

Case Study

Treatment of Melanoma with Human Acellular Dermal Matrix (ADM)

Barysch MJ¹, Bertasi G², Hafner J¹, Läuchli S¹

¹ University of Zurich, Swiss

² University of Padua, Italy

Melanoma originates in the majority of cases from the melanocytes found in the epidermis. Melanocytes are located in the deepest layer of the epidermis and produce melanin, the pigment responsible for skin coloration. Over the last 10 years, new cases of melanoma in the Caucasian population have increased by approximately 5% per year. Melanoma can appear at any age, and it is one of the most frequently occurring cancers in adults aged between 30 and 40 years.

Those most at risk of developing melanoma have one or more of the following characteristics:

- Fair complexion, blue eyes and blond or red hair
- Often clinically atypical
- History of repeated sunburn, especially in adolescence and young age
- Personal history of melanoma
- Nearest relatives suffering from melanoma
- Alterations of immune defenses (for example subject undergoing organ transplant or in immunosuppressive therapy)

Early diagnosis provides the most effective instrument for reducing death associated with melanoma. The prognosis is often considered excellent if the melanoma can be recognized at the initial phase. Surgery is the principal therapeutic modality in the treatment of primary melanomas and metastases to lymph nodes. Surgery also plays an important role in the treatment of locally recurrent tumors and distant metastases in select cases. Wide excision of the primary tumor in stage I malignant melanoma results in a recurrence rate of about 2%. Surgical techniques for lymph node dissection are the same for melanoma as those performed for other malignancies. There is little question that dissection is necessary in obviously involved nodal areas, but controversy remains regarding the value of lymph node dissection of clinically uninvolved regional nodes. A prospective, randomized study by the World Health Organization has shown no improvement in survival for patients who underwent elective regional node dissection.

The following case presentation involves the treatment of a melanoma tumor excision site with a human acellular dermal matrix (ADM), DermACELL AWM®.

Patient

- An 83 year old woman diagnosed with local relapse of a stage IIIB melanoma on the left arm.

Treatment

- The primary tumor was excised with a safety margin 5 years prior, but further staging and further treatment were declined. As the tumor bulk became bothersome due to the size and pain, the patient requested a local excision in an ambulant setting (**Fig. 1**). Besides dementia of a mixed vascular Alzheimer type and an arterial hypertonia, the patient was otherwise healthy.
- The tumor bulk was excised with a safety margin of 1 cm in a lazy-S-shape and directly closed after undermining. The patient was referred to us three days postoperative due to an extensive postoperative hematoma with tension bullae (**Fig 2**). After wound revision, the wound was partly adapted and bandaged with silver-coated alginate wound dressing.
- As the patient refused donation for a large split skin graft or an extensive flap, we opted for the application of a decellularized matrix, DermACELL, for wound closure. Therefore, we freshened the wound edges 5 days after wound revision (**Fig. 3**), and applied the meshed (1:1.5) decellularized matrix on the wound. The matrix was fixed with non-absorbable sutures at the edges and the center (**Fig. 4**). Non-adhesive gauze was applied below a vacuum-assisted wound closure dressing for negative pressure wound therapy with a continuous pressure of 80 mmHg. Postoperatively, cefuroxime was taken for 3 days.
- Four days after application of the decellularized matrix, the first changes of the wound dressing took place (**Fig. 5**). The wound was clean, bleeding had stopped, and the matrix was white in appearance as expected. The wound dressing was changed into a non-adhesive gauze containing nanocrystalline silver ions.

Results

- Postoperatively at eight and eleven days (**Fig. 6**), the wound still revealed the whitish mesh scaffold, but an increase in remodeling was evident. The crusted overlaying edges were removed.
- Fifteen days postoperatively (**Fig. 7**), the whitish scaffold was barely visible, as it was almost completely remodeled. Additionally, the wound showed granulation from the base.
- The wound size continuously decreased until fully closing at the final 8 weeks postoperative wound visit (**Fig. 8**). The wound was completely healed, and remarkably, the scar was thin and much lower in diameter than the original wound. Six months afterwards, the wound was still stable and comparable to the clinical presentation at the eight weeks postoperative visit.

Conclusion

- DermACELL AWM was able to successfully heal the excision site of a melanoma tumor.



Figure 1. Tumor bulk before excision



Figure 2. 3 days postoperative with extensive hematoma



Figure 3. Wound before DermACELL application



Figure 4. DermACELL graft application



Figure 5. 4 days postoperative



Figure 6. 8-11 days postoperative



Figure 7. 15 days postoperative



Figure 8. 8 weeks postoperative

References

1. Attie, N.J., Kafif, R.A.: Melanotic Tumors. Springfield, Illinois, C.C. Thomas Publishers, 1964, p. 193
2. Pack, G., Scharnagel, I., Morfit, M.: The principle of excision and dissection in continuity of primary and metastatic melanoma of the skin. *Surgery*17:849, 1945
3. Veronesi, U., Adamus, J., Bandiera, D.C., Brennhovd, I.O., Caceres, E., Cascinelli, N., Caludio, F., Ikonopisov, R.L., Javorskj, V.V., Kirov, S., Kulakowski, A., Lacour, J., Lejeune, F., Mechl, Z., Morabito, A., Rodé, I., Sergeev, S., van Slooten, E., Szczygiel, K., Trapeznikov, N.N., Wagner, R.I.: Inefficacy of immediate node dissection in stage I melanoma of the limbs. *N. Engl. J. Med.*297:627, 1977
4. Veronesi, U., Bajetta, E., Cascinelli, N., Clemente, C., Rilke, F.: New trends in the treatment of malignant melanoma. In *International Advances in Surgical Oncology*, Vol. 1, Murphy, G.P., editor. New York, A.R. Liss, 1978, pp. 113–156 [Google Scholar](#)
5. Davis, N.C., Little, J.H.: The role of frozen section on the diagnosis and management of malignant melanoma. *Br. J. Surg.*61:505, 1974
6. McGovern, V.J.: Malignant Melanoma: Clinical and Histological Diagnosis. New York, John Wiley and Sons, 1976
7. Cascinelli, N., Balzarini, G.P., Fontana, V., Morabito, A., Orefice, S.: Long-term results of surgical treatment of melanoma of the limbs. *Tumori*62:233, 1976
8. Olsen, G.: Surgical treatment of primary melanoma: some views of the sites and depth of excision. In *Melanoma and Skin Cancer*, McCarthy, W.H., editor. Sydney, V.C.N. Blight, 1972, p. 389
9. Breslow, A., Macht, S.D.: Optimal size of resection margin for thin cutaneous melanoma. *Surg. Gynecol. Obstet.*145:691, 1977
10. Cohen, M.H., Schour, L., Felix, E.L., Bernstein, A.D., Chretien, P.B., Rosenberg, S.A., Ketcham, A.S.: Staging laparotomy in the treatment of metastatic melanoma of lower extremities. *Ann. Surg.*182:710, 1975
11. Fortner, J.G., Schottenfeld, D., MacLean, B.J.: En bloc resection of primary melanoma with regional lymph node dissection. *Arch. Surg.*110:674, 1975
12. Papachristou, D., Fortner, J.G.: Comparison of lymphedema following in-continuity and discontinuity groin dissection. *Ann. Surg.*185:13, 1977 [Google Scholar](#)
13. Southwick, H.W.: Malignant melanoma: role of node dissection reappraised. *Cancer*37:202, 1976
14. Veronesi, U., Cascinelli, N., Balzarini, G.P. et al.: Treatment of regional node metastases. In *Melanoma and Skin Cancer*, McCarthy, W.H., editor. Sydney, V.C.N. Blight, 1972, p. 417
15. Davis, N.: Cutaneous melanoma: the Queensland experience. *Curr. Probl. Surg.*13:1, 1976
16. Wanebo, H.J., Fortner, J.G., Woodruff, J., MacLean, B., Binkowski, E.: Selection of optimum surgical treatment of stage I melanoma by depth of micro-invasion: use of the continued microstage technique (Clark-Breslow). *Ann. Surg.*182:302, 1975
17. Clark, W.H., From, L., Bernardino, E.A., Mihm, M.C.: The histogenesis and biological behavior of primary tumor malignant melanoma of the skin. *Cancer Res.*29:705, 1969
18. Breslow, A.: Tumor thickness, level of invasion, and node dissection in stage I cutaneous melanoma. *Ann. Surg.*182:572, 1975
19. WHO Collaborating Centers for Evaluation of Methods of Diagnosis and Treatment of Melanoma (Breslow, A., Cascinelli, N., Van der Esch, E.P., Morabito, A.): Stage I melanoma of the limbs: assessment of prognosis by levels of invasion and maximum thickness. *Tumori*64:273, 1978
20. Sim, F.H., Taylor, W.F., Ivins, J.C., Pritchard, D.J., Soule, E.H.: A prospective, randomized study of the efficacy of routine elective lymphadenectomy in management of malignant melanoma. Preliminary results. *Cancer*41:948, 1978
21. Paterson, A.H.G., McPherson, T.A., Willans, D.J.: Malignant melanoma (stage IIIB): a pilot study of adjuvant chemo-immunotherapy. *Cancer Treat. Rep.*62:571, 1978
22. WHO Collaborating Centers for Evaluation of Methods of Diagnosis and Treatment of Melanoma (Beretta, G.): Controlled study for prolonged chemotherapy, immunotherapy and chemotherapy plus immunotherapy as an adjuvant to surgery. Adjuvant therapies and markers of post-surgical minimal residual disease. Intermediate report presented in Paris, June 22–24, 1978
23. Fortner, J.G., MacLean, B., Mulcare, R.J.: Treatment of recurrent malignant melanoma. In *Melanoma and Skin Cancer*, McCarthy, W.H., editor. Sydney, V.C.N. Blight, 1972, p. 453
24. WHO Collaborating Centers for Evaluation of Methods of Diagnosis and Treatment of Melanoma (Bufalino, R., Cascinelli, N., Morabito, A., Preda, F.): Register: analysis of 4739 cases, 1977. [Google](#)
25. Mark A Moore, PhD, Decellularization of Human Dermis Using MATRACELL[®] Technology: Process, Preclinical Studies, and Medical Applications.
26. Dr. Eran Rosines, Dr. Qishan Lin, Analysis of the Acellular Matrix, Growth Factors, and Cytokines Present in DermACELL[®] Advanced Wound Management
27. Mark Moore, PhD, Kristen Trost, MSS, Ralph Powers, DDS, Ensuring the Safety of Allograft Dermal Tissue