Hospital-based multidisciplinary lung cancer care in Australia: a survey of the landscape in 2021

Fraser J H Brims,1,2 Chellan Kumarasamy,1 Jessica Nash,3 Tracy L Leong,3,4 Emily Stone,5,6 Henry M Marshall7

ABSTRACT

Introduction. Lung cancer is the leading cause of cancer death in Australia and has the highest cancer burden. Numerous reports describe variations in lung cancer care and outcomes across Australia. There are no data assessing compliance with treatment guidelines and little is known about lung cancer multidisciplinary team (MDT) infrastructure around Australia.

Methods. Clinicians from institutions treating lung cancer were invited to complete an online survey regarding the local infrastructure for lung cancer care and contemporary issues affecting lung cancer.

Results. Responses from 79 separate institutions were obtained representing 72% of all known institutions treating lung cancer in Australia. Most (93.6%) held a regular MDT meeting although recommended core membership was only achieved for 42/73 (57.5%) sites. There was no thoracic surgery representation in 17/73 (23.3%) of MDTs and surgery was less represented in regional and low case volume centres. Specialist nurses were present in just 37/79 (46.8%) of all sites. Access to diagnostic and treatment facilities was limited for some institutions. IT infrastructure was variable and most sites (69%) do not perform regular audits against guidelines. The COVID-19 pandemic has driven most sites to incorporate virtual MDT meetings, with variable impact around the country. Clinician support for a national data-driven approach to improving lung cancer care was unanimous.

Discussion. This survey demonstrates variations in infrastructure support, provision and membership of lung cancer MDTs, in particular thoracic surgery and specialist lung cancer nurses. This heterogeneity may contribute to some of the well-documented variations in lung cancer outcomes in Australia.

INTRODUCTION

Lung cancer is the fifth most commonly diagnosed cancer in Australia and the leading cause of cancer death in Australia.1 There were more than 13,200 cases diagnosed in Australia in 2020. It has a 19% 5-year survival rate4 as most patients present with incurable disease where 42% of cases are diagnosed at stage 4 (most advanced) and 29% of cases are not staged at all.2 Lung cancer accounts for 9.1% of all cancers but is responsible for 18% of deaths from all cancers in Australia and 18.6% of the cancer burden.1

Lung cancer diagnosis and care is becoming increasingly complex, with a greater array of diagnostics and therapies than ever before. There are clear, evidence-based national guidelines which outline the expected standards of care and infrastructure available for patients with lung cancer, through the Optimal Care Pathway for People with Lung Cancer (OCP) and the Lung Cancer Framework Principle of Best Practice.3 4 These guidelines recommend timely access to critical interventions such as positron emission tomography-CT (PET-CT), endobronchial ultrasound (EBUS), thoracic surgeons with expertise in lung cancer and consideration for clinical trials. Due to the complexities of lung cancer management, the guidelines also recommend that every patient be discussed by a lung cancer multidisciplinary team (MDT), regardless of the patient’s geographic location. The OCP specifies MDT membership including a ‘core’ team, who attend all/most meetings either in person or remotely, and an extended team.3

Key messages

► The adequacy, resources and infrastructure for treating lung cancer in Australia is not known and may be associated with variations in patient outcomes.

► This national survey highlights widespread deficiencies in staffing at multidisciplinary team (MDT) meetings and access to critical diagnostics and treatment modalities; just 47% of all institutions have a specialist nurse and 23% have no access to thoracic surgery at MDT.

► This service heterogeneity may contribute to some of the well-documented variations in lung cancer outcome in Australia, highlighting urgent need to address these shortfalls.
Diagnostic and treatment delays worsen outcomes for lung cancer and outcomes vary across the country. Australian and international data indicate wide variances in care between centres even in standard, established therapies. Data from two states in Australia (Victoria and New South Wales) consistently demonstrate similar findings: variations in care and outcomes between Indigenous and non-Indigenous Australians, rural and metropolitan patients, public and private centres and those of differing socioeconomic status. At least some of these variations result from differing infrastructure to investigate and treat lung cancer across Australia’s hospitals.

Identifying gaps in care and ways to improve service delivery requires data to drive changes in practice. There are no Australian data assessing compliance with the guidelines, nor national data on the quality of care or outcomes for lung cancer at the patient level. At the most basic level, understanding the ‘who’ and ‘where’ of lung cancer treatment provision is important, yet little is currently known about the infrastructure of lung cancer care around Australia. This study, therefore, aimed to identify institutions treating patients with lung cancer in Australia, and survey infrastructure support and multidisciplinary care.

METHODS
MDT lead clinicians at institutions from all states and territories across Australia were invited to complete a voluntary online Qualtrics questionnaire survey. We identified institutions and MDT Leads via multiple sources including the Lung Foundation Australia (LFA) website, national professional bodies, lung cancer clinical and research interest groups and personal networks. The survey was open from 22 January 2021 to 7 July 2021.

Only one response per institution was required to avoid duplication. Variables included: institution characteristics (public/private; metropolitan/regional); estimated numbers of new lung cancers seen annually; MDT frequency, MDT team composition (see table 1), meeting set-up (face-to-face; virtual); diagnostic and treatment services on-site, information technology (IT) support for MDT meetings; staff and infrastructure support; impact of the COVID-19 pandemic; attitudes towards a potential Australian lung cancer clinical quality registry (CQR). Responses to statement questions were structured using 0–10 Likert scales. Case volume was defined as low (<59 new cases annually) or high (≥100 new cases annually). A copy of the survey is provided as an online supplemental file.

Patient and public involvement
Consumer representation influenced the concept and planning for this project as part of a wider effort towards identifying and alleviating variations in outcomes for lung cancer in Australia.

### Statistical analysis

Adequate core data were defined as confirming the name of the institution that treats lung cancer and if the institution was regional or metropolitan, and public or private. Descriptive analysis was performed on the continuous variables of interest and frequencies were assessed for the dichotomised variables. The differences in categorical variables were analysed using the \( \chi^2 \) test. Qualitative data were analysed and grouped by themes, for instance, functioning and infrastructure issues for MDT and positive and negative impacts from the COVID-19 pandemic on the running of the MDT. Likert scales were analysed in accordance with standard approaches with the most frequent (mode) and/or median responses identified. Data were analysed using SPSS Statistics, V.24 (SPSS).

### RESULTS

The LFA MDT database identified 86 institutions with a lung cancer MDT. The survey received a total of 96 responses, 13 were excluded due to incomplete survey data (in all cases respondents restarted and completed a separate response) and 4 were removed for duplication (multiple responders from same organisation). This left a total of 79 valid responses from separate institutions across all Australian states and territories. This included 23 institutions that were not included on the LFA database. The possible denominator for lung cancer MDTs in Australia could therefore reach 109 centres, suggesting a survey response rate of at least 72.4% for Australian lung cancer MDTs.

A summary of the data is presented in table 1. The denominator for responses varied as not all questions were answered by every respondent. Fifty-five institutions identified themselves as metropolitan and 24 regional with state breakdown as follows: Victoria 26, New South Wales 23, Queensland 13, Western Australia 10, South...
Australia 2, Tasmania 2, Northern Territory 2 and Australian Capital Territory 1. Every public tertiary hospital in Australia responded. More than two thirds of sites (70.2%) estimated an annual new caseload of ≥ 100.

Five (6.3%) institutions reported no MDT meeting, one institution did not comment on MDT status. Of these, 4/5 were public, 3/5 regional, 1/5 reported a new annual caseload of 100–199 and 4/5 reported a new annual caseload < 50. Most MDTs met weekly, but around half of regional and low volume centres met fortnightly.

Personnel and MDT membership

Lung cancer nurse specialists (LCNS) were employed in 37/79 (46.8%) sites, with a slightly lower proportion employed in regional areas. For those institutions with an LCNS, the median full-time equivalent (FTE) was 0.8 (range 0.2–1.5). The total FTE for LCNS reported for all institutions across Australia was 37 (with 27 FTE within metropolitan sites). Just over a quarter (27.3%) of low case volume sites (<99/year) had less representation from thoracic surgery, nuclear medicine, specialist nurses and administrative support (table 3 and figure 1B).

Table 2 Summary of responses from each Australian institution identified as managing lung cancer patients about the functioning of the local lung cancer multidisciplinary team

<table>
<thead>
<tr>
<th></th>
<th>Total n (%)</th>
<th>Metropolitan n (%)</th>
<th>Regional n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total no institutions*</td>
<td>79</td>
<td>55</td>
<td>24</td>
</tr>
<tr>
<td>Public</td>
<td>65 (82.3)</td>
<td>42 (76.4)</td>
<td>1 (4.2)</td>
</tr>
<tr>
<td>Private</td>
<td>14 (17.7)</td>
<td>13 (23.6)</td>
<td>23 (95.8)</td>
</tr>
<tr>
<td>Has regular MDT</td>
<td>73/78 (93.6)</td>
<td>52/54 (96.3)</td>
<td>21/24 (87.5)</td>
</tr>
<tr>
<td>No regular MDT</td>
<td>5/78</td>
<td>2</td>
<td>3</td>
</tr>
</tbody>
</table>

Estimate annual lung cancer cases for all centres

<table>
<thead>
<tr>
<th>Case Number</th>
<th>Total n (%)</th>
<th>Metropolitan n (%)</th>
<th>Regional n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0–49</td>
<td>8/78 (10.3)</td>
<td>4/54 (7.3)</td>
<td>4/24 (16.7)</td>
</tr>
<tr>
<td>50–99</td>
<td>15/78 (19.2)</td>
<td>8/54 (14.5)</td>
<td>7/24 (29.2)</td>
</tr>
<tr>
<td>100–199</td>
<td>27/78 (34.6)</td>
<td>18/54 (32.7)</td>
<td>9/24 (37.5)</td>
</tr>
<tr>
<td>&gt;200</td>
<td>28/78 (35.9)</td>
<td>24/54 (43.6)</td>
<td>4/24 (16.7)</td>
</tr>
</tbody>
</table>

Specialist lung cancer nurse for all centres

<table>
<thead>
<tr>
<th></th>
<th>Total n (%)</th>
<th>Metropolitan n (%)</th>
<th>Regional n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>If yes FTE (median)</td>
<td>0.8</td>
<td>0.8</td>
<td>0.8</td>
</tr>
</tbody>
</table>

Frequency of MDT

<table>
<thead>
<tr>
<th></th>
<th>Total n (%)</th>
<th>Metropolitan n (%)</th>
<th>Regional n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Weekly</td>
<td>49/71 (69.0)</td>
<td>38/52 (73.1)</td>
<td>11/19 (42.1)</td>
</tr>
<tr>
<td>Fortnightly</td>
<td>22/71 (31.0)</td>
<td>14/52 (26.9)</td>
<td>11/19 (57.9)</td>
</tr>
</tbody>
</table>

Core MDT members per OCP guidelines

<table>
<thead>
<tr>
<th></th>
<th>Total n (%)</th>
<th>Metropolitan n (%)</th>
<th>Regional n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>None</td>
<td>15/71 (21.1)</td>
<td>13/52 (25)</td>
<td>2/19 (10.5)</td>
</tr>
<tr>
<td>Very infrequent</td>
<td>11/71 (15.5)</td>
<td>9/52 (17.3)</td>
<td>2/19 (10.5)</td>
</tr>
<tr>
<td>Yes, ad hoc</td>
<td>23/71 (32.4)</td>
<td>16/52 (30.8)</td>
<td>7/19 (36.8)</td>
</tr>
<tr>
<td>Yes, regularly</td>
<td>22/71 (31.0)</td>
<td>14/52 (26.9)</td>
<td>8/19 (42.1)</td>
</tr>
</tbody>
</table>

QA against National guidelines

<table>
<thead>
<tr>
<th></th>
<th>Total n (%)</th>
<th>Metropolitan n (%)</th>
<th>Regional n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>None</td>
<td>15/71 (21.1)</td>
<td>13/52 (25)</td>
<td>2/19 (10.5)</td>
</tr>
<tr>
<td>Very infrequent</td>
<td>11/71 (15.5)</td>
<td>9/52 (17.3)</td>
<td>2/19 (10.5)</td>
</tr>
<tr>
<td>Yes, ad hoc</td>
<td>23/71 (32.4)</td>
<td>16/52 (30.8)</td>
<td>7/19 (36.8)</td>
</tr>
<tr>
<td>Yes, regularly</td>
<td>22/71 (31.0)</td>
<td>14/52 (26.9)</td>
<td>8/19 (42.1)</td>
</tr>
</tbody>
</table>

All data presented as n=N (%) unless otherwise stated.

*The denominator for responses varied as not all questions were answered by every respondent.

FTE, (fraction of) full time equivalent; MDT, multidisciplinary team; OCP, Optimal Care Pathway; QA, quality assurance.

Figure 1 Per cent of lung cancer MDT attendance representation from subspecialty discipline in Australia, stratified by (A) metropolitan and regional institutions, and (B) high and low volume annual case numbers. MDT, multidisciplinary team.
Clinical infrastructure

All 73 MDTs provided information on availability of diagnostic and treatment services. Medical oncology and palliative care services were available on all sites, with respiratory not available at one institution. PET-CT was available on site for 56/73 (76.7%) and EBUS bronchoscopy for 58/73 (79.5%). EBUS was less available in regional areas (metropolitan vs regional 46/53 vs 12/20, \( \chi^2 \) statistic 6.38, \( p=0.012 \)). Thoracic surgery was on site for 56/73 (76.7%) institutions, external beam radiotherapy 56/73 (76.7%) and stereotactic ablative radiotherapy 48/73 (65.8%). One regional site commented on an ‘ad hoc service with minimal resources’ and a private site reported ‘fragmented care with no nurse’.

IT support

Three MDTs reported no routine data collection and 6/73 (8.2%) reported no routine IT support. For those MDTs which routinely collected data, 37/73 (46.8%) used commercial third-party software and 27/73 (34.1%) used standalone databases (eg, Microsoft Excel or Access). Over one-third of MDTs (30/73, 41.1%) reported no dedicated administrative support and data entry was performed by clinicians (consultants, fellows and registrars) in 53/73 MDTs (72.6%), or by an LCNS in 12/73 MDTs (16.4%).

Multi-institution MDTs

Collaboration across different institutions was evident: 9/73 sites commented on combined MDTs with larger institutions, 12/73 sites relying on other institutions for PET-CT and EBUS services and eight sites referring patients off site for radiation oncology and/or thoracic surgery. Further, 37/73 institutions reported having an LCNS on site, although 43/73 MDTs had LCNS representation.

Impact from the COVID-19 pandemic

There was a bimodal distribution of reported impact from the COVID-19 pandemic on lung cancer care (figure 2). Of 71, 27 (38.0%) MDTs reported instituting fully virtual meetings with 32 (45.1%) initiating hybrid meetings. For fully virtual MDTs, five commented on increased participation by MDT members and one respondent commented that they ‘can see pathology slides and radiology better’; two sites noted reduced educational opportunities and less ‘informal discussion’ among members. There was no clear pattern for sites reporting lower or higher impact by state, location or case volume. Minimal or no change as a result of the pandemic was reported by 15 institutions, with nine sites continuing with face-to-face meetings. The impact of the pandemic was most evident in case presentations and service delivery. Twelve institutions commented on delays in case presentation and two that they had noted a stage shift to more advanced
disease at presentation; one site noted an increase in referrals. Fourteen sites noted impact on service delivery in delays in access to diagnostic investigations and treatment. One institution commented that ‘patients did not attend surgery’ and one Victorian institution reported, ‘significant impact (on services) during the second wave’.

Attitudes towards a national lung cancer CQR
Regarding a national lung cancer CQR, 75 responses were received; there was strong support for a national registry (median score of 10 out of 10). Respondents indicated they felt the need for a national effort to improve lung cancer care is urgent (median score 9 out of 10).

DISCUSSION
This is the first Australia-wide survey of lung cancer services and care, representing up to 72% of all institutions known to treat lung cancer in Australia across all states and territories. Most responding sites were public and 70% reported seeing more than 100 cases per year. The results indicate important variations in the infrastructure for investigating and managing lung cancer.

No institutions had full recommended MDT membership, only 57% fulfilled recommended core membership and five institutions did not host a lung cancer MDT. Commonly missing groups included LCNSs (53.2%), MDT co-ordinators (42.5%), nuclear medicine (35.6%) and thoracic surgery (23.3%). Lung cancer MDT discussion improves a range of outcomes including staging, clinical decision making, better utilisation and uptake of treatments and survival and the quality of such meetings may be a reflection of the functioning of the wider culture of cancer care within a centre. National and international guidelines recommend MDTs in lung cancer care and, and therefore, there is a clear need to improve the membership and infrastructure of lung cancer MDTs in Australia. Mitigating these MDT gaps, we observed a degree of collaboration across sites and note the recent widespread implementation of virtual meeting technologies. Similar to how telehealth has enabled remote patient–practitioner interactions, these technologies provide a mechanism to share resources and knowledge, and increase collaboration and referral pathways between low volume and high volume centres, which ultimately should translate to better patient outcomes. In addition, as discussed below, virtual meeting technologies may foster support and mentoring of newly established MDTs.

The survey identified key gaps in on-site services and staffing. Although fewer than half of all institutions reported having an LCNS, particularly low case-volume institutions, the total FTE for all LCNS was better than previous estimates. Access to an LCNS is associated with increased treatment rates, reduced unplanned hospital admissions and improved addressing of patient needs including complex case navigation and emotional and psychological support. Specialist cancer nurses in Australia are well recognised in breast and prostate cancer with an estimated 400 breast cancer nurses and at least 50 prostate cancer nurses in Australia. Federal commitment to fund five FTE LCNS in the 2021 Budget is welcome, but considering the burden of disease from lung cancer in Australia, the need for a significant uplift in the number of specialist lung cancer nurses remains.

Nearly one-quarter of MDTs (23%) had no thoracic surgery representation despite the direct influence of surgical resection rates on survival in lung cancer. The initial National Lung Cancer Audit reports for England and Wales demonstrated considerable variation in access to thoracic surgery, with subsequent changes leading to notable improvements resection rates. Site location was associated with infrastructure variations. Collaboration across sites appears to address some gaps for MDTs without on-site access to key investigations and treatments, however, regional and low volume centres demonstrated likely clinically important differences in access to EBUS bronchoscopy and thoracic surgery. Low volume sites reported less representation at MDT from thoracic surgery, nuclear medicine, specialist nurses and administrative support, and less frequent MDT meetings. This may worsen outcomes including survival, particularly for early stage lung cancer. Time delays may also affect regional patients due to travel required for investigations and/or curative treatment.

This survey identified 23 institutions not documented in the LFA database. Of these, 70% were public hospitals, 48% low volume centres (0–49 cases a year) and 10 regional centres. This likely, at least in part, represents expanding health services within local health networks across different jurisdictions. In turn, this reflects the dynamic landscape of lung cancer care, as new MDTs are established and grow to serve changing populations. Although intermittent snapshots of MDT care are of interest and paint a broad picture, they are unlikely to drive better patient outcomes in themselves. Rapid advances in lung cancer therapies, superimposed on continuously evolving demographics, mean that a ‘real-time’ national CQR should be the goal. Increased healthcare access for patients with lung cancer is undoubtedly important, and the establishment of new MDTs is welcome. At the same time, healthcare resource planning should be cognisant of research from other areas of medicine (including lung cancer surgery) demonstrating that high case volume is associated with better outcomes across a wide range of procedures and conditions.

Although UK data demonstrates a correlation between quality of lung cancer services and 1-year survival rates, evidence to determine optimal lung cancer MDT case volumes is lacking. A recent consensus statement from Spain describes quality indicator requirements for a lung cancer MDT but does not make recommendations on case volume. This supports the urgent need to assess and ensure the quality of lung cancer services is adequate across all centres in Australia, to be able to track MDT progress.
delivery and outcomes over time and to support newly established MDTs.

The COVID-19 pandemic has driven innovations in models of care. Most centres had transitioned to either hybrid or fully virtual meetings, with only 12 sites continuing face-to-face meetings, some citing benefit and some drawbacks with these approaches. This survey was not designed to investigate the impact in detail, although the free-text comments supported other recent reports that have demonstrated a decline in diagnostic procedures performed for lung cancer along with lower rates of diagnosis, reduction in clinical trial participation and increased 30-day mortality.\(^{14,29}\)

It is perhaps surprising that many centres do not practice regular quality assurance (QA) against accepted standards of care. Fewer than one-third of MDTs reported a regular QA process. This survey identifies a number of areas which could be targeted to improve adherence to national and international standards of care for lung cancer.\(^{3,4}\) For example, dedicated MDT coordinators and commercial databases with IT support would likely improve data integrity. Clinicians clearly recognise the limitations in the data systems at present, and the value that a high-quality, uniform and linked data platform could provide.\(^{30}\) In this survey, the support for a national approach to auditing lung cancer care in Australia was strong, unanimous and urgent.

This survey has important limitations to consider. To reduce the volume of free text, the survey was designed with specific, focused questions, restricting the flexibility and depth of responses. To avoid duplication, the responses were not anonymised, which may have introduced responder bias. The survey did not specifically reference a guideline as a gold standard (but listed the full possible extended MDT membership). The second edition of the OCP for people with lung cancer was published June 2021\(^{31}\) and this survey closed on 7 July 2021, suggesting there will have been little confusion with the updated guidelines. Further, the suggested core MDT membership has not changed in the latest version.

The strength of this survey is its reach - likely representing over at least 70% of all institutions treating lung cancer in Australia. It represents an important first step towards improving lung cancer care. Multidisciplinary models of care overcome barriers to treatment, improve adherence to guidelines and are more likely to improve the quality of care for patients.\(^{31}\) The data from this survey provide a basis for review of Australian lung cancer services to improve collaboration across sites, reduce delays and improve access to timely investigation and treatment. Virtual platforms could improve MDT attendance and fill gaps in subspecialties across sites. The survey has quantified two major gaps in expertise—access to thoracic surgery support at MDTs and the lack of LCNS at almost half of all sites—both require urgent attention.

This survey of front-line clinicians who provide lung cancer care in Australia demonstrates variations in infrastructure support, provision and membership of lung cancer MDTs. This service heterogeneity may contribute to some of the well-documented variations in lung cancer outcome in Australia. Our survey respondents identified critical access issues to thoracic surgery and specialist lung cancer nurses and, for the first time, strong and unanimous support for a national CQR.

**Author affiliations**
1 Curtin Medical School, Curtin University, Perth, Western Australia, Australia
2 Department of Respiratory Medicine, Sir Charles Gairdner Hospital, Nedlands, Western Australia, Australia
3 Department of Respiratory and Sleep Medicine, Austin Health, Heidelberg, Victoria, Australia
4 Personalised Oncology Division, Walter and Eliza Hall Institute of Medical Research, Parkville, Victoria, Australia
5 Department of Respiratory Medicine, St Vincent’s Hospital, Sydney, New South Wales, Australia
6 St Vincent’s Clinical School, University of New South Wales, Sydney, New South Wales, Australia
7 Thoracic Research Centre, University of Queensland, Brisbane, Queensland, Australia

**Contributors** FB: concept, design, data collection, analysis, interpretation, writing and guarantor for overall content; CK: data analysis and writing; JN: data analysis and writing; TLL: concept, design, interpretation, writing; ES: concept, design, interpretation, writing; HMM: concept, design, interpretation, writing.

**Funding** The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

**Competing interests** None declared.

**Patient consent for publication** Not applicable.

**Ethics approval** Curtin University Human Research Ethics approved the project (reference HREC2021-0031).

**Provenance and peer review** Not commissioned; externally peer reviewed.

**Data availability statement** All data relevant to the study are included in the article or uploaded as online supplemental information.

**Supplemental material** This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

**Open access** This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/.

**ORCID iDs**
Fraser J H Brims http://orcid.org/0000-0002-6725-7535
Henry M Marshall http://orcid.org/0000-0002-9626-8014

**REFERENCES**