PDF version

Bita Eslami[1], Sadaf Alipour[1,2], Monireh S Seyyedsalehi[3], Azin Nahvijou[3], Ramesh Omranipour[1,4], Mojtaba V Rajabpour[3] and Kazem Zendehdel[1,5]

[1]Breast Diseases Research Center, Cancer Institute, Tehran University of Medical Science, Tehran, Islamic Republic of Iran (Correspondence to Kazem Zendehdel: kzendeh@tums.ac.ir).
[2]Department of Surgery, Arash Women's Hospital, Tehran University of Medical Science, Tehran, Islamic Republic of Iran.
[3]Cancer Research Center, Cancer Institute, Tehran University of Medical Sciences, Tehran, Islamic Republic of Iran.
[4]Department of Surgical Oncology, Cancer Institute, Tehran University of Medical Sciences, Tehran, Islamic Republic of Iran.
[5]Department of Medical and Surgical Sciences, University of Bologna, Bologna, Italy.

Abstract

Background: Comorbidities have a significant impact on the treatment and outcome of breast cancer. However, data on such comorbidities from low-income countries are limited.

Aim: To evaluate the feasibility and accuracy of comorbidity data extracted from medical records for estimating the prevalence of comorbidities among patients registered in the clinical breast cancer registry of the Islamic Republic of Iran.

Methods: We collected data from the medical records of 400 patients on 30 comorbidities included in the Charlson Comorbidity Index (CCI) and Elixhauser Comorbidity Index (ECI). The sensitivity and specificity of comorbidity data extracted from medical records were calculated

using data from interviews with 97 randomly selected patients. We studied the prevalence of comorbidities using the CCI and ECI. Data were analysed using SPSS version 24.

Results: The mean age of patients was 51.69 SD 12.28 years. The sensitivity and specificity of medical records for detecting any comorbidity data contained in CCI versus non-comorbidity were 93.2% and 69.8%, respectively. However, for the comorbidity data included in ECI, both sensitivity (86.9%) and specificity (44.4%) were lower than in CCI. Hypertension (n = 144, 36.0%) and diabetes without chronic complications (n = 77, 19.3%) were the most prevalent comorbidities. A higher proportion of patients had no comorbidity with CCI (72.2%) than with ECI (44.8%).

Conclusion: It is feasible to construct a comorbidity index using medical records with high accuracy, especially when we extract comorbidities using the CCI. Further studies are needed to understand the association between comorbidity index and breast cancer survival among Iranian patients.

Keywords: breast cancer, cancer registry, Charlson Comorbidity Index, Elixhauser Comorbidity Index, Iran.

Citation: Eslami B, Alipour S, Seyyedsalehi MS, Nahvijou A, Omranipour R, Rajabpour MV, Zendehdel K. Feasibility of measuring comorbidity indices based on clinical breast cancer records. East Mediterr Health J. 2023;29(11):870–876. https://doi.org/10.26719/emhj.23.091
Received: 09/12/22; Accepted: 28/04/23

Copyright © Authors 2023; Licensee: World Health Organization. EMHJ is an open access journal. This paper is available under the Creative Commons Attribution Non-Commercial ShareAlike 3.0 IGO licence (CC BY-NC-SA 3.0 IGO; https://creativecommons.org/licenses/by-nc-sa/3.0/igo).

Introduction

Comorbidity refers to a long-term health condition or disorder that coexists with a primary disease (1), and multimorbidity refers to the presence of ≥ 2 long-term health conditions (2).

Comorbidity is common among older cancer patients, and 4 out of 10 cancer patients have at least one comorbid disease and 15% have multimorbidity (3). Breast cancer is the most common female cancer worldwide and is the main cause of cancer mortality among women (4). The coexistence of breast cancer and other disease has an impact on treatment planning and outcome. A systematic review reported that the prevalence of comorbidity among breast cancer patients ranged widely from 0.4% to 87% in different populations (5). Different comorbidity measures have been developed and the most commonly used indices are the Charlson Comorbidity Index (CCI) and the Elixhauser Comorbidity Index (ECI), or their derivatives (6, 7).

The impact of comorbidity on management and survival, care cost, and ability to predict outcome of breast cancer patients has been evaluated in several studies (8–14). The prevalence of comorbidity and weighted indices vary depending on the target population and the type of cancer. The best strategy for evaluating comorbidity is to develop an index for each type of cancer and a specific weight for comorbidity in the study population (15). To the best of our knowledge, no study has reported the development and adaptation of comorbidity indices in low and middle-income countries, although these indicators are of interest in developed countries (15,16–20).

Accurate recording of comorbidity details in a valid database is the first step in assessing the impact of comorbidity on treatment of cancer patients. The lack of comorbidity data is one of the shortcomings of some registries. Addition of such data extracted from available records in medical centres, such as paper or electronic medical records of inpatients and outpatients, is a cost-effective method if the reliability is confirmed. The Clinical Breast Cancer Registry of Iran was established in 2014 at the Cancer Institute of Iran and extended to other cancer hospitals across the country (21). Trained registrars review patients' medical records and collect detailed clinical data including diagnosis, staging, treatment, and follow-up data, and register them in a web-based system designed specifically for the Clinical Breast Cancer Registry of Iran. At the time of creating the clinical registry, the entry of comorbidity data was not the goal or priority. Therefore, to optimize research on breast cancer patients, the Cancer Institute Registry Team decided to add comorbidity data.

The aims of this study were: (1) to evaluate the feasibility and accuracy of using comorbidity data extracted from medical records of breast cancer patients admitted to the Cancer Institute of Iran, in order to add comorbidity data based on CCI or ECI to the Clinical Breast Cancer Registry; and (2) to estimate the prevalence of comorbidity among registered patients.

Methods

This study was approved by the Research Ethics Committee of Imam Khomeini Hospital (code:

IR.TUMS.IKHC.REC.1399. 191) affiliated with the Tehran University of Medical Sciences. We collected information on all comorbidities included in CCI (n = 16) and ECI (n = 25). According to a systematic review and meta-analysis in 2018, dyslipidaemia is a prevalent comorbidity in the Islamic Republic of Iran (22), and several studies have shown that serum lipid levels are significantly associated with breast cancer risk and mortality (23, 24). Therefore, we added hyperlipidaemia to the list of comorbidities in this study.

Comorbidity information was extracted from the inpatient and outpatient electronic medical records at the Cancer Institute of Iran. There was no single questionnaire containing information on comorbidities in the patients' files. Therefore, initially, we reviewed medical history recorded by physicians; nursing assessment sheets; preoperative cardiac, anaesthesia, or other specialty consultation notes; and results of laboratory and paraclinical assessments. According to the findings of this initial step, we prepared a guideline to help registrars collect comorbidity data from the medical records, and these data were coded based on the International Classification of Diseases 10th Revision (https://icd.who.int/browse10/2019/en). Trained registrars used the guideline to review all the notes containing relevant data and extracted the comorbidity information. The collected comorbidity data were entered into the Clinical Breast Cancer Registry of Iran.

To evaluate the accuracy of the extracted comorbidity data, we randomly selected 132 of the 400 patients for telephone interview to ask about their comorbidities, and 97 responded (73.5% response rate). Most of the patients we could not get response from could not be reached through their telephone numbers.

We used the comorbidity results reported at interview as a gold standard to calculate the sensitivity, specificity, and accuracy of extracting any comorbidity versus noncomorbidity from medical data. The accuracy of extracting the presence or absence of 2 common diseases (diabetes and hypertension) was calculated separately. We studied the prevalence of comorbidities based on CCI and ECI.

The mean (standard deviation) for continuous variables and percentage for categorical variables were calculated using SPSS version 24.

Results

The mean age of the 400 breast cancer patients was 51.7 (12.3) years, with a range of 24–86 years. There were 395 (98.8%) female and 5 (1.3%) male patients.

The accuracy of the comorbidity data extracted from medical records is shown in Table 1 . The sensitivity, specificity, and accuracy of medical records for detecting any comorbidity data contained in CCI versus noncomorbidity was 93.2%, 69.8%, and 80.4%, respectively. The accuracy of extracting the comorbidity data available in ECI from medical records was lower than for CCI (86.9% sensitivity and 44.4% specificity). Diabetes was registered in medical records with sensitivity, specificity, and accuracy of 100.0%, 71.7%, and 82.5%, respectively.

Table 2 shows the prevalence of comorbidities according to the CCI and ECI. Among the Charlson comorbidities, diabetes without chronic complications (n = 77, 19.3%) was the most prevalent. Based on the Elixhauser comorbidities, uncomplicated hypertension (n = 144, 36.0%) was the most prevalent. Our evaluation showed that the prevalence of hyperlipidaemia was 17.0% (n = 68).

According to CCI and ECI, 72.2% and 44.8% of the patients, respectively had no report of any comorbidity (Table 3). Multimorbidity was seen in 6.9% and 19.1% of the patients according to CCI and ECI, respectively. In the subanalysis, there was no report of comorbidity among patients under 30 years of age. At least 1 comorbidity was reported in 11.4% of patients aged

Discussion

We investigated the feasibility of comorbidity data extraction using medical records based on the CCI and ECI in order to add data to the Clinical Breast Cancer Registry of Iran. The sensitivity and specificity for detecting comorbidities in medical records showed that obtaining comorbidity data from medical records was feasible and had sufficient accuracy to improve clinical data for breast cancer studies. We found that entering comorbidity data based on the CCI was more accurate; however, the CCI did not capture all types of comorbidities, such as hypertension and hypothyroidism, that may be relevant to health outcomes and quality of life in breast cancer patients. Therefore, we believe that extraction of comorbidity data from both CCI and ECI should be continued to improve the mortality index in our population.

Several studies have investigated comorbidities and the use of hospital or self-reported data by cancer patients. Similar to our study, in the California Cancer Registry, comorbidity information for breast cancer patients extracted from hospital discharge data was compared with comorbidity scores derived using the Surveillance, Epidemiology, and End Results (SEER)–Medicare database. The authors concluded that the sensitivity of hospital discharge data for detecting any comorbidity versus noncomorbidity was sufficiently high (76.5%) to allow

the construction of a comorbidity index for breast cancer registries (19). The California study and this study confirm that the assessment of comorbidity data using internal data sources, such as paper or electronic medical records in hospitals, is appropriate and reliable. Using and connecting internal data with health databases outside hospitals, such as private laboratories, are cost- and time-consuming and need specific authorization.

In contrast to our study, among women diagnosed with breast cancer, who were part of the California Breast Cancer Survivorship Consortium, the self-reported information and electronic medical records for 4 common comorbidities (diabetes, hypertension, myocardial infarction, and other heart diseases) were compared (25). The concordance rate for myocardial infarction and other heart diseases was not sufficiently high (

Diabetes is important in the CCI and ECI and in our study population; therefore, we assessed diabetes data in medical records and found 82% accuracy for reporting diabetes without complications. Our results are consistent with those from previous studies that reported the high sensitivity and specificity in recording diabetes by different methods; therefore, it seems that history of diabetes is well documented in medical records (19, 25). Although comorbid diabetes was mentioned in patients' records, end-organ damage associated with diabetes was not well recorded. In the present study, chronic complications following diabetes were recorded correctly in only 2 of 7 patients. We should note that chronic complications of diabetes may overlap with other comorbidities such as heart and renal disease. The prevalence of diabetes with complications in our study was similar to that in the study of Klabunde et al. (1.8% vs 1%) (15). However, in another study by Mehta et al., the rate of diabetes with end-organ damage was higher (4.3%), probably because of the older age of the participants (mean 75.8 years) (17).

Based on our results, 72.2% of patients had no comorbidities according to CCI and this was consistent with the California Cancer Registry in 2017, which showed that 75.3% of breast cancer patients had no relevant comorbidities (19). Similar to our results, the SEER study in 123 680 breast cancer patients reported that 67.8% had no comorbidity (3). Multimorbidity was higher in the SEER study (9.8%) than in our study (6.9%), because of the recruitment of older patients aged ≥ 66 years. The difference may also be because of the younger age of onset of breast cancer among Iranian than the European and North American patients (27). Contrary to the studies mentioned above, Fu et al. reported that, among 134 breast cancer patients, 73.8% had at least 1 comorbidity (28). That study evaluated all CCI comorbidities and other conditions mentioned in the open-ended interview questions; therefore, it is better to compare with our results for ECI, which captured more comorbidities. The number of patients with at least 1 comorbidity in the Fu study was higher than in our study (73.8 % vs 55.2%). This difference may be because the Fu study had a smaller sample size and higher average age (56 years) than in our study (52 years).

In our study, the prevalence of comorbidities was assessed using CCI and ECI. Hypertension and diabetes without chronic complications were the most common comorbidities among breast cancer patients. Similar to our study, uncomplicated diabetes and hypertension were the most prevalent comorbidities in studies from the United States of America (17) and Australia (29). In other studies, the reported prevalence of diabetes without complications ranged from 10% to 22% (15, 17, 19, 28, 29) and that of hypertension from 14% to 58% (17, 28, 29) among cancer patients. The rate of these 2 comorbidities in our study was within these reported ranges. It seems that the difference between the reported prevalence of comorbidities, especially hypertension, resulted from the age of the patients at the time of recruitment.

To the best of our knowledge, despite several studies in developed countries, there has been no systematic evaluation of the feasibility of medical data extraction from health systems to measure comorbidity index in developing countries. Considering that the comorbidity index needs to be tailored to specific populations (15, 19, 28), defining the specific comorbidity index and developing weights for comorbidities among breast cancer patients in each population are necessary. Our study is believed to be the first to systematically investigate comorbidities and add them to cancer registration in developing countries, including the Islamic Republic of Iran. The strengths of the study were that it was conducted in the Cancer Institute of Iran and used data from a high-quality registry and a large study sample.

Our study had some limitations. Patients can have anxiety at the time of cancer diagnosis, and may not remember their history of comorbidity, or omit details of the diseases during hospital admission or during interview (recall bias). Furthermore, we used interview data as a gold standard measurement in this study, which was subject to recall bias and misclassification. Women with a long history of comorbidity or who are taking medication are more likely to report their diseases. We tried to mitigate this bias by employing a trained interviewer for this study.

We are aware that medical records are often fragmented across multiple healthcare programmes, posing an obstacle to clinical care and research studies. Therefore, we recommend the development of electronic records and public–private partnerships to integrate entire records and facilitate accurate registration of comorbidity data, specifically for severe comorbidities that may have a high impact on patient outcomes. Data generated by the private sector, such as laboratories and radiology centres, should be combined with data produced by public and university hospitals to allow exchange of data according to ethical standards. Accessibility to medical records and permission to use personal medical information, while protecting privacy of patients is recommended (30).

In further studies we will investigate the prevalence of comorbidities among breast cancer

patients throughout the Islamic Republic of Iran and compare the results between health centres of different geographical regions. We will continue to develop a clinical comorbidity index for Iranian breast cancer patients that can improve prediction of survival. It is hoped that this index will be implemented by other cancer registries across the country to increase the relevance and usefulness of breast cancer registry data.

Conclusion

It is appropriate to use medical records to collect comorbidity information and construct a comorbidity index for breast cancer patients admitted to the Cancer Institute of Iran. Extraction of comorbidity data from medical records considering CCI provides greater accuracy than for ECI. We recommend recording all comorbidities included in CCI and ECI in the CBCR-IR to develop the best comorbidity index for predicting survival among Iranian breast cancer patients. Active data collection and face-to-face interviews with patients and evaluation of medical records are needed for some comorbidities. To overcome the limitation in this type of study and collect comorbidity data accurately, development of integrated electronic medical records is necessary.

Acknowledgement

We would like to thank the Clinical Breast Cancer Registry of Iran for providing the data for this research and registrars working there.

Funding: This study was supported by the Cancer Institute, Tehran University of Medical Sciences (No# 98-3-259-45418).

Conflict of interest: None declared.

Faisabilité de la mesure des indices de comorbidité à partir des dossiers cliniques sur le cancer du sein en République islamique d’Iran

Résumé

Contexte : Les comorbidités ont un impact significatif sur le traitement et l’issue du cancer du sein. Cependant, les données sur les comorbidités provenant de pays à faible revenu sont limitées.

Objectifs : Évaluer la faisabilité et l'exactitude des données de comorbidité extraites des dossiers médicaux pour estimer la prévalence des comorbidités chez les patient(e)s enregistré(e)s dans le registre clinique du cancer du sein en République islamique d'Iran.

Méthodes : Nous avons recueilli des données à partir des dossiers médicaux de 400 patient(e)s sur 30 comorbidités incluses dans l'indice de comorbidité de Charlson (ICC) et l'indice de comorbidité d'Elixhauser (ICE). La sensibilité et la spécificité des données de comorbidité extraites des dossiers médicaux ont été calculées à partir des données provenant d'entretiens menés auprès de 97 patient(e)s sélectionné(e)s de manière aléatoire. Nous avons étudié la prévalence des comorbidités à l'aide de l'ICC et de l'ICE. Les données ont été analysées au moyen du logiciel SPSS version 24.

Résultats : L'âge moyen des patient(e)s était de 51,69 (ET 12,28) ans. La sensibilité et la spécificité des dossiers médicaux pour la détection de toute donnée de comorbidité contenue dans l'ICC par rapport à la non-comorbidité étaient de 93,2 % et de 69,8 %, respectivement. Cependant, pour les données de comorbidité incluses dans l'ICE, la sensibilité (86,9 %) et la spécificité (44,4 %) étaient plus faibles que pour celles comprises dans l'ICC. Les comorbidités les plus fréquentes étaient l'hypertension (n = 144 ; 36,0 %) et le diabète sans complications chroniques (n = 77 ; 19,3 %). La proportion de patient(e)s ne présentant aucune comorbidité avec l'ICC (72,2 %) était plus élevée qu'avec l'ICE (44,8 %).

Conclusion : Il est possible de concevoir un indice de comorbidité en utilisant des dossiers médicaux affichant un degré élevé de précision, en particulier lorsque nous extrayons des données de comorbidité en utilisant l'ICC. Des études plus poussées sont nécessaires pour comprendre le lien entre l'indice de comorbidité et la survie au cancer du sein chez les patient(e)s iranien(ne)s.

[illegible]

[illegible]

[illegible]

[illegible]

.□□□□□□□

□□□□□□□

CA Cancer J Clin. 2016;66(4):337–50. https://doi.org/10.3322/caac.21342 PMID:26891458

3. Edwards BK, Noone AM, Mariotto AB, Simard EP, Boscoe FP, Henley SJ, et al. Annual Report to the Nation on the status of cancer, 1975–2010, featuring prevalence of comorbidity and impact on survival among persons with lung, colorectal, breast, or prostate cancer. Cancer. 2014;120(9):1290–314. https://doi.org/10.1002/cncr.28509 PMID:24343171

4. Ferlay J, Héry C, Autier P, Sankaranarayanan R. Global burden of breast cancer. In: Li C, editor. Breast cancer epidemiology: Springer; 2010:1–19.

5. Lee L, Cheung WY, Atkinson E, Krzyzanowska MK. Impact of comorbidity on chemotherapy use and outcomes in solid tumors: a systematic review. J Clin Oncol. 2011 Jan 1;29(1):106–17. https://doi.org/10.1200/JCO.2010.31.3049 PMID:21098314

6. Charlson ME, Pompei P, Ales KL, MacKenzie CR. A new method of classifying prognostic comorbidity in longitudinal studies: development and validation. J Chronic Dis. 1987;40(5):373–83. https://doi.org/10.1016/0021-9681(87)90171-8. PMID:3558716

7. Elixhauser A, Steiner C, Harris DR, Coffey RM. Comorbidity measures for use with administrative data. Med Care. 1998 Jan:36(1):8–27. https://doi.org/10.1097/00005650-199801000-00004 PMID:9431328

8. Berglund A, Wigertz A, Adolfsson J, Ahlgren J, Fornander T, Wärnberg F, et al. Impact of comorbidity on management and mortality in women diagnosed with breast cancer. Breast Cancer Res Treat. 2012 Aug;135(1):281–9. https://doi.org/10.1007/s10549-012-2176-4 PMID:22829398

9. Land LH, Dalton SO, Jørgensen TL, Ewertz M. Comorbidity and survival after early breast cancer. A review. Crit Rev Oncol Hematol. 2012 Feb;81(2):196–205. https://doi.org/10.1016/j.critrevonc.2011.03.001 PMID:21536452

10. Salas M, Henderson M, Sundararajan M, Tu N, Islam Z, Ebeid M, et al. Use of comorbidity indices in patients with any cancer, breast cancer, and human epidermal growth factor receptor-2-positive breast cancer: a systematic review. Plos One. 2021 Jun 18;16(6):e0252925. https://doi.org/10.1371/journal.pone.0252925 PMID:34143813

11. Yoon SJ, Kim EJ, Seo HJ, Oh IH. The association between Charlson comorbidity index and the medical care cost of cancer: a retrospective study. BioMed Res Int. 2015;2015:259341. https://doi.org/10.1155/2015/259341 PMID:26347086

12. Patterson RE, Flatt SW, Saquib N, Rock CL, Caan BJ, Parker BA, et al. Medical comorbidities predict mortality in women with a history of early stage breast cancer. Breast Cancer Res Treat. 2010 Aug;122(3):859–65. https://doi.org/10.1007/s10549-010-0732-3 PMID:20077000

13. Breccia M, Latagliata R, Stagno F, Luciano L, Gozzini A, Castagnetti F, et al. Charlson comorbidity index and adult comorbidity evaluation-27 scores might predict treatment compliance and development of pleural effusions in elderly patients with chronic myeloid leukemia treated with second-line dasatinib. Haematologica. 2011 Oct;96(10):1457–61. https://doi.org/10.3324/haematol.2011.041251 PMID:21685471

14. Woo HK, Park JH, Kang HS, Kim SY, Lee SI, Nam HH. Charlson comorbidity index as a predictor of long-term survival after surgery for breast cancer: a nationwide retrospective cohort study in South Korea. J Breast Cancer. 2010 Dec;13(4):409–17. https://doi.org/10.4048/jbc.2010.13.4.409.

15. Klabunde CN, Legler JM, Warren JL, Baldwin L-M, Schrag D. A refined comorbidity measurement algorithm for claims-based studies of breast, prostate, colorectal, and lung cancer patients. Ann Epidemiol. 2007 Aug;17(8):584–90. https://doi.org/10.1016/j.annepidem.2007.03.011 PMID:17531502

16. Klabunde CN, Potosky AL, Legler JM, Warren JL. Development of a comorbidity index using physician claims data. J Clin Epidemiol. 2000 Dec;53(12):1258–67. https://doi.org/10.1016/s0895-4356(00)00256-0 PMID:11146273

17. Mehta HB, Sura SD, Adhikari D, Andersen CR, Williams SB, Senagore AJ, et al. Adapting the Elixhauser comorbidity index for cancer patients. Cancer. 2018 May 1;124(9):2018–25. https://doi.org/10.1002/cncr.31269 PMID:29390174

18. Groll DL, To T, Bombardier C, Wright JG. The development of a comorbidity index with physical function as the outcome. J Clin Epidemiol. 2005 Jun;58(6):595–602. https://doi.org/10.1016/j.jclinepi.2004.10.018 PMID:15878473

19. Lichtensztajn DY, Giddings BM, Morris CR, Parikh-Patel A, Kizer KW. Comorbidity index in central cancer registries: the value of hospital discharge data. Clin Epidemiol. 2017;9:601–9. https://doi.org/10.2147/CLEP.S146395 PMID:29200890

20. Brusselaers N, Lagergren J. The Charlson comorbidity index in registry-based research. Methods Inf Med. 2017;56(5):401–6. https://doi.org/10.3414/ME17-01-0051 PMID:2958293

21. Seyyedsalehi MS, Nahvijou A, Rouhollahi M, Teymouri F, Mirjomehri L, Zendehdel K. Clinical Cancer Registry of the Islamic Republic of Iran: Steps for Establishment and Results of the Pilot Phase. J Registry Manag. 2020 Winter;47(4):200–6. PMID:34170898

22. Toori MA, Kiani F, Sayehmiri F, Sayehmiri K, Mohsenzadeh Y, Ostovar R, et al. Prevalence of hypercholesterolemia, high LDL, and low HDL in Iran: a systematic review and meta-analysis. Iran J Med Sci. 2018 Sep;43(5):449–65. PMID:30214097

23. Ni H, Liu H, Gao R. Serum lipids and breast cancer risk: a meta-analysis of prospective cohort studies. PloS One. 2015 Nov 10;10(11):e0142669. https://doi.org/10.1371/journal.pone.0142669 PMID:26554382

24. Borgquist S, Broberg P, Tojjar J, Olsson H. Statin use and breast cancer survival–a Swedish nationwide study. BMC Cancer. 2019 Jan 11;19(1):1–9. https://doi.org/10.1186/s12885-018-5263-z PMID:30634941

25. Vigen C, Kwan ML, John EM, Gomez SL, Keegan TH, Lu Y, et al. Validation of self-reported comorbidity status of breast cancer patients with medical records: the California Breast Cancer Survivorship Consortium (CBCSC). Cancer Causes Control. 2016 Mar;27(3):391–401. https://doi.org/10.1007/s10552-016-0715-8 PMID:26797455

26. Lujic S, Simpson JM, Zwar N, Hosseinzadeh H, Jorm L. Multimorbidity in Australia: comparing estimates derived using administrative data sources and survey data. PloS One. 2017;12(8):e0183817. https://doi.org/10.1371/journal.pone.0183817 PMID:28850593

27. Lei S, Zheng R, Zhang S, Wang S, Chen R, Sun K, et al. Global patterns of breast cancer incidence and mortality: a population-based cancer registry data analysis from 2000 to 2020. Cancer Commun. 2021 Nov;41(11):1183–94. https://doi.org/10.1002/cac2.12207 PMID:34399040

28. Fu MR, Axelrod D, Guth AA, Cleland CM, Ryan CE, Weaver KR, et al. Comorbidities and quality of life among breast cancer survivors: a prospective study. J Pers Med. 2015 Jun 26;5(3):229–42. https://doi.org/10.3390/jpm5030229 PMID:26132751

29. Pule L, Buckley E, Niyonsenga T, Banham D, Roder D. Developing a comorbidity index for comparing cancer outcomes in Aboriginal and non-Aboriginal Australians. BMC Health Serv Res. 2018 Oct 16;18(1):776. https://doi.org/10.1186/s12913-018-3603-y PMID:30326898

30. Mandl KD, Markwell D, MacDonald R, Szolovits P, Kohane IS. Public standards and patients' control: how to keep electronic medical records accessible but private. BMJ. 2001 3;322(7281):283–7. https://doi.org/10.1136/bmj.322.7281.283

Thursday 11th of July 2024 07:25:42 AM