BRIEF REPORT

DNA methylation changes in biomarker loci occur early in cancer progression [version 2; peer review: 2 approved]

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Abstract
Tumor-specific DNA methylation can be used for cancer diagnostics and monitoring. We have recently reported a set of DNA methylation biomarkers that can distinguish plasma samples from lung cancer patients versus healthy controls with high sensitivity and specificity. Furthermore, the DNA methylation signal from the biomarker loci detected in plasma samples correlated with tumor size and decreased after surgical resection of lung tumors. In order to determine the timing of DNA methylation of these loci during carcinogenesis and thus the potential of the biomarkers to detect early stages of the disease we analyzed the DNA methylation of the biomarker loci in five precancerous conditions using available data from the GEO database. We found that the DNA methylation of the biomarker loci is gained early in carcinogenesis since most of the precancerous conditions already have biomarker loci hypermethylated. Moreover, these DNA methylation biomarkers are able to distinguish between precancerous lesions with malignant potential and those that stay benign where data is available. Taken together, the biomarkers have the potential to detect the earliest cancer stages; the only limitation to detection of cancer from plasma samples or other liquid biopsies is the timing when tumors start to shed enough DNA into body fluids.

Keywords
DNA Methylation, Cancer Biomarker, Epigenetics

Open Peer Review

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16 Dec 2019
report
report

1. Keith D. Robertson, Mayo Clinic, Rochester, USA
2. Judd C. Rice, University of Southern California, Los Angeles, USA

Any reports and responses or comments on the article can be found at the end of the article.
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Author roles: Vrba L: Conceptualization, Formal Analysis, Investigation, Methodology, Software, Validation, Visualization, Writing – Original Draft Preparation, Writing – Review & Editing; Futscher BW: Funding Acquisition, Writing – Review & Editing

Competing interests: B. Futscher is a co-founder of DesertDx, LLC

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First published: 16 Dec 2019, 8:2106 https://doi.org/10.12688/f1000research.21584.1
Introduction

Tumor cells have fundamentally different DNA methylation profile from normal cells of origin. Some of these differences are tumor-specific, i.e. do not occur in any normal cell types, and thus could be used for tumor DNA identification. Since tumors shed DNA into bloodstream or other body fluids, the detection of tumor-specific DNA methylation in these liquid biopsies could be utilized for non-invasive cancer diagnostics and monitoring. This initiated a search for cancer-specific DNA methylation biomarker loci and analysis of these loci in plasma samples and other liquid biopsies. We have previously described a large suite of cancer-specific DNA methylation biomarker loci discovered using TCGA and GEO data from over 10,000 tumor and normal samples. Recently, we developed qPCR amplicons specific for a subset of these biomarker loci designed to detect common carcinoma types and tested them on clinical cfDNA samples from healthy individuals and non-small cell lung cancer (NSCLC) patients. We demonstrated that these biomarkers can distinguish between healthy subjects and NSCLC patients with high sensitivity and specificity.

Moreover, in blood samples from lung cancer patients the biomarker DNA methylation signal positively correlates with tumor size. The purpose of the current study was to find how early during carcinogenesis the biomarker loci gain DNA methylation in order to assess their potential as detectors of early stage cancer. To this end, we analyzed DNA methylation of the biomarker loci in publically available data from several precancerous conditions. We found that the biomarker loci gain DNA methylation early in carcinogenesis since they are methylated already in majority precancerous lesions analyzed; in addition, where the data are available, the markers can distinguish lesions with malignant potential from those that stay benign.

Methods

The DNA methylation data from the Illumina Human Methylation450 platform were downloaded from the GEO database (GEO accessions GSE60185, GSE66313, GSE53051, GSE58999, GSE48684, GSE77954, GSE72872, GSE81334, GSE108123 and GSE39279). These DNA methylation data are presented as beta values - numeric values in interval 0.0-1.0. For unmethylated CpGs the beta value approaches zero, for fully methylated CpGs beta approaches 1 and for CpGs methylated in a fraction of the sample 0<beta<1, e.g. a CpG methylated in 50% of the sample will have a beta value of approximately 0.5. All data were analyzed in the R programming environment, version 3.6.1 as follows: The beta values were normalized as described. The normalized beta values for 10 biomarker CpGs (Table 1) were used in further analysis. Boxplots were created using the R function boxplot and the R library beeswarm, version 0.2.3. Since the beta values do not have normal distribution, nonparametric Wilcoxon rank-sum test was used to test differences between the groups. Multidimensional scaling (MDS) plots were constructed using the R function cmdscale on matrices of distances between samples and projected into two dimensions. The ability of the marker set to distinguish between progressive and regressive lung CIS was evaluated using receiver operating characteristic (ROC) analysis on the sums of the beta values from all 10 marker CpG Illumina probes (Table 1). The ROC analysis and AUC calculations were performed using the R library pROC, version 1.15.3.

Results and discussion

We have previously described a set of DNA methylation biomarker loci that are hypermethylated in 10 common carcinoma tumor types and we demonstrated that the level of DNA methylation of these loci can differentiate between plasma samples from lung cancer patients and healthy individuals. To determine the timing of the hypermethylation of these biomarker loci during human carcinogenesis and thus estimate potential of the markers to detect early disease stages we analyzed here the DNA methylation state of the biomarker loci in several premalignant conditions: breast ductal carcinoma in situ (DCIS), colorectal adenomas, Barrett’s esophagus (BE), pancreatic intraductal papillary mucinous neoplasms (IPMNs) and lung carcinoma in situ (CIS) using publically available Illumina HumanMethylation450 datasets from the GEO database.

Ductal carcinoma in situ is a precursor of invasive breast carcinoma (IBC). We analyzed DNA methylation of the biomarker

Table 1. List of 10 DNA methylation biomarker loci used in the study. The first column specifies Illumina CpG probe and the second column shows the genomic position of each biomarker CpG.

<table>
<thead>
<tr>
<th>Illumina CpG.ID</th>
<th>CpG.position (hg19)</th>
</tr>
</thead>
<tbody>
<tr>
<td>cg14416371</td>
<td>chr1:134602847-43602848</td>
</tr>
<tr>
<td>cg08189989</td>
<td>chr2:105459164-105459165</td>
</tr>
<tr>
<td>cg00100121</td>
<td>chr1:169396635-169396636</td>
</tr>
<tr>
<td>cg03306374</td>
<td>chr16:23847325-23847326</td>
</tr>
<tr>
<td>cg01419831</td>
<td>chr2:162283705-162283706</td>
</tr>
<tr>
<td>cg25875213</td>
<td>chr19:38138055-38183056</td>
</tr>
<tr>
<td>cg00339556</td>
<td>chr5:16180048-16180049</td>
</tr>
<tr>
<td>cg01893212</td>
<td>chr7:49813088-49813089</td>
</tr>
<tr>
<td>cg14732232</td>
<td>chr5:528621-528622</td>
</tr>
<tr>
<td>cg07302069</td>
<td>chr7:27196286-27196287</td>
</tr>
</tbody>
</table>

Any further responses from the reviewers can be found at the end of the article.
loci in normal breast tissue samples, DCIS and IBC from three GEO datasets: GSE60185, GSE66313, GSE53051. The results (Figure 1A) show that the biomarker loci are methylated already in DCIS at about the same level as in IBC. The multidimensional scaling (MDS) plot (Figure 1B) shows DCIS samples scattered among IBC samples, indicating comparable levels of DNA methylation of individual markers, while most of the normal samples form a small cluster on a side of the plot. Furthermore, there is no significant increase in the marker methylation during the progression to metastatic disease, as illustrated by data from a cohort (GSE58999) of 44 pairs of primary breast tumors and lymph node metastases (Figure 1A).

Colorectal adenomas are the precursor neoplasms to colorectal cancer. We analyzed biomarker loci in normal colorectal tissue, colorectal adenomas, colorectal carcinomas and metastatic colorectal tumors from three GEO datasets: GSE48684, GSE77954, GSE53051. Similar to DCIS, biomarker loci are already hypermethylated in colorectal adenomas with no further increase in methylation during the progression into invasive colorectal carcinomas or metastatic colorectal cancer (Figure 1C) and again colorectal adenomas on MDS plot are scattered among colorectal carcinomas (Figure 1D).

Barrett’s esophagus is a precancerous precursor of esophageal adenocarcinoma (EAC). We analyzed normal esophagus...
to distinguish cancer from healthy samples. This report shows that the DNA methylation change of the biomarker loci is already present to its full extent in the earliest cancer stages. Thus, the combination of the sensitive detection and the timing of the release of enough tumor DNA into blood or other body fluids are the factors that will set the limit of the biomarkers to detect cancer early.

Finally, we analyzed lung CIS. Lung CIS is a pre-invasive precursor lesion of lung squamous cell carcinoma (SCC), one of the two non-small cell lung cancers that we previously used to demonstrate the capability of the biomarkers to distinguish between clinical plasma samples from cancer patients and healthy subjects. We analyzed DNA methylation of the biomarker loci in lung CIS together with lung SCC and normal lung tissue samples from GEO datasets GSE108123, GSE39279. The advantage of the original lung CIS study (GSE108123) is that the prospective follow-up information is available for CIS samples and thus the samples could be classified as either progressive (those later progressed into invasive cancer) or regressive (these later regressed to normal epithelium or low-grade disease). Our analysis revealed, similar to the other pre-invasive lesions, that the biomarker loci have increased DNA methylation already at the lung CIS stage (Figure 1I). More importantly, when we analyzed progressive and regressive lung CIS samples separately (Figure 1I), we found that the biomarker set is able to distinguish between the two types of premalignant lesions with high sensitivity and specificity (AUC = 0.92, Figure 1I). The majority of the regressive lung CIS samples on the MDS plots cluster close to normal lung controls while all progressive lung CIS samples are scattered among lung SCC samples (Figure 1K). Even when lung SCC samples are sub-grouped into the individual cancer progression stages (I-III) there is no increase in DNA methylation with the stage (Figure 1I). Together, these results show that the gain of DNA methylation of the biomarker loci is an early epigenetic event during human carcinogenesis.

The data presented here show that DNA methylation of the biomarker loci is fundamentally changed early during the malignant progression since it is already observed in precancerous lesions. The data from lung CIS further show that the DNA methylation level of the biomarkers can differentiate between potentially malignant and benign CIS. Together, these findings indicate that the biomarkers are capable, from the qualitative point of view, to detect cancer at its earliest stages. However, the detection of cancer-specific DNA methylation in blood or other body fluids is quantitative in nature and depends on the tumor size and its propensity to shed DNA into bloodstream; e.g., our previous report shows that the DNA methylation signal from this biomarker set in cfDNA samples depends on the NSCLC tumor size. Later disease stages are thus relatively easy to detect since larger tumors of later cancer stages shed a large amount of DNA into bloodstream resulting in high DNA methylation signal. In order to detect the early cancer stages as well, sensitive detection techniques and especially sample processing leading to minimal background DNA methylation signal will be profound
References


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Version 1

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Judd C. Rice
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In this brief report by Vrba and Futscher, the authors sought to leverage their previous discovery of 10 cancer-associated DNA methylation biomarker loci to investigate the clinical potential of these biomarkers in assessing the progression of carcinogenesis. The authors analyzed several publicly available datasets that included normal tissue, precancerous lesions and tumor samples to generate in silico models of breast, colorectal, esophageal, pancreatic and lung cancer progression. The rigor of the study design is high as all datasets utilized the same technology (Illumina HumanMethylation450) with appropriate sample size of each group for analysis, except for the pancreatic samples. Although the data presented support the authors' conclusions, at least by the eyeball test, the inclusion and description of detailed statistical analyses would increase confidence in the conclusions. The overall results indicate that aberrant methylation of the biomarker loci is an early epigenetic event of carcinogenesis, regardless of cancer type. This is an important finding that further supports the clinical potential of DNA methylation biomarkers in early cancer detection.

Is the work clearly and accurately presented and does it cite the current literature? Yes

Is the study design appropriate and is the work technically sound? Yes

Are sufficient details of methods and analysis provided to allow replication by others? Yes

If applicable, is the statistical analysis and its interpretation appropriate? Yes

Are all the source data underlying the results available to ensure full reproducibility?
Yes

**Are the conclusions drawn adequately supported by the results?**
Yes

**Competing Interests:** No competing interests were disclosed.

**Reviewer Expertise:** Epigenetics.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Author Response 11 Feb 2020

**Lukas Vrba**, The University of Arizona, Tucson, USA

Dear Dr. Rice,

Thank you for the evaluation of our study. According to your suggestion and suggestion from the other reviewer we added the p-values to Figure 1. The results now show that all the differences that made the “eyeball test” are also highly statistically significant, including in the pancreatic samples where the n is rather low but the control cohort is consistently unmethylated. We thank you for this suggestion, since the amended version of our manuscript will have higher impact among the readers.

**Competing Interests:** No competing interests were disclosed.

Reviewer Report 08 January 2020

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**Keith D. Robertson**

Department of Molecular Pharmacology and Experimental Therapeutics, Mayo Clinic, Rochester, MN, USA

In this manuscript, Vrba and Futscher examined the DNA methylation status of candidate loci in pre-malignant stages of a variety of cancers (e.g. breast, pancreas, and esophagus). Ultimately, the authors observed that DNA hypermethylation events previously identified in full-blown cancer were already hypermethylated in the precancerous tissues. One particularly interesting outcome was that progressive lung carcinoma in situ demonstrated elevated methylation levels, while regressive CIS more closely resembled normal lung tissue, suggesting that these CpGs may actually be predictive of progression and/or play a role in carcinogenesis.
This is a straight-forward *in silico* analysis study, and seems worthy of indexing since the methylation biomarker field is moving rapidly and showing significant promise.

This reviewer has two concerns, however, that should be addressed:

1. While the methylation trends are clearly qualitatively visible, statistical significance is not given for any of the comparisons.

2. Due to pre-malignant lesions showing similar patterns as the corresponding cancer, does that reduce the efficacy of these biomarkers as the target screening population malignancy is indistinguishable from the cancer? For example, healthy individuals are not routinely screened for esophageal carcinoma, but those with Barrett's esophagus are.

**Is the work clearly and accurately presented and does it cite the current literature?**
Yes

**Is the study design appropriate and is the work technically sound?**
Yes

**Are sufficient details of methods and analysis provided to allow replication by others?**
Yes

**If applicable, is the statistical analysis and its interpretation appropriate?**
Yes

**Are all the source data underlying the results available to ensure full reproducibility?**
Yes

**Are the conclusions drawn adequately supported by the results?**
Yes

**Competing Interests:** No competing interests were disclosed.

I confirm that I have read this submission and believe that I have an appropriate level of expertise to confirm that it is of an acceptable scientific standard.

Author Response 11 Feb 2020

**Lukas Vrba,** The University of Arizona, Tucson, USA

Dear Dr. Robertson,

Thank you for the prompt and generally positive review of our article. We have postponed our response until all reviews were available and a new version of the article was submitted.

As to the first point raised, the other reviewer had the same suggestion and therefore we added p-values for comparisons of individual sample cohorts. All differences between normal tissues and respective premalignant lesions are highly statistically significant, while
there are no significant differences in DNA methylation of the biomarker loci during further malignant progression.

We especially appreciate your second point. The presence of the increased DNA methylation at marker loci may indeed raise doubts about how well the biomarkers may serve for cancer screening of individuals with premalignant conditions. We think that it depends on the type of the samples used for the screening (liquid vs tissue biopsies) and we have added the whole new paragraph discussing this topic. We thank you for raising this important point since it had to be discussed to make the study complete.

**Competing Interests:** No competing interests were disclosed.