

# Pancreatic Adenocarcinoma: An Overview of Clinical Challenges and Advances in Treatment

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## DESCRIPTION

Adenocarcinoma of the pancreas, commonly referred to as pancreatic cancer, is one of the most lethal malignancies worldwide. This type of cancer originates in the glandular cells of the pancreas, particularly in the exocrine part, which produces digestive enzymes. Pancreatic cancer is notorious for its late presentation, rapid progression resistance to treatment, making it a significant public health challenge.

Pancreatic cancer ranks as the fourth leading cause of cancerrelated deaths in the United States and has a rising incidence globally. The American Cancer Society estimates that over 60,000 new cases will be diagnosed annually in the U.S. alone, with approximately 48,000 deaths attributed to the disease. Risk factors include age, smoking, obesity, family history certain genetic syndromes such as *BRCA2* mutations. It is most commonly diagnosed in individuals over the age of 65, with a slight male predominance.

The majority of pancreatic cancers (about 95%) are classified as Pancreatic Ductal Adenocarcinoma (PDAC). This cancer arises from the ductal epithelial cells, where mutations in key genes such as *KRAS*, *TP53*, *CDKN2A* SMAD4 contribute to the tumorigenesis process. The *KRAS* gene, in particular, is mutated in nearly 90% of pancreatic adenocarcinomas, leading to unchecked cellular proliferation and resistance to apoptosis.

A biopsy is often necessary to confirm the diagnosis techniques such as endoscopic Ultrasound (EUS) or percutaneous needle biopsy may be employed.

Staging of pancreatic cancer is critical for determining the appropriate treatment strategy. The American Joint Committee on Cancer (AJCC) staging system is commonly used, which categorizes tumors based on size, lymph node involvement metastasis. Staging helps guide treatment options, ranging from surgical resection to chemotherapy and radiation therapy.

#### **Treatment** options

The treatment of pancreatic adenocarcinoma is challenging and depends on the stage at diagnosis:

**Surgery:** The Whipple procedure (pancreaticoduodenectomy) is the most common surgical approach for respectable tumors. However, only about 15%-20% of patients are candidates for surgery at the time of diagnosis.

**Chemotherapy:** Adjuvant chemotherapy with regimens like FOLFIRINOX (combination of fluorouracil, leucovorin, irinotecan oxaliplatin) or gemcitabine is standard for patients who undergo surgery. For advanced stages, chemotherapy remains the primary treatment option.

**Radiation therapy:** This may be used in combination with chemotherapy for locally advanced disease or as palliative treatment for symptom control.

The prognosis for pancreatic adenocarcinoma is generally poor, with a 5-year survival rate of around 10%. Factors influencing prognosis include tumor stage at diagnosis, histological grade response to treatment. Ongoing research focuses on improving early detection methods, understanding the molecular basis of the disease developing targeted therapies.

### CONCLUSION

Adenocarcinoma of the pancreas remains a impressive challenge in oncology due to its aggressive nature and late diagnosis. Increased awareness of risk factors, early symptoms advancements in diagnostic and therapeutic modalities are important for improving outcomes. As research continues, there is hope for more effective treatments and improved survival rates for patients diagnosed with this devastating disease.

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Received: 30-Aug-2024, Manuscript No. JCM-24-27167; Editor assigned: 02-Sep-2024, PreQC No. JCM-24-27167; Reviewed: 16-Sep-2024, QC No. JCM-24-27167; Revised: 23-Sep-2024, Manuscript No. JCM-24-27167; Published: 30-Sep-2024, DOI: 10.35248/2157-2518.24.15.459

Citation: Kabir A (2024). Pancreatic Adenocarcinoma: An Overview of Clinical Challenges and Advances in Treatment. J Carcinog Mutagen. 15:459.

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