Melanoma imaging and diagnosis: What does the future hold?



Lena von Schuckmann, Leith Banney, H Peter Soyer

Background

In Australia, artificial intelligence (AI) is increasingly being used in the field of melanoma diagnosis. Early diagnosis is arguably the most important prognostic factor for melanoma survival. The use of digital monitoring of naevi, especially dysplastic naevi, might reduce the number of biopsies needed in managing patients at risk of melanoma, especially in patients with high naevi counts.

Objective

This article discusses advances in imaging and early diagnosis including the use of Al in this process.

Discussion

The benefits of performing biopsies must be balanced with the potential to cause harm. Whole-body imaging can assist with more accurate detection of changing lesions and enable clinicians to focus on lesions where change is detected. IN AUSTRALIA, artificial intelligence (AI) is increasingly being used in the field of melanoma diagnosis. Early diagnosis is arguably the most important prognostic factor for melanoma survival.1 A common clinical challenge is identifying changing or unstable pigmented lesions, especially in patients with high naevi counts.2 The benefits of performing biopsies for suspicious pigmented lesions must be balanced with the potential to cause harm.³ AI, in the form of 3D total body imaging with dermoscopy, has been used in research settings in Australia for many years and is now starting to be implemented into clinical settings to assist clinicians in the diagnosis of melanoma.4

3D total body imaging with dermoscopy creates an Avatar of the patient's skin and body map of naevi.5 These machines have in-built AI that have developed AI mechanisms to: (i) determine whether a lesion has reached the threshold for a diagnostic biopsy; and (ii) identify dermatoscopic changes by comparing serial photographs sequentially.4 It has been demonstrated that these AI technologies are able to diagnose a pigmented lesion with an accuracy comparable to a dermatologist.6 Sequential photographs have been shown to increase the sensitivity and specificity of diagnosing melanomas.7 The main challenges with this technology are difficult-to-image

areas (eg acral surfaces, groin, skin folds) and the reliance on melanographers for performing dermoscopy.

A study conducted by an Austrian– Australian research team showed that AI phone applications for categorising pigmented lesions as benign or malignant, matched the diagnostic accuracy of experts and outperformed less-experienced physicians.⁸ In this study, the AI applications more commonly recommended removal of benign lesions, indicating that clinician judgement remains essential for treatment planning to avoid excessive false-positive findings and overtreatment.

Future applications

The future of AI in the diagnosis and treatment of melanoma is expected to bring significant advancements and improvements.

Screening

AI algorithms for differentiating between benign and malignant pigmented lesions are expected to become more sophisticated, allowing for even greater accuracy in diagnosing melanoma. These advancements will likely come from improved machine learning models that can analyse a wider range of data and incorporate information on patient history and even genetics. This technology will likely also become available for the interpretation of lesions imaged with reflectance confocal microscopy (RCM), which is a non-invasive technique enabling visualisation of the epidermis and papillary dermis. Although the resolution is almost comparable to conventional histology, the use of RCM is limited to thin lesions.

As currently used in mammography, AI can exclude normal lesions and reduce the number of lesions needing diagnosis by clinicians. This could greatly reduce the workload of melanoma screening programs and allow clinicians to concentrate on challenging lesions. High-quality imaging also facilitates obtaining second opinions for more difficult lesions.

Early detection

Wearable devices equipped with sensors and AI capabilities aim to detect abnormal biometric markers in enzymes, antibodies, cell receptors or organelles for early detection of diseases such as cancer. Use of wearable biosensors could continuously monitor skin health, alerting users to any changes that might warrant professional evaluation. This would be particularly beneficial for individuals at high risk of melanoma.

Personalised treatment plans

AI might be used to develop more personalised treatment plans for melanoma patients. By using algorithms, AI might help identify the most effective diagnostic tests and treatments for a case. This might incorporate tumour information (histopathology, molecular and genetic factors), as well as patient characteristics (ie age, sex, comorbidities, medication use).

Early diagnosis of recurrence

AI in the form of wearable biosensors, might also be increasingly used for early detection of recurrence before visible symptoms appear and/or are detectable with imaging. Algorithms could also analyse historical data and identify patterns or risk factors that predispose individuals to recurrence, leading to closer monitoring and earlier interventions.

Automation in pathology

AI might automate certain aspects of pathology, such as the initial screening and diagnosis of biopsy specimens. More advanced models might assist with diagnosing borderline pigmented lesions with higher accuracy. This might reduce time delay from biopsy to diagnosis and increase the sensitivity and specificity of melanoma diagnosis.

Telemedicine and remote diagnostics

With the advancement of AI, remote diagnosis of melanoma might become achievable, where melanoma screening is reliably performed via 3D body imaging or AI smartphone applications. Patients in remote areas could receive timely diagnoses without needing to travel to specialised centres.⁹

Conclusion

AI in Australia is increasingly utilised for the diagnosis of melanoma. The most effective use of AI in melanoma diagnosis and treatment now, and in the near future, involves a collaborative approach, where AI supplements the skills and knowledge of healthcare professionals, rather than replacing them.¹⁰ As AI becomes more integrated into healthcare, there will be an increased focus on the regulation of AI tools and ethical considerations, such as patient privacy and the decision-making process in AI-driven technology.⁴

Authors

Lena von Schuckmann BSc, MBBS, MPH, PhD, FRACGP, FACD, Senior Clinical Lecturer, Frazer Institute, The University of Queensland, Dermatology Research Centre, Brisbane, Qld; Senior Medical Officer, Dermatology Department, Princess Alexandra Hospital, Brisbane, Qld; Senior Medical Office, Dermatology Department, Sunshine Coast University Hospital, Sunshine Coast, Qld; Private Practice, Spring Hill, Brisbane and Samford Valley, Moreton Bay, Qld

Leith Banney MBBS, FACD, Senior Clinical Lecturer, Griffith University School of Medicine, Sunshine Coast University Hospital, Sunshine Coast, Qld; Head of Department, Sunshine Coast University Hospital, Sunshine Coast, Qld; Private practice, Dermatology Solutions (online teledermatology service)

H Peter Soyer MD, FACD, FAHMS, Professor in Dermatology, Frazer Institute, The University of Queensland, Dermatology Research Centre, Brisbane, Qld; Senior Staff Specialist, Princess Alexandra Hospital, Brisbane, Qld

Competing interests: HPS is a shareholder of MoleMap NZ Limited and an e-derm consult for GmbH and undertakes regular teledermatological reporting for both companies. HPS is also a Medical Consultant for Canfield Scientific Inc and a Medical Advisor for First Derm.

Funding: None.

Provenance and peer review: Commissioned, externally peer reviewed.

Correspondence to:

Leith.Banney@health.qld.gov.au

References

- Baade PD, Whiteman DC, Janda M, et al. Long-term deaths from melanoma according to tumor thickness at diagnosis. Int J Cancer 2020;147(5):1391–96. doi: 10.1002/ijc.32930.
- Argenziano G, Soyer HP. Dermoscopy of pigmented skin lesions—A valuable tool for early diagnosis of melanoma. Lancet Oncol 2001;2(7):443–49. doi: 10.1016/S1470-2045(00)00422-8.
- Glasziou PP, Jones MA, Pathirana T, Barratt AL, Bell KJ. Estimating the magnitude of cancer overdiagnosis in Australia. Med J Aust 2020;212(4):163–68. doi: 10.5694/mja2.50455.
- Primiero CA, Rezze GG, Caffery LJ, et al. A narrative review: Opportunities and challenges in artificial intelligence skin image analyses using total body photography. J Invest Dermatol 2024:S0022-202X(23)03123-8. doi: 10.1016/j. jid.2023.11.007. Epub ahead of print.
- Koh U, Cust AE, Fernández-Peñas P, et al. ACEMID cohort study: Protocol of a prospective cohort study using 3D total body photography for melanoma imaging and diagnosis. BMJ Open 2023;13(9):e072788. doi: 10.1136/bmjopen-2023-072788.
- Cerminara SE, Cheng P, Kostner L, et al. Diagnostic performance of augmented intelligence with 2D and 3D total body photography and convolutional neural networks in a high-risk population for melanoma under real-world conditions: A new era of skin cancer screening? Eur J Cancer 2023;190:112954. doi: 10.1016/j.ejca.2023.112954.
- Menzies SW, Emery J, Staples M, et al. Impact of dermoscopy and short-term sequential digital dermoscopy imaging for the management of pigmented lesions in primary care: A sequential intervention trial. Br J Dermatol 2009;161(6):1270-77. doi: 10.1111/j.1365-2133.2009.09374.x
- Menzies SW, Sinz C, Menzies M, et al. Comparison of humans versus mobile phone-powered artificial intelligence for the diagnosis and management of pigmented skin cancer in secondary care: A multicentre, prospective, diagnostic, clinical trial. Lancet Digit Health 2023;5(10):e679–91. doi: 10.1016/S2589-7500(23)00130-9.
- Drabarek D, Habgood E, Ackermann D, et al. Perspectives and experiences of patient-led melanoma surveillance using digital technologies from clinicians involved in the MEL-SELF pilot randomized controlled trial: Qualitative interview study. JMIR Dermatol 2022;5(4):e40623. doi: 10.2196/40623.
- 10. Tschandl P, Rinner C, Apalla Z, et al. Humancomputer collaboration for skin cancer recognition. Nat Med 2020;26(8):1229–34. doi: 10.1038/s41591-020-0942-0.

correspondence ajgp@racgp.org.au