Characterization of Chromosomal Aberrations in Neuroblastoma Formalin-Fixed Paraffin-Embedded Specimens with Standard ArrayCGH Procedure - Preliminary Experience

Abstract

Background: The broad spectrum of neuroblastoma (NB) clinical behavior depends on a genomic landscape of tumor cells. The amplification of MYCN oncogene is the most powerful negative prognostic marker in NB. Moreover, segmental chromosomal alterations are also associated with a poor outcome. Therefore, the comprehensive characterization of tumor genetic features is obligatory for NB patients. These features determine the risk stratification and therapeutic decisions in treatment.

Purpose: Our report focuses on a possibility to use standard microarray procedure to demonstrate critical structural chromosomal alteration in archival samples of NB tumors.

Methods and Findings: Formalin-fixed paraffin-embedded tissue samples from 8 NB primary tumors have been analyzed by cytogenetic microarrays. It achieved a very good quality of genomic DNA from fixed samples. Chromosomal abnormalities were detected in 7 out of 8 cases. It was not an incidence of MYCN amplification.

Conclusion: The results demonstrate that it is possible to obtain reliable and high-quality microarray data from archival samples.

Keywords: Advanced neuroblastoma; Chromosomal aberrations; Molecular karyotyping; Tumor genome screening

Introduction

Neuroblastoma (NB) is one of the most common pediatric cancer (6% to 10%) [1]. Approximately 90% of patients are diagnosed before the age of 5 years old [2,3]. The primary form of NB is an extracranial solid tumor arisen from neural crest progenitor cells [1-4]. NB tumors have a very high tendency to form distant metastases [2]. The comprehensive characterization of tumor genetic and biological features is obligatory for NB patients [4]. These features determine the risk stratification and therapeutic decisions in treatment. The most unfavorable genetic factor in NB is MYCN oncogene amplification [2-7]. Moreover, tumor aggressiveness and progression is strongly associated with Segmental Chromosomal Aberrations (SCAs) observed in malignant cells [2,5,8]. Aggressive metastatic tumors are characterized by a high number of SCAs and a low number of Numerical Changes (NCAs). On the whole, genome-wide studies have demonstrated that critical chromosomal damages occur more frequently in older patients and that SCAs accumulate in an age-dependent manner, as supported by Schleiermacher and Coco [8-12].

According to the international guidelines the current gold standard for detecting MYCN oncogene amplification in NB tumor cells is the fluorescence in situ hybridization (FISH) analysis of interphase nuclei [5]. Other, critical copy number changes like 11q23 deletion and 17q gain can also be evaluated by FISH analysis [5,13-17]. However, pan-genomic techniques like high-resolution array-based comparative genomic hybridization (aCGH) or multiplex Ligation-Dependent Probe Amplification (MLPA) provide alternative methods for whole-genome screening and characterizing SCAs and NCAs in NB tumor cells [18-21]. These techniques are equivalent to thousands of FISH
experiments. Numerous aCGH studies had performed on NB primary tumor tissues revealed new, often occurring unfavorable SCAs including deletions in regions: 3p, 4p, 6q, 10q, 14q, 18q, and additional copies: 3p, 12, 18p, as well as cases of the uniparental disomy and the chromothripsis in malignant cells [8,17,21-25].

The vast majority of FISH and aCGH routine analysis in NB have been and currently are performed on fresh tumor tissue samples. However, it is worth emphasizing that both research techniques provide for a genetic evaluation also formalin-fixed paraffin-embedded (FFPE) specimens. Archival, FFPE samples represent an invaluable source of material especially for retrospective molecular studies. The possibility of long-term clinical follow-up of FFPE samples makes them a valuable source to evaluate links between genetic and clinical information.

So far, a few studies have used FFPE-derived DNA for aCGH due to nucleic acid degradation and chemical modifications, and also cross-linking with proteins introduced during the fixation process. In NB, recently reported study confirmed an ability to recognize chromosomal aberrations in FFPE specimens using aCGH but with a new modified microarray method (OncoScan™ FFPE Assay Kit, Affymetrix) [26]. The used protocol gave rise to a significant improvement over most of the previous methods in terms of accuracy in detecting copy number variant in FFPE tissues.

The main purpose of this study is evaluation the practical potential of routine use aCGH platform for detection of numerical and structural chromosomal changes in FFPE-derived DNA samples of NB primary tumor tissues.

### Material and Methods

Diagnostic fixed tissue samples (FFPE specimens) from NB primary tumors have been analyzed in Laboratory of Molecular Genetics, University Children’s Hospital in Krakow since June 2016. FFPE tissues required initial de-waxed with xylene followed by washing with ethanol. Genomic DNA was extracted from the sample with use of a DNA tissue kit (KURABO Industries Ltd, Osaka, JAP) and quantified by OD at 260 nm using the NanoDrop isolation kit. Moreover, there were no obstacles to achieving the required DNA quantity.

The aCGH evaluation was performed for 8 FFPE specimens. It was achieved a very good quality of genomic DNA with the described detection limit of 200 kb for appropriate detection of structural chromosomal abnormalities.

### Results

The aCGH evaluation was performed for 8 FFPE specimens. It was achieved a very good quality of genomic DNA with the described isolation kit. Moreover, there were no obstacles to achieving the required DNA quantity.

Abnormalities in the form of numerous NCAs were detected in 5 out of 8 cases. These tumors were identified as hyperdiploid. The most common NCAs were multiplication of whole chromosomes: 6, 7, 8, 9, 17, 20, 22. In 2 cases we have found also loss of whole chromosomes (Table 1). All patients with the NCAs-positive aCGH profile were younger than 18 months.

Both, SCAs and NCAs were found in 2 out of 8 cases (Figure 1). These patients were older than 18 months, what was an unfavorable factor. Among SCAs we have found critical changes like 1p36 deletion and 17q gain. The size of above structural abnormalities ranged from 5.8 Mb to 49.5 Mb. It was not an incidence of MYCN amplification (Table 1). One patient from the research group had a normal diploid aCGH profile, and was over 18-months-old.

### Discussion

There is a well-known association between patient outcome and tumor ploidy in NB [2-5]. NB tumors with diploid DNA content are more likely to present in older children and with metastatic disease. These diploid tumor’s cells are more likely to harbor MYCN gene amplification or other SCAs. In contrast, near-triploid DNA content is most commonly found in infant tumors that remain localized and are associated with more favorable outcome [2-9,12].

The most common type of SCAs in NB cells beside MYCN amplification is gain of the distal part of long arm of chromosome 17 which occurs in at least 50% of primary NB tumors. This abnormality is associated with poor prognosis, and is frequently coexisted with other biomarkers of aggressive disease such as MYCN amplification, older age, and chromosome 1p36 deletion. Loss of 1p36 region occurs in approximately a quarter of NB tumors and also predicted adverse outcome [2,3,7,11,13]. In addition, a recent INRG report on non-MYCN amplified tumors determined that it is not single genetic markers, but the overall

### Table 1 Microarray analysis results in the research group.

<table>
<thead>
<tr>
<th>Patient number</th>
<th>Patient age (months)</th>
<th>Abnormal aCGH profile</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>&gt;18</td>
<td>arr[hg19] 1p36.13p33x1, 17x3, 18x3, 19p13p11x1</td>
</tr>
<tr>
<td>2</td>
<td>&gt;18</td>
<td>arr[hg19] 1p36.33p36.1x1, 3x1, 4x1, 6x1, 7x3, 9x1, 10x1, 14x1, 15x1, 16x1, 17q21.2q25.3x3, 19x1, 21x1, 22x1</td>
</tr>
<tr>
<td>3</td>
<td>&lt;18</td>
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</tr>
<tr>
<td>4</td>
<td>&lt;18</td>
<td>arr[hg19] 3x1, 6x3, 7x3, 9x3, 16x1, 17x3, 20x3, 21x3</td>
</tr>
<tr>
<td>5</td>
<td>&lt;18</td>
<td>arr[hg19] 6x3, 7x3, 8x3, 9x3, 13x3, 17x3, 20x3, 21x1, 22x3</td>
</tr>
<tr>
<td>6</td>
<td>&lt;18</td>
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<td>7</td>
<td>&lt;18</td>
<td>arr[hg19] 1x3, 2x3, 6x3, 7x3, 8x3, 9x3, 12x3, 17x3, 19x3, 20x3, 22x3</td>
</tr>
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segmental genomic profile of tumors that adds information to patient prognosis [11].

Conclusion

Whole-genome DNA copy number analyses have demonstrated that the overall genomic pattern adds significant prognostic information to clinical algorithms. Integration of molecular profiles and genomic markers that are reflective of tumor biology into future classification systems will allow more precise prognostication and refined treatment assignment.

References

instability phenotype with later onset. Proc Natl Acad Sci USA 107: 4323-4328.


