Prepectoral implant placement in plastic-reconstructive breast surgery – a contribution to the discussion

Präpektorale Implantateinlage in der plastisch-rekonstruktiven Mammachirurgie – ein Diskussionsbeitrag

Authors
Stefan Paepke¹, Ralf Ohlinger², Evelyn Klein¹, Marc Thill³

Affiliations
1 Breast Centre of the Technical University of Munich, Germany
2 Breast Centre of the Ernst Moritz Arndt University of Greifswald, Germany
3 Obstetrics and Gynaecology Clinic, Breast Centre, Agaplesion Markus Hospital, Frankfurt am Main, Germany

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Correspondence
Dr. Stefan Paepke
Chief Physician: Senology
Breast Centre of the Technical University of Munich,
Ismaninger Strasse 22, 81675 Munich
stefan.paepke@lrz.tu-muenchen.de

ZUSAMMENFASSUNG
Die implantatbasierten Rekonstruktionstechniken nehmen in den letzten Jahren national und international kontinuierlich zu und decken ein weites Spektrum rekonstruktiver Anforderungen ab. Die nipple-sparing subcutaneous Mastektomie mit subpektoraler Implantat-Sofortrekonstruktion stellt das derzeitige Standardverfahren dar. Diese Operationstechnik erfordert jedoch den Einsatz heterologen Materials in Form von azeelläreren dermalen Matrices oder synthetischen Netzen, um eine optimale Implantatabdeckung, eine Stabilisierung des unteren Brustpols, die Stabilisierung des Implantats und die Fixierung des Muskels zu erreichen und somit ein opti-
Introduction

Throughout the year, many current aspects of plastic-reconstructive breast surgery have been discussed in a wide range of lectures and debates in various symposia on senology and surgery. Standards have been defined, perspectives offered, and the field has been opened up to possible further developments.

- Indications
- Patient selection
- Anatomically correct surgical techniques with the use of intra-operative sonography to ensure surgical precision
- Complications and the possibility of avoiding or minimising them, as well as problem-solving alternatives
- Classification of surgical techniques
- National study data.

When considering the developments in implant-based reconstructive breast surgery of recent years, the following should be noted:

- Changes in basic oncological techniques, from radical to skin- and nipple-saving mastectomy techniques
- Highly-developed cooperation between professionals (senology, surgery, plastic surgery) in multi-disciplinary teams
- An increase in bilateral interventions, both primary and secondary prophylactic interventions
- An increase in reconstructive interventions in every age group, with an above-average increase in young women
- An increase in the number of implant-based, reconstructive interventions overall
- Widespread and safe use of autologous reconstruction techniques
- The introduction of newly-developed breast implants for reconstructive breast surgery
- The introduction of materials to bridge, substitute or support the tissues in reconstructive breast surgery since around 2008
- Further developments in materials to bridge, substitute or support the tissues, implemented in the German Working Group for Gynaecological Oncology’s treatment recommendations since 2011
- The introduction of minimally invasive surgical techniques
- Additional tools for intraoperative safeguarding of skin perfusion

Implant-based plastic-reconstructive breast surgery

Implant-based techniques constitute approximately 40 – 60% of all breast reconstructions in Europe and approximately 75% in the USA [2 – 4]. It is therefore not surprising that implant-based breast surgery modifications of implant-based reconstructions is a subject of discussion in interdisciplinary knowledge exchange. The performance of a subcutaneous mastectomy within anatomical limits [5 – 7] and the optimal design of the implant bed have been specified. In this context, the surgical separation plane of the pectoralis major muscle, which is used for the cranial covering of the implant, has also been defined [8 – 10].

Subpectoral implant placement was the previous standard method of implant-based reconstruction, either with

- exclusive skin coverage (mostly as an interim solution until the definitive reconstruction is specified)
- complete muscle coverage
- dual-plane method with a de-epithelised corial flap
- caudo-lateral interposition of an acellular matrix
- caudo-lateral interposition of a synthetic mesh
- combination techniques.

Complication types and rates are described in many analyses, although these are mainly retrospective [11, 12]. The few prospective studies largely support the findings of the retrospective analyses. To summarise: it can be said that the use of both synthetic meshes and of acellular dermal and tissue matrices can be considered safe [1, 13 – 15]. The use of synthetic meshes seems to have few complications with regard to seroma formation and implant loss [10]. A single prospective, directly comparative study has shown that titanised polypropylene mesh (Tiloop® Bra) has an advantage over the porcine acellular dermal matrix Strattice™ regarding both the complication rate and the aesthetic result [16]. Newly-developed dermal matrices display a low rate of side effects in first application and will be analysed further in studies and registries.

Besides the question of the optimum material, the question of paramount importance regarding the implant bed is: pre- or sub-pectoral?

A large number of sometimes significant problems are caused by what is regarded as the standard placement of the subpectoral implant (Fig. 1):

- unnatural breast shapes to some extent
- the “jumping breast” phenomenon
- muscle response even with less expansive movements
- muscle pain
- contraction of the upper exterior of the severed pectoralis major muscle with wrinkle formation below the muscle origin.
- fasciculations
- restricted shoulder-arm mobility.

It must be considered, however, that in the past, a prepectoral implant placement exclusively covered by remaining skin led to changes in the subpectoral implant placement due to the high rate of complications. Current data shows the limitations of prepectorally placed implants, which presuppose per se a very thick layer of subcutaneous fat that must be considered critical from the oncological perspective [17]. However, Sigalove et al. [18] report that over 207 patients have been operated on since 2008, all for prepectoral implant placements, ADM covering and lipofilling (“bio-engineered breast concept”) in 353 operations. The complication rate is remarkably low; infections occurred in 4.5% of cases, seromas in only 2% and flap necroses in 2.5%. The authors explain that the contraindications to a prepectoral implant placement arise with thin, poorly perfused or ischaemic skin, in response to previous radiotherapy in conjunction with a scar in the lower pole or in the region of the radiation boost, with a BMI
several years [19, 20]. At this year has also increasingly been used in German-speaking countries for retrospective analyses shows that the prepectoral implant bed categorically applies. The publication of cases, case series and counter-argument against prepectoral implant placement no longer now be compensated for. Thus, in our view, an important coun-

t calibrated procedure that included the formation of a complete implant sheath from an ADM, more specifically from the porcine ADM Strattice® – a procedure which is likely to be problematic from the perspective of health economics. Preference is often given to ADMs when considering prepectoral implant placements, as the benefit of tissue substitution is ascribed to them, resulting in better coverage of the cranial parts of the breast. Meanwhile, different manufacturers have responded to the special require-
ments of prepectoral implant placement by developing new forms using known materials. Pre-formed ADM materials such as Braxton (DecomedSrl, Italy) [21] can be used directly as implant pockets and other ADMs such as Tutomesh or Strat tice and Artia can be used to shape the contours. Other ADMs, to the best of our knowledge, are not shaped but are squarer. H. Becker et al. [22] use both different synthetic meshes and ADMs to cover the im-
plant completely in a prepectoral implant placement.

The authors of this discussion contribution also use synthetic meshes to cover prepectorally positioned implants just as frequently as ADMs and tissue matrices [23]. Similarly, some patients with epipectoral implant placement and Tiloop® Bra coverage were included and evaluated in the PRO Bra-Trial [24, 25]. Observations of possible complications and problems also arise from this experience in individual cases. New materials are also being developed in the field of synthetic meshes. Building on the experi-
ce of D. Casella et al. [26] and M. Rezai, a titaniumised polypropylene implant pocket was developed, considerably simplifying prepectoral implant placement with a full covering of titanium mesh (Fig. 2, 3).

Three different implant pockets are available, enabling incre-
mental implant volumes of <270, <420 and <550 cm³ and im-
plant widths from 11.0 to 15.0 cm with projections of <4.5 to <6.0 cm. In addition to the already frequent use of the Tiloop® Bra pocket in Italy, different breast centres across Germany have used it on approximately 60 occasions (Fig. 4).

To summarise the considerations regarding the individual deci-
sion for a prepectoral implant placement from the authors’ point of view, the following factors present themselves: from the patient’s perspective, her own wishes are naturally paramount. More objective characteristics may be found, on the one hand, in a physically active, non-smoking patient with a BMI of <35, firm or slightly sagging skin with no striae distensae or current long-term systemic cortisone treatment, with mild or moderate ptosis and a resection weight of under 500 g. On the other hand, patients with glandular bodies in subinvolution and very flat cleavage that a

A 34-year-old patient, 7 months post subcutaneous mastectomy (status post primary chemotherapy, status post radiother-
apy). Immediate reconstruction using subpectoral implant place-
ment with caudal mesh interponation. Currently with grade III capsular contracture, implant cranialisation and empty volume in the caudal quadrants.

Fig. 1 A 34-year-old patient, 7 months post subcutaneous mastectomy (status post primary chemotherapy, status post radiotherapy). Immediate reconstruction using subpectoral implant placement with caudal mesh interponation. Currently with grade III capsular contracture, implant cranialisation and empty volume in the caudal quadrants.

Fig. 2 Synthetic mesh specially designed for prepectoral implant placement (Tiloop® Bra Pocket) Source: pfm medical AG, Cologne.
Concluding remarks

When reviewing the current developments in reconstructive breast surgery, it is striking that the basic strategies have been or are covered by retrospective analyses and an increasing number of prospective studies and data from registers, and are reflected in the respective evidence level of the German Working Group for Gynaecological Oncology’s treatment recommendations. This provides a high degree of user safety. Patient satisfaction regarding the cosmetic result reaches 80 – 90 % and the complications rate is absolutely acceptable. Nevertheless, the optimisation of autologous and implant-based reconstructive surgical techniques must remain the focus. The consideration of which implant bed is to be preferred in the context of individual surgery planning will be rendered much simpler by the introduction of advanced materials. Indications and complications should be recorded in the registers.

Conflict of interest

The authors will declare any conflicts of interest (honorary activities, travel and conference support, support for scientific events).

References


