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Management of Early Breast Cancer at an Australian Cancer Centre During the Early Phase of COVID-19 Pandemic

Elissa J Zhang^{a,b}, Kirsty Stuart^{a,c,e}, Rina Hui^{a,c,e}, Rhiannon Mellor^e, Wei Wang^{a,e}, Verity Ahern^{a,c,e}, Farid Meybodi^{a,c}, James French^{a,c}, Elisabeth Elder^{a,c}, Meagan E Brennan^{a,c,d}

^a Westmead Breast Cancer Institute, Westmead Hospital, Westmead, NSW, Australia

^b Faculty of Medicine, University of New South Wales, Kensington, NSW, Australia

^c Westmead Clinical School, Faculty of Medicine and Health, University of Sydney, Westmead NSW, Australia

^d School of Medicine Sydney, University of Notre Dame Australia, Darlinghurst, NSW, Australia

^e Crown Princess Mary Cancer Centre, Westmead Hospital, Westmead, NSW, Australia

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ABSTRACT

Background: This study aimed to prospectively record changes to treatment for early breast cancer patients during the first wave of the COVID-19 pandemic in Australia. The purpose was to assess the impact on breast cancer outcomes and to determine the need for any mitigative actions.

Methods: The study was conducted in the breast cancer unit of a tertiary referral hospital. Patients with early (non-metastatic) breast malignancy discussed in multidisciplinary team meetings between March and June 2020 were included. Patients were newly diagnosed, post-operative or post-neoadjuvant chemotherapy. Standard treatment was defined by Westmead Breast Cancer Institute protocols and any variations related to the pandemic were recorded.

Results: In the study, 145 patients were included (median age 59 years). Pandemic-related changes to management were noted in 13 of 145 (9.0%) patients. Four patients experienced a delay to cancer treatments, four were not offered reconstructive/ symmetrisation surgical procedures, three had altered radiotherapy protocols and two patients were not offered enrolment to a clinical trial. These impacts affected the groups presenting with new cancers (n=7/86, 8.1%), post-operative cases (n=4/25, 16.0%) and post-neoadjuvant chemotherapy cases presenting for surgical planning (n=2/34, 5.9%).

Conclusion: Most patients (91.0%) received standard treatment during the first wave of the pandemic. The minor variations from institutional protocols observed in this study are unlikely to affect local control or survival in this patient cohort, but close follow-up is required. Quality of life may have been affected for four patients who had downgraded or delayed reconstructive procedures.

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* Address for correspondence:

Elissa Zhang, MD

Address: Westmead Breast Cancer Institute, Westmead Hospital, Block F/189 Cnr, Hawkesbury & Darcy Rd, Westmead NSW 2145, Australia

Email: elissa.zhang@health.nsw.gov.au

INTRODUCTION

During the early phase of the COVID-19 pandemic in Australia (March to June 2020), restrictions were placed on the treatment of early breast cancer following public health recommendations to contain the spread of the novel coronavirus.¹ In the state of New South Wales (NSW), the national population screening program



BreastScreen closed between late March and early May, and elective surgery was suspended between 25 March and 1 July of 2020.² Professional bodies in Australia and world-wide released guidelines for cancer treatment due to the pandemic (Table 1).³⁻⁶ The type and extent of breast cancer operations were minimized to reduce hospital admission days and risk of complications. Surgery for low grade DCIS, re-excision of involved margins, complex oncoplastic procedures, contralateral risk-reducing or symmetrising procedures and most breast reconstruction were recommended against.^{1, 4} Radiotherapy and chemotherapy restrictions mandated considering the risks and benefits of treatment to minimize the numbers of patients attending hospital and numbers of immunosuppressed patients in the

community. This included staged plans to reduce radiotherapy if staff illness reduced capacity, using hypofractionation, omitting boost treatments, and omitting treatment for DCIS and invasive cancer in older women.⁷ Systemic treatment recommendations included the use of G-CSF with chemotherapy, neoadjuvant or adjuvant endocrine therapy without chemotherapy for patients with low risk ER-positive/HER2-negative breast cancer, trastuzumab and paclitaxel without anthracycline-based chemotherapy for node negative small size HER2-positive cancer, delay of routine follow-up echocardiograms, and bone mineral density scans.⁸⁻¹⁰ Many consultations during treatment and follow-up were moved to telehealth.

Table 1. COVID-19 recommendations and restrictions in NSW.

Recommending body	Recommendation
	<u><i>Elective surgery during COVID-19</i></u>
NSW Health- Government (Health Directive) ¹	Surgery restricted to Category 1 (most urgent) only. No non-urgent surgery. Implemented 25/3/2020; staged return to normal from 17/4/2020. (Relevance for breast service: No breast reconstruction, risk reducing surgery, complex oncoplastic procedures or contralateral procedures during this time.)
	<u><i>Guidelines from BreastSurgANZ Council relating to the COVID-19 pandemic</i></u>
Breast Surgeons of Australia and New Zealand (Guidelines) ⁴	Surgery should be restricted to Category 1 cases. Defer surgery for low and intermediate grade DCIS. Limit complexity of surgery- consider deferral of immediate breast reconstruction and contralateral risk reducing mastectomy. Reconsider need and delivery of chemotherapy. Consider neoadjuvant endocrine therapy when surgery is delayed. Rigorous MDT* discussion and documentation are required.
	<u><i>Position statement regarding breast reconstruction during the COVID-19 pandemic</i></u>
Australian Society of Plastic Surgeons and Breast Surgeons of Australia and New Zealand (Position statement) ³	In general, breast reconstruction should be delayed. Delayed breast reconstruction and planned secondary or revision breast reconstruction should be postponed. Immediate autologous flap reconstruction for breast reconstruction should be delayed where possible. Immediate tissue expander or direct to implant reconstruction can be evaluated on a case-by-case basis.
	<u><i>Principles for Radiation Oncology Practices</i></u>
Royal Australian and New Zealand College of Radiologists (Guidance) ⁷	In summary, limiting transmission where possible, segregation of teams, segregation of well patients from those with or with suspected COVID-19, maximising communication between staff members, and for radiotherapy protocols consideration of shorter fractionation, delay in commencement of treatment, very low risk disease deferment.
	<u><i>COVID-19 Clinical Response Plan for Radiation Oncology</i></u>
Peter MacCallum Cancer Centre, Victoria, Australia (Department of Radiation Oncology, Clinical Response Plan) ⁵	Advising patients should stay away if unwell with COVID-19 symptoms, staff quarantine after exposure, reduction of transmission on hard surfaces, hand sanitisation, reduction in visitor access, increased telehealth facilities, segregation of teams, working from home where possible, avoidance of non-essential contact. The plan also discusses service impact levels correlating with proportion of working staff available.



Recommending body	Recommendation
Medical Oncology Group of Australia ⁹	<p><i>Practical Considerations for Treating Patients With Cancer in the COVID-19 Pandemic</i></p> <p>In summary, strongly consider the use of primary prophylactic G-CSF with second- and third-generation adjuvant regimens, identify patients for no requirement of chemotherapy or less intensive regimens where possible, e.g. neoadjuvant and adjuvant endocrine therapy without chemotherapy for low risk ER positive/HER2 negative cancer, paclitaxel with trastuzumab without AC for small size, node negative, HER2 positive cancer and conversion to telehealth consultations where possible.</p>

From May 2020, there was a staged easing of restrictions in public hospitals in NSW after a six-week period of 'lockdown'. Social distancing rules, mask wearing and limitations on hospital visitors remained until March 2021. Overall NSW experienced a low COVID-19 caseload, peaking at 213 cases in a single day in late March 2020 (Figure 1), with a cumulative total of 56 deaths.^{11, 12}

Nationally subsidised breast cancer related imaging including mammogram, 3D-tomosynthesis and MRI decreased by 37% from March to April 2020, before fully returning to pre-COVID service numbers in June 2020.¹³ Mastectomies, breast lesion excisions and axillary lymph node procedures in public hospitals across Australia remained stable in March to April before decreasing by 33% in May, reflecting the flow-on effects of reductions in imaging and subsequent cancer diagnoses.¹³

Impacts on early breast cancer treatments during this period may have resulted in sub-optimal treatment, placing them at risk of recurrence or other complications. For example, if a patient did not have an involved margin re-excised, or had altered radiotherapy and chemotherapy regimes, there might be a higher risk of local or distant relapse. Quality of life might be adversely affected if oncoplastic procedures or breast reconstructions were not offered. Understanding alterations to management will help to plan future care and surveillance for patients treated during this time.

The aim of this study was to prospectively document and analyze any changes to treatment for early (non-metastatic) breast cancer during the first wave of the COVID-19 pandemic in NSW, compared to evidence-based protocols.

Methods

This study was conducted at the breast cancer unit of a tertiary referral hospital (the NSW Westmead Hospital Breast Cancer Institute) between March 15 to June 15, 2020 during the most severe restrictions on hospital treatment. Inclusion criteria were: female patients who were either newly diagnosed DCIS or non-metastatic invasive cancer, post-operative, or finishing neo-adjuvant chemotherapy (NACT), who

were being discussed in MDT meetings held three times a week during the study period. Patients with benign lesions or recurrent/metastatic breast cancer were excluded. Patient demographic and cancer data were collected. Each case was allocated a category of "new cancer", "post-op cancer" and "post-NACT", corresponding to the reason for their first MDT discussion during the study period. Therefore, new cancer cases referred before the study period were classified as 'post-operative' or 'post-neoadjuvant' when re-discussed at MDT meeting. Management recommended by the MDT was recorded and any changes from local protocols ('ideal' management) caused by COVID-19 restrictions were documented.

Data were collected in an Excel spreadsheet and analysed in IBM SPSS v26. Cases where the actual (MDT-recommended) management differed from the 'ideal' management were identified, examined in detail and were flagged for recall for closer follow-up post COVID-19 restrictions easing.

Ethical Approval

The study was registered with the Australian New Zealand Clinical Trials Registry (ACTRN126-20000869976). Approval was obtained from the Western Sydney Local Health District Human Research Ethics Committee (Project ID 2020/PID00900).

Results

There were 344 new referrals to the institute (86 new malignancies; the remainder atypical or benign). This compares to 456 cases (109 new malignancies) for the same three-month period in 2019, a 24.6% reduction in total new referrals and a 21.1% reduction in new malignancy referrals.

Overall, 145 eligible cases of early breast cancer (DCIS or invasive cancer) were discussed across the MDT meetings. The demographics of the study population are shown in Table 2. All patients were female. Median age was 59 years.

All patients had core biopsies of the breast performed prior to referral or organised through the institute. The majority of patients (n=123, 84.8%) had invasive cancer, and 22 (15.2%) had DCIS on biopsy results.

Table 2. Participant and cancer demographics.

Variables		N (%)	
Age (years)	Mean	57.7 (SD 11.8)	-
Age groups (years)	30-39		11 (7.6)
	40-49		23 (15.9)
	50-59		46 (31.7)
	60-69		40 (27.6)
	70-79		20 (13.8)
	80+		5 (3.4)
Cancer Demographics		Core Biopsy (n=145) N (%)	Surgical Histology (n=127) N (%)
DCIS		22 (15.2)	15 (11.8)
Grade	Low	2 (9.1)	0
	Intermediate	11 (50.0)	10 (66.7)
	High	7 (31.8)	5 (33.3)
	Missing	2 (9.1)	
Invasive cancer		123 (84.8)	109 (86.5)
Histological Type	No specific type	88 (71.5)	77 (70.6)
	Infiltrating lobular carcinoma	18 (14.6)	18 (16.5)
	Invasive other	13 (10.6)	8 (7.3)
	Atypical **	3 (2.4)	-
	pCR	NA	6 (5.5)
	Missing	1 (0.8)	-
Histological Grade	1	17 (13.8)	23 (21.1)
	2	65 (52.8)	53 (48.6)
	3	25 (20.3)	24 (22.0)
	Missing	16 (13.0)	9 (8.3)
Receptor status	ER positive	97 (78.9)	90 (82.6)
	HER2 positive	18 (14.6)	11 (10.1)
	Triple negative	13 (10.6)	7 (6.4)

*Including invasive tubular, invasive mucinous, invasive papillary, invasive carcinoma

** Atypical including atypical apocrine, atypical hyperplasia na=not applicable

Details of presentation (reason for discussion by the MDT) are shown in Table 3. Of the sample, 86 (59.3%) were new cancer cases, 25 (17.2%) were post-operative cancer cases and 34 (23.4%) were cases returning for surgical planning after neoadjuvant chemotherapy. Of the 86 new cancer referrals, up-front surgery was recommended in 69 (80.2%) cases and neoadjuvant therapies in 17 (19.8%) cases.

COVID-related changes to treatment

Changes to management related to COVID-19 were recorded in 13 of 145 (9.0%) patients. Of these, four patients experienced a delay to cancer treatments, four were not offered their ideal reconstructive or

symmetrisation procedures, three had altered radiotherapy protocols and two were not recruited to eligible clinical trials. These impacts mostly affected new cancer patients (n=7, 54%), followed by post-operative (n=4, 31%), then post-NACT (n=2, 15%). (Table 4).

Changes to surgery

Six patients experienced COVID-related changes to surgical management:

- Two patients with intermediate grade DCIS less than 20mm in size had their surgery deferred. They showed no microinvasion on surgical



histopathology and were both treated with breast conservation surgery when restrictions were lifted. Patient C had surgery delayed by three months, and Patient E by only ten days.

- Patients B, F and L were denied immediate contralateral symmetrising breast reduction, breast

reconstruction for low grade DCIS, and contralateral prophylactic mastectomy, respectively. Delayed surgical procedures were planned. One patient requested autologous reconstruction, which was changed to implant-based reconstruction.

Table 3. Reason for discussion at MDT and treatment recommendations during COVID-19 period of restriction (N=145)

Reason and Recommendations	N (%)	
Group 1: New cancer	86 (59.3)	
Upfront surgery	69 (80.2)	
<u>Breast</u>		
- Wide local excision (WLE) +/- local flap	53 (76.8)	
- WLE + contralateral symmetrisation	1 (1.4)	
- Simple mastectomy	9 (13.0)	
- Mastectomy and immediate reconstruction	4 (5.8)	
- Excision of atypical lesions	2 (2.9)	
<u>Axilla</u>		
- Sentinel lymph node biopsy alone	54 (78.3)	
- Axillary lymph node dissection	3 (4.3)	
- No axillary surgery (DCIS)	12 (17.4)	
Neoadjuvant therapy	17 (19.8)	
- Chemo/targeted	13 (76.5)	
- Endocrine only	4 (23.5)	
COVID-related changes	7 (8.1)	See Table 4; Cases A-G
Group 2: Post-operative Cancer	25 (17.2)	
Further breast surgery	6 (24.0)	
Further axillary surgery	3 (12.0)	
COVID-related change	4 (16.0)	See Table 4; Cases H-K
Group 3: Post NACT surgical planning	34 (23.4)	
Breast		
- WLE	22 (64.7)	
- Simple mastectomy	7 (20.6)	
- Mastectomy and immediate reconstruction	5 (14.7)	
Axilla		
- Sentinel lymph node biopsy alone	14 (41.2)	
- Axillary lymph node dissection	12 (35.3)	
- Targeted axillary dissection	8 (23.5)	
COVID-related change	2 (5.9)	See Table 4; Cases L-M

**Table 4.** COVID-19 related changes to treatment

Case	Age	Category	Cancer information	Treatment Changed	Details
A	71	New cancer	pT1cN0M0 Inv NST G1 ER+PR+HER2-	Trial	Not recruited for clinical trial due to trial recruitment shut down for COVID (EXPERT candidate) * Would have considered NACT but surgery was recommended to avoid multiple trips to hospital with immunosuppression.
B	57	New cancer	pT2N0M0 Inv NST G3 Triple negative; WLE and SNB with adjuvant radiotherapy and chemotherapy	Surgery/NACT	Would have elected immediate contralateral reduction but was delayed due to COVID policy to avoid non-urgent surgery with potential complications. DCIS surgery deferred due to COVID. Followed up after 3 months –WLE performed at that time- restrictions lifted. Radiotherapy not initiated immediately post op - delayed 3 months. Commenced on endocrine therapy in interim. Operation in March. Radiotherapy completed in Aug
C	58	New cancer	DCIS	Surgery	DCIS surgery cancelled, but only delayed by 10 days- restrictions lifted. No reconstruction offered post mastectomy (low grade DCIS). Referred to plastic surgery team for delayed reconstruction.
D	71	New cancer	PT1cN0M0 Inv mucinous G1 ER+PR+HER2-	Radiotherapy	Not recruited for clinical trial due to trial recruitment shut down for COVID (EXPERT candidate) *
E	67	New cancer	DCIS	Surgery	Not offered free flap despite patient preference. Had implant-based reconstruction.
F	47	New cancer	DCIS	Surgery	Boost to tumour bed omitted.
G	54	New cancer	pT1cN0M0 Inv NST G2 ER+PR+HER2neg	Trial	Boost to tumour bed omitted.
H	61	Post op cancer	pT2N0M0 Inv NST G3 ER-PR-HER2pos	Surgery	Radiotherapy delayed until 4 months post operatively. Commenced on Endocrine therapy in interim.
I	52	Post op cancer	DCIS 12mm intermediate grade	Radiotherapy	
J	59	Post op cancer	DCIS 16mm intermediate grade	Radiotherapy	
K	61	Post op cancer	pT1cN0M0 Inv mucinous G2 ER+PR+HER2equiv	Radiotherapy	



Case	Age	Category	Cancer information	Treatment Changed	Details
L	35	Post NACT	pT1bN0M0 Inv NST G3 Triple positive	Surgery	Requested bilateral mastectomy but only offered unilateral.
M	56	Post NACT	IDC triple negative, node positive, RCB-II	Radiotherapy	Treated with hypofractionated radiotherapy rather than conventional fractionated post-mastectomy radiotherapy

*The EXPERT Trial is a randomised phase III trial of adjuvant radiation therapy versus observation following breast conserving surgery and endocrine therapy in patients with molecularly characterised luminal A early breast cancer.¹⁴

NACT=neoadjuvant chemotherapy

Changes to chemotherapy

Of 145 patients, 104 (71.7%) involved a medical oncologist in their breast cancer management. Of the 104 patients, 30 (28.8%) completed neoadjuvant systemic treatment and proceeded to surgery, 19 (18.3%) commenced neoadjuvant or adjuvant chemotherapy +/- anti-HER2 treatment, 17 (16.3%) proceeded to adjuvant endocrine therapy, 5 (4.8%) did not receive any systemic treatment and 33 (31.7%) had their medical oncology consultations after the study period.

One patient (B) experienced COVID-19 related changes to chemotherapy:

- Neoadjuvant chemotherapy was ideally recommended due to her grade 3 triple negative cancer with high proliferative index of 70%. Ultimately up-front surgery was recommended to avoid multiple hospital visits and immunosuppression. She promptly underwent wide local excision and sentinel node biopsy (11 days after MDT discussion), followed by adjuvant chemotherapy (47 days after surgery) and radiotherapy.

There were no pandemic related changes to adjuvant chemotherapy regimens or dosing. However, prophylactic addition of pegfilgrastim to doxorubicin/cyclophosphamide (AC) chemotherapy was implemented in March 2020. None of the 16 patients who started AC experienced febrile neutropenia during the study period, compared to an average of two patients per month prior to this period.

The time intervals between MDT discussion and Medical Oncology consultation (nine days for neoadjuvant chemotherapy and 12 days for adjuvant chemotherapy) and subsequent first cycle of chemotherapy (12 days) were comparable to outside this time period without delay.

Patients on clinical trials continued treatment as per protocol without interruption, but telehealth safety monitoring was instigated during the study period. There was no new recruitment to clinical trials during this period.

Changes to radiotherapy

There were 124 of 145 women (85.5%) recommended to receive radiotherapy. Of these, 99 (79.8%) were consented for radiotherapy during the study period, 19 (15.3%) were awaiting consent at the end of the study period, two (1.6%) declined radiotherapy, one (0.8%) was deemed unsuitable due to prior radiotherapy, and three (2.4%) had their treatment at a different centre. COVID-related radiotherapy consent alterations only occurred in the first two months of the first wave of the pandemic. During this period, ultra-hypofractionation regimens such as the FAST Forward trial were not implemented in this centre.¹⁵

Five patients experienced a change to radiotherapy due to COVID-19.

- Two patients (D, K) had a delay to treatment start. Both were over 60 years of age with either a grade 1 or 2 tumour and were commenced on adjuvant endocrine therapy starting radiotherapy three months later.
- Three patients had changes to their radiotherapy protocols:
 - Initially, a change in prescription to shorten the course of radiotherapy was recommended in five patients. However, between consent and initiation of treatment (2-5 weeks), two prescriptions were reverted to standard treatment. This involved one patient where a boost was to be omitted but was delivered, and one patient who was consented for a hypofractionated protocol but changed to conventional fractionation for whole breast and comprehensive nodal irradiation during the radiotherapy planning process as COVID-19 cases in NSW declined.
 - Two patients had a boost to the surgical cavity omitted to shorten treatment course duration
 - One patient received hypofractionated chest wall radiotherapy rather than conventional fractionated post-mastectomy radiotherapy post-NACT.



DISCUSSION

These results show that in the context of low pandemic burden, a minority of early breast cancer patients (9.0%) received altered management, with the majority still receiving ideal treatment regimens. These findings provide reassurance that minimal patients may experience future recurrence as a result of ‘less than ideal’ management during the pandemic. The modest changes recorded are likely influenced by factors including reduced new presentations, institutional mitigative strategies and comparatively low COVID-19 transmission. These observations can provide insights in minimising compromises to oncological care for other institutions with improving COVID-19 rates and for future pandemics.

The changes to surgery, radiotherapy, and chemotherapy treatments were unlikely to impact significantly on risk of recurrence. The patient who was treated with up-front surgery rather than neoadjuvant chemotherapy is unlikely to have adverse effects from the change in sequencing. For other chemotherapy patients, the use of prophylactic G-CSF reduced the incidence of febrile neutropenia in neoadjuvant or adjuvant treatments and resulted in hospitalisations reducing substantially. Approximately 20% of patients may experience adverse events with bone pain, which can often be controlled with analgesia. Primary prophylaxis using pegylated G-CSF with AC may become routine treatment for the post-COVID-19 era.

Radiotherapy changes include two DCIS cases treated with whole-breast radiation with omission of a boost to the tumour bed. The benefit of a boost to the tumour bed for patients with non-low risk DCIS was not known until the publication of the TROG DCIS trial results in December 2020.¹⁶ One case of low-grade mucinous cancer experienced an intentional three-month delay to her radiotherapy treatment and commenced endocrine therapy in the interim. The likelihood of this significantly increasing risk of recurrence is considered very small.

Two patients were not referred for an eligible trial study during MDT discussions. Recommendation for trial therapy was not primarily discussed during these meetings and possibly many more patients referred for eligible trials at other points of management were not captured by this study. This has impacted the recruitment and progression of clinical trials suspended during this period.

Institutions abroad facing higher COVID-19 prevalence implemented stricter precautions to minimize the risk of infecting vulnerable cancer patients and experienced higher levels of resource competition.¹⁷⁻¹⁹ Strategies to cope included “hub and spoke” models, where a central facility coordinates care that is carried out in designated “COVID-19

free” satellite locations.¹⁷⁻¹⁹ Similarly in this study, most new referrals recommended for upfront surgery underwent their operations at COVID-free facilities within the same public hospital district or at locally affiliated private hospitals. This allowed the institution to focus on COVID management, protect surgical cancer patients from viral exposure, and utilize available resources in the private sector, which experienced a massive reduction in work-load during the suspension of elective surgeries. Breast cancer specific services and personnel remained largely operational during the study period, and were not significantly impacted by resource limitations. This was facilitated by a comparatively low burden of local COVID-19 cases in NSW. No patients in the study cohort were found to be COVID-19 positive although there may have been a low infection risk from isolated events of community transmission to healthcare staff who tested positive during this period.

Telemedicine is also being increasingly utilized for breast cancer management and follow-up appointments. Long term consequences may still emerge such as potentially missing recurrences and early signs of complications due to the omission of clinical examination. Alternatively, it may encourage saving of healthcare resources and increase patient convenience.²⁰

The full impact of COVID-19 on early breast cancer may take years to understand as delayed presentations manifest. This study was conducted over the three-month period of most severe lockdowns and restrictions in NSW and showed a 25% reduction in new cancer referrals during the study period. This correlates to when the national screening program was closing down.² Furthermore, national data demonstrated a decrease of 30–50% in Medicare benefits (Australian universal healthcare scheme) claimed for breast cancer diagnostic and treatment services in April/May 2020, with some recovery in later months.¹³ Significant reductions in new breast cancer referrals observed during lockdown period in Australia are reflected in other countries such as the UK where urgent new cancer referrals dropped by up to 80% in early 2020, attributed to disrupted screening programs, reduced primary care presentations, and more cautious health-seeking behaviours.^{21, 22} Diagnostic delays may be associated with a backlog of patients who could present with more advanced disease requiring more complex treatment courses involving systemic therapies, upgraded surgical procedures, and poorer outcomes.^{21, 23} The national BreastScreen program has already observed a surge of 12000 screening mammograms in July to September 2020 following re-opening, compared to the same period in 2018.²⁴ Modelling of the impact of diagnostic delays for



patients presenting in a 12 month period under different COVID-19 scenarios estimates an increased number of deaths due to breast cancer by 7.9-9.6% in the ensuing 5 year period.²²

These flow-on effects affect not only new patients at diagnosis and primary management but cause delays to “non-urgent” treatments, such as reconstructive procedures. After non-urgent surgeries were suspended between March and July 2020 in the NHS, there is now a backlog of over 1500 patients awaiting post mastectomy breast reconstructions.²⁵

This study has some strengths and limitations. The strengths include the prospective documentation of ‘ideal’ treatment plans and COVID-related changes through robust data collection by the MDT. This was limited by excluding post-operative patients who were already receiving adjuvant treatment during the study period as they had treatment plans decided pre-COVID. This study also did not measure the psychological impact of treatment changes, which may be significant. Several patients were unable to have their preferred plans for breast reconstruction even though the importance of choice for these treatments on psychological well-being has been documented.²⁶ Further research is needed to explore the impacts on quality of life by changes to treatment related to COVID-19, and the psychological impacts of a cancer diagnosis at this time of heightened overall distress in the community. A further limitation is that patient-reported outcomes were not collected. The study did not continue beyond mid-2020 to see if there was a ‘catch-up’ in referrals, as would be expected according to national data.¹³ Also, follow-up data is not available at this stage, and this will be required to evaluate the impact of changes to treatment.

Much of the research on COVID-19 impacts on breast cancer care comprises observational, single centre data, to which this paper contributes perspectives from a low pandemic burden institution. For healthcare systems integrating pandemic-related strategies into mid and long-term practice, larger multicentre studies and systematic reviews are required to establish evidence-based protocols. As

vaccination programs roll out, countries entering recovery status may take precedent from countries with low pandemic burden when accounting for surging capacity in routine screening and treatment. Similarly, countries with rising cases related to easing lockdowns and spread of new COVID strains may more quickly adopt proven guidelines to protect cancer patients from infection while preserving ideal oncological outcomes. Approximately 12 months from the lifting of the initial lock-down, Sydney experienced a second wave of COVID-19 caused by the Delta variant, so the experience obtained from the first wave was put into practice again.

CONCLUSION

Despite the temporary cessation of breast cancer screening, restrictions on surgery and modified radiotherapy and chemotherapy recommendations, 91.0% of patients at our institution received standard care for their breast cancer during the first wave of the COVID-19 pandemic in NSW. This was possible due to the relatively small COVID-19 incidence in Australia, combined with partnerships between public and private hospitals to facilitate the provision of urgent cancer surgery. Review of the cases experiencing a change to treatment shows that the modifications are unlikely to be of oncological significance. It is possible, however, that the restricted access to breast reconstruction and contralateral surgery options may have quality of life implications which will be addressed in the follow-up of these

CONFLICT OF INTEREST

RH has participated in advisory boards for AstraZeneca, Bristol Myers Squibb, Eli Lilly, Merck, Merck Sharp and Dohme, Novartis, Oncosec, Pfizer, Roche and Seagen and has received speaker honoraria from Merck Sharp and Dohme, Novartis and Roche. EE has participated in advisory board for Merck Sharp and Dohme. There are no conflicts of interests to declare for the other authors. There were no funding sources to declare. All authors confirm that they had full access to all the data in the study.

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