Management of neuroblastoma: a study of first- and second-line chemotherapy responses, a single institution experience

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Abstract

Neuroblastoma is a high-grade malignancy of childhood. It is chemotherapeutically and radio-sensitive but prone to relapse after initial remission.

The aim of the current study was to study the results of the first- and second-line chemotherapy on the short-term response and long-term survival of children, and to further describe the side effects of treatment. Ninety-five children with advanced neuroblastoma were included in the study, divided into two groups according to the treatment strategy: 65 were treated by first-line chemotherapy alone, and 30 children who were not responding or relapsed after first-line chemotherapy were treated by second-line chemotherapy. External beam radiotherapy was given to bone and brain secondary cancers when detected. Staging workup was performed before, during and after management. Response was documented after surgery for the primary tumor. Median follow-up was 32 months (range 24-60 months). Chemotherapy was continued until toxicity or disease progression occurred, indicating interruption of chemotherapy. Patients received a maximum of 8 cycles. Toxicity was mainly myelo-suppression, with grade II-III severity in 60% of the first-line and 70% of the second-line chemotherapy patients. Median total actuarial survival was nearly 51 months for the first-line chemotherapy group and 30 months for the second-line line group, with a statistically significant difference between the two groups (P<0.01).

Introduction

Neuroblastoma is the most common extra-cranial solid tumor in childhood; over 75% of patients are in an advanced stage at diagnosis and prognosis is poor. It is chemotherapeutically sensitive; 70% of patients with high-risk neuroblastoma could achieve complete remission after comprehensive therapy, including chemotherapy, surgery, radiotherapy, bone marrow transplantation, and biotherapy. However, the remission time interval is short, and the majority of patients die of tumor relapse. The 5-year survival of patients with high-risk NB is only approximately 30%.\textsuperscript{1} At diagnosis, the defining characteristics of high-risk neuroblastoma include age over 1.5 years, advanced stage, amplification of the N-MYC oncogene, and histological findings.\textsuperscript{2,4}

Progress in the treatment of high-risk neuroblastoma may have been due to the use of higher doses of chemotherapy\textsuperscript{1} and improved supportive care. Moreover, vitamin A (retinol) plays a critical role in normal neural crest development. Intracellular retinol is metabolized to all-trans retinoic acid (ATRA). Exposure of human neuroblastoma cell lines to supraphysiological doses of ATRA caused a reduction in cell growth and induction of neurite differentiation that was similar to normal neural cells. This property of vitamin A directed oncologists to use fenretinide, a vitamin A derivative, in the treatment of pediatric neuroblastoma in combination with chemotherapy.\textsuperscript{5,6} Further still, a coumarin derivative RKS262 belongs to a new class of potential anti-tumor agents. RKS262 was identified by structural optimization of nifurtimox, which is currently undergoing phase II clinical trials to treat high-risk neuroblastoma.\textsuperscript{7,8} In another attempt to improve the results of treatment of high-risk neuroblastoma cases, patients have received metronomic chemotherapy. Recent experimental studies have suggested that frequent administration of certain cytotoxic agents at low doses, known as metronomic chemotherapy, increases the anti-angiogenic activity of certain drugs. The advantage of this strategy is lower toxicity and risk of emergence of drug-resistant tumor cells than conventional administration.\textsuperscript{9,11} Monoclonal antibody therapy for neuroblastoma is also an attractive investigational treatment option. GD2-disialoganglioside is expressed on the surface of childhood neuroblastoma cells. Because of this tumor selective expression, it is an attractive target for tumor specific therapy with monoclonal antibodies. Over the last two decades, several anti-GD2 antibodies have been developed and investigated for the therapy and consolidation of pediatric neuroblastoma.\textsuperscript{12,14} For patients presenting with a large tumor burden at the time of treatment, 131-iodine-mIBG therapy is usually recommended.\textsuperscript{15} The aim of the current study was to study the outcome of first- and second-line conventional chemotherapy on the short-term response, and long-term survival of high-risk neuroblastoma cases.

Materials and Methods

Between June 2005 and December 2009, 95 neuroblastoma patients were diagnosed at the Pediatric Unit of the Oncology Department, Faculty of Medicine, Cairo University, Egypt. The median follow-up period was 32 months (range 24-60 months). Cases were either children
with Evans stage III or IV who were newly diagnosed and treated with chemotherapy alone (65 cases) or those who failed initial successful therapeutic modalities (30 cases). The patients were younger than 16 years with a Karnovsky performance status of more than 30%. Patients with a lower performance status were excluded from the study due to the lowered tolerance to chemotherapy. Whereas, with treatment, an improvement in performance status would be expected in responders. Sixty-five children were included in the first-line chemotherapy group alone and all were treated with 6-8 cycles of alternating courses of OPEC/OJEC every three weeks as follows: vincristine 1.5 mg/m² Day 1, etoposide 200 mg/m² Day 1, cyclophosphamide 600 mg/m² Day 1, and cisplatin 80 mg/m² Day 1 or carboplatin 500 mg/m² Day 1; these treatments alternating with each other every 21 days if there was hematologic recovery. Second-line therapy was given to 30 children and all were treated with 6-8 cycles according to the French SFOP studies protocol. This consisted of alternating courses of CAdO (cyclophosphamide 300 mg/m² Day 1, vincristine 1.5 mg/m² Day 1, and doxorubicin 60 mg/m² Day 1) and CE (carboplatin 40 mg/m² Day 1-5 and etoposide 100 mg/m² Day 1-5 every three weeks). All children were subjected to the same chemotherapy (CT) for 6-8 cycles till remission, grade 3 or 4 toxicity, or disease progression occurred. Radiological evaluation of response to therapy was performed every 2 cycles before administration of the forthcoming cycle. External beam radiotherapy was given to bone and brain secondary cancers when detected irrespective of group.

Diagnostic work up
- A full clinical examination with documentation of all measurable disease.
- Performance status using K.I and body weight in kgs.
- Laboratory investigations included: CBC, BUN, serum creatinine, LDH, ESR, and urinary VMA.
- Bone marrow aspiration and tumor biopsy.
- Radiological imaging included: chest X-ray, CT scan of the tumor site and abdomino-pelvic sonography.
- Radionuclide imaging included: bone scan and 131I-MIBG scans.

Response criteria were as follows:18
- Complete response (CR): complete disappearance of disease.
- Partial response (PR): 50-90% decrease in tumor volume.
- Stationary disease (SD): <50% decrease in tumor volume.
- No response (NR): no change in size of tumor.
- Disease progression (DP): increase in size of tumor.

Statistical analysis

Independent proportions were compared using χ² and Fisher’s exact tests. Total actuarial survival curves were plotted with log rank test for survival time and ANOVA was used to compare groups. P<0.05 was considered statistically significant.

Results

Patients’ characteristics are summarized in Table 1. In the first-line chemotherapy group, bone marrow involvement was found in 39 cases (60%) and lymph node metastases in 30 (46.2%) of those 39. Of those cases with bone marrow involvement, 38 of the 39 also had a positive bone scan and the remaining patient also had secondary brain tumors. Two cases with bone marrow and bone involvement also had soft tissue extension. In the second-line chemotherapy group, lymph node metastases were found in 15 cases (50%), and all of those cases also had bone marrow involvement. Eleven of those 15 also had also a positive bone scan, 7 had secondary brain tumors, and the remaining 4 had soft tissue extension. Both performance status and body weight improved significantly after therapy (Figures 1 and 2, respectively). Mean Karnovsky Index ± SD was 66.2%±7.2% and 68.9%±6.5% pre-therapy, and increased significantly to 84.9%±9.73% and 82.4%±4.1% post therapy in the first-line and second-line chemotherapy groups, respectively. The increase in the body weight in the groups was statistically significant (P=3.72e-9 and P=8.63e-15, respectively).

Short-term response assessment

Response in the primary tumors is summarized in Table 2. Outcome of metastatic cases at presentation is summarized in Table 3. The toxicity of each treatment group is summarized in Table 4.

Long-term survival assessment

Figure 3 shows the total actuarial 5-year survival of the studied groups during the follow-up period. Median survival was 51 months for the first-line chemotherapy group and 33 months for the second-line chemotherapy group (P<0.01 for both groups). At the end of the follow-up period (median follow-up 32 months, range 2-5 years), 40% of patients in the first-line chemotherapy group and 14% in the second-line chemotherapy group were still alive (P<0.01).

Discussion

Prognosis for high-risk neuroblastoma patients has improved over the last decades. However, even after highly intensive treatment, only
a few patients become long-term survivors. Most high-risk patients relapse after initial response to induction treatment. The optimum treatment for high-risk neuroblastoma has still not been established and the results of most of the protocols regarding, CR, event free and overall survival rates are unsatisfactory.

Our study is one of the few studies to analyze the results of conventional treatment of stage 3 and 4 neuroblastoma cases applied in the majority of cancer centers with unsatisfactory long-term outcome. It was noted that a small percentage of the cases could be cured with chemotherapy. These were cases with: low tumor burden, negative n-myc amplification, younger age at presentation, and no distant metastases.

It was observed that the majority of cases were abdominal and thoracic; this leads us to screen cases by abdomen-pelvic ultrasound and chest X-ray. Metastases were more common in bone marrow, followed by bone and lastly lymph nodes, confirming the importance of primary baseline bone marrow biopsies and isotopic bone scans for all cases. Moreover, early diagnosis is of utmost importance as it indicates better treatment outcomes, underlining the importance of screening programs, especially for high-risk groups.19

The current study showed that the initial treatment of neuroblastoma cases carries a better chance of obtaining CR that is reflected on overall survival. This indicates the importance of optimizing the primary treatment to induce the highest possible CR.

Even in cases with advanced disease, it is possible to obtain a remission rate of 70% or more with chemotherapy and surgery.20 However, the relapse rate is high among these patients, and many of them die because of the refractory recurrent tumor.2,3,5 In the current study, the majority of the chemotherapy-alone treated patients showed stationary disease (35.4%) rather than CR (21.5%) or PR (24.6%).

Concerning survival, in the Spanish Neuroblastoma Group, the probability of survival with stage IV disease was 0.24 at five years.20 Moreover, Philip and colleagues21 reported an outcome of survival rate for patients given a highly intensive investigational megatherapy. The toxicity-related death rate in this group of 33 patients was 24%. In the European registry, overall survival after five years was no more than 33% for stage 3 and stage 4 patients.22 The current study showed a 40% survival in the first-line chemotherapy group that is very similar to the European study22 and that of Philip et al.,21 and higher than that of the Spanish group.20 However, in the current study, survival figures were markedly lower to 14.1% at five years in the second-line chemotherapy group. This was due to toxicity build up, mainly in the form of myelosuppression, with a poor performance status and impaired vital organ reserve reducing the tolerance to chemotherapy and consequently survival.16,17

Sixty percent of neuroblastomas in young children reported by the literature are Stage 4 (undifferentiated and widely disseminated) at

<table>
<thead>
<tr>
<th>Response</th>
<th>First-line chemotherapy</th>
<th>Second-line chemotherapy</th>
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<tbody>
<tr>
<td>CR</td>
<td>3/14, 21.4%</td>
<td>0.0018, 2/6, 33.3%</td>
</tr>
<tr>
<td>PR</td>
<td>9/16, 56.3%</td>
<td>7/18, 38.9%</td>
</tr>
<tr>
<td>SD</td>
<td>15/23, 65%</td>
<td>2/2, 100%</td>
</tr>
<tr>
<td>NR</td>
<td>5/5, 100%</td>
<td>0.0016, 1/1, 100%</td>
</tr>
<tr>
<td>DP</td>
<td>7/7, 100%</td>
<td>3/3, 100%</td>
</tr>
</tbody>
</table>

*P value comparison between first and second groups.*

<table>
<thead>
<tr>
<th>Toxicity type</th>
<th>First-line chemotherapy</th>
<th>Second-line chemotherapy</th>
</tr>
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<tbody>
<tr>
<td>Grade II-III myelosuppression</td>
<td>39/65, 60%</td>
<td>21/30, 70%</td>
</tr>
<tr>
<td>Grade II-III vomiting</td>
<td>50/65, 77%</td>
<td>22/30, 73%</td>
</tr>
<tr>
<td>Grade III alopecia</td>
<td>65/65, 100%</td>
<td>30/30, 100%</td>
</tr>
<tr>
<td>Abdominal distension</td>
<td>0</td>
<td>20/30, 66.6%</td>
</tr>
<tr>
<td>Massive hemorrhage</td>
<td>0</td>
<td>1/30, 3%</td>
</tr>
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Figure 1. Comparison between the mean values of Karnofsky scale pre and post treatment.

Figure 2. Comparison between the mean values of body weight pre and post treatment.

Figure 3. Comparison between total actuarial survival of the studied groups.
diagnosis; that is very close to the percentage reported in the current study with poor treatment outcomes. Novel treatment methods are required to deal with this problem. One of the interesting new modalities is the inclusion in disease management of 131I-mIBG therapy because of the specificity of the agent and the radio-sensitivity of the primary and metastatic disease. Another investigational treatment is differentiation therapy with retinoids. Because neuroblastomas are classified as embryonal tumors arising from immature cells of the neural crest, the induced differentiation of neuroblastoma cells using retinoids either alone or in combination with other treatments has become a part of therapeutic protocols. This differentiating effect is achieved by all-trans retinoic acid (ATRA). All-trans retinoic acid is a major mediator of the effects of vitamin A via activation of a number of RAR and RXR nuclear receptors that heterodimerize and regulate gene transcription. Neuroblastomas show a high level of multidrug resistance during chemotherapy. In 2009, Choudhry and colleagues explored the effect of a combination of sorafenib and genistein on growth inhibition of neuroblastoma cells. They found that this combination abolishes the expression of MDR in both neuroblastoma SK-N-DZ and SH-SY5Y cell lines.

In an attempt to treat neuroblastoma by monoclonal antibodies, it was found that neuroblastoma cells are the tumor cells expressing the most GD2. This expression of tumor selection makes it an attractive target for tumor specific therapies, such as antibody therapy. Several anti-GD2 antibodies have been developed and used either alone or in combination with other agents in the treatment of resistant and refractory neuroblastoma. Immunotherapy may be particularly effective for low levels of minimal residual disease that is responsible for disease relapse after initial remission. Prevention of these relapses with additional conventional chemotherapy is limited because of cumulative toxicity. Thus, additional treatments to chemotherapy, surgery, and radiotherapy have to be sought. Monoclonal antibodies directed against GD2 have offered another promising avenue for treating minimal residual disease. Moreover, metronomic low-dose chemotherapy was thought to have the potential to prevent relapses with acceptably low toxicity. Therefore, an oral chemotherapy for consolidation of remission with cyclophosphamide, etoposide and melphalan was introduced in trial NB90.

Finally, we would like to add that among the problems we faced during this study, was the fact that the majority of parents were unwilling to agree that their children receive any investigational therapy that had no reliable outcome. These parents, for cultural and religious reasons, would not accept any further treatment after failure of conventional therapies.

Conclusions

Even today, the treatment of advanced neuroblastoma has a very high failure rate, although some there has been some decrease in both progression and relapse rates over time. Despite this progress, neuroblastoma remains a challenging disease for both clinicians and researchers due to the lack of acceptable cure rates in high-risk neuroblastoma. Clearly, new therapeutic front-line strategies are needed to significantly increase survival. In the unfortunate case of recurrence or in the case of tumor progression, experimental therapies can be proposed. The patient’s quality of life should also be carefully discussed and evaluated, and strategies for appropriate care when cure appears unlikely should receive strong consideration.

Studies using a combination of radionuclide therapy with conventional chemotherapy offer a tempting treatment option for high-risk cases of neuroblastoma, but extensive research is needed to confirm impact on survival and late effects. Future efforts should be directed to design protocols that have a multimodality treatment approach.

References