A comparative study of Watson for Oncology and tumor boards in breast cancer treatment

Dohoon Kim1, Yun Young Kim1,2, Joon-Hyop Lee1,2, Yoo Seung Chung1,2, Sangtae Choi1,2, Jin Mo Kang1,2, Heung Kyu Park1,2, Yong Soon Chun1,2

1Department of Surgery, Gachon University Gil Medical Center, Incheon; 2Department of Surgery, Gachon University College of Medicine, Incheon, Korea

Purpose: Watson for Oncology (WFO) is a computing system for considering treatment option with patients. The aim of this study is to determine the concordance rate of WFO and tumor board in treatment options of breast cancer.

Methods: One hundred and seventy breast cancer patients who were treated at Gachon University Gil Medical Center (GMC) from December 2016 to March 2018 were investigated. “Concordance” is defined that treatment provided by GMC tumor board and WFO-provided treatments (Recommend or For Consideration) were coincide. “Discordance” is defined that treatment provided by GMC tumor board and WFO-provided treatments (Not Recommended or Not Available) were coincide.

Results: In chemotherapy, of the total 170 patients, 23 patients were excluded, and the concordance rate was compared in 147 patients. Concordance rate was 93% (136/147). In the 11 patients who were discordant, seven patients complied with the treatment proposed by GMC tumor board due to insurance problems and age, and two patients did not follow the treatment proposed by WFO due to pregnancy and patient’s choice, and one patient chose a treatment determined by WFO, last one patient was not treated with chemotherapy because it was not considered to be clinically necessary. In radiotherapy, the overall agreement rate was as high as 99% (147/148).

Conclusion: Through this study, we found that many of the treatment proposals provided by WFO are highly reliable. Although the overall agreement is high in the radiotherapy regimen, the indication for radiotherapy by WFO is more aggressive and hypofractionated high dose intensity radiotherapy is growing trend by WFO.

Keywords: Watson for Oncology, Breast neoplasms, Treatment option, Concordance, Tumor board

INTRODUCTION

The treatment methods for breast cancer are evolving rapidly. New therapies such as radiation therapy, chemotherapy, and endocrine therapy, have been developed with the improvements in science and technology, and now molecular genetic methods have been developed to greatly affect the treatment and prognosis of breast cancer. As of October 2017, there were 69 drugs approved by the Food and Drug Administration for breast cancer treatment [1].

In a situation where technology in various fields is developed [2], the guidelines for the diagnosis and treatment of breast cancer and the development of treatment plans are improving at an accelerating rate [3]. Therefore, it is necessary to choose a treatment that suits the individual characteristics of a patient, but there are too many factors for a physician to synthesize and determine the knowledge in all fields.

Clinical decision-support systems can be a great solution to
solve these difficulties because they can help collect and analyze patient information as well as assist the clinicians in treatment decisions [4].

Watson for Oncology (WFO) is a cognitive computing system designed by Memorial Sloan Kettering clinicians and IBM that considers various treatment options with patients. The system uses natural language, evaluates cases with evolving machine-learned models, and processes large volumes of data rapidly [5]. When WFO was introduced to Gachon University Gil Medical Center (GMC) in September of 2016, it had already learned more than 300 medical journals, more than 200 medical textbooks, and 1,500 pages of medical information [6]. These characteristics can be of great help in presenting a range of treatments for patients.

Because of the benefits of WFO, WFO is now being used to assist in the treatment of cancer patients at GMC. This study examined whether WFO can be an alternative to clinically useful treatment options by investigating the concordance rate between WFO and GMC tumor boards for patients with breast cancer.

**METHODS**

**Patients**

A total of 170 patients treated with breast cancer from December 2016 to March 2018 at GMC were included. GMC tumor boards composed of six multidisciplinary doctors using WFO discussed all patients, and treatment options presented by WFO and GMC tumor boards were compared. If the treatment options of WFO and GMC tumor boards are not consistent, treatment methods are determined by a discussion of members of GMC tumor board to reflect patient’s preference. In 170 patients, 14 patients were excluded. Six patients refused chemotherapy and radiotherapy and histology results of four patients were unavailable from WFO. Two patients were diagnosed of other cancers in the past and WFO does not cover the patients with multiple cancers. One patient was lost in follow up and one case was required neoadjuvant chemotherapy but the patient requested surgery first. All patients had ductal, lobular and mixed histology of breast cancer. In 156 patients, 147 patients underwent adjuvant chemotherapy after surgical treatment, and nine patients experienced neoadjuvant chemotherapy.

**Concordance determination**

WFO suggests treatment options that consist of three categories (Recommend, For Consideration, and Not Recommended). In this study, “concordance” was defined as when the treatment options provided by GMC tumor boards and WFO-provided treatment options (Recommend or For Consideration) coincide. “Disconcordance” was defined as when the treatment options provided by GMC tumor boards and WFO-provided treatment options did not coincide (Not Recommended or Not Available). The concordance rate was only for chemotherapy and radiotherapy, not endocrine therapy.

**RESULTS**

Table 1 lists the patients’ characteristics.

**Chemotherapy**

The Oncotype Dx test is an important factor in determining the treatment plan for chemotherapy in WFO. On the other hand, GMC has performed the Endopredict test instead of Oncotype Dx. Therefore, Oncotype Dx and Endopredict tests were assessed as they are the same. Among the 147 patients who received adjuvant chemotherapy, the concordance rate was 93% (136/147) (Fig. 1). Six patients were not concordant due to the insurance problems. One patient who was not concordant was recommended CMF (cyclophosphamide+methotrexate+5-FU) regimen by WFO but chemotherapy was not performed due to old age and history of heart disease. One patient with stage I breast cancer was not
Radiotherapy

Radiotherapy as proposed by WFO has three methods: hypofractionated whole breast, conventionally fractionated whole breast with/without comprehensive regional nodes, and conventionally fractionated chest wall with comprehensive regional nodes. There are four types of radiotherapy in the GMC: breast only, breast with supraclavicular fossa, breast with supraclavicular fossa and posterior axilla boost, or breast with supraclavicular fossa, posterior axilla boost and internal mammary lymph nodes. Of the 155 patients, two patients failed to undergo radiotherapy due to follow-up loss after chemotherapy and nine patients had neoadjuvant chemotherapy. As a result, the data of 144 patients were examined.

The concordance rate of radiotherapy between WFO and GMC was 99% (143/144); one patient was unable to perform radiotherapy due to pregnancy. The overall agreement rate was as high as 99%, but there were some differences in the implementation.

WFO and GMC tumor boards. WFO recommended palbociclib but due to weakly estrogen receptor (ER) positive, progesterone receptor (PR) negative, GMC oncologist recommended AC (doxorubicin+cyclophosphamide) followed by paclitaxel and the patients chose GMC treatment.

Of a total 156 patients, nine patients underwent neoadjuvant chemotherapy, and seven of them had a concordance rate of 78% which is in agreement with the treatment options proposed by WFO and the GMC tumor boards. In nine cases of neoadjuvant chemotherapy, two cases were discordant with the proposal of

Table 2. Discordant cases in chemotherapy

<table>
<thead>
<tr>
<th>Case</th>
<th>Stage</th>
<th>Pathology</th>
<th>ER/PR/her2</th>
<th>WFO</th>
<th>GMC</th>
<th>Implement</th>
<th>Cause</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>T1N0M0</td>
<td>IDC</td>
<td>+/-/-/+</td>
<td>TH+H</td>
<td>CMF+H</td>
<td>CMF+H</td>
<td>Insurance</td>
</tr>
<tr>
<td>2</td>
<td>T1N0M0</td>
<td>IDC</td>
<td>+/-/+/-</td>
<td>AC+T+H</td>
<td>CMF+H</td>
<td>CMF+H</td>
<td>Insurance</td>
</tr>
<tr>
<td>3</td>
<td>T1N0M0</td>
<td>IDC</td>
<td>+/-/-/-</td>
<td>TH+H</td>
<td>CMF+H</td>
<td>CMF+H</td>
<td>Insurance</td>
</tr>
<tr>
<td>4</td>
<td>T1N0M0</td>
<td>IDC</td>
<td>+/-/+/-</td>
<td>TH+H</td>
<td>CMF+H</td>
<td>CMF+H</td>
<td>Insurance</td>
</tr>
<tr>
<td>5</td>
<td>T2N0M0</td>
<td>ILC</td>
<td>+/-/-/-</td>
<td>AC+T</td>
<td>CMF</td>
<td>CMF</td>
<td>Insurance</td>
</tr>
<tr>
<td>6</td>
<td>T1N0M0</td>
<td>IDC</td>
<td>+/-/-/-</td>
<td>AC+T</td>
<td>CMF</td>
<td>CMF</td>
<td>Insurance</td>
</tr>
<tr>
<td>7</td>
<td>T1N0M0</td>
<td>ILC</td>
<td>+/-/-/-</td>
<td>CMF</td>
<td>No chemotherapy</td>
<td>No chemotherapy</td>
<td>Oncologist’s opinion</td>
</tr>
<tr>
<td>8</td>
<td>T1N0M0</td>
<td>ILC</td>
<td>+/-/-/-</td>
<td>CMF</td>
<td>No chemotherapy</td>
<td>No chemotherapy</td>
<td>Oncologist’s opinion</td>
</tr>
<tr>
<td>9</td>
<td>T2N0M0</td>
<td>ILC</td>
<td>+/-/-/-</td>
<td>AC+T</td>
<td>CMF</td>
<td>No chemotherapy</td>
<td>Patient’s choice</td>
</tr>
<tr>
<td>10</td>
<td>T2N0M0</td>
<td>ILC</td>
<td>+/-/-/-</td>
<td>AC+T</td>
<td>AC</td>
<td>AC</td>
<td>Pregnancy</td>
</tr>
<tr>
<td>11</td>
<td>T1N1M0</td>
<td>IDC</td>
<td>+/-/-/-</td>
<td>AC+T</td>
<td>TC</td>
<td>AC+T</td>
<td>Patient’s choice</td>
</tr>
</tbody>
</table>

ER, estrogen receptor; PR, progesterone receptor; her2, human epidermal growth factor receptor 2; WFO, Watson for Oncology; GMC, Gachon University Gil Medical Center; IDC, invasive ductal carcinoma; ILC, invasive lobular carcinoma; TH, paclitaxel+trastuzumab; H, trastuzumab; AC, doxorubicin+cyclophosphamide; T, paclitaxel; CMF, cyclophosphamide+methotrexate+5-FU; TC, taxol+cyclophosphamide.

Fig. 2. Discordant cases in chemotherapy.
Discussions

In this study, the concordance rate of chemotherapy was high. In the 11 patients who were not concordant, six patients complied with the treatment proposed by GMC due to insurance problems, and one patient complied with the GMC treatment because of old age. Another patient did not follow the treatment proposed by WFO due to pregnancy and one patient was not treated with chemotherapy because it was not considered to be clinically necessary. One patient selected the treatment option of WFO and one selected the GMC treatment options.

One of the drawbacks of comparing the chemotherapy concordance rates in this study was the synchronization of the recurrence risk. Oncotype Dx was required in WFO but we are using Endopredict test in GMC. Therefore, they were synchronized and used in the study [7]. According to a study comparing Endopredict test and Oncotype Dx risk groups, the agreement rate was moderate to 76%. In addition, patients in a low-risk group of Oncotype Dx accounted for a high risk in Endopredict test in 18% [8]. Since the risk group according to Endopredict test was synchronized with the risk group of Oncotype Dx, we could consider the high-risk patients by the Oncotype Dx were not downgraded to the low-risk patients. The problem should be resolved later if the WFO includes the Endopredict test in the patient information setting.

Compared to adjuvant chemotherapy, the concordance rate of neoadjuvant chemotherapy was lower than that of adjuvant chemotherapy because the chemotherapy regimens proposed by WFO in advanced cases are more not covered by insurance.

In the case of radiotherapy, hypofractionated whole breast radiotherapy was less frequently performed in GMC. Therefore, when the treatment of WFO and GMC were concordant, most cases were considered as “For Consideration” options proposed by WFO.

The WFO decision-making process, which presents the treatment options as soon as we input patient’s data related to the disease could be more economically efficient, rather than the tumor board, which is composed of multiple doctors. However, the roles of WFO are a proposal for treatment and assisting a doctor. Future artificial intelligence services will be developed to fill doctor’s deficiencies and make better choices. At present, the high concordance rate of GMC and WFO is not the ultimate success of artificial intelligence, but it shows the possibility of development of better artificial intelligence.

Through this study, we found most treatment options proposed by WFO were highly reliable. If the patient’s personal information input is upgraded, it may be possible to obtain better results for individualized treatment. The weakness of this study is that the number of patients included in the study is small and the study period is short. Hence the recurrence rate could not be confirmed because the WFO had just started. Further studies considering these points will be needed.

Conflict of Interest

No potential conflict of interest relevant to this article was reported.

References