeking advice is a major and burdensome problem that often leads to hurt feelings, frustration, or anger. What is routine for the physician is an entirely new life situation for the individual seeking advice. What is professional work for the physician is personal fate for the individual seeking advice. The most important instrument for guidance is the guides themselves. In a consultation conversation it is not only the medical expertise of the counsellor that is needed but also their communicative competence as a significant qualitative determinant. Appropriate communication requires mastery of situation-appropriate conversation techniques as well as interpersonal skills. In recent years the concept of patient-centered communication has gained prominence, emphasizing understanding and considering the

patient's perspective at its core [24]. Patient-centered communication includes providing understandable information to the individual seeking advice regarding examination techniques such as genetic analysis in cases of possible familial disposition, diagnosis, and treatment options, as well as patient-centered, empathetic communication even for unfavorable news. The concept of patient-centered communication does not only incorporate the individual experiences, expectations, and fears of the patient or individual seeking advice into the physician-patient interaction but also aims to perceive and understand patients/individuals seeking advice within their own psychosocial reference frames. Patient-oriented dialogue necessitates mutual listening and sharing. The concept of auxiliary (help-related) communication implements these basic assumptions by not only providing factual information but also attempting to comprehend the individual's reference system and experience, including its impact on their life. The guiding attitude is patient-oriented rather than centered on the perspective of the advisor or expert. The advising physician does not need to agree with or find everything the individual seeking advice expresses to be good or right. What is crucial is that the advisor "does not find" it at all – neither positively nor negatively. This non-evaluative stance is a prerequisite for the individual seeking advice to open up without fearing rejection. Furthermore, neutrality ensures a neutral stance of the advisor towards the individual seeking advice. If the advisor manages to accept the individuals seeking advice with all their pain and fear and withstand it; it will also be easier for the individuals seeking advice to accept themselves. The advising physician is a companion who presents alternatives or suggestions for solutions. However the final decision always remains with the patient.

Resolution of the Case Presentation

After receiving counselling Julia decided to start by estimating her individual lifetime risk. The IBIS analysis based on the Tyrer-Cuzick model revealed that Julia has a 10-year risk of 3% and a lifetime risk of 24% for developing breast cancer. This indicates a risk around 3 times higher than that of a woman with a normal risk. This assumption considers that her sister's BRCA status is unknown.

However if her sister were found to have the BRCA1 gene Julia's 10-year risk would be nearly 18% and her lifetime risk would be close to 48% indicating a nearly fivefold increased risk of breast cancer. Due to her sister's breast cancer diagnosis at the age of 33 Julia meets the criteria for undergoing genetic testing herself.

Consequently Julia is confronted with the questions formulated in Tab. 2 which we as the advising physician can help her address by providing factual information and appropriate communication techniques. It is ultimately her decision whether she wants to proceed with genetic testing, opt for intensified surveillance, consider prophylactic mastectomy, or choose not to undergo any of these measures.

Conclusion for Clinical Practice

Asymptomatic women who are concerned about an increased risk of breast cancer due to family history should first receive comprehensive and detailed counselling including a discussion of the benefits and risks of diagnostic and preventive measures. Since these individuals are not "patients" but healthy women seeking guidance the principle of non-maleficence takes precedence.

In cases of high-risk situations besides prophylactic mastectomy, intensified

surveillance, following clearly structured recommendations of the relevant professional societies is an established approach. These measures should be carried out at specialized centers within the frame of quality-assured programs. The focus should be on considering the individual preferences of the women seeking advice.

Conflict of interest:
The authors declare that there were no conflicts of interest within the meaning of the recommendations of the International Committee of Medical Journal Editors when the article was written.

Correspondence address:
Prof. Dr. med. Florian Dammann, MD
University Institute of Diagnostic, Interventional and Paediatric Radiology
Inselspital, University Hospital Bern
University of Bern
Freiburgstrasse 18
CH-3010 Bern
Switzerland
Email: f.dammann@gmx.com

Dr. med. Susanne Ditz, MD
Psychooncology and Psychosomatics Dept.
University Women's Hospital Heidelberg
University Hospital, University of Heidelberg
Im Neuenheimer Feld 440
D-69120 Heidelberg
Germany
Email: ditz@psychosomatische-frauenheilkunde.de

References

1. S3-Leitlinie Früherkennung, Diagnose, Therapie und Nachsorge des Mammakarzinoms. Version 4.3. 2020. AWMF-Registernummer: 032-045OL. (http://www.leitlinienprogramm-onkologie.de/leitlinien/mammakarzinom). search from 03.11.2022

2. Arbeitsgemeinschaft Gynäkologische Onkologie e. V. Früherkennung und Diagnostik. (https://www.ago-online.de/leitlinien-empfehlungen/leitlinien-empfehlungen/kommission-mamma). search from 03.11.2022

3. Deutsches Konsortium Familiärer Brust- und Eierstockkrebs. Konsensusempfehlungen des Konsortiums (Stand 2022). (https://www.konsortium-familiaerer-brustkrebs.de/konsensusempfehlung/). search from 03.11.2022

4. Robert Koch-Institut, Zentrum für Registerdaten. Brustkrebs (Mammakarzinom). (https://www.krebsdaten.de/Krebs/DE/Content/Krebsarten/Brustkrebs/brustkrebs_node.html#:~:text=Brustkrebs%20(Mammakarzinom)&text=Mit%20zuletzt%20rund%20071.375%20Neuerkrankungen,Prozent%20aller%20Neuerkrankungen%20betri%EF%AC%80t%20M%C3%A4nner.). search from 03.11.2022

5. Antoniou A et al. Average risks of breast and ovarian cancer associated with BRCA1 or BRCA2 mutations detected in case Series unselected for family history: a combined analysis of 22 studies. Am J Hum Genet 2003; 72: 1117–30

6. Goldberg JI & Borgen PI. Breast cancer susceptibility testing: past, present and future. Expert Rev Anticancer Ther 2006; 6: 1205–14

7. Engel C et al. Breast cancer risk in BRCA1/2 mutation carriers and noncarriers under prospective intensified surveillance. Int J Cancer 2020; 146: 999–1009

8. Fachal L et al. Fine-mapping of 150 breast cancer risk regions identifies 191 likely target genes. Nat Genet 2020; 52: 56–73

9. Dorling L et al. Breast Cancer Association Con-sortium. Breast Cancer Risk Genes - Association Analysis in More than 113,000 Women. N Engl J Med 2021; 384: 428–39

10. Hu C et al. A Population-Based Study of Genes Previously Implicated in Breast Cancer. N Engl J Med 2021; 384: 440–51

11. Yang X et al. Ovarian and Breast Cancer Risks Associated With Pathogenic Variants in RAD51C and RAD51D. J Natl Cancer Inst 2020; 112: 1242–50

12. Alm El-Din MA et al. Breast cancer after treatment of Hodgkin's lymphoma: risk factors that really matter. Int J Radiat Oncol Biol Phys 2009; 73: 69–74

13. Preston DL et al. Radiation effects on breast cancer risk: a pooled analysis of eight cohorts. Radiat Res 2002; 158: 220–35

14. Travis LB et al. Cumulative absolute breast cancer risk for young women treated for Hodgkin lymphoma. J Natl Cancer Inst 2005; 97: 1428–1437

15. Tyrer J, Duffy SW, Cuzick J. A breast cancer pre-diction model incorporating familial and personal risk factors. Stat Med 2004; 23: 1111–30

16. Ikonopedia. IBIS (International Breast Cancer Intervention Study). Online Tyrer-Cuzick model breast cancer risk evaluation tool – IBIS risk assessment tool v8 .0b. (https://ibis.ikonopedia.com/). search from 03.11.2022

17. Bundesministerium für Gesundheit. Krebsfrüherkennungsuntersuchung in der gesetzlichen Krankenversicherung (§ 25 Abs. 2 SGB V) / Mammographie. Informationsblatt Nr. 315-02. (https://www.bundesgesundheitsministerium.de/fileadmin/Dateien/3_Downloads/M/Mammografie/Infoblatt_Krebsfrueherkennung_Mammographie.pdf). search from 03.11.2022

18. Deutsches Konsortium Familiärer Brust- und Eierstockkrebs. Intensivierte Früherkennung Früherkennung durch moderne Bildgebung. (https://www.konsortium-familiaerer-brustkrebs.de/betreuungskonzept/praevention/intensivierte-frueherkennung/). search from 03.11.2022

19. Jakub JW et al. Oncologic Safety of Prophylactic Nipple-Sparing Mastectomy in a Population With BRCA Mutations: A Multi-institutional Study. JAMA Surg 2018; 153: 123–9

20. Jeffers L et al. Interventions to improve psycho-social well-being in female BRCA-mutation carriers following risk-reducing surgery. Cochrane Database Syst Rev 2019; 10: CD012894

21. Scheepens JCC et al. Contralateral prophylactic mastectomy: A narrative review of the evidence and acceptability. Breast 2021; 56: 61–9

22. Bick U et al. German Consortium for Hereditary Breast and Ovarian Cancer (GC-HBOC). High-risk breast cancer surveillance with MRI: 10-year experience from the German consortium for hereditary breast and ovarian cancer. Breast Cancer Res Treat 2019; 175: 217–28

23. Gödde E. Erbliche Krebsdisposition: Familienplanung zwischen Angst und Verantwortung. EHK 2010; 59: 189–98

24. Langewitz W. Patientenzentrierte Kommunikation. In: Adler RH, Herzog W, Joraschky P, Köhle K, Langewitz W, Söllner W, Wesiak W, Hrsg. Uexküll, Psychosomatische Medizin. Theoretische Modelle und klinische Praxis. München: Urban & Fischer; 2011. S. 338–47