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However, the NetAnalyzer's report performance might be a real problem under 10.4.2. There are a few problems like constant rewriting of 'Modified' field in the report. In order to overcome these issues, I. IP-verifier is a free service of the IPVanish VPN. A randomized phase II clinical trial to assess the efficacy of cisplatin plus gemcitabine in unresectable pancreatic carcinoma: a single-center study. The aim of this study was to evaluate the efficacy of cisplatin and gemcitabine in the treatment of patients with pancreatic cancer. We conducted a prospective randomized, single-center phase II clinical trial. Fifty-two eligible patients were randomly assigned into two groups: 36 patients were treated with gemcitabine (1000 mg/m²) for 7 days plus cisplatin (25 mg/m²) for day 8 every 3 weeks, and 16 patients were treated with gemcitabine (1000 mg/m²) for 7 days alone. The response rate in the two groups was 34% and 16.7% respectively. The median time to progression was 4 months in the gemcitabine plus cisplatin arm compared to 2 months in the gemcitabine alone arm (P = 0.09) and the median survival was 8 months versus 6 months respectively (P = 0.37). Toxicities were moderate. Gemcitabine alone is as effective and well tolerated as the combination in the treatment of patients with pancreatic cancer. Effects of reperfusion on energy metabolism in mitochondria of guinea pig heart. Metabolism in isolated mitochondria of guinea pig heart was studied in the closed system with 1 mM external substrates. Perfusion of mitochondria with oxygen-deficient Krebs-Henseleit solution resulted in the accumulation of ADP in the incubation medium. The respiration rate of the isolated mitochondria decreased after the addition of the Krebs-Henseleit solution. A rapid consumption of the accumulated ADP followed by the reoxidation of CoASH was noted. The consumption of ADP was accompanied by a small generation of lactate in the incubation medium. The generation of H₂O₂ was not observed. The accumulation of ADP was due to the ATPase activity of the oxygen-deficient mitochondria. The consumption of ADP was due to the transport of adenylates into the mitochondria

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