Attenuated regimen of biweekly gemcitabine/nab-paclitaxel in patients aged 65 years or older with advanced pancreatic cancer

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Abstract

Background: Treatment with gemcitabine/nab-paclitaxel confers a survival benefit over gemcitabine monotherapy in patients with advanced pancreatic cancer (APC). However, such treatment can be associated with significant toxicities especially in older patients and carries practical disadvantages related to a weekly schedule along with financial cost. We retrospectively analyzed patients > 65 years of age with APC who received a modified biweekly regimen of gemcitabine/nab-paclitaxel to evaluate efficacy and toxicity.

Methods: Patients aged > 65 years with chemo-naïve APC with Eastern Cooperative Oncology Group performance status ≤ 2 were studied. Patients were treated with a modified regimen of gemcitabine 1000 mg/m² and nab-paclitaxel 125 mg/m² every 2 weeks on days 1 and 15 of a 28-day cycle. Patients were evaluated for progression-free survival (PFS) and overall survival (OS) with analyses performed using the Kaplan–Meier method. Adverse events were recorded on the day of chemotherapy. Cancer antigen 19.9 was measured in every cycle and restaging scans were performed every two cycles.

Results: A total of 73 patients (median age: 73 years; range: 66–93) were treated with biweekly gemcitabine/nab-paclitaxel as first-line treatment. The median OS and PFS were 9.1 months and 4.8 months, respectively. Around 66% of patients received growth-factor support based on American Society of Clinical Oncology guidelines and no patient developed neutropenic fever. The incidences of grade ≥ 3 toxicity for neutropenia, anemia, thrombocytopenia, and neurotoxicity were 2%, 7%, 3%, and 5%, respectively. Dose reductions of gemcitabine/nab-paclitaxel were required in 10% and 4% patients, respectively.

Conclusion: In patients older than > 65 years of age with APC, a modified regimen of biweekly gemcitabine/nab-paclitaxel was found to be effective when compared with the historical control from the MPACT study. This regimen allowed for fewer dose reductions, reduced healthcare costs from additional appointments, travel-related cost, as well as a favorable side-effect profile while maintaining efficacy. Though retrospective in nature, this study underlines the need for further investigation, particularly in elderly patients with poor performance status, such as those with pancreatic cancer, and in order to combine with a third agent, such as a targeted treatment or immunotherapy.

Keywords: advanced pancreatic cancer, attenuated regimen, biweekly, chemotherapy, elderly, improved toxicity

Received: 16 June 2020; revised manuscript accepted: 29 October 2020.
Introduction

Despite various revolutions in the field of oncology, advanced pancreatic cancer (APC) has a grim prognosis and is now the third leading cause of cancer-related deaths in the USA. It is estimated that pancreatic cancer will become the second leading cause of cancer-related deaths in the USA by 2030. Approximately 56,770 Americans will be diagnosed with pancreatic cancer and more than 45,750 will die of the disease yearly, highlighting how lethal this disease has become. Most patients present with advanced disease leading to the dismal 5-year survival rate of less than 10%. Furthermore, it is estimated that the number of elderly patients with pancreatic cancer will continue to rise due to the aging population. Management of these patients is quite limited and further hindered by underrepresentation of this population in clinical trials.

Gemcitabine historically was considered the standard of care for metastatic pancreatic cancer as it provided a survival benefit as well as alleviation of symptoms compared with fluorouracil (5-FU). Many trials after this yielded disappointing results as they failed to show improvement on this landmark trial including combination treatments with novel agents and other chemotherapy agents in addition to gemcitabine monotherapy.

More recently, two chemotherapy combination regimens have shown superiority in patients with metastatic disease in randomized phase III trials. In the PRODIGE/ACCORD trial, a combination of 5-FU with oxaliplatin and irinotecan showed an overall survival benefit with a median overall survival (OS) of 11.1 months compared with 6.8 months in the gemcitabine arm. Of note, patients older than 75 years of age were excluded in this trial. In the MPACT trial, the combination of gemcitabine/nab-paclitaxel showed superiority over gemcitabine with a median OS of 8.5 months in the combination arm versus 6.7 months in the monotherapy arm. Treatment was administered with gemcitabine (1000 mg/m²) and nab-paclitaxel (125 mg/m²) given on days 1, 8, and 15 of a 4-week cycle. Although these results were promising and effective, they came at a cost of increased toxicity as well as practical disadvantages for the patients. Prior studies have suggested a modified regimen of biweekly gemcitabine/nab-paclitaxel could lessen adverse effects of the regimen while maintaining efficacy. Although promising, this regimen has not been readily studied in the elderly population and it is unclear how to best manage these patients in the setting of advanced disease. Therefore, we adopted an attenuated regimen of biweekly gemcitabine/nab-paclitaxel for patients older than 65 years of age as treatment for APC and present the efficacy, side-effect profile, and cost-analysis data of our study.

Methods

Patients and study design

This study was approved with written consent by our institution’s Institutional Review Board. Ethics approval was not sought as this was a retrospective review. We performed a retrospective analysis of patients older than 65 years of age with chemo-naïve APC from 2015 to 2017. All patients were treated at our institution and monitored for efficacy as well as toxicity. All patients who were treated with this modified regimen were included in the analysis during this time period.

Eligibility

Patients older than 65 years of age with locally advanced/unresectable or metastatic pancreatic adenocarcinoma were eligible. Diagnosis through tissue biopsy was required. All patients had to have Eastern Cooperative Oncology Group performance status of 0 or 1. Patients were treated with this regimen based on the discretion of the treating physician. Decisions were based upon open discussions with the patients and mostly related to patient preference for not wanting to come to the treatment room for weekly treatments. There were no limitations based on comorbidities as long as performance status was adequate. Patients were chemo-naïve as no previous treatment for pancreatic cancer was allowed.

Treatment schedule

Chemotherapy was administered on days 1 and 15 of a 28-day cycle. Patients received 125 mg/m² of nab-paclitaxel followed by 1000 mg/m² of gemcitabine on both days. Pre-medications and antiemetics were ordered according to American Society of Clinical Oncology (ASCO) guidelines. Granulocyte-colony stimulating factor was also administered according to ASCO guidelines. Treatment was continued until evidence of disease progression or unacceptable toxicity.
Comprehensive physician office visits were conducted every 2 weeks prior to initiation of treatment and between each cycle of treatment. Detailed history and physical examination were performed at each clinic visit. Toxicities were assessed using the National Cancer Institute Common Terminology Criteria for Adverse Events, version 5. Computed tomography imaging of the chest, abdomen, and pelvis was used to assess disease status every two cycles or as clinically indicated. Baseline blood work was obtained prior to initiating treatment as well as prior to each subsequent treatment. Laboratory values included a complete blood count, comprehensive metabolic panel, and cancer antigen 19.9 levels. Baseline information to assess disease status was also evaluated including demographic information, metastatic sites of disease, number of metastatic sites, biliary stent placement, and subsequent lines of treatment. All information was recorded in each patient’s secure online medical electronic record.

Statistical analyses
Study endpoints that were evaluated included progression-free survival (PFS), OS, and toxicity profile. Descriptive statistics were used to evaluate demographic information, subsequent lines of treatment, as well as toxicity profile. The Kaplan–Meier method was used for the calculation of PFS and OS. PFS was calculated from the start of chemotherapy until disease progression or death, whichever came first. OS was obtained from the initiation of chemotherapy until death from any cause.

Results
A total of 73 patients who received this attenuated regimen of gemcitabine/nab-paclitaxel for APC as first-line treatment were evaluated. Demographic data are presented in Table 1. All patients were older than 65 years of age with a median of age 73 years. The median OS was 9.1 months and the median PFS was 4.8 months. All patients were able to receive second-line treatments once progression of disease was noted. Standard-of-care subsequent treatments were utilized including capcitabine monotherapy or other 5-FU-based regimens as outlined in Figure 1.

Most toxicities were rated as grade 2 or less and related to fatigue, neuropathy, skin rash, or nausea. Grade 3 or higher toxicities were rare in our cohort. The incidences of grade 3 or 4 hematologic toxicities for neutropenia, anemia, and thrombocytopenia were minimal, resulting in 2%, 10%, and 3%, respectively. Around 66% of patients received growth-factor support based on the ASCO guidelines. Although the standard to receive primary prophylaxis includes regimens where the risk of neutropenia is >20%, many patients in our analysis received growth-factor support largely due to older age and comorbidities. It has been shown that older patients aged greater than 65 years may
be more vulnerable to chemotherapy-related febrile neutropenia. Furthermore, having more advanced cancer and comorbidities carries a higher risk of complications from chemotherapy. In our analysis, no patients developed neutropenic fever. Furthermore, rates of grade 3 or 4 nonhematologic toxicity were also low. For example, grade 3 or 4 neurotoxicity was found in 5% of patients. Dose reductions of gemcitabine/nab-paclitaxel were required in only 10% and 3% of patients, respectively.

**Figure 1.** Second-line treatments after progression of disease.
5-FU, fluorouracil; CAPOX, capecitabine and oxaliplatin; FOLFI, folic acid, 5-FU, and irinotecan; FOLFIRINOX, 5-FU, irinotecan, oxaliplatin, and folic acid; FOLFOX, folic acid, 5-FU, and oxaliplatin; LCV, leucovorin.

The side-effect profile is outlined in Figure 2 and is compared with the historical MPACT trial in Table 2.

**Discussion**

Our study showed that an alternative biweekly treatment regimen with gemcitabine/nab-paclitaxel appears to have similar efficacy when compared with the historical control from the MPACT trial. Although this trial showed statistically significant results and improved upon the standard of gemcitabine monotherapy, it came at a cost of increased toxicity, dose reductions, and delays in treatment. In this trial, nab-paclitaxel dosing was administered at 125 mg/m² on days 1, 8, and 15 of a 28-day cycle. Gemcitabine was also administered on days 1, 8, and 15 at 1000 mg/m². Dose reduction was required with the use of nab-paclitaxel and gemcitabine in 41% and 47% of patients, respectively. This compares with a rate of 3% and 10% in our cohort. A total of 22% of patients permanently discontinued treatment in the MPACT study. Furthermore, rates of hematologic and nonhematologic toxicities were much lower with the biweekly attenuated regimen.

Previous trials have shown that biweekly gemcitabine monotherapy or combination therapy may improve the side-effect profile while maintaining efficacy in various settings compared with weekly dosing. Previous retrospective studies have also shown that a modified schedule of treatment with gemcitabine/nab-paclitaxel on days 1 and 15 may maintain efficacy while improving upon the toxicity profile of these agents in this setting of APC.

**Figure 2.** Side-effect profile.
However, these studies did not focus on primarily elderly patients. Pancreatic cancer is known to be a disease of older adults, with the median age of 71 years at diagnosis in the USA. In the MPACT study, only 41% of patients were aged 65 years or older. A subgroup analysis showed that there was a benefit of PFS in this population with combination therapy; however, there was no difference in OS with a hazard ratio of 0.81 (0.63–1.03). The older population remains a heterogeneous group and difficult to treat based on unclear tolerability of treatments as well as exclusion from some clinical trials.

The findings from our study suggested that the attenuated regimen maintains relative efficacy with considerable improvement in side-effect profile. Continuing treatments on schedule without dose reductions and interruptions may also help explain why survival was unchanged. Very few patients required dose modifications, and no patients had to discontinue treatment altogether. There were also no instances of neutropenic fever with only 2% of patients experiencing grade 3 or 4 neutropenia. This may explain why survival data are slightly improved in our cohort. However, definite conclusions cannot be made in this retrospective analysis and would require randomized data. It should be noted, however, most patients received growth-factor support due to advanced age in line with current recommendations.

In addition to an improved toxicity profile, the modified biweekly regimen of gemcitabine/nab-paclitaxel also has a significant impact on healthcare costs. By eliminating day 8 of treatment, financial toxicity can be greatly reduced factoring in costs of chemotherapeutic agents, infusion related costs, and personnel need at the cancer center. Furthermore, there is increasing awareness of quality-of-life measures in patients with cancer. By reducing treatment room visits, patients may have considerable improvement in these measures related to convenience factors as well as transportation needs. This may also encompass more quality time with loved ones at home, which is of paramount importance for most patients with APC. We did not include a quality-of-life assessment in our study, but this could present an interesting area of research in this population.

Lastly, improvement in these toxicities while producing similar results may allow for additional treatments. By preserving toxic side effects from treatment, further investigation can be performed with the addition of a third agent to this combination. Management of advanced disease is evolving with ongoing research into various targeted treatments and immunotherapies in APC, but mortality rate remains high. With the continued advancements in oncology, there is a new focus on precision medicine and the identification of novel biomarkers. With the emergence of immunotherapy and these targeted treatments, there is a research need to explore further these associations. With our attenuated regimen of gemcitabine/nab-paclitaxel, we present an opportunity to further improve outcomes while minimizing toxicities in this historically lethal disease.

We acknowledge the limitations that accompany our study. This was carried out in a retrospective nature which included inherent biases such as the possibility of selection bias. Cross-over comparison from different studies also lends to bias as different populations were evaluated. There is no direct comparison of demographic data and lack of a control arm could limit generalizability of results. A relatively small sample size is also a limitation as this was conducted solely at our institution as a single-center retrospective analysis. However,

| Table 2. Cross comparison with historical control. |
|-----------------------------------------------|
| **Survival data**                              |
| Progression-free survival, median (months)     |
| Our study: n (%)                              |
| MPACT trial: n (%)                            |
| Overall survival, median (months)             |
| 4.8                                          |
| 5.5                                          |
| **Grade 3 or 4 toxicities**                    |
| Growth-factor support                         |
| 48 [66%]                                      |
| 110 [26%]                                     |
| Anemia                                        |
| 7 [10%]                                       |
| 53 [13%]                                      |
| Neutropenia                                   |
| 2 [2%]                                        |
| 153 [38%]                                     |
| Neutropenic fever                             |
| 0 [0%]                                        |
| 14 [3%]                                       |
| Thrombocytopenia                              |
| 3 [3%]                                        |
| 52 [13%]                                      |
| Neurotoxicity                                 |
| 5 [5%]                                        |
| 70 [17%]                                      |
| **Dose reduction in patients**                |
| Nab-paclitaxel                                |
| 3 [3%]                                        |
| 41%                                          |
| Gemcitabine                                   |
| 7 [10%]                                       |
| 47%                                          |
keeping these points in mind, we believe our patients represented a reasonably similar profile to the general population in this advance disease setting. All patients had a good performance status at the discretion of the treating physician and were given standard treatment doses. Patients were monitored as they would be in any clinical setting. In the setting of an elderly population, our results showed that a biweekly regimen of gemcitabine/nab-paclitaxel allowed for a favorable side-effect profile while maintaining efficacy in APC.

Conclusion
An attenuated regimen of biweekly gemcitabine/nab-paclitaxel for the treatment of APC in elderly patients showed similar results to historical precedents. Using this regimen minimized toxicities associated with these agents, reduced healthcare costs, and perhaps improved quality-of-life measures in these patients. Better tolerability of these agents may allow for combination with a third agent, such as a targeted treatment or immunotherapy, especially as we enter the era of precision medicine in oncology. Though retrospective in nature, this study underlines the need for further investigation, particularly in elderly patients with APC to optimize outcomes while minimizing toxicities and preserving quality of life.

Conflict of interest statement
The authors declared the following potential conflicts of interest with respect to the research, authorship, and/or publication of this article: Dr. Saif had grant funding from Celegene. The rest of the authors have no conflicts of interest to declare.

Funding
The authors received no financial support for the research, authorship, and/or publication of this article.

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