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Remnant: From the Ashes. Letos Lab Update v220617-CODEX. Remnant From The Ashes Letos Lab Update v220617-CODEX. □□□□□□□ by : whitestorm of MEGA Linux. Letos Lab PC-CODEX. Remnant.From.The.Ashes.Subject.2923.Update.v248587-CODEX. Released: 2019-11-06.Efficacy and safety of continuous-infusion paclitaxel in patients with advanced breast cancer: a phase I-II study. The limited activity of paclitaxel given as a bolus infusion has led to investigation of new methods of administration. We performed a phase I-II study to determine the maximum tolerated dose (MTD), the dose-limiting toxicities (DLTs), and the preliminary antitumor activity of continuous-infusion (CI) paclitaxel. Patients with advanced breast cancer, previously untreated with chemotherapy and who had not received radiotherapy, were enrolled in the phase I part of the study. To decrease toxicity, a 30% dose reduction was planned if one or more DLTs occurred. A maximum of six patients were enrolled in the phase II part of the study. Paclitaxel was infused at a constant rate of 5 mg/m(2)/h for 48 hours. MTD was defined as the dose at which > or =2 of the 6 planned cycles of treatment were administered. Twenty-eight patients were entered in the phase I part of the study. DLTs included grade 3 leukopenia (n = 4) and grade 4 neutropenia (n = 4), one of which developed subsequently into grade 5 sepsis. No grade 2 or grade 3 nonhematological toxicities were observed. One patient with a bilobular breast cancer underwent recurrence of disease three months after study entry and another patient developed a nonmelanoma skin cancer of the trunk. Nine patients were entered in the phase II part of the study. The median number of treatment cycles administered was 4 (range 2 to 6). There was one patient with a complete response and 14 patients with a partial response for an overall objective response rate of 60%. In patients previously treated with chemotherapy and/or radiotherapy, CI paclitaxel was well tolerated, with only grade 3-4 hematologic toxicities. The preliminary antitumor activity was encouraging. A phase III trial of

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