Original Article

Use of immunohistochemistry for detection of epidermal growth factor receptor mutation status in lung adenocarcinoma

Niharika Shah¹, Helmut Popper², Smriti Karki¹, Deebya Raj Mishra¹

¹Department of Pathology, B.P. Koirala Institute of Health Sciences, Dharan, Nepal
²Medical University of Graz, Graz, Austria

Keywords:
Adenocarcinoma; Epidermal growth factor receptor; Immunohistochemistry;

ABSTRACT

Background: Since the advent of targeted therapy in lung cancer, in settings where it is not possible to send for molecular testing of lung adenocarcinoma, immunohistochemistry for EGFR mutation-specific antibodies can be used as an alternative for detection of EGFR mutation.

Materials and Methods: 50 lung adenocarcinoma cases were screened at the Medical University of Graz. 19 cases in which molecular test as well as immunohistochemistry were positive for EGFR mutation. Cases where immunohistochemistry results and molecular test for the E-746-A750 deletion and the L858 mutation were positive, were considered true positives. Similarly, false positives, true negatives, and false negatives were determined.

Results: The mean age of the patients was 78.6 yrs. Among 19 cases, 7 were positive for E-746-A750 deletion (7/19, 36%), among which 4 cases also showed positivity with IHC. 4 were positive for the L858 mutation (4/19, 21%), among which 3 also showed positivity with immunohistochemistry. The rest of the 8 cases were positive for EGFR mutation in other loci. The sensitivity and specificity of the immunohistochemistry test of antibody specific to E-746-A750 exon 19 deletion was 100% and 80% respectively. The sensitivity and specificity of the IHC test of Ab specific to L858 mutation was 75%, and 100% respectively.

Conclusion: Our results have been comparable to previous studies. However, our sample size was a limitation. It can still be concluded that immunohistochemistry can be a diagnostic option in low resource settings, and can aid in ensuring that patients with a positive antibody test get targeted therapy.

INTRODUCTION

The advent of targeted therapy has revolutionized the landscape of lung cancer management. Tyrosine kinase receptors selective for epidermal growth factor (EGFR) receptor mutations in Non-small cell lung cancers (NSCLC) being an example.¹ Molecular testing for these specific mutations is emerging as the standard clinical practice.⁵ 10-50% of NSCLCs harbor an activating mutation in the tyrosine kinase domain of EGFR,⁶ among which approximately 90% include deletions of exon 19 and point mutations involving exon 21.⁷ This can be detected by DNA sequencing and real-time polymerase chain reaction (RT-PCR) but are limited by their low sensitivity, increased cost...
and turnaround time, and procedure complexity.\textsuperscript{5,6}

In settings where it is not possible to send for molecular testing or is unaffordable, immunohistochemistry (IHC) for EGFR mutation-specific antibodies can be used as an alternative for detection of EGFR status in NSCLC.\textsuperscript{10,11}

Jain D et al in their study included 206 biopsies of primary lung ADC on which EGFR mutation-specific antibodies against del E746-A750 and L858R were used. 26.6\% of patients showed positive IHC results and resolution melting analysis (HRM) results were available in 14 patients which showed EGFR mutations in correspondence with del E746-750 or L858R in 64.2\%. A concordance of 85.7\% was established between molecular mutation and IHC which proved that although the number tested was small, the concordance observed was good between molecular EGFR mutation and IHC expression.\textsuperscript{4}

Seo AN et al\textsuperscript{4} in their study enrolled a cohort of 240 consecutive patients with surgically resected lung adenocarcinomas on whom mutant EGFR protein expression was assessed by IHC using specific antibodies to the 2 major forms of EGFR mutations. Both antibodies (anti-EGFR E746-A750 del antibody and anti-EGFR L858R antibody) showed high specificity (99.0\% and 89.7\%, respectively) and sensitivity (70.6\% and 80.4\%, respectively). Although each antibody showed relatively high specificity, some EGFR-mutant cases were not detected by the mutation-specific antibodies. Various forms of exon 19 deletions, except E746 A750, were rarely detected by the mutant-specific antibody. They thus concluded that IHC-negative cases require further molecular analysis to confirm the presence of EGFR mutations.\textsuperscript{7}

\textbf{MATERIAL AND METHODS}

50 cases of biopsy and immunohistochemistry-proven lung adenocarcinoma (TTF-1 positive, p40 negative) were initially screened at the Medical University of Graz. Ethical clearance was obtained from the institutional ethical review board Medical University of Graz.

From these 50 cases, we selected cases in which genetic mutation analysis was done and the test result was positive for EGFR mutation. IHC analyses of EGFR mutation-specific antibodies were done on these cases on paraffin blocks cut to a thickness of 4 microns for immunostaining. Two primary antibodies delE746-A750 mutation-specific monoclonal antibody (EGF receptor, E746-A750del Specific, D6B6, XP Rabbit mAb, Cell Signal) and L858R mutation-specific monoclonal antibody (EGF Receptor, L858 mutant specific, 43B2, Rabbit mAb, Cell Signal) were used for immunostaining. The analysis was done on a tissue microarray and cases with tissues that were lost, or necrosed were not included.

The IHC staining was reported as either positive, negative, or inconclusive (depending on whether there was any neoplastic lung tissue or not or if the sample used for IHC was necrosed). The result of the IHC tests was assessed by two pathologists independently. Eventually, we included the 19 cases in which genetic mutation analysis was done and the test results were positive for EGFR mutation and IHC for mutation-specific antibody was also performed. Among these 19 cases, in some, the IHC results were inconclusive with either the antibody for E746-A750 deletion or with the antibody for the L858R mutation. However, in all of these cases, at least one antibody showed conclusive results.

The statistical analysis was performed based on these 19 cases. Collected data were entered into Microsoft Office Excel software. Data analysis was done by using SPSS (Statistical Package for Social Sciences) version 20. The data was appropriately coded. For Descriptive Statistics: Percentage (%), ratio, mean and standard deviation were calculated. For inferential statistics: the Chi-square test and proportion test were used to find out the significant association between the variables. For diagnostic measurement: sensitivity, specificity, positive predictive value, negative predictive value was calculated.

Cases in which the IHC results and the molecular test both for the E-746-A750 deletion were positive, were considered true positives and the same for the L858 mutation. Cases in which the IHC result was positive with the antibody for E-746-A750 deletion but molecular test for mutation was negative was considered as false positives, and the same for the L858 mutation. Similarly, the true negatives and false negatives were also determined.

\textbf{RESULTS}

We screened 50 cases of lung adenocarcinoma and selected 19 cases suitable for the study between November 5 2018 to November 30, 2018 (fig 1). The mean age of the patients was 78.6 yrs. Out of the 19 cases that were positive for EGFR mutation, the histopathological diagnosis of Adenocarcinoma with a predominant acinar pattern was most common (13/19).

Among the 19 cases, 7 were positive for the E-746-A750 deletion (7/19, 36\%) (fig. 3) and 4 were positive for the L858R mutation (4/19, 21\%) (fig. 4) by molecular tests (MT). The rest of the 8 cases were positive for the EGFR mutation in other loci. Out of 7 cases (7/19, 36\%) that were positive for E-746-A750 exon 19 deletion on genetic analysis, 4 cases also showed positivity with antibodies specific to this genetic defect (true positives). In the remaining 3, IHC results were inconclusive due to no neoplastic lung tissue or necrosis. All 7 cases were negative on IHC with an antibody specific to the L858R mutation. Out of 4 cases (4/19, 21\%) that were positive for L858R exon 21 mutation on genetic
analysis, 3 also showed positivity with antibodies specific to this genetic defect (true positives). However, one case showed negative IHC results despite the positive mutational status of genetic analysis (false negative). All 4 of these cases were negative for IHC with an antibody specific to E746-A750 deletion. (Table 1)

Out of the 8 cases that were positive for EGFR mutation at other loci, 2 cases showed positivity for antibodies specific to E746-A750 del (2 false positives). However, none of these cases with EGFR mutation positivity at other loci showed positivity with antibodies specific to L858 mutation (no false positives). 8 cases were negative for E-746-A750 deletion by MT and negative on IHC by E-746-A750 deletion specific antibody (true Negatives). 13 cases were negative for L858 mutation by MT and negative on IHC by L858 mutation-specific antibody (true Negatives). (Table 2 and 3)

The sensitivity of the IHC test of Ab specific to E-746-A750 exon 19 deletion was 100%, Specificity of the IHC test of Ab specific to E-746-A750 exon 19 deletion was 80%. The positive predictive value was 66.7%, and the Negative Predictive Value was 100%. The sensitivity of the IHC test of Ab specific to L858 mutation was 75%, Specificity of the IHC test of Ab specific to E-746-A750 exon 19 deletion was 100%. The positive predictive value was 100%, and the Negative Predictive Value was 93%.

DISCUSSION

Two types of mutations account for approximately 90% of mutated cases: a specific point mutation, L858R, that occurs in exon 21 and short in-frame deletions in exon 19.12,13 We observed in our study that the most common pattern seen was a predominant acinar pattern among all cases of Adenocarcinoma with EGFR mutation-positive status (13/19, 68%). Similar findings were found in some other studies.14 This could indicate that perhaps the acinar predominant pattern is associated with a positive EGFR mutation status more often and holds an intermediate prognosis.15

In low resource settings, molecular genetic tests are generally not available to patients. Getting test results by sending samples abroad is very expensive. Thus, it becomes imperative to develop a routine diagnostic test that is much more accessible and affordable. The IHC test of Ab specific to E746-A750 deletion and L858R exon 21 mutation offers a promising alternative. The high sensitivity and specificity of this test make it a valuable tool for the diagnosis of EGFR mutations, allowing for earlier identification of patients who may benefit from targeted therapies.

Table 1: IHC results with Antibodies specific to E746-A750 exon 19 deletion and L858R exon 21 mutation

<table>
<thead>
<tr>
<th>Histopathological diagnosis</th>
<th>IHC with E746-A750 exon 19 deletion specific Ab</th>
<th>IHC with L858R exon 21 mutation-specific Ab</th>
<th>Mutation status</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adenocarcinoma, acinar, papillary</td>
<td>Positive</td>
<td>Negative</td>
<td>E746-A750 del exon 19</td>
</tr>
<tr>
<td>Adenocarcinoma, acinar, solid</td>
<td>Negative</td>
<td>Negative</td>
<td>L858R exon 21</td>
</tr>
<tr>
<td>Adenocarcinoma</td>
<td>Positive</td>
<td>Negative</td>
<td>E746-A750 exon 19</td>
</tr>
<tr>
<td>Adenocarcinoma, acinar</td>
<td>Inconclusive</td>
<td>Negative</td>
<td>E746-A750 exon 19</td>
</tr>
<tr>
<td>Adenocarcinoma, acinar, solid</td>
<td>Inconclusive</td>
<td>Negative</td>
<td>E746-A750 exon 19</td>
</tr>
<tr>
<td>Adenocarcinoma, acinar</td>
<td>Negative</td>
<td>Positive</td>
<td>L858R exon 21</td>
</tr>
<tr>
<td>Adenocarcinoma, acinar, papillary</td>
<td>Negative</td>
<td>Positive</td>
<td>L858R exon 21</td>
</tr>
<tr>
<td>Adenocarcinoma</td>
<td>Positive</td>
<td>Negative</td>
<td>E746-A750 exon 19</td>
</tr>
<tr>
<td>Adenocarcinoma, acinar</td>
<td>Positive</td>
<td>Negative</td>
<td>E746-A750 exon 19</td>
</tr>
<tr>
<td>Adenocarcinoma, acinar, papillary</td>
<td>Negative</td>
<td>Positive</td>
<td>L858R exon 21</td>
</tr>
<tr>
<td>Adenocarcinoma, lepidic, tubular, papillary</td>
<td>Inconclusive</td>
<td>Negative</td>
<td>E746-A750 exon 19</td>
</tr>
</tbody>
</table>

Data expressed as mean ± SD
more cost-effective, gives a relatively rapid diagnosis, and is also able to aid in making faster treatment decisions.

IHC has been used in pathology laboratories for a long time and helps the pathologist make a rapid diagnosis, is easy to interpret, and is cost-effective as well.14,15 Thus, IHC tests could at least be used to screen for mutations in the EGFR receptor in NSCC in low resource settings. However, IHC for total EGFR seems to correlate poorly with the presence of the mutation in the same, and is thus not recommended for selection of treatment with a tyrosine kinase inhibitor.18 Though the sample size for this study is small, it nevertheless shows that mutation-specific antibodies can be used for rapid screening of the 2 most common EGFR mutations, E-746-A750 exon 19 deletion, and L858 exon 21 mutation. Clone D6B6 was used to detect E746-A750 exon 19 deletion with a sensitivity of 75%, a specificity of 100%, PPV of 100%, and NPV of 93%. Clone 43B2 was used to detect L858 exon 21 mutation with a sensitivity of 100%, a specificity of 80%, PPV of 67%, and NPV of 100%.

Most other studies show comparable results, however, the specificity of the present study with regards to a mutation-specific antibody for L858 mutation seems to be slightly on the lower side. This could be due to the small sample size of the study. Only 11 cases showed the 2 most common mutations, among which only 4 cases showed L858 mutation by genetic analysis, out of which 3 were positive by mutation-specific antibodies. However, sensitivity was extremely high (100%), as none of the cases that were negative for the mutation by genetic analysis showed a positive IHC test result by the mutation-specific antibody.

One of the disadvantages of these mutation-specific antibodies is that they can detect only specific mutations and will miss the other EGFR mutations, as can be seen in Table 2. However, in low resource settings, it could still be used as a screening tool. Our study shows a very high sensitivity for L858 mutations which is in contrast to other studies, and though the sample size is small it can still justify its use in lower-income countries with a high prevalence of lung cancer, like in Nepal. One of the caveats we can apply is, if both the mutation-specific antibodies are negative, we could strongly advocate for molecular genetic testing. The high specificity (100%) observed with the E746-A750 deletion specific antibody is comparable to the previous studies.19–21 We have not commented on the pattern of the IHC positivity in our study, whereas some other studies have
recommended that, unless a strong and homogenous membranous and cytoplasmic positivity was seen, it did not correlate well with the presence of the specific mutation. Some studies have mentioned that a staining pattern of 1+ is associated with more false positives, and a staining pattern of 2+ and 3+ is indicative of true positives. The two false-positive cases that were seen in our study with the E746-A750 deletion specific antibody could probably be because of that.

CONCLUSIONS

Several studies have been done previously to evaluate the sensitivity and specificity of these mutation-specific antibodies to detect the EGFR mutation status in a cost-effective manner. The present study more or less has come up with comparable results. However, the sample size was quite small, and the specificity of IHC with an antibody specific for the L858 mutation was quite low as opposed to other studies, at the same time the sensitivity for the same was very high (there were no false positives). We were also unable to follow patients for treatment response. A larger sample size is needed to give a stronger recommendation.

Acknowledgement

The authors would like to acknowledge the Union for International Cancer Control (UICC) for financial help.

Conflict of interest: None

REFERENCES


DOI: 10.3126/jpn.v10i2.30170