## ORIGINAL ARTICLE

## Resection Status, Age and Nodal Involvement Determine Survival among Patients Receiving Adjuvant Chemoradiotherapy in Pancreatic Adenocarcinoma

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#### ABSTRACT

**Context** Pancreas cancer can potentially be cured by resection, but the role of adjuvant chemotherapy and/or chemoradiation has been controversial. **Objectives** To better define clinicopathological factors that may serve as predictive and/or prognostic variables. **Patients** Between 1984 and 2006, we retrospectively analyzed 91 patients with pancreas cancer treated with pancreaticoduodenectomy or total pancreatectomy followed by adjuvant 5-fluorouracil-based chemoradiation at the University of Pennsylvania. Final pathological coding including margin status was confirmed by a pathologist. **Interventions** Patients were treated with 48.6 to 63.0 Gy, and 96.7% completed their prescribed radiation dose. **Main outcome measures** The prognostic significance of demographic factors, stage, year of surgery, tumor location, grade, resection status, and number of positive lymph nodes on overall survival were examined. **Results** With a median follow-up of 6.5 years, the overall median survival was 2.3 years (95% CI 1.5-3.2 years), and the 5-year overall survival was 28.9%. In multivariate analysis, completeness of resection (P<0.001), fewer number of positive lymph nodes (0 *vs.* 1-2 *vs.* 3 or more) (P=0.004), and age less than, or equal to, 60 years (P=0.006) were all independently associated with improved overall survival. The overall survival reported in this study compares favorably with the results of other single-institution studies and with the RTOG 97-04 trial. **Conclusions** Adjuvant 5-FU-based chemoradiation following radical pancreatectomy can be delivered safely and results in comparatively good overall survival. The results of this analysis underscore the importance of resection status, number of involved lymph nodes and patient age as prognostic characteristics. These factors may be considered stratification variables for future post-pancreatectomy adjuvant therapy trials.

#### INTRODUCTION

Although surgical resection is the only treatment modality that provides the potential for cure in patients with pancreatic adenocarcinoma, postoperative patients receiving no further treatment remain at a very high risk for local and distant relapse. Starting with the Gastrointestinal Tumor Study Group (GITSG) study [1] in 1985 and then followed by Whittington *et al.* [2] in 1991, improved survival has been demonstrated with adjuvant chemoradiotherapy. Over the subsequent years, multiple reports have evaluated the role of

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adjuvant therapy and have shown improved clinical outcomes, including overall survival, with adjuvant therapy when compared with surgery alone [3, 4, 5, 6]. Nonetheless, controversies abound regarding the exact role and benefit of adjuvant chemoradiotherapy [7, 8, 9, 10, 11, 12], and clinicopathological factors that may serve as predictive and/or prognostic variables are not well-defined [13, 14, 15, 16, 17, 18, 19]. In this retrospective analysis, we report the experience with adjuvant chemoradiation after pancreaticoduodenectomy or total pancreatectomy at the University of Pennsylvania and provide an analysis of commonly accepted clinicopathological factors to determine their prognostic value for favorable survival in this population.

#### METHODS

#### **Study Population**

We created a secure database of all pancreatic cancer patients consecutively treated at the Hospital of the University of Pennsylvania from 1981 to 2006. After cross-referencing with the Department of Radiation Oncology medical record system, a total of 91 patients with adenocarcinoma of the pancreas were identified that were treated with adjuvant chemoradiation, and represent the patient populations for this analysis.

#### **Data Collection**

All T and N stages were included; stage coding was by AJCC 7<sup>th</sup> edition criteria. Data collected included demographics (age, gender, race), location of pancreatic tumors (head *vs.* body/tail), and pathologic features (size, histologic grade, nodal status, and resection status). Microscopic *versus* gross residual disease was verified by careful review of both operative and pathology reports. Final pathological coding was confirmed with a pathologist specializing in gastrointestinal malignancies (E.E.F.). Time points included in this analysis consisted of dates of surgery, discharge and adjuvant treatment. Radiation Oncology Department clinic charts were reviewed to collect treatment details. Death dates were determined via the Social Security Death Index.

#### Treatment

Twelve surgeons performed all pancreatic resections at our institution during the study period. However, a single surgeon (E.F.R.) performed a majority (twothird) of these resections. The 30- and 60-day crude mortality rates for all resections of malignant lesions from the pancreaticoduodenobiliary tract during the study period were 1.8% and 3.4%, respectively (n=572).

The use of concurrent chemoradiotherapy in the postoperative setting became routine at the University of Pennsylvania in 1984, and therefore, only patients treated during or after 1984 are included. While detailed chemotherapy data were not available for some patients, documented records in the Radiation Oncology clinic notes confirmed that all patients in this report received a 5-FU-based regimen of concurrent chemoradiotherapy. Prior to 1986, our institutional policy was to treat patients with bolus 5-FU (500 mg/m<sup>2</sup>/day) on the first three and last three days of radiation. This regimen evolved to the use of a 96 h 5-FU infusion (1,000 mg/m<sup>2</sup>/day) during the first and fifth weeks of radiation. In 1991, the treatment policy

changed to 5-FU by protracted infusion (250  $\text{mg/m}^2/\text{day}$ ) continuously throughout the course of radiation. Patients who tolerated radiation and completed treatment with a performance status of 1 or 2 generally received maintenance chemotherapy after completion of adjuvant chemoradiotherapy with bolus 5-FU, 5-FU with low dose folinic acid, or gemcitabine. All patients were treated with high-energy (megavoltage) photons delivered with a linear accelerator, and CT-defined clinical target volumes routinely included adjacent nodal beds. Patients were treated to 48.6 or 50.4 Gy, but some patients with R1 or R2 resections were treated to higher doses (54.0, 59.4 or 63.0 Gy). Eighty-eight patients (96.7%) completed their prescribed radiation dose.

### ETHICS

The secure database was created after approval from the University of Pennsylvania Institutional Review Board.

The written informed consent was obtained for all investigations made according to the usual clinical practice. The study protocol was approved by the institutional review committee. The study protocol conforms to the ethical guidelines of the World Medical Association Declaration of Helsinki.

#### STATISTICS

Overall survival was defined from the date of surgery to the date of death due to any cause, or censored at last patient contact. The 5-year overall survival rate was estimated by the Kaplan-Meier method and was reported together with the standard error (SE). The hazard ratio and statistical significance of each clinical factor was assessed by Cox regression analysis. To determine whether clinical factors were significant independent prognostic factors, multivariate modeling was performed using Cox regression analysis using a backward elimination selection strategy. Prior to modeling, tumor variables that were significant by univariate regression analysis (T stage, N stage, number of lymph nodes involved and resection status) were tested for collinearity by a linear trend test. These results were used to arrive at a parsimonious final model. All analyses were performed in SPSS version 17.0 (SPSS Inc., Chicago, IL, USA), and two-tailed P

|                         | University of Penn series<br>(present study) | Mayo Clinic series [1] | Johns Hopkins series [2] | RTOG 97-04 trial [3] |
|-------------------------|--|------------------------|--------------------------|----------------------|
| Total subjects          | 91   | 466                    | 271                      | 451                  |
| Median age              | 60 years                                     | 67 years               | 64 years                 | 61-62 years          |
| Male                    | 60.4% (n=55)                                 | 54%                    | 54%                      | 57%                  |
| Fumor in head           | 86.8% (n=79)                                 | NA                     | NA                       | 86%                  |
| 3 or T4 tumor           | 74.1% (63/85)                                | 52%                    | NA                       | 75%                  |
| N1                      | 54.9% (n=50)                                 | 48%                    | 80%                      | 66%                  |
| High grade <sup>a</sup> | 46.4% (39/84)                                | 69%                    | 43%                      | 27% <sup>b</sup>     |
| Margins positive        | 38.5% (n=35)                                 | NA                     | 45%                      | 34%                  |

NA: Data not available

<sup>a</sup> High grade is defined as moderately poor to poor

<sup>b</sup> Data missing on 21 patients

| Variable                                    | No. of patients | 5-year overa |         | Hazard ratio | 95% confidence | P value |
|---|-----------------|--------------|---------|--------------|----------------|---------|
|   |                 | Rate SE      |         | _            | interval       |         |
| Pancreas cancer patients                    | 91              | 28.9%        | 5.5%    |              |                |         |
| Gender:                                     |                 |              |         |              |                | 0.962   |
| - Male                                      | 55              | 29.5%        | 7.1%    | Reference    | -              |         |
| - Female                                    | 36              | 28.1%        | 9.0%    | 1.0          | 0.6-1.7        |         |
| Race:                                       |                 |              |         |              |                | 0.026   |
| - Caucasian                                 | 72              | 23.2%        | 6.1%    | Reference    | -              | -       |
| - African-American                          | 16              | 58.4%        | 13.1%   | 0.4          | 0.2-0.9        | 0.025   |
| Other                                       | 3               | 0.0          | 0.0     | 1.8          | 0.5-5.7        | 0.344   |
| Age:  |                 |              |         |              |                | 0.006   |
| 60 years or less                            | 45              | 46.6%        | 8.7%    | Reference    | -              |         |
| More than 60 years                          | 46              | 14.1%        | 5.9%    | 2.1          | 1.2-3.6        |         |
| Date of surgery:                            |                 |              |         |              |                | 0.542   |
| 1980-1984                                   | 1               | 0.0          | 0.0     | 0.4          | 0.05-3.3       | 0.409   |
| 1985-1989                                   | 15              | 12.4%        | 10.7%   | 0.2          | 0.03-1.8       | 0.163   |
| 1990-1994                                   | 25              | 38.8%        | 9.9%    | 0.2          | 0.04-2.2       | 0.229   |
| 1995-1999                                   | 19              | 25.3%        | 10.2%   | 0.3          | 0.04-2.6       | 0.222   |
| 2000-2004                                   | 21              | 34.3%        | 11.2%   | 0.2          | 0.02-2.0       | 0.167   |
| 2005-2006                                   | 10              | NR           | -       | Reference    | -              | -       |
| Days from surgery to discharge:             | 10              | INK          |         | Reference    |                | 0.827   |
| 14 days or less                             | 39              | 36.1%        | 9.6%    | Reference    | -              | 0.027   |
| More than 14 days                           | 27              | 31.0%        | 9.0%    | 1.1          | 0.6-2.0        |         |
| Days from surgery to start of radiotherapy: | 21              | 51.070       | 9.270   | 1.1          | 0.0-2.0        | 0.261   |
|   | 46              | 20.7%        | 6.7%    | Reference    | _              | 0.201   |
| 60 days or less                             | 40              |              |         |              | 0.5-1.2        |         |
| More than 60 days                           | 45              | 41.0%        | 8.1%    | 0.8          | 0.3-1.2        | 0.200   |
| fumor location:                             | -               | 0.6 70/      | 5 50/   | D (          |                | 0.306   |
| Head  | 79              | 26.7%        | 5.7%    | Reference    | 0.0.1.5        |         |
| Distal                                      | 12              | 50.9%        | 16.3%   | 0.7          | 0.3-1.5        |         |
| Grade:                                      |                 | 2.5.10       | 4.4.004 | 5.0          |                | 0.749   |
| Well to moderately-well                     | 14              | 36.4%        | 14.0%   | Reference    | -              | -       |
| Moderate                                    | 31              | 27.0%        | 9.5%    | 1.1          | 0.5-2.4        | 0.720   |
| Moderately-poor to poor                     | 39              | 21.6%        | 8.0%    | 1.3          | 0.6-2.6        | 0.466   |
| fumor size:                                 |                 |              |         |              |                | 0.233   |
| 2 cm or less                                | 17              | 42.5%        | 13.0%   | Reference    | -              |         |
| Greater than 2 cm                           | 60              | 22.2%        | 6.8%    | 1.5          | 0.8-2.9        |         |
| ſ stage:                                    |                 |              |         |              |                | 0.048   |
| T1-T2                                       | 22              | 34.7%        | 12.8%   | Reference    | -              |         |
| T3-T4                                       | 63              | 24.0%        | 6.2%    | 1.8          | 1.0-3.4        |         |
| N stage:                                    |                 |              |         |              |                | 0.036   |
| NO  | 41              | 45.2%        | 8.8%    | Reference    | -              |         |
| N1  | 50              | 15.2%        | 6.2%    | 1.7          | 1.0-2.8        |         |
| Resection status:                           |                 |              |         |              |                | <0.001  |
| Complete                                    | 56              | 35.3%        | 7.8%    | Reference    | -              | -       |
| Microscopic                                 | 24              | 25.0%        | 9.4%    | 1.7          | 0.9-2.9        | 0.079   |
| Gross disease                               | 11              | 0.0          | 0.0     | 6.7          | 3.0-15.0       | < 0.001 |
| No. of positive lymph nodes:                |                 |              |         |              |                | 0.004   |
| 0   | 41              | 45.2%        | 8.8%    | Reference    | -              | -       |
| 1-2   | 35              | 23.1%        | 8.6%    | 1.3          | 0.8-2.3        | 0.323   |
| 3 or more                                   | 15              | 0.0          | 0.0     | 3.4          | 1.8-6.8        | < 0.001 |

NR: not reached

<sup>a</sup> Cox regression analysis

values less than 0.05 were considered statistically significant.

#### RESULTS

#### **Patient and Disease Characteristics**

A comparison of these characteristics with other large series of pancreatic cancer patients treated with adjuvant chemoradiotherapy is presented in Table 1. Demographic and disease characteristics of the patients in this analysis are shown in Table 2. The majority of patients were Caucasian (72 patients, 79.1%); men comprised 60.4% of the study population (55 patients). The median age was 60 years. Tumors were predominately located in the head of the pancreas (79 patients, 86.8%). The majority (63 patients, 69.2%) were T3/T4 and over half (50 patients, 54.9%) had nodal involvement. High grade tumors (39/84 patients,

46.4%) and positive resection margins (35 patients, 38.5%) were seen in less than half of the cases.

#### **Survival Outcomes**

The median follow-up was 6.5 years for all patients. Of the 91 patients in the series, 64 (70.3%) are deceased. The median survival from the date of surgery was 2.3 years (95% confidence interval 1.5-3.2 years). The overall survival rate ( $\pm$ SE) at 2 and 5 years was 56.2 $\pm$ 5.4% and 28.9 $\pm$ 5.5%, respectively (Figure 1). For the 27 patients alive at last contact, the median followup was 2.8 years.

#### **Characteristics Associated with Survival**

Gender was not significantly associated with overall survival (P=0.962), but race was (P=0.026), with 5-year overall survival rates ( $\pm$ SE) of 58.4 $\pm$ 13.1% for African-American and 23.2 $\pm$ 6.1% for Caucasian

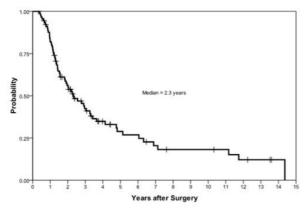


Figure 1. Overall survival after surgical resection for pancreatic adenocarcinoma. Median follow up was 6.5 years, and the median overall survival was 2.3 years. Crosses represent censored patients.

patients (Table 2). Age, dichotomized at the median (60 years), was also found to be significantly associated with overall survival (P=0.006). The 5-year overall survival rate for patients older than 60 years was  $14.1\pm5.9\%$  and for patients up to 60 years was  $46.6\pm8.7\%$ , with a corresponding hazard ratio (HR) of 2.1 (Figure 2a).

Patients with T1-T2 lesions had significantly better overall survival than patients with T3-T4 lesions (5year survival 34.7±12.8% vs. 24.0±6.2%, P=0.048). Both the American Joint Committee on Cancer (AJCC) nodal status (N1 vs. N0) (HR 1.7, P=0.036), and the number of positive lymph nodes showed a statistically significant association with overall survival. We defined cutoffs for number of positive lymph nodes as 0, 1-2 or more than 2. The 5-year overall survival rates for these three groups were 45.2±8.8%, 23.1±8.6% and 0%, respectively (P=0.004, Figure 2b). Resection status was also strongly associated with overall survival (P<0.001, Figure 2c). The 5-year overall survival rates were 35.3±7.8%, 25.0±9.4% and 0% for R0, R1 and R2 patients, respectively. Tumor location within the pancreas, histological grade and tumor size dichotomized at 2 cm were not significantly associated with overall survival.

Given ongoing improvements in surgical technique and perioperative care, we evaluated the impact of treatment era and stratified patients by 5-year increments according to year of surgery. There was no association between overall survival and study period (P=0.542). Five-year overall survival rates were consistently at or above 25% from 1990 onward. Duration of hospital stay and interval from surgery to the initiation of adjuvant therapy, each dichotomized at the median value, were also not significantly associated with survival (P=0.827 and P=0.261, respectively).

#### **Multivariate Analysis**

To determine whether the factors found to be significant in univariate analyses were independent prognostic factors, a multivariate analysis was undertaken, taking into account collinearity among variables. T stage was found to be strongly associated with resection status (P=0.003) and was thus not retained in the final model. There was a weak association between resection status and the number of positive lymph nodes (P=0.106) with a fewer nodepositive patients who had R0 resections. Resection status, age (dichotomized at 60 years) and the number of positive lymph nodes (in 3 groups, as defined earlier) were found to be independent predictors of survival from surgery (Table 3). Resection status stood out as the most strongly associated characteristic, with gross residual disease (R2) having an adjusted hazard ratio of 7.8 (95% CI: 3.4-17.8), when compared to no residual disease (R0).

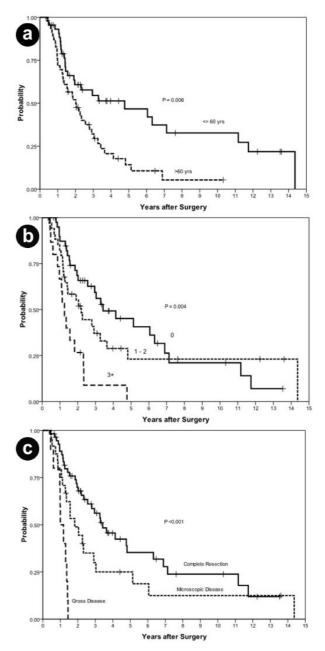


Figure 2. Overall survival after surgical resection for pancreatic adenocarcinoma by age group (a.), number of positive lymph nodes (b.), and resection status (c.).

| Table 3. Multivariate | analysis of surviva | l from date of surgerv | (Cox regression analysis). |
|-----------------------|---------------------|------------------------|----------------------------|
|                       |                     |                        |                            |

| Variable                                   | Univariate results of candidate factors |          |         | Multivariate results (final model) |          |         |
|--|---|----------|---------|------------------------------------|----------|---------|
|  | Unadjusted HR                           | 95% CI   | P value | Adjusted HR                        | 95% CI   | P value |
| Race:                                      |   |          | 0.026   | -                                  | -        | -       |
| - African-American vs. Caucasian           | 0.4                                     | 0.2-0.9  | 0.025   |                                    |          |         |
| - Other vs. Caucasian                      | 1.8                                     | 0.5-5.7  | 0.344   |                                    |          |         |
| Age: More than 60 years vs. up to 60 years | 2.1                                     | 1.2-3.6  | 0.006   | 2.2                                | 1.3-3.8  | 0.004   |
| <b>T stage:</b> T3-T4 vs. T1-T2            | 1.8                                     | 1.0-3.4  | 0.048   | -                                  | -        | -       |
| N stage: N1 vs. N0                         | 1.7                                     | 1.0-2.8  | 0.036   | -                                  | -        | -       |
| Resection status:                          |   |          | <0.001  |                                    |          | < 0.001 |
| - Microscopic (R1) vs. complete (R0)       | 1.7                                     | 0.9-2.9  | 0.079   | 1.4                                | 0.8-2.5  | 0.224   |
| - Gross disease (R2) vs. complete (R0)     | 6.7                                     | 3.0-15.0 | < 0.001 | 7.8                                | 3.4-17.8 | < 0.001 |
| No. of positive lymph nodes:               |   |          | 0.004   |                                    |          | 0.003   |
| - 1-2 vs. 0                                | 1.3                                     | 0.8-2.3  | 0.323   | 1.3                                | 0.7-2.3  | 0.362   |
| - 3+ <i>vs</i> . 0                         | 3.4                                     | 1.8-6.8  | < 0.001 | 3.4                                | 1.7-6.8  | 0.001   |

HR: hazard ratio

#### DISCUSSION

# Adjuvant Therapy Outcomes and Prognostic Variables

At the University of Pennsylvania, we have offered postoperative chemoradiotherapy routinely for over two decades to all patients with pancreatic cancer who have undergone potentially curative surgical resection. In 91 patients treated, a 5-year overall survival rate of 28.9% and a median survival of 2.3 years has been achieved. These results compare favorably with similar reports of postoperative adjuvant chemoradiotherapy, where 5-year overall survival rates range from 16.5% to 28.0% and median survival ranges from 1.49 to 1.84 years (Table 4). The EORTC study reported a 5-year overall survival rate of 20% and a median survival of 1.43 years for pancreatic head tumor patients [20]. In addition, the 5-year overall survival results reported by ESPAC-1 were 13% and 7% in patients who received postoperative chemoradiotherapy with or without maintenance chemotherapy, respectively [5]. Also, in the American College of Surgeons Oncology Group (ACOSOG) Z05031 multi-institutional phase II study, which continues to report some of the most favorable results presented to date, 89 patients treated with postoperative radiotherapy to 50.4 Gy with concurrent protracted venous infusion of 5-FU, cisplatin and interferon-alpha achieved a median survival time of 2.26 years [21, 22].

In an attempt to identify independent prognostic factors that may be associated with survival after adjuvant chemoradiotherapy, in univariate we found that gender, primary tumor location (pancreatic head versus distal site), and histologic grade, as well as race in the successive multivariate analysis, all failed to demonstrate associations with survival in this series. Our results, however, indicate that age (dichotomized at 60 years), resection status and number of positive lymph nodes were strongly, and independently, associated with survival. Although T stage and resection status were strongly associated, only resection status was retained in the final multivariate model and thus should be viewed as a superior predictor of survival. In most other major series, resection status, T and N stages, and tumor grade, in

various combinations, are the variables that have been consistently shown to be associated with survival outcomes [5, 14, 15, 20]. While our results are similar. they indicate that resection status may be the primary determinant of outcomes, and that the number of positive lymph nodes may be a better prognosticator than a simple binary lymph node variable (N0 vs. N1). Keeping in line with this observation, the planned subset analysis by the ACOSOG Z05031 study also failed to identify a prognostic value for T1-T2 vs. T3-T4, tumor size (less than 3 cm), and nodal status [21]. Resection of T4 tumors is only partial by nature, and the lack of long term survivors with R2 resections emphasizes the importance of achieving complete resections when the intent is curative. Although not all series agree on the same age cutoff, we found that age less than 60 years was strongly associated with survival.

In our current report, 97% of patients were able to safely complete their prescribed radiation dose. We believe adequate nutritional support plays an important role and have provided a separate analysis from our experience elsewhere [23]. This includes close attention to nutritional concerns, including the common practice of leaving in place a feeding jejunostomy tubes by at least one surgeon (E.F.R.).

#### Study Limitations

The authors acknowledge that retrospective collection of data from a single institution limits interpretation of

|   | Five-year<br>overall<br>survival <sup>a</sup> | Median<br>survival      |
|---|---|-------------------------|
| University of Penn series (present study) | 28.9%   | 2.30 years              |
| Mayo Clinic series [1]                    | 28.0%   | 1.80 years <sup>b</sup> |
| Johns Hopkins series [2]                  | 20.1%   | 1.49 years <sup>b</sup> |
| RTOG 97-04 trial [3]                      | 16.5%   | 1.84 years <sup>c</sup> |

<sup>a</sup> Survival data for all series are presented for the adjuvant chemoradiotherapy arms only

<sup>b</sup> Mayo Clinic and Johns Hopkins series: survival from date of surgery

<sup>c</sup> RTOG 97-04: survival from date of randomization

our findings. Many patients following resection sought their adjuvant therapy outside of our institution, and the long-term outcomes for these patients were not analyzed. Yet, it should be emphasized that our study population was inclusive of all patients who completed trimodality therapy at our institution with a broad age range (32-81 years) and included 11 patients with grossly positive margins and 10 patients with invasion of the superior mesenteric artery or celiac axis. There has long been a uniform policy to offer trimodality therapy to patients with both favorable and unfavorable pathologic factors, and this is easily identifiable in our dataset with inclusion of pT1N0 lesions with negative margins as well as pT4 lesions, which by definition have grossly positive margins. CA 19-9 levels, which have also been shown to be prognostic in other series, were not available for all of these patients due to the treatment era.

Ultimately, a prospective randomized trial of postoperative chemotherapy with and without radiation is needed to help resolve the controversy regarding the role of adjuvant therapy and prognostic variables for pancreatic cancers. A recently conducted randomized phase II/III study of adjuvant gemcitabine with and without radiation therapy for pancreatic cancer was reported. The study has not yet been unblinded and the pooled analysis (all patients combined) reported a 1year overall survival of 79.4% [24]. Chemoradiation has largely been abandoned by European practitioners, but the Radiation Therapy Oncology Group (RTOG) has recently opened an intergroup trial to address this important question. Also, traditional prognostic and predictive variables may be supplemented with genetic profiles to determine radiosensitivity, and detection of patients with subclinical metastatic disease [25, 26, 27, 281.

#### CONCLUSION

Numerous institutions currently employ a strategy of trimodality therapy with postoperative chemoradiotherapy and the literature is now replete with suggestions of favorable outcomes with this treatment approach [1, 2, 14, 15, 20, 22, 29, 30, 31, 32, 33]. However, definitive information is lacking and until randomized trials establish uniformly accepted standards, results such as those presented in this report can continue to help guide treatment decisions in the adjuvant setting and provide further evidence for the efficacy of adjuvant chemoradiation following resection of pancreatic cancers. The results of this study also underscore the importance of resection status, number of involved lymph nodes and patient age as prognostic characteristics and they may be considered stratification variables for future postpancreatectomy adjuvant therapy trials.

Abbreviations ACOSOG: American College of Surgeons Oncology Group; AJCC: American Joint

Committee on Cancer; RTOG: Radiation Therapy Oncology Group

**Conflicts of interest** The authors have no potential conflicts of interest

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