High prevalence of skin cancers and actinic keratoses in lung transplant recipients

DOI:
10.1016/j.healun.2017.11.016

Document Version
Accepted author manuscript

Link to publication record in Manchester Research Explorer

Citation for published version (APA):

Published in:
Journal of Heart and Lung Transplantation

Citing this paper
Please note that where the full-text provided on Manchester Research Explorer is the Author Accepted Manuscript or Proof version this may differ from the final Published version. If citing, it is advised that you check and use the publisher's definitive version.

General rights
Copyright and moral rights for the publications made accessible in the Research Explorer are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

Takedown policy
If you believe that this document breaches copyright please refer to the University of Manchester’s Takedown Procedures [http://man.ac.uk/04Y6Bo] or contact uml.scholarlycommunications@manchester.ac.uk providing relevant details, so we can investigate your claim.
HIGH PREVALENCE OF SKIN CANCERS AND ACTINIC KERATOSES IN LUNG TRANSPLANT RECIPIENTS

Azadeh Sahebian\textsuperscript{1,2}, Nirmala Pandeya\textsuperscript{3}, Daniel C Chambers\textsuperscript{4,5}, H Peter Soyer\textsuperscript{1,2}, Adele C Green\textsuperscript{6,7}

\textsuperscript{1}Dermatology Research Centre, The University of Queensland; The University of Queensland Diamantina Institute, Translational Research Institute, Brisbane, Australia; \textsuperscript{2}Department of Dermatology, Princess Alexandra Hospital, Brisbane, Australia; \textsuperscript{3}School of Public Health, The University of Queensland, Brisbane, Australia; \textsuperscript{4}Queensland Lung Transplant Service, The Prince Charles Hospital, Brisbane, Australia; \textsuperscript{5}School of Medicine, The University of Queensland, Brisbane, Australia; \textsuperscript{6}Population Health Department, QIMR Berghofer Medical Research Institute, Brisbane, Australia; \textsuperscript{7}Cancer Research UK Manchester Institute and Institute of Inflammation and Repair, University of Manchester, Manchester, United Kingdom

ORCID:
H. Peter Soyer: 0000-0002-4770-561X
Adele C. Green: 0000-0002-2753-4841

Corresponding author:
Adele C Green, QIMR Berghofer Medical Research Institute, Cancer and Population studies, Brisbane, QLD, Australia; T: +61 7 3362 0235; F: +61 7 3845 3503
Adele.Green@qimrberghofer.edu.au

Word count= 1000
Increased risk of malignancy is a major complication of lung transplantation. Keratinocyte cancers (KC), namely cutaneous squamous cell cancers (SCCs) and basal cell cancers (BCCs), are a particular risk, and cause high morbidity and some mortality among lung transplant recipients (LTRs) (1). Actinic keratoses (AKs) are potentially premalignant skin lesions strongly associated with SCC although the AK burden in LTR has not been specifically measured. The heightened risks of skin cancer and AKs are due to two main factors, high-dose immunosuppression and sun exposure, but their relative contributions are unknown. To assess the burden of skin cancer and AKs in LTRs at a point in time in a high sun exposure region, we therefore conducted a cross-sectional study and estimated the prevalence of skin cancer and AKs in LTRs in the state of Queensland, Australia.

The study was approved by Institutional Ethics Committees. We invited LTRs aged over 17 to participate during routine clinics (July 2012-July 2013) at The Prince Charles Hospital, the only treatment centre for LTR in the state. Participants provided information via a self-completed questionnaire (2) about standard risk factors for skin cancer (age, sex, skin phenotype, past sun exposure, smoking, history of previous skin cancer). They underwent a full skin examination by a dermatology-trained physician who mapped the location of all lesions clinically suspicious for malignancy and of AKs, and visually graded the severity of neck solar elastosis. LTRs were referred to their doctor for histopathologic diagnosis of suspicious lesions (usually available within 3 months). Information on date of transplant, current immunosuppressive therapy and use of voriconazole (risk factor for SCC (3)) were obtained by medical chart review. We calculated period prevalences of skin cancers based on malignant lesions histopathologically-confirmed within 3 months of examination. Site-specific AK counts were summed to give total AK prevalence at baseline. Logistic regression was used to assess associations with the composite outcome of having either skin cancer or >5AKs present on the skin in a multivariable model adjusting for all potential confounding factors.

Of 191 LTRs aged over 17 in Queensland, 159 were approached and 122 (77%) participated with informed, written consent (Supp Figure). Non-participants were similar to participants in age and sex but had longer mean time from transplantation (6.8 years vs 4.7 years). Average age of LTRs
studied was 50 years and 93% used triple therapy immunosuppression (calcineurin inhibitor, antimetabolite, corticosteroid). Half the participants had fair skin, 22% had moderate or severe solar elastosis and 44% reported past skin cancer (15% pre-transplant).

There were 44 confirmed skin cancers (7 BCCs, 20 SCCs, 16 intraepithelial carcinomas (IECs), 1 melanoma) in the 3-month baseline period and 9 LTRs (7%) had multiple (33) BCCs, SCCs or IECs present on the skin (Table 1). Corresponding prevalences of BCC, SCC and total skin cancer were 5%, 8% and 13% respectively. Of the 122 LTRs examined, 74 (61%) had one or more AKs present, and a third had >5 AKs. In a multivariable model adjusted for all potential confounding factors, we found that categories of increasing severity of solar elastosis of the neck and a history of previous KC were associated with having either skin cancer or >5AKs present on the skin at examination (Table 2). While odds of KC or >5AK almost doubled with voriconazole treatment (adjusted OR 1.8) this was not statistically significant. Odds of KC or >5AK increased with increasing transplant duration (Table 2) (and this was true for AK and KC separately, though sparse numbers limited the precision of estimated ORs).

One previous study of KC in LTRs showed similar transplant-related risk factors but it assessed incidence over time (rather than prevalence that indicates necessary outlay of treatment services) and did not assess sun exposure (1). No other generalizable, population-based data on skin cancer and AK prevalence and role of sun exposure are available for LTRs, though small numbers of prevalent KC limited our statistical power. The composite endpoint may have diluted KC associations. Skin cancer prevalence of 13% in Queensland LTRs is far more than the 5% in the general population aged 20-69 (4). The 61% AK prevalence alone is notably higher than 42% with AK or skin cancer in referred French organ transplant recipients (5). Still our figures may substantially underestimate the burden in Queensland LTRs because of longer transplant duration in non-participants, and because the study did not count as outcome events the non-histologically confirmed skin tumors that were destructively treated (eg cryotherapy and 5-flurouracil).
Our results provide further evidence of the association between sun exposure and skin carcinogenesis in LTRs and we therefore vigorously encourage LTRs’ use of sun protection. We also support the establishment of dedicated, specialist skin cancer surveillance clinics and curative treatment of AKs.

Financial conflict of interest disclosure

All authors declare no conflict of interest. This study was funded in part by the EPIDERM Foundation, who played no role in the collection of data, its analysis and interpretation, or in the right to approve or disapprove publication of the finished manuscript

Acknowledgements

Our thanks to Dr Peter Hopkins, Dr Conrad Morze, Michelle Grant and Stephanie Yerkovich for their assistance with data collection
References: