Overexpression of MicroRNA 21 in Cervical Carcinoma with Lymph Node Metastasis

N.W. Armerinayanti¹, I.M. Bakta², I.G. Alit Artha³, I.A.I. Wahyuniari⁴ and Samuel Widodo¹

¹Anatomic Pathology Department, Faculty of Medicine, Warmadewa University, Indonesia.
²Internal Medicine Department, Faculty of Medicine Udayana University, Indonesia.
³Anatomic Pathology Department, Faculty of Medicine Udayana University, Indonesia.
⁴Histologic Department, Faculty of Medicine Udayana University, Indonesia.
*Corresponding Author E-mail: armerinayantipranata@gmail.com

https://dx.doi.org/10.13005/bpj/2628

(Received: 05 December 2022; accepted: 16 January 2023)

Cervical carcinoma being the second common cancer in women in Indonesia, as well as in Bali, and mostly patients are diagnosed at an advanced stage with having metastases. The lymphatic pathway (lymph nodes) is the most frequent route for cervical cancer metastases. MicroRNA is a novel invention for predicting the biological behavior of cervical carcinoma and has the potential to act as the foundation for targeted therapy for cervical cancer. Several microRNA profiles, including microRNA 21, microRNA 126, and microRNA 143, were discovered to regulate the biological activity of cervical cancer. However, no studies have established a correlation between the expression of one of these microRNAs and the incidence of lymph node metastases in cervical cancer. This study aims to analyze whether overexpression of microRNA 21 is a risk factor for lymph node metastases in cervical cell carcinoma. Collected data was descriptively analyzed using the chi-square test with a p-value<0.05 and 95% CI. The results showed that microRNA 21 was significantly overexpressed in cervical carcinomas with lymph node metastases compared to those without lymph node metastases, representing a 19-fold increased risk for lymph node metastases. This can be influenced by the activity of microRNA 21 on several signaling pathways, such as Phosphatase and Tensin Homolog (PTEN), Programmed Cell Death Protein 4 (PDCD4), and Tissue inhibitor of metalloproteinase 3 (TIMP3), that affect the progression, invasion capacity, and metastasis of tumor cells.

Keywords: Cervical Carcinoma; lymph node; metastases; microRNA 21 Overexpression; Risk Factor.
In 2008, the Indonesian Cancer Foundation (YKI) reported that 52 million Indonesian women were at risk for cervical carcinoma and that 36% of all cancer patients were women with cervical carcinoma.\textsuperscript{1,2} Despite undergoing treatment, the mortality rate for cervical carcinoma remains about 90%, with the majority of patients dying of metastases. Metastasis is alarming for cancer patients since it represents the tumor’s aggressiveness and is related to a poor prognosis. The lymphatic pathway (lymph nodes) is the most common route for cervical carcinoma metastases.\textsuperscript{2,3}

In cervical cancer, determining the status of the inguinal lymph nodes is crucial because it serves as a prognostic indicator of distant metastases, recurrence, and appropriate therapy. However, the status of the lymph nodes cannot always be determined because clinicians do not always perform lymphadectomy. Determining the prognostic characteristics related with metastases is critical, as this will affect future treatment. Classic prognostic variables used to predict the occurrence of metastases include tumor size, grade, stage, lymphovascular invasion, and resection margins, with researchers obtaining varying findings. Therefore, new measures that correlate with the risk of metastasis in cervical cancer are required. Metastases of the lymph nodes can only be identified through histopathological examination of radical surgical tissue.\textsuperscript{1,3,4} Predicting the possibility of lymph node metastases can help with the prevention of lymph node metastases and also in determining whether the clinician will perform surgery and if the patient requires radical action.\textsuperscript{5,6}

Several parameters have been developed that could also assist clinicians in predicting the possibility of lymph node metastasis development.\textsuperscript{7,8} One of the most investigated biomolecular parameters is the expression profile of microRNAs (miRNAs), a class of endogenous non-coding RNAs that act as negative regulators/suppressors of protein-coding gene expression by selectively binding to the base complement 3 untranslated region (3-UTR) of messenger RNAs (mRNAs). Several miRNA profiles, including miR-21, miR-126, and miR-143, have been identified as playing a role in determining the biological behavior of cervical carcinoma.\textsuperscript{9,10}

A recent study has investigated the important role of miR-21 in cervical carcinoma. In cervical carcinoma, miR-21 is classified as an oncogene miR because it stimulates cell proliferation and migration. The expression levels of miR-21 and ZEB1 (Zinc finger E-box binding homeobox 1 transcription factor) were significantly higher in cervical carcinoma tissue with lymph node metastases than in normal cervical tissue, according to previous research. In a recent study, it was discovered that the upregulation of miR-21 in cervical carcinoma tissue modulates EMT (Epithelial-Mesenchymal Transition) activation, which in turn induces lymphatic metastasis. In 2020, Sánchez conducted opinion on the overexpression of miR-21 in cervical cancer cell lines HeLa and SiHa, which led to the upregulation of mesenchymal cell markers vimentin and N-cadherin and the decreased expression of epithelial cell marker E-cadherin.\textsuperscript{11} The EMT is the ability of epithelial cells to transform from immobile to motile mesenchymal progenitor cells. This pathway is essential for development (type 1), normal tissue regeneration or pathological fibrosis (type 2), and cancer cell metastatic transformation (type 3). Type 3 EMT is essential for tumor progression to metastasis, and both reactivation and dedifferentiation as well as activated cancer cells are generated to have an invasive and motile phenotype.\textsuperscript{4,10}

In addition, previous studies discovered a significant correlation between miR-21 overexpression with muscular infiltration, and parametrial invasion. Multiple studies have shown that elevated miR-21 expression is related to cell proliferation, tumor invasion, metastasis, angiogenesis, immunological response, and apoptosis inhibition in cervical cancer cases.\textsuperscript{5,6} The aim of this study was to investigate the role of miR-21 overexpression of lymph node metastases in cervical carcinoma.

**MATERIALS AND METHODS**

This study used a case-control study design to explore the correlation between miR-21 overexpression and the presence of lymph node metastases in cervical carcinoma. In addition,
RESULTS AND DISCUSSION

Case Distribution Based on Patient Clinical Data

The majority of cervical carcinoma cases (63.3%) were diagnosed in women younger than 50 years old, according to research data. This is congruent with findings from the International Agency for Research on Cancer (IARC) indicating that cervical carcinoma is one of the top three cancers among women aged 45. This may be due to the fact that sexual activity, a risk factor for cervical carcinoma, is most prevalent in this age group, which corresponds to the reproductive age range. Epidemiological studies indicate that HPV infection is most prevalent in women in their twenties, as determined by the detection of HPV DNA. It takes between 10 to 20 years from the presence of HPV infection to the development of cervical carcinoma, hence the prevalence of cervical cancer in the third and fourth decades is considerable.1,4,5

According to table 1, 70% of the tumors in this study group were larger than or equal to four centimeters. This relates to the selection of study samples that use radical surgical tissue, which is typically obtained from cervical carcinomas that are apparent to the naked eye (above stage IB1). In developing nations, such as Indonesia, cervical cancer is still diagnosed at a more advanced stage, where the size of the tumor exceeds 4 centimeters and already causes serious clinical symptoms such as profuse bleeding and spread to surrounding tissues such as the vagina and bladder.

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>n=30 (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (year)</td>
<td></td>
</tr>
<tr>
<td>&lt;50</td>
<td>19 (63.3%)</td>
</tr>
<tr>
<td>&gt;=50</td>
<td>11 (36.7%)</td>
</tr>
<tr>
<td>Tumor Size (cm)</td>
<td></td>
</tr>
<tr>
<td>&lt;4</td>
<td>9 (30%)</td>
</tr>
<tr>
<td>&gt;=4</td>
<td>21 (70%)</td>
</tr>
<tr>
<td>Lymph node metastases group</td>
<td></td>
</tr>
<tr>
<td>1) positive metastases</td>
<td>15 (50%)</td>
</tr>
<tr>
<td>2) negative metastases</td>
<td>15 (50%)</td>
</tr>
</tbody>
</table>

The data was descriptively analyzed using the chi-square test to determine the correlation between miR-21 overexpression and lymph node metastases in cervical carcinoma. The significance test was determined at $p<0.05$. Data precision is determined by a 95% Confident Interval (CI).
Correlation between Overexpression of MicroRNA-21 with Patient Clinical Data

Quantitative real-time PCR showed no significant association between miR-21 overexpression and the patient’s age or tumor size ($p>0.05$). This is consistent with findings from prior studies indicating that miR-21 expression is unaffected by age or the aging process. The miR-21 is a small non-coding RNA molecule whose biological control is correlated to its role in immunological activity, cell differentiation, and carcinogenesis in diverse types of tumors.\textsuperscript{12,13}

The contrary association was noticed between tumor growth and miR-21 overexpression ($p<0.05$). This is consistent with the findings of numerous studies indicating that miR-21 expression is implicated in several hallmarks of carcinogenesis, including tumor proliferative activity, tumor development, and the capacity to invade and metastasis.\textsuperscript{2,3}

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>n=30 (%)</th>
<th>miR-21 Overexpression (n)</th>
<th>$p$ value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (year)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;50</td>
<td>19 (63.3%)</td>
<td>12</td>
<td>0.592</td>
</tr>
<tr>
<td>$\geq50$</td>
<td>11 (36.7%)</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>Tumor Size (cm)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;4</td>
<td>9 (30%)</td>
<td>6</td>
<td>0.037</td>
</tr>
<tr>
<td>$\geq4$</td>
<td>21 (70%)</td>
<td>14</td>
<td></td>
</tr>
</tbody>
</table>

Table 2. Correlation between miR-21 overexpression and research sample clinical data

The contrary association was noticed between tumor growth and miR-21 overexpression ($p<0.05$). This is consistent with the findings of numerous studies indicating that miR-21 expression is implicated in several hallmarks of carcinogenesis, including tumor proliferative activity, tumor development, and the capacity to invade and metastasis.\textsuperscript{2,3}

Graph 1. Distribution of relative multiples of miR-21 expression throughout samples compared to internal control
Correlation Between Overexpression of MicroRNA-21 with Lymph Node Metastasis

In this study, miR-21 expression was evaluated in both sample groups using quantitative real-time PCR (cervical carcinoma with lymph node metastases and those without lymph node metastases). Normal cervical tissue samples were used as the negative control to analyze the multiple expression of miR-21 between the two groups in this investigation, with U6 sRNA serving as the internal control.

According to graph 1, both sample groups had a considerably higher multiple of expression compared to negative internal controls. In addition, miR-21 was detected in both sets of study samples, including those with lymph node metastases and those without lymph node metastases. It is known that the carcinogenesis of cervical carcinoma is consistently related to previous infections with high-risk HPV types 16 and 18. The miR-21 genomic locus resides in the FRA17B locus of chromosome 17q 23.2, which is one of the HPV 16 integration sites. MiR-21 expression in Cervical carcinoma is a marker for HPV 16 integration factor involvement. Because miR-21 is involved in the carcinogenesis process of cervical cancer, it may be deduced that, in general, cases of cervical cancer will have higher miR-21 expression than normal cervical tissue.

The activation of the PI3K and its target, AKT, and mTOR signalling pathways, which regulate differentiation, proliferation, and cell survival, play a role in the continuation of the cervical carcinoma carcinogenesis process. By activating the PI3K/AKT/mTOR signalling pathway, tumor cell proliferation is increased.

The expression of microRNAs increased significantly in the cervical carcinoma group with lymph node metastases compared to the group without lymph node metastases ($p=0.000; p<0.05$). The relative expression multiples in samples with lymph node metastases were higher than five times the expression of the reference control, hence they were classified as overexpressed.
miR-21 overexpression increases the risk of cervical carcinoma lymph node metastasis by 19-fold. This is congruent with results regarding the molecular mechanism of miR-21’s participation in inducing invasion and metastasis in a variety of solid carcinomas; thus, miR-21 is recognized as a metasmiR. Involved molecular mechanisms may include suppression of the activation of PDCD4, TIMP3, TPM1, SERPINB5, coding for Maspin, and PTEN, among others. If PDCD4 is blocked, there will be a decrease in JNK activation, resulting in an increase in BCL2-like (anti-apoptotic) proteins, which prevents apoptosis and eventually contributes to tumor cell proliferation and migration.14,15,16

Graph 2 further shows that PTEN activity can suppress the PI3K/PTEN/AKT signalling pathway and target AKT to modulate cellular processes such as cell growth, differentiation, proliferation, and migration. Overexpression of miR-21 can downregulate PTEN, hence maintaining the activation of the PI3K/PTEN/AKT signalling pathway, which affects tumor proliferation and progression. PTEN can also inhibit MMP-9 and MMP-2 expression by dephosphorylating FAK. Thus, inhibition of PTEN activity would result in the overexpression of matrix metalloproteinases such as MMP2 and MMP9, which promote migration, cellular invasion, and metastasis, including lymph node metastasis.1,2,5,10,17 In this study, the overexpression of miR-21 in lymph node metastasis samples is most likely influenced by the inhibition of this PTEN pathway.

CONCLUSION

It can be concluded that, miR-21 expression was significantly higher in cervical carcinomas with lymph node metastases compared to cervical carcinomas without lymph node metastases, furthermore, cervical carcinomas with lymph node metastases showed overexpression of miR-21. In cervical carcinoma, overexpression of miR-21 multiplies the risk of lymph node metastasis by 19-fold. This relates to the wide range of molecular roles of miR-21 in various signalling pathways of tumor progression, migration, and metastasis, such as inhibition of PTEN, which triggers anti-apoptotic activity, and activation of metalloproteinases such as MMP-9, which affect the emergence of EMT phenotypes in cervical carcinoma, thereby facilitating deeper and deeper invasion. In addition, this study also showed that miR-21 overexpression has an impact on tumor size, since there was a significant difference between tumors with a size of >4 cm and those with a size of <4 cm with regard to miR-21 overexpression.

ACKNOWLEDGEMENTS

We would like to express our deepest appreciation to the Biomolecular Laboratory of the Faculty of Medicine and Health Sciences at Warmadewa University and The Anatomical Pathology Laboratory at Balimed Denpasar Hospital, which has given the infrastructure and facilities necessary for this research to be conducted.

Conflict of Interest

There are no Conflict of Interest

Funding Source

This research was supported by grants from institutions provided by the Faculty of Medicine and Health Sciences at Warmadewa University. The grant number is 098/Unwar/FKIK/UP2M/UR-21/IV/2020.

REFERENCES


