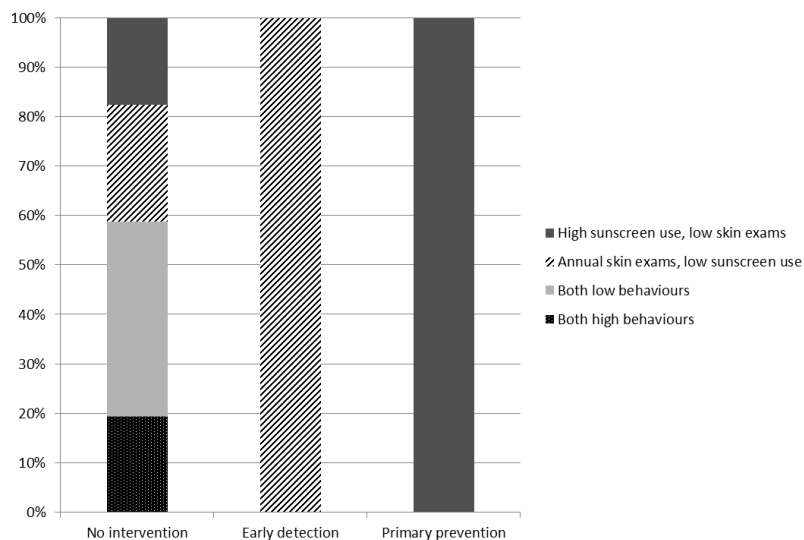


SUPPLEMENTARY FILE

Contents

Figure S1: Proportions of health behaviours for each strategy	2
Further details of model inputs	3
Model validation.....	4
Figure S2: Tornado Plot: Incremental cost-effectiveness ratio, primary prevention vs early detection	4
Figure S3: Scatterplot of incremental costs and QALYs for Primary prevention versus Early detection.	5
Figure S4: Scatterplot of mean costs and QALYs for all three strategies	5
Figure S5: Cost-effectiveness acceptability curve for 3 strategies	6
Data inputs	7
Distribution of melanoma stage	7
Melanoma mortality	7
Incidence of melanoma	7
Incidence of Keratinocyte cancers.....	8
Incidence of Multiple keratinocyte cancers	8
Background mortality	9
References	10

Figure S1: Proportions of health behaviours for each strategy



Definitions: high sunscreen use = >50% time outdoors, low sunscreen use <50% time outdoors, skin exams – any clinical skin exam within past 3 years

Early detection strategy – skin exams annually + low sunscreen use

Primary prevention strategy – High sunscreen use + low skin exams

No intervention – mix of behaviours according to QSkin analysis of n=38638

In the 'No intervention' strategy, the baseline behaviours of sunscreen use and skin exams were estimated and represent the real-world scenario of current 'no intervention'. Using QSKIN data, the behaviour probabilities were 17% for high sunscreen use and no skin exams, 24% for low sunscreen use and skin exams, 19% for both high behaviours, and 40% for both low behaviours. Under the early detection strategy, 'skin exams and low sunscreen use' was 100% and in the primary prevention scenario, the 'high sunscreen use and low skin exams category probability was 100%. The behaviours were not explicitly modelled as data inputs in the model but rather the consequences from them.

Further details of model inputs

The estimate for solar keratosis incidence was 24% [18] (used for 'benign skin lesions' health state). The effect of increased skin exams was a 50% increase in all skin malignancies as seen in the Skin Awareness Trial [1]. This increase was applied to the age-specific Queensland population incidence of melanoma, KCs and benign lesions [19, 20]. The effect of early detection is through an 18% reduction in thicker melanomas (>1mm) over three years [9].

A weighted average was calculated for thicker melanomas which combined stages II, III and IV. Five year survival for stage I melanoma was 98% versus 32% for stage IIID (the latter was assumed the same for stage IV in our model). This is supported by increasing metastatic mortality rates from pharmacotherapy trial evidence at 4-years for ipilimumab and nivolumab (53% survival). Ten year survival for stage I melanoma was 95% versus 24% for stage IIID[7]. We did not apply excess mortality to individuals with multiple melanomas as evidence is unclear and similarly, for KCs, deaths from KCs are uncommon in the general population and are lower than background mortality rates [22]. Background mortality rates were based on life tables for the Queensland population [8] to reflect that individuals may die at any age from any cause

Health Utilities. Utility scores (similar to health-related quality of life scores) were assigned to the health states where appropriate. Scores ranged from 0 (indicating death) to 1 (indicating best possible health) and applied to survival to generate 'quality-adjusted life years' (QALYs). Utility scores for melanoma stages, including patients with advanced melanoma receiving targeted therapies, were obtained from a meta-analysis (Table 1) [13]. Robust evidence on health utilities for patients with KC or benign skin lesions is limited, but there is an appreciable quality of life effect where some individuals face disfiguring, multiple cancers, anxiety and other symptoms [23]. From Seidler *et al.* (2009), we assigned a utility score of 0.984 to patients with KCs (0.95 and 0.99 in sensitivity analysis) and a utility reduction of -0.03 each time an individual had an additional KC [24].

Resource Use and Costs. Short-term costs associated with implementing promotional campaigns or intervention programs were excluded. Individuals having any clinical skin exam were assumed to visit their primary physician once per year. Costs were included for resources consumed in the routine diagnosis, treatment and follow-up of skin cancers and skin lesions in Australia [14, 15] and included new therapies for treating advanced melanoma [15] (Table 1).

Model validation

In the 'no intervention' strategy, the model predicted 70.5% were alive at age 80 which aligns to 67.4% of 50-year olds in the general population[8]. The modelled 10-year survival in the no intervention strategy was 96.4% compared with 97.4% survival in Queenslanders with thin melanoma from Green *et al.* 2012.

Figure S2: Tornado Plot: Incremental cost-effectiveness ratio, primary prevention vs early detection

Simple expected values analysis

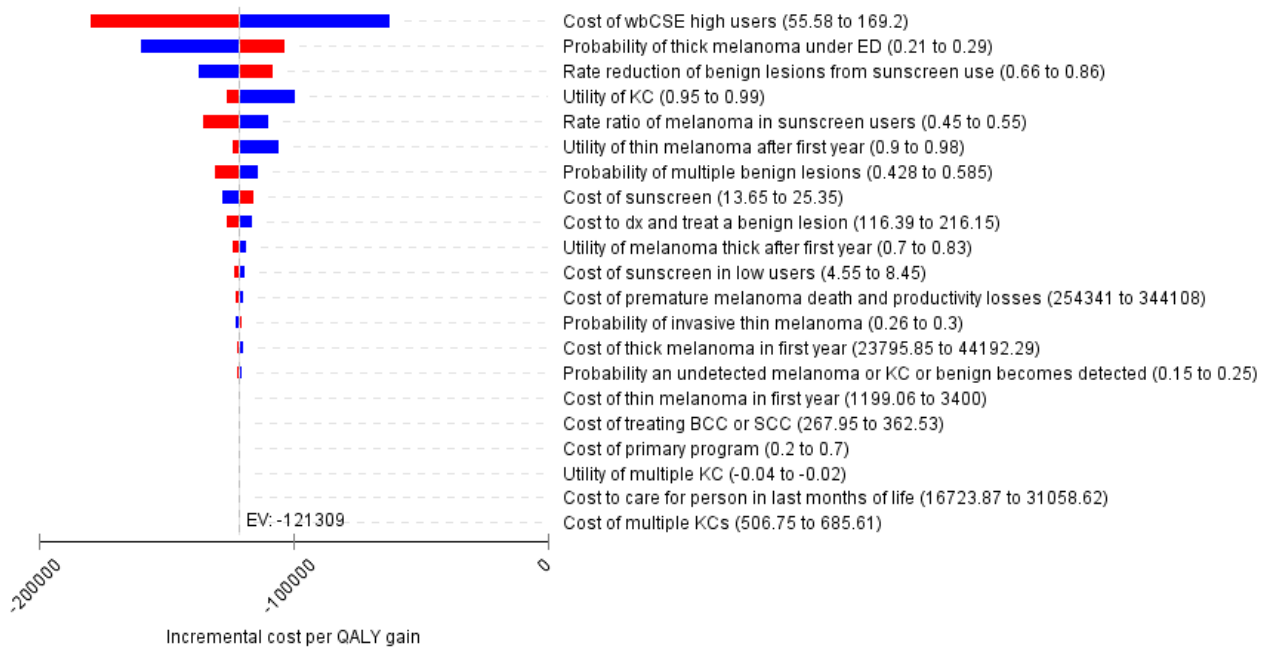
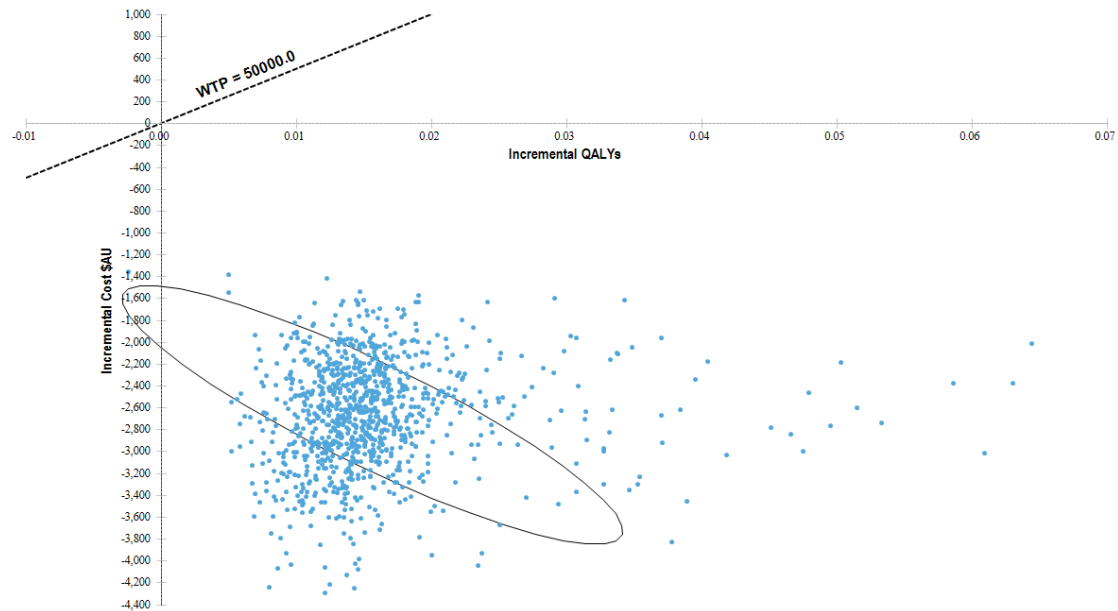


Figure S3: Scatterplot of incremental costs and QALYs for Primary prevention versus Early detection.



Note: Each dot represents an incremental cost and incremental QALY pairing, using the assigned distributions around each model parameter, selected randomly during 5000 iterations. Dots falling below the diagonal line (the willingness-to-pay threshold of AU\$50,000 per QALY) are considered cost-effective. The proportion of simulations considered cost-effective is 100%. The oval is the 95% ellipse.

Figure S4: Scatterplot of mean costs and QALYs for all three strategies

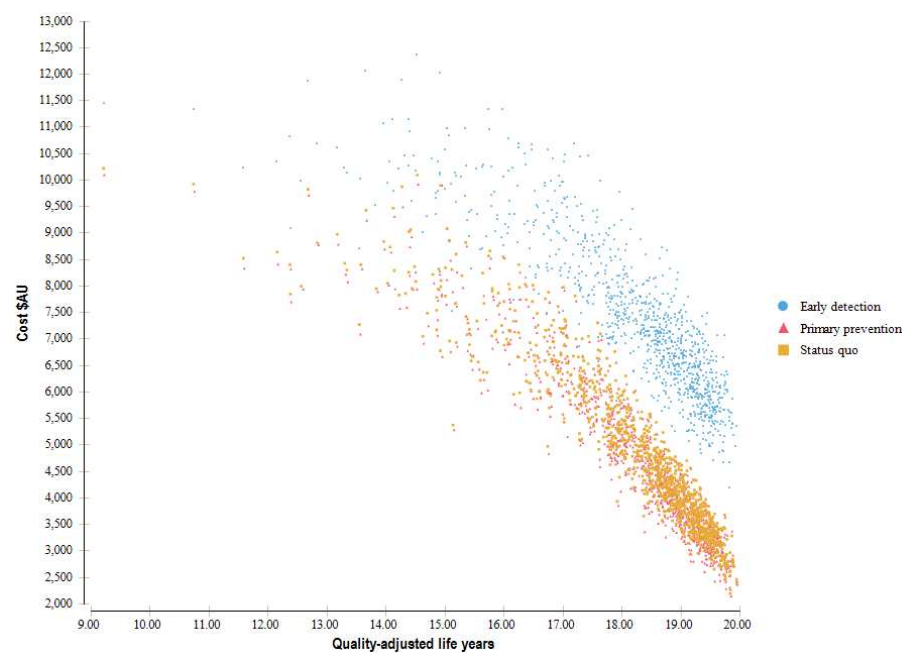
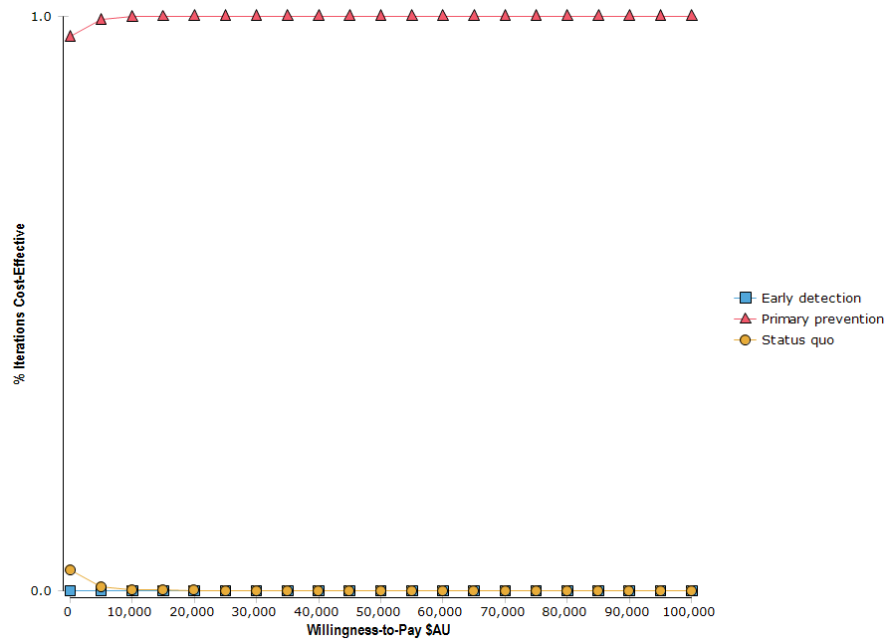
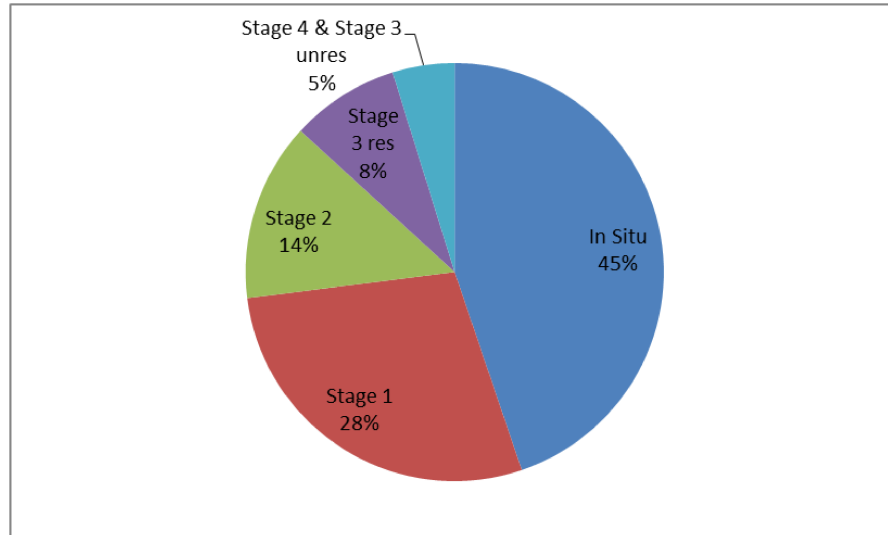


Figure S5: Cost-effectiveness acceptability curve for 3 strategies



Data inputs

Distribution of melanoma stage



Sources: Aitken 2018[5] & NSW data

Melanoma mortality

- a) For invasive melanomas, thin melanomas were Stage I, thicker melanomas were Stages II-IV and weighted by stage II (60%), III (22%), IV (18%). Mortality rate was the complement of survival and rate to probability calculations applied. Stage IIID in Gershenwald 2017 was assumed to be the same for Stage IV.

Year since diagnosis	Thin (0<1mm)	Thicker (>1mm)
5	0.02	0.233
10	0.05	0.301

Source: Gershenwald 2017 [7]

- b) For in situ melanomas – no mortality possible, background mortality rates applied

Incidence of melanoma

-in Queenslanders by Age, all persons, rate to prob calculation

Age	Prob of mel
4	0.00000100
9	0.00000100
14	0.00000900
19	0.00005000
24	0.00011799

29	0.00025897
34	0.00035594
39	0.00047189
44	0.00060582
49	0.00077770
54	0.00098551
59	0.00118829
64	0.00170255
69	0.00210877
74	0.00243503
79	0.00278013
84	0.00308822
100	0.00297856

Source: Queensland Cancer Statistics On-Line, 2017. Viertel Cancer Research Centre, Cancer Council Queensland (qcsol.cancerqld.org.au). Based on data released by the Queensland Cancer Registry (1982-2014; released January 2017).[3]

Incidence of Keratinocyte cancers

-in Queenslanders by Age, all persons, rate to prob calculation

Age	Incid KC
19	0.0027
39	0.0180
59	0.0565
79	0.0709
99	0.0900

Source: Pandeya 2017[2], Age 99 was estimated

Incidence of Multiple keratinocyte cancers

-in Queenslanders by Age, all persons, rate to prob calculation

Index	Incid multiple KC
20	0.0007
40	0.0057
60	0.0246
80	0.0342

Source: Pandeya 2017[2]

Background mortality

These were for Queenslanders as reported in government life tables for 2015-2017.

Index	All
0	0.004008
1	0.000289
2	0.000180
3	0.000150
4	0.000125
5	0.000105
6	0.000090
7	0.000080
8	0.000075
9	0.000075
10	0.000075
11	0.000080
12	0.000100
13	0.000125
14	0.000160
15	0.000210
16	0.000264
17	0.000329
18	0.000373
19	0.000413
20	0.000442
21	0.000462
22	0.000477
23	0.000497
24	0.000512
25	0.000527
26	0.000542
27	0.000566
28	0.000591
29	0.000621
30	0.000661
31	0.000706
32	0.000746
33	0.000801

34	0.000850
35	0.000910
36	0.000960
37	0.001015
38	0.001074
39	0.001144
40	0.001224
41	0.001313
42	0.001408
43	0.001508
44	0.001612
45	0.001717
46	0.001831
47	0.001966
48	0.002105
49	0.002255
50	0.002434
51	0.002623
52	0.002837
53	0.003080
54	0.003343
55	0.003626
56	0.003924
57	0.004256
58	0.004623
59	0.005020
60	0.005441
61	0.005876
62	0.006356
63	0.006886
64	0.007489
65	0.008167
66	0.008949
67	0.009823
68	0.010806

69	0.011895
70	0.013126
71	0.014517
72	0.016088
73	0.017877
74	0.019853
75	0.022084
76	0.024636
77	0.027555
78	0.030891
79	0.034707
80	0.039141
81	0.044251
82	0.050041
83	0.056585
84	0.064097
85	0.074709
86	0.084204
87	0.094600
88	0.105923
89	0.118217
90	0.131459
91	0.145525
92	0.160324
93	0.175518
94	0.190727
95	0.203434
96	0.217400
97	0.231974
98	0.246855
99	0.264965
100	0.281137

Source: Life Tables
Queensland[8]

References

1. Janda M, Youl P, Neale R, *et al.* Clinical skin examination outcomes after a video-based behavioral intervention: analysis from a randomized clinical trial. *JAMA Dermatol.* 2014;150(4):372-9. doi: 10.1001/jamadermatol.2013.9313.
2. Pandeya N, Olsen CM, Whiteman DC. The incidence and multiplicity rates of keratinocyte cancers in Australia. *Med J Aust.* 2017;207(8):339-343.
3. Queensland Cancer Statistics On-Line. *Viertel Cancer Research Centre, Based on data released by the Queensland Cancer Registry (1982-2014 released January 2017).* qcsol.cancerqld.org.au
4. Youlden DR, Baade PD, Soyer HP, *et al.* Ten-Year Survival after Multiple Invasive Melanomas Is Worse than after a Single Melanoma: a Population-Based Study. *J Invest Dermatol.* 2016;136(11):2270-2276. doi: 10.1016/j.jid.2016.03.014. Epub 2016 Mar 24.
5. Aitken JF, Youlden DR, Baade PD, *et al.* Generational shift in melanoma incidence and mortality in Queensland, Australia, 1995-2014. *Int J Cancer.* 2018;142(8):1528-1535. doi: 10.1002/ijc.31141. Epub 2017 Nov 21.
6. Aitken JF, Janda M, Elwood M, *et al.* Clinical outcomes from skin screening clinics within a community-based melanoma screening program. *J Am Acad Dermatol.* 2006;54(1):105-14. doi: 10.1016/j.jaad.2005.08.072. Epub 2005 Nov 28.
7. Gershenwald JE, Scolyer RA, Hess KR, *et al.* Melanoma staging: Evidence-based changes in the American Joint Committee on Cancer eighth edition cancer staging manual. *CA Cancer J Clin.* 2017;67(6):472-492. doi: 10.3322/caac.21409. Epub 2017 Oct 13.
8. Australian Bureau of Statistics. *Life Tables 2015-2017.* In. Canberra: Commonwealth of Australia; 2018.
9. Aitken JF, Elwood M, Baade PD, *et al.* Clinical whole-body skin examination reduces the incidence of thick melanomas. *Int J Cancer.* 2010;126(2):450-8. doi: 10.1002/ijc.24747.
10. Green AC, Williams GM, Logan V, *et al.* Reduced melanoma after regular sunscreen use: randomized trial follow-up. *J Clin Oncol.* 2011;29(3):257-63. doi: 10.1200/JCO.2010.28.7078. Epub 2010 Dec 6.
11. van der Pols JC, Williams GM, Pandeya N, *et al.* Prolonged prevention of squamous cell carcinoma of the skin by regular sunscreen use. *Cancer Epidemiol Biomarkers Prev.* 2006;15(12):2546-8. doi: 10.1158/1055-9965.EPI-06-0352. Epub 2006 Nov 28.
12. Seidler AM, Bramlette TB, Washington CV, *et al.* Mohs versus traditional surgical excision for facial and auricular nonmelanoma skin cancer: an analysis of cost-effectiveness. *Dermatol Surg.* 2009;35(11):1776-87.
13. Tran AD, Fogarty G, Nowak AK, *et al.* A systematic review and meta-analysis of utility estimates in melanoma. *Br J Dermatol.* 2018;178(2):384-393. doi: 10.1111/bjd.16098. Epub 2018 Jan 17.
14. Gordon LG, Elliott TM, Olsen CM, *et al.* Multiplicity of skin cancers in Queensland and their cost burden to government and patients. *Aust N Z J Public Health.* 2018;42(1):86-91. doi: 10.1111/1753-6405.12738. Epub 2017 Nov 22.
15. Elliott TM, Whiteman DC, Olsen CM, *et al.* Estimated Healthcare Costs of Melanoma in Australia Over 3 Years Post-Diagnosis. *Appl Health Econ Health Policy.* 2017;15(6):805-816. doi: 10.1007/s40258-017-0341-y.
16. Reeve R, Srasuebku R, Langton JM, *et al.* Health care use and costs and the end of life: a comparison of elderly Australian decedents with and without a cancer history. *BMC Palliative Care.* 2018;17(1):DOI 10.1186/s12904-017-0213-0
17. Carter HE, Schofield DJ, Shrestha R. The Productivity Costs of Premature Mortality Due to Cancer in Australia: Evidence from a Microsimulation Model. *PLoS One.* 2016;11(12):e0167521. doi: 10.1371/journal.pone.0167521. eCollection 2016.

18. Darlington S, Williams G, Neale R, *et al*. A randomized controlled trial to assess sunscreen application and beta carotene supplementation in the prevention of solar keratoses. *Arch Dermatol*. 2003;139(4):451-5. doi: 10.1001/archderm.139.4.451.
19. (AIHW) AloHaW. ACIM (Australian Cancer Incidence and Mortality) Books. In. Canberra: AIHW; 2012.
20. National Cancer Control Initiative. The 2002 national non-melanoma skin cancer survey. A report by the NCCI Non-melanoma Skin Cancer Working Group. . In: Staples MP, (ed). Melbourne: NCCI 2003.
21. Helgadottir H, Tuominen R, Olsson H, *et al*. Cancer risks and survival in patients with multiple primary melanomas: Association with family history of melanoma and germline CDKN2A mutation status. *J Am Acad Dermatol*. 2017;77(5):893-901. doi: 10.1016/j.jaad.2017.05.050. Epub 2017 Aug 14.
22. Australian Institute of Health and Welfare (AIHW). Australian Cancer Incidence and Mortality (ACIM) books. In. . Canberra: Australian Institute of Health and Welfare; 2017.
23. Gaulin C, Sebaratnam DF, Fernandez-Penas P. Quality of life in non-melanoma skin cancer. *Australas J Dermatol*. 2015;56(1):70-6. doi: 10.1111/ajd.12205. Epub 2014 Sep 8.
24. Seidler AM, Bramlette TB, Washington CV, *et al*. Mohs versus traditional surgical excision for facial and auricular nonmelanoma skin cancer: an analysis of cost-effectiveness. *Dermatol Surg*. 2009;35(11):1776-87. doi: 10.1111/j.1524-4725.2009.01291.x. Epub 2009 Sep 8.
25. van der Pols JC, Williams GM, Neale RE, *et al*. Long-term increase in sunscreen use in an Australian community after a skin cancer prevention trial. *Prev Med*. 2006;42(3):171-6. doi: 10.1016/j.ypmed.2005.10.007. Epub 2005 Dec 2.