The incidence and correlates of non-adherence in adolescents receiving chemotherapy

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Abstract

A retrospective chart review was conducted to investigate the prevalence of non-adherence to chemotherapy among adolescents with cancer.

Chemotherapy treatment protocols were identified as Treatment I or Treatment II depending on treatment phase and disease progression. Treatment I was defined as the series of drugs given during the induction and maintenance phases of treatment. Treatment II was a different series of drugs, offered if the patient failed to achieve remission during induction or if experienced relapse at any time during treatment. Each treatment was analyzed separately and independently. Data were extracted from a total of 49 charts. Forty-eight charts had information available from the time of diagnosis and one chart had information only from the time of relapse because the patient received initial treatment at another center.

Of the 48 patients who entered Treatment I, one patient refused chemotherapy and 47 accepted. The one patient who refused an initial offer of chemotherapy accepted an equivalent treatment of total nodal radiation. He was not considered non-adherent; however, factors that led to his decision to refuse chemotherapy will be examined.

Of the 47 patients who received chemotherapy in Treatment I, one patient dropped out, four patients modified their treatment and five delayed treatment, for a total non-adherence rate of 35% (n = 10). There were a total of 16 patients in Treatment II which was composed of 15 of the initial 47 patients in Treatment I and one patient transferred from another hospital. Of the 16 patients in Treatment II, two patients dropped out and three patients modified treatment for a non-adherence rate of 31% (n = 3).

The major reason for modification or delay of treatment was the inability to tolerate nausea and vomiting. Antineoplastics were refused by 13 patients (27%) due to the unpleasant side effects of these drugs. Other factors contributing to non-adherence and implications of this study for future research are discussed.

Introduction

At the time of diagnosis, adolescent cancer patients and their families receive information about the nature of their illness and are advised of treatment protocols which will improve their chances of long-term survival. Although the protocol improves the prognosis, it results in unpleasant side effects, inconvenience, and restrictions in activities of daily living. Often, decisions about following protocols involve making trade-offs between quality of life and longevity. For many patients, or their families, the decision is straightforward; survival takes priority in decision-making and quality-of-life factors are secondary. For others, the treatment is considered worse than the disease. Either in anticipation of the toxicity and complications, or after experiencing them for a period of time, patients may decide to drop out of treatment, or have treatment modified because the restrictions imposed on their quality of life outweigh the possible benefit of reduced morbidity and mortality.

The purpose of this study was to examine the prevalence and factors associated with non-adherence to chemotherapy in a group of adolescents. Through a retrospective chart audit, decisions made by the adolescent to refuse, modify, delay, or discontinue treatment were identified and factors contributing to these decisions were noted. The chart audit was chosen as a means to give direction to a more extensive, prospective study that would examine, in more detail, patient decision-making about non-adherence.

The prevalence of non-adherence across a variety of conditions in adult and pediatric populations has been shown to be significant. Approximately 50% of all patients, regardless of age or condition, have difficulty following prescribed medical regimens (Sackett, & Snow, 1979). In a review of 11 adherence studies with chronically ill children, Lit and Caskey (1968) found non-adherence rates ranging from 12 to 88 per cent. When adolescents were separated out, they were found to be less adherent than younger children. Factors, regardless of age, that have been found to adversely affect adherence include:

1. the length and complexity of the prescribed regimen (Haynes, Taylor, & Sackett, 1979),
2. side effects (Koch, Fine, & Negrette, 1978; Smith, Rosen, Trueworthy, & Lowman, 1979),
3. the degree of difficulty in managing side effects (Richardson, Marks & Levine, 1988) and,
4. the degree to which the child’s normal developmental function is interrupted (Koocher 1986; Friedman et al. 1986).

The complex nature of cancer treatment protocols may make them particularly difficult to follow. Adherence, for the adolescent, means time.
spent in hospital away from family, school and peers; numerous clinic visits; painful diagnostic and evaluation procedures and living with the toxic effects of the chemotherapy drugs. Drug side effects that are visible such as weight gain, acne, and hair loss are most disturbing to adolescents (Klopowitz, & Trueworthy 1985) as this altered appearance conflicts with their need to conform to the image of their peers. At a time when peer acceptance, body image and emerging independence are important issues for the adolescent, cancer treatment forces them to be more dependent on their parents and sets them visibly apart from their peers.

Drug therapy for the treatment of cancer is considered to be instrumental in achieving disease-free survival (Finkel 1976), yet little detailed information is available about the consequences of delay, dosage modification, or refusal of drugs within a protocol. Most clinicians would agree that labelling an individual as non-adherent has negative implications, however this must be weighed against the benefits of having targeted an individual for added support and intervention that might help them to comply with therapy.

Varying definitions of what behaviours constitute adherence, and diverse techniques for measuring adherence make it problematic to compare results across studies. Indirect measures such as pill counts, appointment keeping, prescription renewal rates and patient testimony do not always give an accurate picture of patient adherence (Sackett, & Snow 1979). Direct methods involving serum drug levels or urine metabolites are more accurate but are expensive and can involve invasive procedures. Depending on the extent to which adherence was operationally defined and reliably measured, study results must be interpreted with caution.

Smith, Rosen, Trueworthy, & Lowman (1979) were the first investigators to use a direct measure of medication adherence. Adherence to prednisone therapy was determined using a random, urine 17- ketogenic steroid assay. Thirty-three per cent of the total assays performed were below the value set for adherence and of the 17 assays performed on eight patients over 11 years, 59% were in the non-adherence range.

Lansky, Smith, Cairns, N. and Cairns, G. (1983) used the same method to measure prednisone adherence in 31 leukemia patients under 15 years of age. The overall non-adherence rate was 45% with no patient found to be 100% adherent. While both of these studies used a rigorous method for determining adherence, they are limited to prednisone therapy which is only a part of the treatment for cancer. It is speculative to generalize adherence behaviour with other drugs in the protocol based on the adolescent's adherence to prednisone therapy.

Tebbi, Cummings, Zevon, Smith, Richards, & Mallon (1986) measured self-reported non-adherence with prednisone and crosscorrelated the testimony with serum bioassays. They found that children over 10 years of age had a greater adherence problem than younger children. An interesting finding from this study was the observation that patients' parents gave accurate accounts of missed dosages which were supported by the results of the serum bioassays. A limitation of the study was the definition of non-adherence which included patients missing from one to greater than three doses. In terms of the impact on survival, missing one dose of a drug probably does not have the same implications as missing many doses or taking the drugs sporadically. It seems unrealistic given the complexity and prolonged length of cancer treatment protocols, to classify patients as non-adherent if they miss one dose of medication.

In studies that used indirect methods to measure adherence, such as physicians' perceptions (Dolgin, Katz, Doctors, & Siegel, 1986), clinic attendance (Levanthal, & Boek, 1978; MacDougall, Wilson, Cohn, Shuenyane, & McElligott, 1989), and patient self-report (MacDougall et al., 1989), non-adherence was a significant problem, with adolescents generally less adherent than younger children.

Two studies have established a link between prednisone adherence and survival. Trueworthy (1982) categorized 17 children with lymphoblastic leukemia as adherent or non-adherent based on urinary assays. The 12 patients in the adherent group had no incidence of relapse whereas four of the five patients in the non-adherent group had relapsed. Richardson, Shelton, Kraio and Levine (1990) used serum drug and metabolic levels to assess adherence to allopurinol and prednisone in 94 adults. They found a significant correlation between adherence with oral medication and survival.

The results of adherence research suggests that adolescents have a difficult time following treatment protocols. However, it is difficult to know the exact scope of the problem and the implications for survival. In order to be meaningful, the problem of adherence needs to be studied from a broader perspective, taking into account different phases of treatment eg. induction, maintenance, after relapse, and examining provider and self-administered chemotherapy. Most of the studies focused on one drug and none mentioned at what point in treatment the non-adherence behaviour occurred. Non-adherence during induction may have different implications for survival than non-adherence after relapse. This study was conducted to examine the incidence and factors associated with treatment refusal, delay, modification or discontinuation of provider-administered chemotherapy during induction and maintenance (Treatment I) and after relapse (Treatment II).

Method

Procedure

A retrospective chart review was conducted at a children's hospital in Ontario. The centre cares for approximately 40 newly diagnosed cancer patients per year and has a total active caseload of 200 patients per year. Chart audits were conducted to examine treatment decisions for adolescents offered chemotherapy over a ten-year period (1979-1988). Individuals ranging in age from 11 to 18 years were classified as adolescents. This age span covers early, middle and late adolescence as described by Felice (1985).

All diagnostic categories were considered eligible to be included in the study. Treatment protocols were classified as Treatment I or Treatment II depending on whether the patient entered at the time of diagnosis (I) or at treatment failure/relapse (II).

Charts of all patients who were offered chemotherapy over a 10-year period were identified using the Canadian Classification of Diagnostic, Therapeutic and Surgical Procedures code for injection or infusion of cancer chemotherapy substances. Of the 231 charts screened for inclusion in the study, 49 were between the ages of 11 and 18, and were no longer receiving chemotherapy. Forty-eight charts selected were from patients treated at the time of diagnosis (Treatment I) and one chart was from a patient transferred to the centre at the time of relapse (Treatment II), having finished the initial treatment elsewhere. (see Figure One). Information concerning chemotherapy adherence was extracted from medication and treatment records, physicians', nurses' and allied health workers' notes. The notes for each clinic visit were examined to ascertain if the adolescent received the treatment that was planned at the previous visit. For example, if the physician noted read "return to clinic in one week for Vincristine 2 mg IV" it was then possible to evaluate the next visit to see if the patient kept the appointment and received the drug as planned. If the drug was not given or the dosage was modified the reasons were classified as patient or physician determined. Physician determined reasons were mainly related to drug toxicities such as decreased white blood count, infection, stomatitis, diarrhea etc. Examples of patient determined reasons for modifying or delaying chemotherapy were...
cancelled appointments, pain, nausea, refusal etc. We did not attempt to analyze how much, or what type of input parents had into initiating changes in treatment because, for the most part, it was not possible to obtain this information retrospectively from the charts.

Results of each chart review were validated with the principle investigator and two research assistants. The nurse in the oncology clinic was consulted to clarify information in the charts when needed. If the patient entered more than one treatment protocol, which was the case for all patients that relapsed, each treatment was analyzed separately. Rates of non-adherence are reported for Treatment I, which is the series of chemotherapy drugs given for the induction and maintenance phases of treatment, and Treatment II, which begins at the time of remission or failure of Treatment I to put the patient into remission. These treatments were analyzed separately and independently of each other. Other variables extracted from the medical record included: Diagnosis, age at diagnosis, sex, type of treatment protocol, types of health professionals involved with the patient, and side effects related to chemotherapy.

Analysis
Rates of adherence and non-adherence were calculated based on patients’ decisions. Three decision categories were established: Adolescents that delayed or modified treatment on three or more occasions or patients that refused/dropped out of treatment were considered non-adherent. In an effort to recognize that some flexibility exists within treatment protocols, adherence was defined as less than three episodes of chemotherapy delay or modification. Due to the lengthy and complex nature of cancer treatment protocols, it did not seem realistic to classify an adolescent as non-adherent if there were infrequent modifications or delays in chemotherapy administration. Three episodes was arbitrarily chosen as a reasonable dividing line between adherence and non-adherence. There is nothing in the literature to provide guidelines as to how much a protocol can be altered before treatment success or survival is affected.

Descriptive statistics were used to describe the sample. Chi-square analysis and Fisher’s exact test were used to test the relationship between demographic and clinical variables and adherence categories.

Results
The 49 patients selected for inclusion in the study ranged in age at time of diagnosis from 11.5 to 18 years with a mean age of 13.82 (SD 1.7). Twenty-eight (57%) patients were 13 to 15 years of age. There were 25 males and 24 females in the study. Lymphoma was the largest diagnostic category with 16 (33%) patients. The second largest category was sarcoma with 15 (31%) patients. Outcome data indicated that 10 (21%) of the patients lived at the time of the chart review and 19 (39%) of the patients had died. Figure Two shows adherence status in Treatment I by age and sex; and Figure Three shows adherence status by diagnosis.

The modal patient in this sample was male, diagnosed at 12 years of age with a diagnosis of lymphoma or sarcoma. He followed only one protocol (Treatment I), a T10 or a MOPP (Demithe, Serpick, & Carbone 1970) and was alive at follow-up. In addition to nurses and physicians, he had a social worker and a child life worker involved in his care.

Among those patients entering Treatment I, the non-adherence rate was 18%, with a 95% confidence interval of 9 to 33. This non-adherence rate included nine modifier/delayers, and one drop-out. Although the non-adherence rate for Treatment II was 31%, it was not considered meaningful due to the small number of patients (n = 16) and the resultant wide confidence intervals.

It was noted that 13 (27%) patients refused IV and oral antiemetics sometime during their treatment. Refusal was linked to side-effects of the antiemetics. Statements such as "they make me feel out of control," or "I don't like the way they make me feel," were typical of patients' reasons for refusing the antiemetics. The relationship between adherence status and refusal of antiemetics is presented in Figure Four.

Refusers/drop-out rates
Of the 48 patients offered chemotherapy at the time of diagnosis (Treatment I), one patient refused, and one patient dropped out of treatment after completing 12 months of a 18-month protocol. The patient who refused chemotherapy was male, 17 years of age at the time of diagnosis, with Stage II, Hodgkin's lymphoma. In conjunction with the oncologist and his parents, he opted for total nodal radiation (Stanford protocol) with the understanding that he might have to undergo chemotherapy if radiotherapy proved inadequate. He was not entered into the statistical analysis as non-adherent because he accepted an alternative to chemotherapy that did not jeopardize his chances of survival. Reasons given for refusal of
Chemotherapy were related to hair loss and the fear of becoming sterile as a result of chemotherapy.

The patient who dropped out of treatment was also male, and diagnosed at 12 years with non-Hodgkin's lymphoma. He was noted to have refused chemotherapy due to vomiting and/or fear of needles at least nine times before finally dropping out of treatment. Currently he is alive and in remission.

Sixteen of the 49 patients entered Treatment II; of this group, two (10%) patients dropped out. Both were males, one diagnosed at 13 years with malignant schwannoma and the other diagnosed at 16 years with Ewing's sarcoma of the right wrist. Both patients were adherent in Treatment I, relapsed and were offered a second series of drugs (Treatment II). The patient with malignant schwannoma refused chemotherapy, opted for radiotherapy and subsequently died. The patient with Ewing's sarcoma indicated many times that he did not want any more chemotherapy before finally choosing not to have further treatment. He did return for chemotherapy at another centre for a short time before his death.

Delay/modification rates

In Treatment I (n = 47), four patients were categorized as modifiers and five patients were categorized as delays for a combined rate of 19%. In Treatment II (n = 16), two patients delayed treatment and one patient modified treatment for a delay/modification rate of 19%.

The most frequently cited explanations for non-adherence were nausea and vomiting, missing appointments, pain and a nonspecific complaint of "feeling unwell." Overall, the reasons for non-adherence can be categorized into drug side-effects, psychological and social variables. Drug side-effects that contributed to non-adherence included pain, nausea and vomiting, loss of appetite, dizziness, and generally feeling "unwell." The psychological factors most frequently cited in the charts were depression, fatigue/tension, fear of hair loss, anxiety, and fear of needles. The social variables that contributed to treatment delay or modification included family vacations, school-related activities, cancelled appointments and one instance of medication being withheld by the mother, as a punishment.

Correlates of non-adherence

Several variables had more than 25% of the cells with expected frequencies less than five. Therefore Fisher's exact test was used to replace Chi square analyses where appropriate. Adherence was considered in view of its possible association with age, sex, diagnosis, outcome, treatment regimen and number of health professionals involved other than nurses or physicians. (See Table One) The significance level was adjusted to p<.008 to account for the increased chance of obtaining a type I error. There was no statistically significant association between adherence status and any of the demographic, diagnostic and treatment variables. The lack of statistical correlation between selected variables and adherence may be due to small sample sizes. In at least three variables (sex, diagnosis and outcome) absolute differences in proportions of 15% were noted.

Discussion

Overall, the non-adherence rate of 21% appears to fall within the range of rates reported in the literature and was considered clinically significant. It is difficult to compare adherence behaviour across studies because of diverse methodologies and measurement techniques. Criteria for non-adherence are often different, not clearly articulated, or subjectively reported. However, this study supports the premise that adolescents have difficulty following treatment regimens. This is not surprising given the fact that adolescents are at an age where control over their lives is an important issue. Non-adherence behaviour becomes a vehicle, albeit maladaptive, to exercise an emerging sense of power and autonomy within the structured, often inflexible routines of hospitals and clinics.

A limitation of this study was the fact that it was a retrospective chart review. The amount of information in the chart, the logic and clarity of the notes tended to vary depending on the author. However, it was still possible to get a sense of how well or poorly the patient was coping with treatment. When drugs were refused or dosages modified there was a notation on the chart giving some explanation. It was then possible to classify the modifications or delays as patient or physician-directed. It was also apparent that many of the adherent patients had great difficulty coping with treatment side-effects, school absences, and the overall stress of having cancer.

The development of new treatments and the success of existing treatment protocols have dramatically increased long-term survival rates for adolescents with cancer. However, the efficacy of a treatment protocol is, in part, dependent on how well the patient is able to follow it. Therefore, it is important to be able to predict which adolescents may not comply with treatment and which treatments are particularly difficult to follow. Predictors variables could be identified and tested in a prospective study. Patients identified as being at risk for non-adherence could be targeted for intervention before a problem develops. Variables identified from this study which may be useful to include in a prospective study are severity of nausea and vomiting, refusal of antinecics, and support provided by family, and by health professionals.

The refuser/dropout rate of 2%, in Treatment I, was composed of one patient who dropped out before treatment was completed. This is in contrast to the 19% (n = 9) of patients in Treatment I who delayed or modified their treatment regimens when objective toxicity or treatment failure was noted. Although modification and delay of treatment is not as extreme as refusing or dropping out of treatment, it is not known to what extent this may compromise treatment outcome. A prospective study would permit evaluation of this more moderate non-adherence in terms of its effect on outcome. It would provide information about the extent to which flexibility exists within a specific protocol. For example, how often and to what extent can drugs be delayed or omitted and dosages modified, and still expect optimum long-term survival rates from a protocol?

Thirteen patients (27%) refused antinecics for all or part of their treatment. Of the 15 patients that were non-adherent across Treatment I and II, five were also antinecic refusers, for a total of 33% of the non-adherent

<table>
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<th>Variable</th>
<th>Adherent (N=37)</th>
<th>Non-adherent (N=10)</th>
<th>X²</th>
<th>P Value</th>
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<td>&lt; 14</td>
<td>N=21</td>
<td>17 (46%)</td>
<td>4 (40%)</td>
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<td>≥ 14</td>
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<tr>
<td>Male</td>
<td>N=25</td>
<td>21 (57%)</td>
<td>4 (40%)</td>
<td>.17</td>
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<tr>
<td>Female</td>
<td>N=22</td>
<td>16 (43%)</td>
<td>6 (60%)</td>
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<td>Lymphoma</td>
<td>N=15</td>
<td>10 (27%)</td>
<td>5 (50%)</td>
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<td>12 (32%)</td>
<td>3 (30%)</td>
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<td>Leukemia</td>
<td>N=11</td>
<td>9 (24%)</td>
<td>2 (20%)</td>
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<td>Type of protocol</td>
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<tr>
<td>MOPP</td>
<td>N=9</td>
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<td>2 (20%)</td>
<td>.45</td>
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<td>T-9</td>
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<td>7 (19%)</td>
<td>1 (10%)</td>
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<tr>
<td>Others</td>
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<td>23 (62%)</td>
<td>7 (70%)</td>
<td></td>
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<td>Number of health professionals involved in additional to physicians and nurses</td>
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<tr>
<td>&lt; 2</td>
<td>N=17</td>
<td>13 (35%)</td>
<td>4 (45%)</td>
<td>**</td>
</tr>
<tr>
<td>≥ 2</td>
<td>N=30</td>
<td>24 (65%)</td>
<td>6 (35%)</td>
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<td>Outcome</td>
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<tr>
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<td>22 (59%)</td>
<td>8 (75%)</td>
<td>**</td>
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<tr>
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<td>15 (41%)</td>
<td>2 (25%)</td>
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* These were the three largest categories
** Fisher's exact test used

| CONJ: 2/1/92 | 4 |
| RCNO: 2/1/92 | 6 |
group. A few patients made specific statements about side-effects of the antiepileptic drugs that they found intolerable. Because one of the major reasons for non-adherence was nausea and vomiting, it would be useful to determine factors contributing to antiepileptic refusal and how this affects adherence.

The difficulties that adolescents have coping with the side-effects of antiepileptic therapy suggest the need for drugs that can control nausea and vomiting without the drowsiness, extrapyramidal effects and perceptual alterations that are common with current therapy. Onondesertin hydrochloride (Zofran), a promising new antiemetic, seems to be very effective in controlling nausea and vomiting without undesirable side-effects (Marty, Pouillart, School, et al. 1990; Cubeddu, Hoffman, Fuenmayor, & Finn 1990; Schmoll, 1989). At this point, research on ondansetron is limited, due to its relatively recent development. It would be useful to evaluate how acceptable this drug is to the adolescents and what impact alleviation of nausea and vomiting has on adherence.

In addition to drug therapy, it would be interesting to look at alternative approaches to the treatment of chemotherapy-induced nausea and vomiting and subsequent effects on adherence. Some success in alleviating post-treatment and anticipatory nausea and vomiting has been reported with relaxation techniques such as guided imagery, hypnosis (Cotanch, Hockenberry & Herman, 1985), systematic desensitization (Moore & Morell, 1982), progressive muscle relaxation and biofeedback (Burish, Shariner & Lyles, 1981). These techniques are relatively easy to learn, can be controlled by the patient, and have no side-effects. They might prove especially useful for adolescents who refuse antiepileptic drugs.

The extent to which social support influences adherence is an area that could use further study. There was anecdotal evidence in the charts indicating many of the patients who modified and delayed treatment also had chaotic family relationships. In one instance, a mother failed to renew her daughter’s prescription for chemotherapy for three weeks as a punishment for bad behaviour. In other instances, it was mentioned that parents were having difficulties coping with their own lives and were not able to offer much support to the patient. It is not known to what degree family stability or support is influential in adherence in an adolescent cancer population, nor to what extent health care professionals can aid in a family situation where social support is lacking. Further investigation may provide more understanding in this area.

Summary

In summary, non-adherence is a complex phenomenon and, in adolescents, one we know very little about. The significant rate of non-adherence observed in this Canadian adolescent population warrants a prospective study. Patient interviews would give us information about the specific factors that lead to the decision to delay or modify treatment. Treatment delay and modification needs to be examined as it relates to long-term survival. The severity and persistence of nausea and vomiting and the inability to tolerate antiepileptic needs to be examined in relation to treatment delay and modification.

Acknowledgement

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References


