GEMCITABINE THYMOORGAN®

Gemcitabine (as HCl)

WARNINGS

Caution - Prolongation of the infusion time beyond 60 minutes and more frequent than weekly dosing have been shown to increase toxicity.
Hematology - GEMCITABINE THYMOORGAN can suppress bone marrow function as manifested by leukopenia, thrombocytopenia, and anemia (see ADVERSE REACTIONS), and myelosuppression is usually the dose-limiting toxicity. Patients should be monitored for myelosuppression during therapy. See DOSAGE AND ADMINISTRATION for recommended dose adjustments.
Pulmonary - Pulmonary toxicity has been reported with the use of GEMCITABINE THYMOORGAN. In cases of severe lung toxicity, GEMCITABINE THYMOORGAN therapy should be discontinued immediately and appropriate supportive care measures instituted (see Pulmonary under Single-Agent Use and under Post-marketing experience in ADVERSE REACTIONS).
Renal - Hemolytic Uremic Syndrome (HUS) and/or renal failure have been reported following one or more doses of GEMCITABINE THYMOORGAN. Renal failure leading to death or requiring dialysis, despite discontinuation of therapy, has been rarely reported. The majority of the cases of renal failure leading to death were due to HUS (see Renal under Single - Agent Use and under Post-marketing experience in ADVERSE REACTIONS).
Hepatic - Serious hepatotoxicity, including liver failure and death, has been reported very rarely in patients receiving GEMCITABINE THYMOORGAN alone or in combination with other potentially hepatotoxic drugs (see Hepatic under Single-Agent Use and under Post-marketing experience in ADVERSE REACTIONS).
Pregnancy - Pregnancy Category D. GEMCITABINE THYMOORGAN can cause fetal harm when administered to a pregnant woman. GEMCITABINE THYMOORGAN is embryotoxic causing fetal malformations (cleft palate, incomplete ossification) at doses of 1.5 mg/kg/day in mice (about 1/200 the recommended human dose on a mg/m2 basis). GEMCITABINE THYMOORGAN is fetotoxic causing fetal malformations (fused pulmonary artery, absence of gall bladder) at doses of 0.1 mg/kg/day in rabbits (about 1/600 the recommended human dose on a mg/m2 basis). Embryotoxicity was characterized by decreased fetal viability, reduced live litter sizes, and developmental delays. There are no studies of GEMCITABINE THYMOORGAN in pregnan women. If GEMCITABINE THYMOORGAN is used during pregnancy, or if the patient becomes pregnant while taking GEMCITABINE THYMOORGAN, the patient should be apprised of the potential hazard to the fetus.

ACTION

GEMCITABINE THYMOORGAN (gemcitabine HCl) is a nucleoside analogue that exhibits antitumor activity.
GEMCITABINE THYMOORGAN exhibits cell phase specificity, primarily killing cells undergoing DNA synthesis (S-phase) and also blocking the progression of cells through the G1/S-phase boundary. GEMCITABINE THYMOORGAN is metabolized intracellularly by nucleoside kinases to the active diphosphate (dFdCDP) and triphosphate (dFdCTP) nucleosides. The cytotoxic effect of GEMCITABINE THYMOORGAN is attributed to a combination of two actions of the diphosphate and the triphosphate nucleosides, which leads to inhibition of DNA synthesis. First, GEMCITABINE THYMOORGAN diphosphate inhibits ribonucleotide reductase, which is responsible for catalyzing the reactions that generate the deoxynucleoside triphosphates for DNA synthesis. Inhibition of this enzyme by the diphosphate nucleoside causes a reduction in the concentrations of deoxynucleotides, including dCTP. Second, GEMCITABINE THYMOORGAN triphosphate competes with dCTP for incorporation into DNA. The reduction in the intracellular concentration of dCTP (by the action of the diphosphate) enhances the incorporation of GEMCITABINE THYMOORGAN triphosphate into DNA (self-potentiation). After the GEMCITABINE THYMOORGAN nucleotide is incorporated into DNA, only one additional nucleotide is added to the growing DNA strands. After this addition, there is inhibition of further DNA synthesis. DNA polymerase epsilon is unable to remove the GEMCITABINE THYMOORGAN nucleotide and repair the growing DNA 2 strands (masked chain termination). In CEM T lymphoblastoid cells, GEMCITABINE THYMOORGAN induces internucleosomal DNA fragmentation, one of the characteristics of programmed cell death.
GEMCITABINE THYMOORGAN demonstrated dose-dependent synergistic activity with cisplatin in vitro. No effect of cisplatin on GEMCITABINE THYMOORGAN triphosphate accumulation or DNA double-strand breaks was observed. In vivo, GEMCITABINE THYMOORGAN showed activity in combination with cisplatin against the LX-1 and CALU-6 human lung xenografts, but minimal activity was seen with the NCI-H460 or NCI-H520 xenografts. GEMCITABINE THYMOORGAN was synergistic with cisplatin in the Lewis lung murine xenograft. Sequential exposure to GEMCITABINE THYMOORGAN 4 hours before cisplatin produced the greatest interaction.

INDICATIONS

Ovarian Cancer: Gemcitabine in combination with carboplatin is indicated for the treatment of patients with advanced ovarian cancer that has relapsed at least 6 months after completion of platinum-based therapy.
Breast Cancer: Gemcitabine in combination with paclitaxel is indicated for the first-line treatment of patients with metastatic breast cancer after failure of prior anthracycline-containing adjuvant chemotherapy, unless anthracyclines were clinically contraindicated.
Non-Small Cell Lung Cancer: Gemcitabine is indicated in combination with cisplatin for the first-line treatment of patients with inoperable, locally advanced (Stage IIIA or IIIB), or metastatic (Stage IV) non-small cell lung cancer.
Pancreatic Cancer: Gemcitabine is indicated as first-line treatment for patients with locally advanced (nonresectable Stage II or Stage III) or metastatic (Stage IV) adenocarcinoma of the pancreas. Gemcitabine is indicated for patients previously treated with 5-FU.

DOSAGE AND ADMINISTRATION

Gemcitabine is for intravenous use only.

Adults

Single-Agent Use:
Pancreatic Cancer: Gemcitabine should be administered by intravenous infusion at a dose of 1000 mg/m2 over 30 minutes once weekly for up to 7 weeks (or until toxicity necessitates reducing or holding a dose), followed by a week of rest from treatment. Subsequent cycles should consist of infusions once weekly for 3 consecutive weeks out of every 4 weeks.
Dose Modifications: Dosage adjustment is based upon the degree of hematologic toxicity experienced by the patient. Clearance in women and the elderly is reduced and women were somewhat less able to progress to subsequent cycles.
Patients receiving Gemcitabine should be monitored prior to each dose with a complete blood count (CBC), including differential and platelet count. If marrow suppression is detected, therapy should be modified or suspended according to the guidelines in Table 1.

Dosage Reduction Guidelines

Absolute granulocyte count (x 106/L)		Platelet count (x 106/L)	% of full dose
≥1000	and	≥100,000	100
500-999	or	50,000-99,999	75
<500	or	<50,000	Hold

Laboratory evaluation of renal and hepatic function, including transaminases and serum creatinine, should be performed prior to initiation of therapy and periodically thereafter. Gemcitabine should be administered with caution in patients with evidence of significant renal or hepatic impairment as there is insufficient information from clinical studies to allow clear dose recommendation for these patient populations.
Patients treated with Gemcitabine who complete an entire cycle of therapy may have the dose for subsequent cycles increased by 25%, provided that the absolute granulocyte count (AGC) and platelet nadirs exceed 1500 x 106/L and 100,000 x 106/L, respectively, and if non-hematologic toxicity has not been greater than WHO Grade 1. If patients tolerate the subsequent course of Gemcitabine at the increased dose, the dose for the next cycle can be further increased by 20%, provided again that the AGC and platelet nadirs exceed 1500 x 106/L and 100,000 x 106/L, respectively, and that non-hematologic toxicity has not been greater than WHO Grade 1.
Combination Use:
Non-Small Cell Lung Cancer: Two schedules have been investigated and the optimum schedule has not been determined. With the 4-week schedule, Gemcitabine should be administered intravenously at 1000 mg/m2 over 30 minutes on Days 1, 8, and 15 of each 28-day cycle. Cisplatin should be administered intravenously at 100 mg/m2 on Day 1 after the infusion of Gemcitabine. With the 3-week schedule, Gemcitabine should be administered intravenously at 1250 mg/m2 over 30 minutes on Days 1 and 8 of each 21-day cycle. Cisplatin at a dose of 100 mg/m2 should be administered intravenously after the infusion of Gemcitabine on Day 1. See prescribing information for cisplatin administration and hydration guidelines.
Dose Modifications: Dosage adjustments for hematologic toxicity may be required for Gemcitabine and for cisplatin. Gemcitabine dosage adjustment for hematological toxicity is based on the granulocyte and platelet counts taken on the day of therapy. Patients receiving Gemcitabine should be monitored prior to each dose with a complete blood count (CBC), including differential and platelet counts. If marrow suppression is detected, therapy should be modified or suspended according to the guidelines in Table 1. For cisplatin dosage adjustment. In general, for severe (Grade 3 or 4) non-hematological toxicity, except alopecia and nausea/vomiting, therapy with Gemcitabine plus cisplatin should be held or decreased by 50% depending on the judgment of the treating physician. During combination therapy with cisplatin, serum creatinine, serum potassium, serum calcium, and serum magnesium should be carefully monitored (Grade 3/4 serum creatinine toxicity for Gemcitabine plus cisplatin was 5% versus 2% for cisplatin alone).
Breast Cancer: Gemcitabine should be administered intravenously at a dose of 1250 mg/m2 over 30 minutes on Days 1 and 8 of each 21-day cycle. Paclitaxel should be administered at 175 mg/m2 on Day 1 as a 3-hour intravenous infusion before Gemcitabine administration. Patients should be monitored prior to each dose with a complete blood count, including differential counts. Patients should have an absolute granulocyte count ≥1500 x 106/L and a platelet count ≥100,000 x 106/L prior to each cycle.
Dose Modifications: Gemcitabine dosage adjustments for hematological toxicity is based on the granulocyte and platelet counts taken on Day 8 of therapy. If marrow suppression is detected, Gemcitabine dosage should be modified according to the guidelines in Table 2.

Day 8 Dosage Reduction Guidelines for Gemcitabine in Combination with Paclitaxel

Absolute granulocyte count (x 106/L)		Platelet count (x 106/L)	% of full dose
≥1200	and	>75,000	100
1000-1199	or	50,000-75,000	75
700-999	and	≥50,000	50
<700	or	<50,000	Hold

In general, for severe (Grade 3 or 4) non-hematological toxicity, except alopecia and nausea/vomiting, therapy with Gemcitabine should be held or decreased by 50% depending on the judgment of the treating physician. For paclitaxel dosage adjustment.
Ovarian Cancer — Gemcitabine should be administered intravenously at a dose of 1000 mg/m2 over 30 minutes on Days 1 and 8 of each 21-day cycle. Carboplatin AUC 4 should be administered intravenously on Day 1 after Gemcitabine administration. Patients should be monitored prior to each dose with a complete blood count, including differential counts. Patients should have an absolute granulocyte count ≥1500 x 106/L and a platelet count ≥100,000 x 106/L prior to each cycle.
Dose Modifications — Gemcitabine dosage adjustments for hematological toxicity within a cycle of treatment is based on the granulocyte and platelet counts taken on Day 8 of therapy. If marrow suppression is detected, Gemcitabine dosage should be modified according to guidelines in Table 3.

Day 8 Dosage Reduction Guidelines for Gemcitabine in Combination with Carboplatin

Absolute granulocyte count (x 106/L)		Platelet count (x 106/L)	% of full dose
≥1500	and	≥100,000	100
1000-1499	and/or	75,000-99,999	50
<1000	and/or	<75,000	Hold

In general, for severe (Grade 3 or 4) non-hematological toxicity, except nausea/vomiting, therapy with Gemcitabine should be held or decreased by 50% depending on the judgment of the treating physician. For carboplatin dosage adjustment.
Dose adjustment for Gemcitabine in combination with carboplatin for subsequent cycles is based upon observed toxicity. The dose of *Gemcitabine in subsequent cycles should be reduced to 800 mg/m2 on Days 1 and 8 in case of any of the following hematologic toxicities:*
- Absolute granulocyte count <500 x 106/L for more than 5 days.
- Absolute granulocyte count <100 x 106/L for more than 3 days.
- Febrile neutropenia.
- Platelets <25,000 x 106/L.
- Cycle delay of more than one week due to toxicity.

If any of the above toxicities recur after the initial dose reduction, for the subsequent cycle, Gemcitabine should be given on Day 1 only at 800 mg/m2.
Gemcitabine may be administered on an outpatient basis.
Instructions for Use/Handling: The recommended diluent for reconstitution of Gemcitabine is 0.9% Sodium Chloride Injection without preservatives. Due to solubility considerations, the maximum concentration for Gemcitabine upon reconstitution is 40 mg/mL. Reconstitution at concentrations greater than 40 mg/mL may result in incomplete dissolution, and should be avoided.
To reconstitute, add 5 mL of 0.9% Sodium Chloride Injection to the 200-mg vial or 25 mL of 0.9% Sodium Chloride Injection to the 1-g vial. Shake to dissolve. These dilutions each yield a gemcitabine concentration of 38 mg/mL which includes accounting for the displacement volume of the lyophilized powder (0.26 mL for the 200-mg vial or 1.3 mL for the 1-g vial). The total volume upon reconstitution will be 5.26 mL or 26.3 mL, respectively. Complete withdrawal of the vial contents will provide 200 mg or 1 g of gemcitabine, respectively. The appropriate amount of drug may be administered as prepared or further diluted with 0.9% Sodium Chloride Injection to concentrations as low as 0.1 mg/mL.
Reconstituted Gemcitabine is a clear, colorless to light straw-colored solution. After reconstitution with 0.9% Sodium Chloride Injection, the pH of the resulting solution lies in the range of 2.7 to 3.3. The solution should be inspected visually for particulate matter and discoloration, prior to administration, whenever solution or container permit. If particulate matter or discoloration is found, do not administer.
When prepared as directed, Gemcitabine solutions are stable for 24 hours at controlled room temperature 20° to 25°C (68° to 77°F). Discard unused portion. Solutions of reconstituted Gemcitabine should not be refrigerated, as crystallization may occur.
The compatibility of Gemcitabine with other drugs has not been studied. No incompatibilities have been observed with infusion bottles or polyvinyl chloride bags and administration sets.
Unopened vials of Gemcitabine are stable until the expiration date indicated on the package when stored at controlled room temperature 20° to 25°C (68° to 77°F).
Caution should be exercised in handling and preparing Gemcitabine solutions. The use of gloves is recommended. If Gemcitabine solution contacts the skin or mucosa, immediately wash the skin thoroughly with soap and water or rinse the mucosa with copious amounts of water. Although acute dermal irritation has not been observed in animal studies, 2 of 3 rabbits exhibited drug-related systemic toxicities (death, hypoactivity, nasal discharge, shallow breathing) due to dermal absorption.
Procedures for proper handling and disposal of anti-cancer drugs should be considered. Several guidelines on this subject have been published.1-5 There is no general agreement that all of the procedures recommended in the guidelines are necessary or appropriate.

CONTRAINDICATIONS

Gemcitabine is contraindicated in those patients with a known hypersensitivity to the drug.

WARNINGS AND PRECAUTIONS

WARNINGS

Caution: Prolongation of the infusion time beyond 60 minutes and more frequent than weekly dosing have been shown to increase toxicity.
Hematology: Gemcitabine can suppress bone marrow function as manifested by leukopenia, thrombocytopenia, and anemia, and myelosuppression is usually the dose-limiting toxicity. Patients should be monitored for myelosuppression during therapy.
Pulmonary: Pulmonary toxicity has been reported with the use of Gemcitabine. In cases of severe lung toxicity, Gemcitabine therapy should be discontinued immediately and appropriate supportive care measures instituted.
Renal: Hemolytic Uremic Syndrome (HUS) and/or renal failure have been reported following one or more doses of Gemcitabine. Renal failure leading to death or requiring dialysis, despite discontinuation of therapy, has been rarely reported. The majority of the cases of renal failure leading to death were due to HUS.
Hepatic: Serious hepatotoxicity, including liver failure and death, has been reported very rarely in patients receiving Gemcitabine alone or in combination with other potentially hepatotoxic drugs.
Pregnancy: Pregnancy Category D. Gemcitabine can cause fetal harm when administered to a pregnant woman. Gemcitabine is embryotoxic causing fetal malformations (cleft palate, incomplete ossification) at doses of 1.5 mg/kg/day in mice (about 1/200 the recommended human dose on a mg/m2 basis). Gemcitabine is fetotoxic causing fetal malformations (fused pulmonary artery, absence of gall bladder) at doses of 0.1 mg/kg/day in rabbits (about 1/600 the recommended human dose on a mg/m2 basis). Embryotoxicity was characterized by decreased fetal viability, reduced live litter sizes, and developmental delays. There are no studies of Gemcitabine in pregnant women. If Gemcitabine is used during pregnancy, or if the patient becomes pregnant while taking Gemcitabine, the patient should be apprised of the potential hazard to the fetus.

PRECAUTIONS

General: Patients receiving therapy with Gemcitabine should be monitored closely by a physician experienced in the use of cancer chemotherapeutic agents. Most adverse events are reversible and do not need to result in discontinuation, although doses may need to be withheld or reduced. There was a greater tendency in women, especially older women, not to proceed to the next cycle.
Laboratory Tests: Patients receiving Gemcitabine should be monitored prior to each dose with a complete blood count (CBC), including differential and platelet count.
Suspension or modification of therapy should be considered when marrow suppression is detected.
Laboratory evaluation of renal and hepatic function should be performed prior to initiation of therapy and periodically thereafter.
Carcinogenesis, Mutagenesis, Impairment of Fertility: Long-term animal studies to evaluate the carcinogenic potential of Gemcitabine have not been conducted. Gemcitabine induced forward mutations in vitro in a mouse lymphoma (L5178Y) assay and was clastogenic in an in vivo mouse micronucleus assay. Gemcitabine was negative when tested using the Ames, in vivo sister chromatid exchange, and in vitro chromosomal aberration assays, and did not cause unscheduled DNA synthesis in vitro. Gemcitabine IP doses of 0.5 mg/kg/day (about 1/700 the human dose on a mg/m2 basis) in male mice had an effect on fertility with moderate to severe hypospermatogenesis, decreased fertility, and decreased implantations. In female mice, fertility was not affected but maternal toxicities were observed at 1.5 mg/kg/day IV (about 1/200 the human dose on a mg/m2 basis) and fetotoxicity or embryolethality was observed at 0.25 mg/kg/day IV (about 1/1300 the human dose on a mg/m2 basis).
Pregnancy: Category D.
Nursing Mothers — It is not known whether Gemcitabine or its metabolites are excreted in human milk. Because many drugs are excreted in human milk and because of the potential for serious adverse reactions from Gemcitabine in nursing infants, the mother should be warned and a decision should be made whether to discontinue nursing or to discontinue the drug, taking into account the importance of the drug to the mother and the potential risk to the infant.
Elderly Patients: Gemcitabine clearance is affected by age. There is no evidence, however, that unusual dose adjustments (i.e., other than those already recommended in DOSAGE AND ADMINISTRATION) are necessary in patients over 65, and in general, adverse reaction rates in the single-agent safety database of 979 patients were similar in patients above and below 65. Grade 3/4 thrombocytopenia was more common in the elderly. In the randomized clinical trial of Gemcitabine in combination with carboplatin for recurrent ovarian cancer, 125 women treated with Gemcitabine plus carboplatin were <55 years and 50 were ≥65 years. Similar effectiveness was observed between older and younger women. There was significantly higher Grade 3/4 neutropenia in women 65 years of age or older. Overall, there were no substantial differences in toxicity profile of Gemcitabine plus carboplatin based on age.
Gender — Gemcitabine clearance is affected by gender. In the single-agent safety database (N=979 patients), however, there is no evidence that unusual dose adjustments (i.e., other than those already recommended in DOSAGE AND ADMINISTRATION) are necessary in women. In general, in single-agent studies of Gemcitabine, adverse reaction rates were similar in men and women, but women, especially older women, were more likely not to proceed to a subsequent cycle and to experience Grade 3/4 neutropenia and thrombocytopenia.
Pediatric Patients — The effectiveness of Gemcitabine in pediatric patients has not been demonstrated. Gemcitabine was evaluated in a Phase 1 trial in pediatric patients with refractory leukemia and determined that the maximum tolerated dose was 10 mg/m2/min for 360 minutes three times weekly followed by a one-week rest period. Gemcitabine was also evaluated in a Phase 2 trial in patients with relapsed acute lymphoblastic leukemia (22 patients) and acute myelogenous leukemia (10 patients) using 10 mg/m2/min for 360 minutes three times weekly followed by a one week rest period. Toxicities observed included bone marrow suppression, febrile neutropenia, elevation of serum transaminases, nausea, and rash/desquamation, which were similar to those reported in adults. No meaningful clinical activity was observed in this Phase 2 trial. Patients with Renal or Hepatic Impairment — Gemcitabine should be used with caution in patients with preexisting renal impairment or hepatic insufficiency as there is insufficient information from clinical studies to allow clear dose recommendation for these patient populations. Administration of Gemcitabine in patients with concurrent liver metastases or a preexisting medical history of hepatitis, alcoholism, or liver cirrhosis may lead to exacerbation of the underlying hepatic insufficiency.
Drug Interactions — No specific drug interaction studies have been conducted. For information on the pharmacokinetics of Gemcitabine and cisplatin in combination.
Radiation Therapy — A pattern of tissue injury typically associated with radiation toxicity has been reported in association with concurrent and non-concurrent use of Gemcitabine. Non-concurrent (given >7 days apart) — Analysis of the data does not indicate enhanced toxicity when Gemcitabine is administered more than 7 days before or after radiation, other than radiation recall. Data suggest that Gemcitabine can be started after the acute effects of radiation have resolved or at least one week after radiation. Concurrent (given together or ≤7 days apart) — Preclinical and clinical studies have shown that Gemcitabine has radiosensitizing activity. Toxicity associated with this multimodality therapy is dependent on many different factors, including dose of Gemcitabine, frequency of Gemcitabine administration, dose of radiation, radiotherapy planning technique, the target tissue, and target volume. In a single trial, where Gemcitabine at a dose of 1000 mg/m2 was administered concurrently for up to 6 consecutive weeks with therapeutic thoracic radiation to patients with non-small cell lung cancer, significant toxicity in the form of severe, and potentially life-threatening mucositis, especially esophagitis and pneumonitis was observed, particularly in patients receiving large volumes of radiotherapy [median treatment volumes 4795 cm3]. Subsequent studies have been reported and suggest that Gemcitabine administered at lower doses with concurrent radiotherapy has predictable and less severe toxicity. However, the optimum regimen for safe administration of Gemcitabine with therapeutic doses of radiation has not yet been determined in all tumor types.

DRUG INTERACTIONS

When Gemcitabine (1250 mg/m2 on Days 1 and 8) and cisplatin (75 mg/m2 on Day 1) were administered in NSCLC patients, the clearance of gemcitabine on Day 1 was 128 L/hr/m2 and on Day 8 was 107 L/hr/m2. The clearance of cisplatin in the same study was reported to be 3.94 mL/min/m2 with a corresponding half-life of 134 hours. Analysis of data from metastatic breast cancer patients shows that, on average, Gemcitabine has little or no effect on the pharmacokinetics (clearance and half-life) of paclitaxel and paclitaxel has little or no effect on the pharmacokinetics of Gemcitabine. Data from NSCLC patients demonstrate that Gemcitabine and carboplatin given in combination does not alter the pharmacokinetics of Gemcitabine or carboplatin compared to administration of either single-agent. However, due to wide confidence intervals and small sample size, interpatient variability may be observed.

SIDE EFFECTS

Gemcitabine has been used in a wide variety of malignancies, both as a single-agent and in combination with other cytotoxic drugs.
Single-Agent Use: Myelosuppression is the principal dose-limiting toxicity with Gemcitabine therapy. Dosage adjustments for hematologic toxicity are frequently needed and are described in DOSAGE AND ADMINISTRATION.
The data in Table 4 are based on 979 patients receiving Gemcitabine as a single-agent administered weekly as a 30-minute infusion for treatment of a wide variety of malignancies. The Gemcitabine starting doses ranged from 800 to 1250 mg/m2. Data are also shown for the subset of patients with pancreatic cancer treated in 5 clinical studies. The frequency of all grades and severe (WHO Grade 3 or 4) adverse events were generally similar in the single-agent safety database of 979 patients and the subset of patients with pancreatic cancer. Adverse reactions reported in the single-agent safety database resulted in discontinuation of Gemcitabine therapy in about 10% of patients. In the comparative trial in pancreatic cancer, the discontinuation rate for adverse reactions was 14.3% for the Gemcitabine arm and 4.8% for the 5-FU arm.
All WHO-graded laboratory events are listed in Table 4, regardless of causality. Non-laboratory adverse events listed in Table 4 or discussed below were those reported, regardless of causality, for at least 10% of all patients, except the categories of Extravasation, Allergic, and Cardiovascular and certain specific events under the Renal, Pulmonary, and Infection categories. Table 5 presents the data from the comparative trial of Gemcitabine and 5-FU in pancreatic cancer for the same adverse events as those in Table 4, regardless of incidence.

Selected WHO-Graded Adverse Events in Patients Receiving Single-Agent Gemcitabine WHO Grades (% incidence)*

	All Patients[a]			Pancreatic Cancer Patients[b]			Discontinuations (%)[c]
	All Gardes	Garde 3	Garde 3	All Gardes	Garde 3	Garde 3	All Patients
Laboratory[d]							
Hematologic							
Anemia	68	7	1	73	8	2	<1
Leukopenia	62	9	<1	64	8	1	<1
Neutropenia	63	19	6	61	17	7	
Thrombocytopenia	24	4	1	36	7	<1	<1
Hepatic							
ALT	68	8	2	72	10	1	<1
AST	67	6	2	78	12	5	
Alkaline Phosphatase	55	7	2	77	16	4	
Bilirubin	13	2	<1	26	6	2	
Renal							
Proteinuria	45	<1	0	32	<1	0	
Hematuria	35	<1	0	23	0	0	<1
BUN	16	0	0	15	0	0	
Creatinine	8	<1	0	6	0	0	
None-laboratory[d]							
Nausea and Vomiting	69	13	1	71	10	2	<1
Pain	48	9	<1	42	6	<1	<1
Fever	41	2	0	38	2	0	<1
Rash	30	<1	0	28	<1	0	<1
Dyspnea	23	3	<1	10	0	<1	<1
Constipation	23	1	<1	31	3	<1	0
Diarrhea	19	1	0	30	3	0	0
Hemorrhag	17	<1	<1	4	2	<1	<1
Infection	16	1	<1	10	2	<1	<1
Alopecia	15	<1	0	16	0	0	0
Stomatitis	11	<1	0	10	<1	0	<1
Somnolence	11	<1	<1	11	2	<1	<1
Paresthesias	10	<1	0	10	<1	0	0

* Grade based on criteria from the World Health Organization (WHO).
[a] N= 699-974; all patients with laboratory or non-laboratory data.
[b] N= 161-241; all pancreatic cancer patients with laboratory or non-laboratory data.
[c] N= 979.
[d] Regardless of causality.
[e] Laboratory data with incidence for all patients ≥10%. For approximately 60% of the patients, non-laboratory events were graded only if assessed to be possibly drug-related.

Selected WHO-Graded Adverse Events From Comparative Trial of Gemcitabine and 5-FU in Pancreatic Cancer WHO Grades (% incidence)*

	Gemcitabine[a]			5-FU[b]		
	All Grades	Grade 3	Grade 4	All Grades	Grade 3	Grade 4
Laboratory[c]						
Hematologic						
Anemia	65	7	3	45	0	0
Leukopenia	71	10	0	15	2	0
Neutropenia	62	19	7	18	2	3
Thrombocytopenia	47	10	0	15	2	0
Hepatic						
ALT	72	8	2	38	0	0
AST	72	10	2	52	2	0
Alkaline Phosphatase	71	16	0	64	10	3
Bilirubin	16	2	2	25	6	3
Renal						
Proteinuria	10	0	0	2	0	0
Hematuria	13	0	0	0	0	0
BUN	8	0	0	10	0	0
Creatinine	2	0	0	0	0	0
Non-laboratory[c]						
Nausea and Vomiting	64	10	3	58	5	0
Pain	10	2	0	7	0	0
Fever	30	0	0	16	0	0
Rash	24	0	0	13	0	0
Dyspnea	6	0	0	3	0	0
Constipation	10	3	0	11	2	0
Diarrhea	24	2	0	31	5	0
Hemorrhage	0	0	0	2	0	0
Infection	8	0	0	3	2	0
Alopecia	18	0	0	16	0	0
Stomatitis	14	0	0	15	0	0
Somnolence	5	2	0	7	2	0
Paresthesias	2	0	0	2	0	0

[a] Grade based on criteria from the World Health Organization (WHO).
[b] N=58-63; all Gemcitabine patients with laboratory or non-laboratory data.
[c] N=61-63; all 5-FU patients with laboratory or non-laboratory data.
[d] Regardless of causality.
[e] Non-laboratory events were graded only if assessed to be possibly drug-related.

Hematologic: In studies in pancreatic cancer myelosuppression is the dose-limiting toxicity with Gemcitabine, but <1% of patients discontinued therapy for either anemia, leukopenia, or thrombocytopenia. Red blood cell transfusions were required by 19% of patients. The incidence of sepsis was less than 1%. Petechiae or mild blood loss (hemorrhage), from any cause, was reported in 16% of patients; less than 1% of patients required platelet transfusions. Patients should be monitored for myelosuppression during Gemcitabine therapy and dosage modified or suspended according to the degree of hematologic toxicity.
Gastrointestinal — Nausea and vomiting were commonly reported (69%) but were usually of mild to moderate severity. Severe nausea and vomiting (WHO Grade 3/4) occurred in <15% of patients. Diarrhea was reported by 19% of patients, and stomatitis by 11% of patients.
Hepatic — In clinical trials, Gemcitabine was associated with transient elevations of one or both serum transaminases in approximately 70% of patients, but there was no evidence of increasing hepatic toxicity with either longer duration of exposure to Gemcitabine or with greater total cumulative dose. Serious hepatotoxicity, including liver failure and death, has been reported very rarely in patients receiving Gemcitabine alone or in combination with other potentially hepatotoxic drugs.
Renal — In clinical trials, mild proteinuria and hematuria were commonly reported. Clinical findings consistent with the Hemolytic Uremic Syndrome (HUS) were reported in 6 of 2429 patients (0.25%) receiving Gemcitabine in clinical trials. Four patients developed HUS on Gemcitabine therapy, 2 immediately posttherapy. The diagnosis of HUS should be considered if the patient develops anemia with evidence of microangiopathic hemolysis, elevation of bilirubin or LDH, reticulocytosis, severe thrombocytopenia, and/or evidence of renal failure (elevation of serum creatinine or BUN). Gemcitabine therapy should be discontinued immediately. Renal failure may not be reversible even with discontinuation of therapy and dialysis may be required.
Fever — The overall incidence of fever was 41%. This is in contrast to the incidence of infection (16%) and indicates that Gemcitabine may cause fever in the absence of clinical infection. Fever was frequently associated with other flu-like symptoms and was usually mild and clinically manageable.
Rash — Rash was reported in 30% of patients. The rash was typically a macular or finely granular maculopapular pruritic eruption of mild to moderate severity involving the trunk and extremities. Pruritus was reported by 13% of patients.
Pulmonary — In clinical trials, dyspnea, unrelated to underlying disease, has been reported in association with Gemcitabine therapy. Dyspnea was occasionally accompanied by bronchospasm. Pulmonary toxicity has been reported with the use of Gemcitabine. The etiology of these effects is unknown. If such effects develop, Gemcitabine should be discontinued. Early use of supportive care measures may help ameliorate these conditions.
Edema — Edema (13%), peripheral edema (20%), and generalized edema (<1%) were reported. Less than 1% of patients discontinued due to edema.
Flu-like Symptoms — "Flu syndrome" was reported for 19% of patients. Individual symptoms of fever, asthenia, anorexia, headache, cough, chills, and myalgia were commonly reported. Fever and asthenia were also reported frequently as isolated symptoms. Insomnia, rhinitis, sweating, and malaise were reported infrequently. Less than 1% of patients discontinued due to flu-like symptoms.
Infection — Infections were reported for 16% of patients. Sepsis was rarely reported (<1%).
Alopecia — Hair loss, usually minimal, was reported by 15% of patients.
Neurotoxicity — There was a 10% incidence of mild paresthesias and a <1% rate of severe paresthesias.
Extravasation — Injection-site related events were reported for 4% of patients. There were no reports of injection site necrosis. Gemcitabine is not a vesicant.
Allergic — Bronchospasm was reported for less than 2% of patients. Anaphylactoid reaction has been reported rarely. Gemcitabine should not be administered to patients with a known hypersensitivity to this drug.
Cardiovascular — During clinical trials, 2% of patients discontinued therapy with Gemcitabine due to cardiovascular events such as myocardial infarction, cerebrovascular accident, arrhythmia, and hypertension. Many of these patients had a prior history of cardiovascular disease.
Combination Use in Non-Small Cell Lung Cancer: In the Gemcitabine plus cisplatin versus cisplatin study, dose adjustments occurred with 35% of Gemcitabine injections and 17% of cisplatin injections on the combination arm, versus 6% on the cisplatin-only arm. Dose adjustments were required in greater than 90% of patients on the combination, versus 16% on cisplatin. Study discontinuations for possibly drug-related adverse events occurred in 15% of patients on the combination arm and 8% of patients on the cisplatin arm. With a median of 4 cycles of Gemcitabine plus cisplatin treatment, 94 of 262 patients (36%) experienced a total of 149 hospitalizations due to possibly treatment-related adverse events. With a median of 2 cycles of cisplatin treatment, 61 of 260 patients (23%) experienced 78 hospitalizations due to possibly treatment-related adverse events.
In the Gemcitabine plus cisplatin versus etoposide plus cisplatin study, dose adjustments occurred with 20% of Gemcitabine injections and 16% of cisplatin injections in the Gemcitabine plus cisplatin arm compared with 20% of etoposide injections and 15% of cisplatin injections in the etoposide plus cisplatin arm. With a median of 5 cycles of Gemcitabine plus cisplatin treatment, 15 of 69 patients (22%) experienced 15 hospitalizations due to possibly treatment-related adverse events. With a median of 4 cycles of etoposide plus cisplatin treatment, 18 of 66 patients (27%) experienced 22 hospitalizations due to possibly treatment-related adverse events. In patients who completed more than one cycle, dose adjustments were reported in 81% of the Gemcitabine plus cisplatin patients, compared with 68% on the etoposide plus cisplatin arm. Study discontinuations for possibly drug-related adverse events occurred in 14% of patients on the Gemcitabine plus cisplatin arm and in 8% of patients on the etoposide plus cisplatin arm. The incidence of myelosuppression was increased in frequency with Gemcitabine plus cisplatin treatment (~90%) compared to that with the Gemcitabine monotherapy (~60%). With combination therapy Gemcitabine dosage adjustments for hematologic toxicity were required more often while cisplatin dose adjustments were less frequently required.
Table 6 presents the safety data from the Gemcitabine plus cisplatin versus cisplatin study in non-small cell lung cancer. The NCI Common Toxicity Criteria (CTC) were used. The two-drug combination was more myelosuppressive with 4 (1.5%) possibly treatment-related deaths, including 3 resulting from myelosuppression with infection and one case of renal failure associated with pancytopenia and infection. No deaths due to treatment were reported on the cisplatin arm. Nine cases of febrile neutropenia were reported on the combination therapy arm compared to 2 on the cisplatin arm. More patients required RBC and platelet transfusions on the Gemcitabine plus cisplatin arm. Myelosuppression occurred more frequently on the combination arm, and in 4 possibly treatment-related deaths myelosuppression was observed. Sepsis was reported in 4% of patients on the Gemcitabine plus cisplatin arm compared to 1% on the cisplatin arm. Platelet transfusions were required in 21% of patients on the combination arm and <1% of patients on the cisplatin arm. Hemorrhagic events occurred in 14% of patients on the combination arm and 4% on the cisplatin arm. However, severe hemorrhagic events were rare. Red blood cell transfusions were required in 39% of the patients on the Gemcitabine plus cisplatin arm, versus 13% on the cisplatin arm. The data suggest cumulative anemia with continued Gemcitabine plus cisplatin use. Nausea and vomiting despite the use of antiemetics occurred slightly more often with Gemcitabine plus cisplatin therapy (78%) than with cisplatin alone (71%). In studies with single-agent Gemcitabine, a lower incidence of nausea and vomiting (58% to 69%) was reported. Renal function abnormalities, hypomagnesemia, neuromotor, neurocortical, and neurocerebellar toxicity occurred more often with Gemcitabine plus cisplatin than with cisplatin monotherapy. Neurohearing toxicity was similar on both arms. Cardiac dysrhythmias of Grade 3 or greater were reported in 7 (3%) patients treated with Gemcitabine plus cisplatin compared to one (<1%) Grade 3 dysrhythmia reported with cisplatin therapy. Hypomagnesemia and hypokalemia were associated with one Grade 4 arrhythmia on the Gemcitabine plus cisplatin combination arm. Table 7 presents data from the randomized study of Gemcitabine plus cisplatin versus etoposide plus cisplatin in 135 patients with NSCLC for the same WHO-graded adverse events as those in Table 5. One death (1.5%) was reported on the Gemcitabine plus cisplatin arm due to febrile neutropenia associated with renal failure which was possibly treatment-related. No deaths related to treatment occurred on the etoposide plus cisplatin arm. The overall incidence of Grade 4 neutropenia on the Gemcitabine plus cisplatin arm was less than on the etoposide plus cisplatin arm (28% versus 56%). Sepsis was experienced by 2% of patients on both treatment arms. Grade 3 anemia and Grade 3/4 thrombocytopenia were more common on the Gemcitabine plus cisplatin arm. RBC transfusions were given to 29% of the patients who received Gemcitabine plus cisplatin versus 21% of patients who received etoposide plus cisplatin. Platelet transfusions were given to 3% of the patients who received Gemcitabine plus cisplatin versus 8% of patients who received etoposide plus cisplatin. Grade 3/4 nausea and vomiting were also more common on the Gemcitabine plus cisplatin arm. On the Gemcitabine plus cisplatin arm 7% of participants were hospitalized due to febrile neutropenia compared to 12% on the etoposide plus cisplatin arm. More than twice as many patients had dose reductions or omissions of a scheduled dose of Gemcitabine as compared to etoposide, which may explain the differences in the incidence of neutropenia and febrile neutropenia between treatment arms. Flu syndrome was reported by 3% of patients on the Gemcitabine plus cisplatin arm with none reported on the comparator arm. Eight patients (12%) on the Gemcitabine plus cisplatin arm reported edema compared to one patient (2%) on the etoposide plus cisplatin arm.

Selected CTC-Graded Adverse Events From Comparative Trial of Gemcitabine Plus Cisplatin Versus Single-Agent Cisplatin in NSCLC CTC Grades (% incidence)*

	Gemcitabine plus Cisplatin[a]			Cisplatin[a]		
	All Grades	Grade 3	Grade 4	All Grades	Grade 3	Grade 4
Laboratory[c]						
Hematologic						
Anemia	89	22	3	67	6	1
RBC Transfusion[e]	39			13		
Leukopenia	82	35	11	25	2	1
Neutropenia	79	22	35	20	3	1
Thrombocytopenia	85	25	25	13	3	1
Platelet Transfusions[e]	21			<1		
Lymphocytes	75	25	18	51	12	5
Hepatic						
Transaminase	22	2	1	10	1	0
Alkaline Phosphatase	19	1	0	13	0	0
Renal						
Proteinuria	23	0	0	18	0	0
Hematuria	15	0	0	13	0	0
Creatinine	38	4	<1	31	2	<1
Other Laboratory						
Hyperglycemia	30	4	0	23	3	0
Hypomagnesemia	30	4	3	17	2	0
Hypocalcemia	18	2	0	7	0	<1
Non-laboratory[d]						
Nausea	93	25	2	87	20	<1
Vomiting	78	11	12	71	10	9
Alopecia	53	1	0	33	0	0
Neuro Motor	35	12	0	15	3	0
Constipation	28	3	0	21	0	0
Neuro Hearing	25	6	0	21	6	0
Diarrhea	24	2	2	13	0	0
Neuro Sensory	23	1	0	18	1	0
Infection	18	3	2	12	1	0
Fever	16	0	0	5	0	0
Neuro Cortical	16	3	1	9	1	0
Neuro Mood	16	1	0	10	1	0
Local	15	0	0	6	0	0
Neuro Headache	14	0	0	7	0	0
Stomatitis	14	1	0	5	0	0
Hemorrhage	14	1	0	4	0	0
Dyspnea	12	4	3	11	3	2
Hypotension	12	1	0	7	1	0
Rash	11	0	0	3	0	0

* Grade based on Common Toxicity Criteria (CTC). Table includes data for adverse events with incidence ≥10% in either arm.
[a] N=217-253; all Gemcitabine plus cisplatin patients with laboratory or non-laboratory data. Gemcitabine at 1000 mg/m2 on Days 1, 8, and 15 and cisplatin at 100 mg/m2 on Day 1 every 28 days.
[b] N=213-248; all cisplatin patients with laboratory or non-laboratory data. Cisplatin at 100 mg/m2 on Day 1 every 28 days.
[c] Regardless of causality.
[d] Percent of patients receiving transfusions. Percent transfusions are not CTC-graded events.
[e] Non-laboratory events were graded only if assessed to be possibly drug-related.

Selected WHO-Graded Adverse Events From Comparative Trial of Gemcitabine Plus Cisplatin Versus Etoposide Plus Cisplatin in NSCLC WHO Grades (% incidence)*

	Gemcitabine plus Cisplatin[a]			Etoposide plus Cisplatin[b]		
	All Grades	Grade 3	Grade 4	All Grades	Grade 3	Grade 4
Laboratory[c]						
Hematologic						
Anemia	88			77		
RBC Transfusions[e]	29	22	0	21	13	2
Leukopenia	86			87		
Neutropenia	88	26	3	87	36	7
Thrombocytopenia	81	36	28	45	20	56
Platelet Transfusions[e]	3	39	16	8	8	5
Hepatic						
ALT	6	0	0	12	0	0
AST	3	0	0	11	0	0
Alkaline Phosphatase	16	0	0	11	0	0
Bilirubin	0	0	0	0	0	0
Renal						
Proteinuria	12	0	0	5	0	0
Hematuria	22	0	0	10	0	0
BUN	6	0	0	4	0	0
Creatinine	2	0	0	2	0	0
Non-laboratory[d]						
Nausea and Vomiting	96	35	4	86	19	7
Fever	6	0	0	3	0	0
Rash	10	0	0	3	0	0
Dyspnea	1	0	1	3	0	0
Constipation	17	0	0	15	0	0
Diarrhea	14	1	1	13	0	2
Hemorrhage	9	0	3	3	0	3
Infection	28	3	1	21	8	0
Alopecia	77	13	0	92	51	0
Stomatitis	20	4	0	18	2	0
Somnolence	3	0	0	3	2	0
Paresthesias	38	0	0	16	2	0

* Grade based on criteria from the World Health Organization (WHO).
[a] N=67-69; all Gemcitabine plus cisplatin patients with laboratory or non-laboratory data. Gemcitabine at 1250 mg/m2 on Days 1 and 8 and cisplatin at 100 mg/m2 on Day 1 every 21 days.
[b] N=57-63; all cisplatin plus etoposide patients with laboratory or non-laboratory data. Cisplatin at 100 mg/m2 on Day 1 and IV etoposide at 100 mg/m2 on Days 1, 2, and 3 every 21 days.
[c] Regardless of causality.
[d] Percent of patients receiving transfusions. Percent transfusions are not WHO-graded events.
[e] Non-laboratory events were graded only if assessed to be possibly drug-related.
[f] Pain data were not collected.

Combination Use in Breast Cancer: In the Gemcitabine plus paclitaxel versus paclitaxel study, dose reductions occurred with 8% of Gemcitabine injections and 5% of paclitaxel injections on the combination arm, versus 2% on the paclitaxel arm. On the combination arm, 7% of Gemcitabine doses were omitted and <1% of paclitaxel doses were omitted, compared to <1% of paclitaxel doses on the paclitaxel arm. A total of 18 patients (7%) on the Gemcitabine plus paclitaxel arm and 12 (5%) on the paclitaxel arm discontinued the study because of adverse events. There were two deaths on study or within 30 days after study drug discontinuation that were possibly drug-related, one on each arm. Table 8 presents the safety data occurrences of ≥10% (all grades) from the Gemcitabine plus paclitaxel versus paclitaxel study in breast cancer.

Adverse Events From Comparative Trial of Gemcitabine Plus Paclitaxel Versus Single-Agent Paclitaxel in Breast Cancera CTC Grades (% incidence)

	Gemcitabine plus Paclitaxel (N=262)			Paclitaxel (N=259)		
	All Grades	Grade 3	Grade 4	All Grades	Grade 3	Grade 4
Laboratory[c]						
Hematologic						
Anemia	69	6	1	51	3	<1
Neutropenia	69	31	17	31	4	7
Thrombocytopenia	26	5	<1	7	<1	<1
Leukopenia	21	10	1	12	2	0
Hepatobiliary						
ALT	18	5	<1	6	<1	0
AST	16	2	0	5	<1	0
Non-laboratory[d]						
Alopecia	90			92		
Neuropathy-sensory	64	14	4	58	19	3
Nausea	50	5	<1	31	3	0
Fatigue	40	1	0	28	2	0
Myalgia	33	6	<1	33	1	<1
Vomiting	29	4	0	15	3	<1
Arthralgia	24	2	0	22	2	0
Diarrhea	20	3	0	13	2	<1
Anorexia	17	3	0	12	2	0
Neuropathy-motor	15	0	0	12	<1	0
Stomatitis/pharyngitis	13	2	<1	10	<1	0
Fever	13	1	<1	8	<1	0
Constipation	11	<1	0	3	0	0
Bone pain	11	<1	0	12	0	0
Pain-other	11	2	0	10	<1	0
Rash/desquamation	11	<1	0	8	<1	0
		<1	<1	5	0	0

* Grade based on Common Toxicity Criteria (CTC) Version 2.0 (all grades ≥10%).
[a] Regardless of causality.
[b] Non-laboratory events were graded only if assessed to be possibly drug-related.
The following are the clinically relevant adverse events that occurred in ≥1% and <10% (all grades) of patients on either arm. In parentheses are the incidences of Grade 3 and 4 adverse events (Gemcitabine plus paclitaxel versus paclitaxel): febrile neutropenia (5.0% versus 1.2%), infection (0.8% versus 0.8%), dyspnea (1.9% versus 0), and allergic reaction/hypersensitivity (0 versus 0.8%). No differences in the incidence of laboratory and non-laboratory events were observed in patients 65 years or older, as compared to patients younger than 65.

Combination Use in Ovarian Cancer: In the Gemcitabine plus carboplatin versus carboplatin study, dose reductions occurred with 10.4% of Gemcitabine injections and 1.8% of carboplatin injections on the combination arm, versus 3.8% on the carboplatin alone arm. On the combination arm, 13.7% of Gemcitabine doses were omitted and 0.2% of carboplatin doses were omitted, compared to 0% of carboplatin doses on the carboplatin alone arm. There were no differences in discontinuations due to adverse events between arms (10.9% versus 9.8%, respectively).
Table 9 presents the adverse events (all grades) occurring in ≥10% of patients in the ovarian cancer study.

Adverse Events From Comparative Trial of Gemcitabine Plus Carboplatin Versus Single-Agent Carboplatin in Ovarian Cancera CTC Grades (% incidence)

	Gemcitabine plus Carboplatin (N=175)			Carboplatin (N=174)		
	All Grades	Grade 3	Grade 4	All Grades	Grade 3	Grade 4
Laboratory[c]						
Hematologic						
Neutropenia	90	42	29	58	11	1
Anemia	86	22	6	75	9	2
Leukopenia	86	48	5	70	6	<1
Thrombocytopenia	78	30	5	57	10	1
RBC Transfusions[e]	38			15		
Platelet Transfusions[e]	9			3		
Non-laboratory[d]						
Nausea	69	6	0	61	3	0
Alopecia	49	0	0	17	0	0
Vomiting	46	6	0	36	2	<1
Constipation	42	6	1	37	3	0
Fatigue	40	3	<1	32	5	0
Neuropathy-sensory	29	1	0	27	2	0
Diarrhea	25	3	0	14	<1	0
Stomatitis/pharyngitis	22	<1	0	13	0	0
Anorexia	16	1	0	13	0	0

* Grade based on Common Toxicity Criteria (CTC) Version 2.0 (all grades ≥10%).
[a] Regardless of causality.
[b] Percent of patients receiving transfusions. Transfusions are not CTC-graded events. Blood transfusions included both packed red blood cells and whole blood.
In addition to blood product transfusions as listed in Table 9, myelosuppression was also managed with hematopoetic agents. These agents were administered more frequently with combination therapy than with monotherapy (granulocyte growth factors: 23.6% and 10.1%, respectively; erythropoietic agents: 7.3% and 3.9%, respectively).
The following are the clinically relevant adverse events, regardless of causality, that occurred in >1% and <10% (all grades) of patients on either arm. In parentheses are the incidences of Grade 3 and 4 adverse events (Gemcitabine plus carboplatin versus carboplatin): AST or ALT elevation (0 versus 1.2%), dyspnea (3.4% versus 2.9%), febrile neutropenia (1.1% versus 0), hemorrhagic event (2.3% versus 1.1%), hypersensitivity reaction (2.3% versus 2.9%), motor neuropathy (1.1% versus 0.6%), and rash/desquamation (0.6% versus 0). No differences in the incidence of laboratory and non-laboratory events were observed in patients 65 years or older, as compared to patients younger than 65.

Post-marketing experience: The following adverse events have been identified during post-approval use of Gemcitabine. These events have occurred after Gemcitabine single-agent use and Gemcitabine in combination with other cytotoxic agents. Decisions to include these events are based on the seriousness of the event, frequency of reporting, or potential causal connection to Gemcitabine.
Cardiovascular — Congestive heart failure and myocardial infarction have been reported very rarely with the use of Gemcitabine. Arrhythmias, predominantly supraventricular in nature, have been reported very rarely.
Vascular Disorders — Clinical signs of peripheral vasculitis and gangrene have been reported very rarely.
Skin — Cellulitis and non-serious injection site reactions in the absence of extravasation have been rarely reported. Severe skin reactions, including desquamation and bullous skin eruptions, have been reported very rarely.
Hepatic — Increased liver function tests including elevations in aspartate aminotransferase (AST), alanine aminotransferase (ALT), gamma-glutamyl transferase (GGT), alkaline phosphatase, and bilirubin levels have been reported rarely. Serious hepatotoxicity including liver failure and death has been reported very rarely in patients receiving Gemcitabine alone or in combination with other potentially hepatotoxic drugs.
Pulmonary — Parenchymal toxicity, including interstitial pneumonitis, pulmonary fibrosis, pulmonary edema, and adult respiratory distress syndrome (ARDS), has been reported rarely following one or more doses of Gemcitabine administered to patients with various malignancies. Some patients experienced the onset of pulmonary symptoms up to 2 weeks after the last Gemcitabine dose. Respiratory failure and death occurred very rarely in some patients despite discontinuation of therapy.
Renal — Hemolytic-Uremic Syndrome (HUS) and/or renal failure have been reported following one or more doses of Gemcitabine. Renal failure leading to death or requiring dialysis, despite discontinuation of therapy, has been rarely reported. The majority of the cases of renal failure leading to death were due to HUS.
Injury, Poisoning, and Procedural Complications — Radiation recall reactions have been reported.

OVERDOSAGE

There is no known antidote for overdoses of Gemcitabine. Myelosuppression, paresthesias, and severe rash were the principal toxicities seen when a single dose as high as 5700 mg/m2 was administered by IV infusion over 30 minutes every 2 weeks to several patients in a Phase 1 study. In the event of suspected overdose, the patient should be monitored with appropriate blood counts and should receive supportive therapy, as necessary.

STORAGE

Store below 25°C. Do not refrigirate.

PRESENTATION

Vials
GEMCITABINE THYMOORGAN 200 mg: Gemcitabine (as HCL) 200 mg
GEMCITABINE THYMOORGAN 1000 mg: Gemcitabine (as HCL) 1000 mg
Excipients: Mannitol, sodium acetate, hydrochloric acid and/or sodium hydroxide.

THIS IS A MEDICAMENT

- A medicament is a product which affects your health, and its consumption contrary to instructions is dangerous.
- Follow the doctor's prescription strictly, the method of use and the instructions of the pharmacist who sold the medicament.
- The doctor and the pharmacist are experts in medicine, its benefits and risks.
- Do not by yourself interrupt the period of treatment prescribed for you.
- Do not repeat the same prescription without consulting your doctor.

THYMOORGAN Pharmazie GmbH

Manufactured by:
Thymoorgan Pharmazie GmbH, Germany
For:
Hikma Pharmaceuticals, Amman - Jordan

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