Alternative polyadenylation is associated with lower expression of PABPN1 and poor prognosis in non-small cell lung cancer

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Key words
Alternative polyadenylation, lung cancer, PABPN1, prognosis, rmodel

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Funding Information
This work was supported by grants from the Takeda Science Foundation, and JSPS KAKENHI (22249017, 24692027, 25830098, 24791455, 23249045, 22249017, 24390326).

Received May 14, 2014; Revised June 16, 2014; Accepted June 24, 2014

Cancer Sci 105 (2014) 1135–1141
doi: 10.1111/cas.12472

Alternative polyadenylation (APA), which induces shortening of the 3′UTR, is emerging as an important phenomenon in gene regulation. APA is involved in development, cancer and cell proliferation. APA may lead to disruption of microRNA-mediated gene silencing in cancer cells via detachment of microRNA binding sites. We studied the correlation between the APA profile and the tumor aggressiveness in cases of lung cancer. We selected the top 10 genes showing significant 3′UTR shortening in lung cancer, using the package of the Bioconductor for probe-level analyses of expression microarrays. We established and evaluated the APA score by quantitative RT-PCR in 147 clinical specimens of non-small cell lung cancer and compared the results with the clinical outcomes and expression levels of APA-related genes, including PABPN1, CPEB1, E2F1 and proliferation markers (MKI67, TOP2A and MCM2). High APA scores were correlated with an advanced tumor stage and a poor prognosis (P < 0.001). Multivariate analysis identified the APA score as an independent prognostic factor (hazard ratio, 3.0; P = 0.03). Both lower expression of PABPN1 and higher expression of the proliferation markers were correlated with high APA scores and a poor prognosis, with suppression of PABPN1 exerting its influence independent of gain of the proliferation markers. Moreover, the APA score was correlated with the maximum standardized uptake value of the tumors on positron emission tomography (r = 0.53; P < 0.001). Our results indicate that the loss of PABPN1, a suppressor of APA, might promote tumor aggressiveness by releasing the cancer cells from microRNA-mediated gene regulation.

In addition to genetic alterations in cancer cells, post-transcriptional regulation of gene expression has recently been shown to play many important roles in human cancer. MicroRNAs are broadly conserved small non-coding RNAs that bind to elements within the 3′UTR of the target mRNAs and regulate gene expression by inhibiting mRNA translation or inducing mRNA degradation. MicroRNA expression is altered in human cancers. While some microRNAs have oncogenic functions, in most cases, loss of microRNA expression has been reported to be correlated with cancer aggressiveness. Suppression of microRNA biosynthesis is one of the ways by which cells may escape microRNA regulation. Decreased expressions of genes involved in microRNA biosynthesis, such as Drosha, Dicer and TARBP2, are associated with cancer aggressiveness. A few reports have suggested that silencing of microRNAs by DNA methylation is associated with tumor invasiveness. Alternative polyadenylation (APA), which induces detachment of microRNA binding sites, is another way by which cells escape microRNA-mediated gene silencing (Fig. 1). APA is emerging as an important player in gene regulation. More than half of mammalian genes use APA to generate multiple mRNA isoforms with different lengths of the 3′UTR. Besides escape from microRNA-mediated regulation, the shortening of the 3′UTR can influence mRNA nuclear export, localization in cytoplasm and non-microRNA-mediated changes in mRNA stability and translational efficiency. It has been shown that proliferating cells, embryonic cells, induced pluripotent stem cells and cancer cells express shorter transcripts. The correlation between the APA profile and tumor malignancy grade is still unclear.

Several factors have been reported to be involved in the regulation of APA. While E2F and CPEB1 enhance APA, PABPN1 has been shown to suppress APA. Therefore, we have evaluated whether the NSCLC specific APA profile and expression levels of APA-related genes, including proliferation markers such as MKI67, TOP2A and MCM2, in clinical tumor specimens might be correlated with a poor prognosis of non-small cell lung cancer (NSCLC).

Materials and Methods

Tumor samples. Clinical lung cancer samples were collected from NSCLC patients who underwent surgical resection with...
A curative intent at the University of Tokyo Hospital between June 2005 and May 2009. Total RNA was isolated using RNAiso Plus (TaKaRa, Shiga, Japan). The diagnoses were based on pathological evidence and the tumors were classified according to the seventh edition of the TNM classification for NSCLC.(32) Patients who underwent preoperative therapy were excluded. Normal lung tissue specimens were collected from non-smoking stage I NSCLC patients without recurrence. Informed consent was obtained from all the patients, and the study was conducted with the approval of the Institutional Ethics Review Committee.

For all patients, medical records were reviewed to extract the data on the clinicopathological characteristics. A disease status census was carried out every 6 months at our outpatient department. We chose freedom from recurrence (FFR) as the primary endpoint of the study. FFR was measured from the date of surgery until documentation of disease recurrence or metastasis, and in cases without relapse, any deaths due to causes other than lung cancer were censored.

In silico selection of alternative polyadenylation-indicator genes. In order to select genes showing significant 3'UTR shortening specifically in lung cancer, we used the Bioconductor package for probe-level analysis of expression microarrays (rmodel), as in Salisbury et al.(33) Rmodel divides a probe set into segments representing the sequence boundaries of transcribed regions that change by different amounts when two sample groups are compared. Microarray data obtained on Affymetrix (Santa Clara, CA, USA) HG-U133_Plus_2 array were downloaded from the Gene Expression Omnibus resistor. We used GSE10245 containing the expression profiles of 40 lung adenocarcinomas and 18 lung squamous cell carcinomas,(34) and GSM415386-8 containing the expression profiles of 3 normal lung tissues.(35) These profiles of normal lung tissue were selected from the dataset GSE16538 that consists of profiles of tissue samples of pulmonary sarcoidosis and normal lung tissue.(35) We reviewed the result of rmodel analysis and excluded clearly inadequate candidates (i.e. genes with no significant change of probe level and genes with probes that were mismapped).

Analysis of the alternative polyadenylation status. The APA status in the in silico-selected APA-indicator genes was analyzed by quantitative RT-PCR (qRT-PCR) using THUNDERBIRD SYBR qPCR Mix (Toyobo, Osaka, Japan), performed in triplicate. We calculated the proportion of the full-length transcripts to the total transcripts using the ddCT method. We designed primer set 1 based on the coding sequence and primer set 2 on the distal side of the 3'UTR. Primer set 2 was designed distal to the boundary between the proximal probe set and the distal probe set divided by the rmodel (Fig. 2). APA score was defined as the total number of the genes in which the proportion of the full-length transcripts was less than half of that in the normal lung tissue. Expressions of the APA-related genes (PABPN1, CPEB1, E2F1, and the three proliferation markers MKI67, TOP2A and MCM2) were also determined by qRT-PCR. HPRT1 was used as the internal control, and the primer sequences are shown in Tables S1 and S2.

Statistical analysis. Statistical analysis was carried out using the JMP 10 software (SAS Institute, Cary, NC, USA). The cutoff value of the APA score was determined by receiver-operating characteristic curve analysis for lung cancer recurrence. The relationships between the APA score, mRNA expression levels of the APA-related genes and the clinicopathological characteristics were analyzed using Student’s t-test, Welch’s method and the χ²-test. The relationship between the APA
scores and the maximum standardized uptake values (SUVmax) was evaluated by calculation of the Pearson correlation coefficient. FFR was estimated using the Kaplan–Meier method and compared by the log-rank test. A multivariate analysis of various prognostic factors was carried out using the Cox proportional hazards regression model.

Results

Identification of the alternative polyadenylation-indicator genes in non-small cell lung cancer. Using the rmodel, we selected 16 genes that showed 3'UTR shortening in more than 20 of the 40 lung adenocarcinoma specimens included within the GSE10245 dataset (Table S3). All 16 genes showed 3'UTR shortening in more than 10 of the 18 lung squamous cell carcinoma specimens within the GSE10245 dataset (Table S3). The APA status of the 16 genes was confirmed in 7 clinical lung cancer specimens and 8 normal lung tissue specimens (Fig. 3), and the top 10 genes which showed more marked APA in lung cancer were selected as the APA-indicator genes (C1orf52, DIEXF, ESYT2, HN1L, MUC20, NDFIP2, RBM33, SCAMP1, SMCA1 and SSR1). For subsequent analysis, we defined the APA score as the total number of genes that showed 3'UTR shortening among the 10 APA-indicator genes.

Correlation between high alternative polyadenylation score and a poor prognosis. We determined the APA scores of the 147 primary NSCLC specimens. The specimens were divided into two groups, the training set and the test set, to determine and validate the appropriate cutoff value of the APA score. The training set comprised 47 lung cancer samples and 4 normal lung tissue samples. The test set comprised 100 lung cancer samples and 6 normal lung tissue samples. The median follow-up period was 78 months for the training set and 58 months for the test set. The clinicopathological characteristics of the corresponding patient groups are shown in Table S4.

The most appropriate cutoff value of the APA score was determined to be 5 from the training set by receiver-operating characteristic curve analysis, with the same value verified as appropriate in the test set. The frequency of advanced disease, non-adenocarcinoma and adjuvant chemotherapy was higher in...
the high APA score group (Table 1). The high APA score was significantly associated with the presence of lymph node metastasis and vascular invasion as well as with the advanced T factor. These observations indicate that abnormal APA status accumulates during the course of tumor progression and the acquisition of an invasive phenotype. The prognosis of the patients in the high APA score group was significantly poorer (Fig. 4a). Univariate analysis using the log-rank test revealed the APA score, histological type, T factor, N factor, presence/absence of lymphatic invasion, presence/absence of vascular invasion and history of adjuvant chemotherapy as prognostic factors (Table 2). Multivariate analysis identified the APA score, N factor and age as independent prognostic factors (Table 3).

Correlation between high alternative polyadenylation status score and low PABPN1 expression. The APA scores were inversely correlated with the relative expression of PABPN1 (r = 0.53; P < 0.001), with the SUVmax values being higher in the high APA score group than those in the low APA score group (11.8 ± 8 vs. 5.6 ± 4.8, P < 0.001).

Table 1. APA score and clinicopathological characteristics

| Age (years) | High APA | Low APA | P | High APA | Low APA | P |
|-------------|----------|---------|---|----------|---------|---|
| <70         | 7        | 19      | NS | 13       | 35      | NS |
| ≥70         | 10       | 11      | 0.01 | 16       | 36      | 0.01 |
| Sex         |          |         |    |          |         |    |
| Male        | 16       | 16      | 0.01 | 19       | 33      | NS |
| Female      | 1        | 14      | 0.33 | 10       | 38      |    |
| Smoking habit |        |         |    |          |         |    |
| Negative history | 1       | 12      | 0.03 | 21       | 38      | NS |
| Positive history | 16      | 18      | 0.33 | 8        | 33      |    |
| Histological type |      |         |    |          |         |    |
| Adenocarcinoma | 11      | 25      | NS | 21       | 65      | 0.03 |
| Others      | 6        | 5       | 8   | 6        | 6       |     |
| T factor    |          |         |    |          |         |    |
| T1          | 4        | 16      | NS | 9        | 46      | 0.004 |
| T2, T3      | 13       | 14      | 0.01 | 20       | 25      |    |
| N factor    |          |         |    |          |         |    |
| N0          | 9        | 24      | NS | 18       | 66      | <0.001 |
| N1, N2      | 8        | 6       | 11  | 5        | 6       |     |
| Lymphatic invasion |      |         |    |          |         |    |
| Negative    | 10       | 24      | NS | 20       | 61      | NS |
| Positive    | 7        | 6       | 9   | 10       |         |     |
| Vascular invasion |      |         |    |          |         |    |
| Negative    | 6        | 21      | 0.04 | 12       | 57      | <0.001 |
| Positive    | 11       | 9       | 17  | 14       |         |     |
| History of adjuvant chemotherapy |      |         |    |          |         |    |
| Negative    | 10       | 23      | NS | 18       | 59      | 0.04 |
| Positive    | 7        | 7       | 11  | 12       |         |     |

APA, alternative polyadenylation; NS, not significant.

Table 2. Results of univariate analyses carried out to identify prognostic factors in patients with resected non-small cell lung cancer

| Training set (n = 47) | Test set (n = 100) |
|----------------------|-------------------|
| APA score ≥5         | 3.9               | 5.4               |
| Age ≥70 years        | 1.1               | 0.6               |
| Male                 | 3.5               | 1.7               |
| Positive smoking history | 2.6           | 2.3               |
| Non-adenocarcinoma   | 3.1               | 2.8               |
| T2, T3               | 13.7              | <0.001            |
| N1, N2               | 2.9               | 9.7               |
| Lymphatic invasion   | 3.3               | 5.8               |
| Vascular invasion    | 1.9               | 8.2               |
| Adjuvant chemotherapy | 2.5               | 4.8               |
| Loss of PABPN1†       | 3.7               | 5.3               |
| Loss of CPEB1†        | 0.7               | 3.4               |
| Gain of MKI67†        | 1.5               | 4.2               |
| Gain of TOP2A†        | 1.5               | 5.3               |
| Gain of MCM2†         | 2.9               | 2.2               |
| Gain of E2F1†         | 1.0               | 2.9               |

APA, alternative polyadenylation; NS, not significant.

Table 3. Results of multivariate analyses carried out to determine the independent prognostic factors in patients with resected non-small cell lung cancer

| Test set (n = 100) |
|-------------------|
| APA score ≥5       | 3.0               |
| T2, T3             | 2.6               |
| N1, N2             | 4.7               |
| Age ≥70 years      | 0.3               |
| Male               | 1.0               |
| Positive smoking history | 2.2           |
| Non-adenocarcinoma | 1.6               |

APA, alternative polyadenylation; NS, not significant.

Table 3. Results of multivariate analyses carried out to determine the independent prognostic factors in patients with resected non-small cell lung cancer

| Test set (n = 100) |
|-------------------|
| Loss of PABPN1†    | 3.5               |
| Loss of CPEB1†     | 2.1               |
| Gain of MKI67†     | 1.7               |
| Gain of TOP2A†     | 2.1               |
| Gain of MCM2†      | 0.4               |
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| Gain of E2F1†      | 3.9               |

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higher expression of the proliferation markers (Fig. 5b). Recent studies have shown that E2F (24) and CPEB1 (25) enhance APA. We did not observe any significant relation between the APA scores and E2F1 expression. The high APA score group tended to show lower expression levels of CPEB1...
than the low APA score group. Moreover, the expression of CPEB1 was markedly suppressed in the lung cancer specimens (Fig. 5a).

We analyzed the correlations between the expression levels of the APA-related genes and the patient prognosis. The prognosis of the patients in the low PABPN1 expression group was significantly poorer (Fig. 4b). Univariate analysis revealed that loss of PABPN1 and CPEB1 and gain of MKI67, TOP2A, and MCM2 were correlated with a poor prognosis (Table 2). Multivariate analysis identified loss of PABPN1 and gain of E2F1 as independent prognostic factors (Table 3).

Discussion

In our study, high APA scores were correlated with advanced tumor stages and a poor prognosis of the patients. We also determined that loss of PABPN1 was correlated with high APA scores and a poor prognosis, independent of the expression of proliferation markers. Moreover, the APA scores were correlated with the SUVmax values of the tumors.

The mechanism of APA is currently under intense investigation. The 3'UTR shortening associated with APA may be the result of enhanced cell proliferation or deregulation of APA-regulator genes, such as PABPN1 and CPEB1, although it remains unclear whether the proposed mechanisms work in cancer cells. In our lung cancer samples, both lower expression of PABPN1 and higher expression of the proliferation markers were correlated with the APA scores and a poor patient prognosis. Importantly, suppression of PABPN1 was independent of the expression of the proliferation markers, suggesting that the pathway of APA induced by the loss of PABPN1 is independent of the pathway of APA in proliferating cells. Low expression of CPEB1 was correlated with high APA scores. This result is in conflict with a previous report which revealed that CPEB1 mediates APA. CPEB1 expression was markedly suppressed in the lung cancer specimens. Thus, it is likely that CPEB1 does not induce APA, at least in lung cancer.

The finding of a correlation between the SUVmax and the APA score is of interest. The SUVmax evaluated by PET represents the tumor uptake of fluorodeoxyglucose and reflects the tumor cell activity. Further interventional research is needed to clarify the causal relationship between APA and enhancement of the tumor cell proliferative activity.

Until date, there have been no consistent reports on how APA might affect the expression levels of the APA-regulated genes. Theoretically, mRNAs with shorter lengths of the 3'UTR are depleted of microRNA binding sites, resulting in increased mRNA expression. This is true for some APA-regulated genes with let-7 microRNA binding sites. In our study, however, expression levels of the APA-indicator genes tended to be lower in the lung cancer samples than in the normal lung tissue samples (Fig. 6). There are three possibilities to explain this discrepancy. First, the expression of the APA-indicator genes

![Fig. 4.](image)

| Training set | Test set |
|--------------|----------|
| ![Graph](image) | ![Graph](image) |

### Discussion

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was relatively suppressed as a result of upregulation of other overexpressed genes in the cancer cells. It should be noted that the APA-indicator genes are mere indicator genes, and are not oncogenes by themselves. Next, the mRNA expressions might be suppressed by negative feedback as a result of the significant increase of the protein production from mRNAs with shorter lengths of the 3'UTR. Finally, some upregulated microRNAs (oncomiRs) in the cancer cells may bind only to the target gene transcripts with full-length 3'UTR, and may result in increasing the proportion of target gene transcripts with short 3'UTR.

In most recent studies on the APA, the APA profile has been evaluated by expression microarrays or high-throughput sequencing. Correlation of shorter lengths of the 3'UTR with a poor prognosis has been shown in breast and lung cancers by analysis of Affymetrix chips. (36) Certainly, analysis of a few transcripts is insufficient to estimate APA, which has a genome-wide influence; however, these methods are still too expensive for clinical application. To resolve this dilemma, we selected APA-indicator genes, using the rmodel, that showed significant 3'UTR shortening specifically in lung cancer. We could evaluate the APA profile easily and efficiently by performing qRT-PCR of these selected genes. The APA profile of the 10 genes (KRAS, N Ras, CCND1, CCND2, E2F1, FGFR2, IMP1, MAPKAPK3, PDRG1 and PDXK) selected based on the results of the past studies (22, 26, 37) were evaluated in the same way (Fig. 3). These genes were associated with a lower frequency of 3'UTR shortening as compared to the APA-indicator genes and were not useful as indices of APA in the clinical lung cancer samples. It has been reported that CCND1, FGFR2 and IMP1 show APA in cancer cell lines, including lung cancer cell lines. (22) PDXK exhibits marked APA in clinical colorectal cancer, (37) and CCND2, E2F1, MAPKAPK3 and PDRG1 show APA in an osteosarcoma cell line with PABPN1 knockdown. (26) Genes showing marked 3'UTR shortening may differ considerably depending on the primary site and on whether the sample is a cell line or a primary tumor. This result is consistent with a recent study carried out using the rmodel, which revealed that the characteristic signatures of 3'UTR shortening can distinguish tumor subtypes. (38) Rmodel-based selection of the APA-indicator genes showing 3'UTR shortening specifically in the target disease is very useful for simple and efficient evaluation of APA.
In conclusion, we revealed significant correlations between high APA scores, loss of PABPN1 and a poor patient prognosis in cases of lung cancer. Because PABPN1 is a recently identified suppressor of APA, this observed link suggests that loss of PABPN1 might promote tumor aggressiveness by releasing the tumor cells from microRNA-mediated gene regulation. The prognostic significance of APA and PABPN1 should be validated in a large independent cohort.

Acknowledgments
This work was supported by grants from the Takeda Science Foundation and JSPS KAKENHI (23390147, 24659267, 25830098, 24791455, 23249045, 22249017 and 24390326).

Disclosure Statement
The authors have no conflict of interest to declare.

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Supporting Information
Additional supporting information may be found in the online version of this article:
Table S1. PCR primer sequences of the selected alternative polyadenylation (APA)-indicator genes.
Table S2. PCR primer sequences of the alternative polyadenylation (APA)-related genes and HPRT1.
Table S3. Alternative polyadenylation (APA) profile of 16 genes that showed frequent 3'UTR shortening in lung cancer.
Table S4. Patient characteristics.