Drug Interaction Results

Selected Drugs: Acetaminophen, Acetylcysteine, Albuterol, Amikacin, Carbamazepine, Cefazolin, Cefepime, Clindamycin HCI, Codeine, Dexamethasone, Dipyrone, Enoxaparin Sodium, Fentanyl, Fludrocortisone Acetate, Fluticasone, Furosemide, Gabapentin, Hydrochlorothiazide, Hydrocortisone, Ipratropium Bromide, Irbesartan, Ketamine HCI, Labetalol, Lactulose, Levetiracetam, Mannitol, Meropenem, Methyldopa, Metoclopramide, Midazolam, NIFEdipine, Neostigmine, Nimodipine, Norepinephrine, Omeprazole, Phenytoin, Propofol, Valproic Acid, Vancomycin HCI, Vasopressin, Vecuronium Bromide, traMADol HCI

Severity: All

Documentation: All

Interaction Type: Drug-Drug

Drugs:	Severity:	Documentation:	Summary:
CARBAMAZEPINE NIFEDIPINE	Contraindicated	<u>Excellent</u>	Concurrent use of NIFEDIPINE and STRONG CYP3A4 INDUCERS may result in reduced NIFEdipine exposure and reduced efficacy of NIFEdipine.
NIFEDIPINE PHENYTOIN	Contraindicated	<u>Excellent</u>	Concurrent use of NIFEDIPINE and STRONG CYP3A4 INDUCERS may result in reduced NIFEdipine exposure and reduced efficacy of NIFEdipine.
ACETAMINOPHEN CARBAMAZEPINE	S <u>Major</u>	Good	Concurrent use of ACETAMINOPHEN and CARBAMAZEPINE may result in an increased risk of acetaminophen hepatotoxicity and reduced acetaminophen exposure.
ALBUTEROL FUROSEMIDE	S <u>Major</u>	Fair	Concurrent use of ALBUTEROL and POTASSIUM-DEPLETING DIURETICS may result in an increased risk of ECG changes or hypokalemia.
ALBUTEROL HYDROCHLOROTHIAZIDE	S <u>Major</u>	Fair	Concurrent use of ALBUTEROL and POTASSIUM-DEPLETING DIURETICS may result in an increased risk of ECG changes or hypokalemia.
ALBUTEROL LABETALOL HYDROCHLORIDE	S <u>Major</u>	Fair	Concurrent use of ALBUTEROL and BETA-ADRENERGIC BLOCKERS may result in reduced efficacy of albuterol.
AMIKACIN SULFATE FUROSEMIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of AMIKACIN and FUROSEMIDE may result in increased amikacin plasma and tissue concentrations and additive ototoxicity and/ or nephrotoxicity.
AMIKACIN SULFATE MANNITOL	S <u>Major</u>	Fair	Concurrent use of AMIKACIN and IV MANNITOL may result in alteration of serum and tissue amikacin concentrations.
AMIKACIN SULFATE VANCOMYCIN HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of AMIKACIN and VANCOMYCIN may result in additive ototoxicity and/or nephrotoxicity.

AMIKACIN SULFATE VECURONIUM BROMIDE	S <u>Major</u>	Good	Concurrent use of AMINOGLYCOSIDES and NONDEPOLARIZING NEUROMUSCULAR BLOCKERS may result in enhanced and/ or prolonged neuromuscular blockade which may lead to respiratory depression and paralysis.
CARBAMAZEPINE CLINDAMYCIN HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and CYP3A4 SUBSTRATES may result in reduced CYP3A4 substrate exposure and reduced efficacy of CYP3A4 substrate.
CARBAMAZEPINE CODEINE SULFATE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and CODEINE may result in reduced opioid efficacy, precipitation of opioid withdrawal, an increased risk of serotonin syndrome, an increased risk of respiratory depression, profound sedation, hypotension and syncope and reduced codeine exposure.
CARBAMAZEPINE DEXAMETHASONE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and DEXAMETHASONE may result in reduced dexAMETHasone exposure and reduced carBAMazepine exposure.
CARBAMAZEPINE FENTANYL	S <u>Major</u>	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and FENTANYL may result in an increased risk of respiratory depression, profound sedation, hypotension and syncope, reduced exposure of fentaNYL and an increased risk of serotonin syndrome.
CARBAMAZEPINE FLUDROCORTISONE ACETATE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and CORTICOSTEROIDS may result in reduced corticosteroids exposure and reduced efficacy of corticosteroids.
CARBAMAZEPINE FLUTICASONE PROPIONATE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and CORTICOSTEROIDS METABOLIZED BY CYP3A may result in reduced corticosteroids exposure and reduced efficacy of corticosteroids.
CARBAMAZEPINE GABAPENTIN	S <u>Major</u>	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and CNS DEPRESSANTS may result in an increased risk of CNS depression.
CARBAMAZEPINE HYDROCORTISONE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and HYDROCORTISONE may result in reduced hydrocortisone exposure and reduced efficacy of hydrocortisone.
CARBAMAZEPINE IRBESARTAN	S <u>Major</u>	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and CYP2C9 SUBSTRATES may result in reduced CYP2C9 substrate exposure and reduced efficacy of CYP2C9 substrate.

CARBAMAZEPINE KETAMINE HYDROCHLORIDE	S Major	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and
	<u></u>		CNS DEPRESSANTS THAT ARE CYP2B6 SUBSTRATES may result in reduced CYP2B6 substrate exposure and reduced efficacy of CYP2B6 substrate and an increased risk of respiratory depression, profound sedation, hypotension and syncope.
CARBAMAZEPINE LABETALOL HYDROCHLORIDE	S Major	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and CYP2C19 SUBSTRATES may result in reduced CYP2C19 substrate exposure and reduced efficacy of CYP2C19 substrate.
CARBAMAZEPINE MIDAZOLAM	S <u>Major</u>	Good	Concurrent use of CARBAMAZEPINE and MIDAZOLAM may result in reduced midazolam exposure and reduced efficacy of midazolam, an increased risk of respiratory depression, profound sedation, hypotension and syncope.
CARBAMAZEPINE NIMODIPINE	S <u>Major</u>	Good	Concurrent use of CARBAMAZEPINE and NIMODIPINE may result in reduced nimodipine plasma concentrations and reduced nimodipine efficacy.
CARBAMAZEPINE OMEPRAZOLE	S <u>Major</u>	Good	Concurrent use of CARBAMAZEPINE and OMEPRAZOLE may result in increased carBAMazepine exposure, an increased risk of carBAMazepine toxicity, reduced omeprazole exposure and reduced efficacy of omeprazole.
CARBAMAZEPINE PHENYTOIN	S <u>Major</u>	Excellent	Concurrent use of CARBAMAZEPINE and FOSPHENYTOIN OR PHENYTOIN may result in reduced phenytoin or fosphenytoin (prodrug of phenytoin) exposure and reduced carBAMazepine exposure.
CARBAMAZEPINE PROPOFOL	S <u>Major</u>	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and CNS DEPRESSANTS may result in an increased risk of CNS depression.
CARBAMAZEPINE VALPROIC ACID	S <u>Major</u>	Good	Concurrent use of CARBAMAZEPINE and VALPROIC ACID may result in reduced carBAMazepine exposure, increased carBAMazepine-10,11-epoxide exposure and reduced valproate exposure.
CARBAMAZEPINE VECURONIUM BROMIDE	S Major	Excellent	Concurrent use of CARBAMAZEPINE and VECURONIUM may result in reduced efficacy of vecuronium.
CARBAMAZEPINE TRAMADOL HYDROCHLORIDE	S Major	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and TRAMADOL may result in increased risk of serotonin syndrome and reduced traMADol exposure and an increased risk of respiratory depression, profound sedation, hypotension and syncope.

CEFEPIME HYDROCHLORIDE FUROSEMIDE	S <u>Major</u>	Good	Concurrent use of CEFEPIME and DIURETICS may result in an increased risk of nephrotoxicity.
CEFEPIME HYDROCHLORIDE HYDROCHLOROTHIAZIDE	S <u>Major</u>	Good	Concurrent use of CEFEPIME and DIURETICS may result in an increased risk of nephrotoxicity.
CODEINE SULFATE DEXAMETHASONE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CODEINE and CYP3A4 INDUCERS may result in reduced codeine and morphine exposure, reduced opioid efficacy and precipitation of opioid withdrawal.
CODEINE SULFATE DIPYRONE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CODEINE and CYP3A4 INDUCERS may result in reduced codeine and morphine exposure, reduced opioid efficacy and precipitation of opioid withdrawal.
CODEINE SULFATE FENTANYL	S <u>Major</u>	<u>Fair</u>	Concurrent use of CODEINE and SEROTONERGIC CNS DEPRESSANTS may result in increased risk of respiratory and CNS depression and an increased risk of serotonin syndrome.
CODEINE SULFATE FUROSEMIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CODEINE and DIURETICS may result in reduced diuretic efficacy.
CODEINE SULFATE GABAPENTIN	S <u>Major</u>	Good	Concurrent use of GABAPENTIN and CNS DEPRESSANTS may result in an increased risk of respiratory depression.
CODEINE SULFATE HYDROCHLOROTHIAZIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CODEINE and DIURETICS may result in reduced diuretic efficacy.
CODEINE SULFATE IPRATROPIUM BROMIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CODEINE and ANTICHOLINERGICS may result in an increased risk of paralytic ileus.
CODEINE SULFATE KETAMINE HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CODEINE and CNS DEPRESSANTS may result in an increased risk of respiratory and CNS depression.
CODEINE SULFATE METOCLOPRAMIDE HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of METOCLOPRAMIDE and CNS DEPRESSANTS may result in increased risk of CNS depression.
CODEINE SULFATE MIDAZOLAM	S <u>Major</u>	<u>Fair</u>	Concurrent use of CODEINE and BENZODIAZEPINES may result in an increased risk of hypotension, respiratory depression, profound sedation, coma, and death.
CODEINE SULFATE PHENYTOIN	S <u>Major</u>	<u>Fair</u>	Concurrent use of CODEINE and CYP3A4 INDUCERS may result in reduced codeine and morphine exposure, reduced opioid efficacy and precipitation of opioid withdrawal.
CODEINE SULFATE PROPOFOL		<u>Fair</u>	Concurrent use of CODEINE and CNS

	S <u>Major</u>		DEPRESSANTS may result in an increased risk of respiratory and CNS depression.
CODEINE SULFATE TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of CODEINE and SEROTONERGIC CNS DEPRESSANTS may result in increased risk of respiratory and CNS depression and an increased risk of serotonin syndrome.
DEXAMETHASONE DIPYRONE	S <u>Major</u>	<u>Fair</u>	Concurrent use of NSAIDS and ORAL CORTICOSTEROIDS may result in increased risk of gastrointestinal ulcer or bleeding.
DEXAMETHASONE FENTANYL	S <u>Major</u>	Good	Concurrent use of FENTANYL and CYP3A4 INDUCERS may result in reduced exposure of fentaNYL, reduced efficacy of fentaNYL and/ or onset of opioid withdrawal symptoms.
DEXAMETHASONE NIFEDIPINE	S <u>Major</u>	Good	Concurrent use of NIFEDIPINE and CYP3A4 INDUCERS may result in reduced NIFEdipine exposure.
DEXAMETHASONE NIMODIPINE	S <u>Major</u>	<u>Fair</u>	Concurrent use of DEXAMETHASONE and NIMODIPINE may result in reduced nimodipine plasma concentrations and reduced nimodipine efficacy.
DEXAMETHASONE PHENYTOIN	S <u>Major</u>	<u>Fair</u>	Concurrent use of DEXAMETHASONE and STRONG CYP3A4 INDUCERS may result in reduced dexAMETHasone exposure.
DEXAMETHASONE TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of TRAMADOL and CYP3A4 INDUCERS may result in reduced traMADol exposure and reduced efficacy of traMADol.
DIPYRONE ENOXAPARIN SODIUM	S <u>Major</u>	Good	Concurrent use of LOW- MOLECULAR-WEIGHT HEPARINS and NSAIDS may result in an increased risk of bleeding.
DIPYRONE FENTANYL	S <u>Major</u>	Good	Concurrent use of FENTANYL and CYP3A4 INDUCERS may result in reduced exposure of fentaNYL, reduced efficacy of fentaNYL and/ or onset of opioid withdrawal symptoms.
DIPYRONE FUROSEMIDE	S <u>Major</u>	Good	Concurrent use of FUROSEMIDE and NSAIDS may result in reduced diuretic effectiveness and possible nephrotoxicity.
DIPYRONE HYDROCHLOROTHIAZIDE	S <u>Major</u>	Good	Concurrent use of NSAIDS and THIAZIDE DIURETICS may result in reduced diuretic effectiveness and possible nephrotoxicity.
DIPYRONE HYDROCORTISONE	S <u>Major</u>	<u>Fair</u>	Concurrent use of NSAIDS and ORAL CORTICOSTEROIDS may result in increased risk of gastrointestinal ulcer or bleeding.

DIPYRONE NIFEDIPINE	S <u>Major</u>	Good	Concurrent use of NIFEDIPINE and CYP3A4 INDUCERS may result in reduced NIFEdipine exposure.
DIPYRONE TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of TRAMADOL and CYP3A4 INDUCERS may result in reduced traMADol exposure and reduced efficacy of traMADol.
FENTANYL FUROSEMIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of FENTANYL and DIURETICS may result in reduced diuretic efficacy.
FENTANYL GABAPENTIN	S <u>Major</u>	Good	Concurrent use of GABAPENTIN and CNS DEPRESSANTS may result in an increased risk of respiratory depression.
FENTANYL HYDROCHLOROTHIAZIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of FENTANYL and DIURETICS may result in reduced diuretic efficacy.
FENTANYL IPRATROPIUM BROMIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of FENTANYL and ANTICHOLINERGICS may result in increased risk of urinary retention and/ or server constipation, which may lead to paralytic ileus.
FENTANYL KETAMINE HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of FENTANYL and CNS DEPRESSANTS may result in increased risk of profound sedation, respiratory depression, hypotension, coma and death.
FENTANYL METOCLOPRAMIDE HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of METOCLOPRAMIDE and CNS DEPRESSANTS may result in increased risk of CNS depression.
FENTANYL MIDAZOLAM	S <u>Major</u>	<u>Fair</u>	Concurrent use of FENTANYL and CNS DEPRESSANTS may result in increased risk of profound sedation, respiratory depression, hypotension, coma and death.
FENTANYL NIFEDIPINE	S <u>Major</u>	<u>Fair</u>	Concurrent use of FENTANYL and NIFEDIPINE may result in an increased risk of severe hypotension and an increased fluid volume requirements.
FENTANYL PHENYTOIN	S <u>Major</u>	Good	Concurrent use of FENTANYL and CYP3A4 INDUCERS may result in reduced exposure of fentaNYL, reduced efficacy of fentaNYL and/ or onset of opioid withdrawal symptoms.
FENTANYL PROPOFOL	S <u>Major</u>	<u>Fair</u>	Concurrent use of FENTANYL and CNS DEPRESSANTS may result in increased risk of profound sedation, respiratory depression, hypotension, coma and death.
FENTANYL TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of FENTANYL and SEROTONERGIC CNS DEPRESSANTS may result in increased risk of respiratory and CNS depression and an increased risk of serotonin syndrome.

FLUDROCORTISONE ACETATE PHENYTOIN	S <u>Major</u>	Good	Concurrent use of FOSPHENYTOIN OR PHENYTOIN and CORTICOSTEROIDS may result in reduced corticosteroid exposure.
FLUTICASONE PROPIONATE PHENYTOIN	S Major	Good	Concurrent use of FOSPHENYTOIN OR PHENYTOIN and CORTICOSTEROIDS may result in reduced corticosteroid exposure.
FUROSEMIDE IRBESARTAN	S <u>Major</u>	<u>Fair</u>	Concurrent use of FUROSEMIDE and ANGIOTENSIN RECEPTOR BLOCKERS may result in severe hypotension and deterioration in renal function, including renal failure.
FUROSEMIDE NOREPINEPHRINE BITARTRATE	S <u>Major</u>	<u>Fair</u>	Concurrent use of FUROSEMIDE and NOREPINEPHRINE may result in decreased arterial responsiveness (vasoconstricting effect) to norepinephrine.
FUROSEMIDE PHENYTOIN	S <u>Major</u>	<u>Fair</u>	Concurrent use of FOSPHENYTOIN OR PHENYTOIN and FUROSEMIDE may result in reduced furosemide exposure.
FUROSEMIDE TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of TRAMADOL and DIURETICS may result in reduced efficacy of diuretics.
GABAPENTIN KETAMINE HYDROCHLORIDE	S <u>Major</u>	Good	Concurrent use of GABAPENTIN and CNS DEPRESSANTS may result in an increased risk of respiratory depression.
GABAPENTIN MIDAZOLAM	S <u>Major</u>	Good	Concurrent use of GABAPENTIN and CNS DEPRESSANTS may result in an increased risk of respiratory depression.
GABAPENTIN PROPOFOL	S <u>Major</u>	Good	Concurrent use of GABAPENTIN and CNS DEPRESSANTS may result in an increased risk of respiratory depression.
GABAPENTIN TRAMADOL HYDROCHLORIDE	S <u>Major</u>	Good	Concurrent use of GABAPENTIN and CNS DEPRESSANTS may result in an increased risk of respiratory depression.
HYDROCHLOROTHIAZIDE TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of TRAMADOL and DIURETICS may result in reduced efficacy of diuretics.
HYDROCORTISONE PHENYTOIN	S <u>Major</u>	Good	Concurrent use of FOSPHENYTOIN OR PHENYTOIN and CORTICOSTEROIDS may result in reduced corticosteroid exposure.
IPRATROPIUM BROMIDE TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of TRAMADOL and ANTICHOLINERGICS may result in increased risk of urinary retention and/ or severe constipation leading to paralytic ileus.
KETAMINE HYDROCHLORIDE METOCLOPRAMIDE HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of METOCLOPRAMIDE and CNS DEPRESSANTS may

result in increased risk of CNS depression.

			CNS depression.
KETAMINE HYDROCHLORIDE MIDAZOLAM	S <u>Major</u>	<u>Fair</u>	Concurrent use of KETAMINE and CNS DEPRESSANTS may result in increased risk of respiratory and CNS depression.
KETAMINE HYDROCHLORIDE PROPOFOL	S <u>Major</u>	<u>Fair</u>	Concurrent use of KETAMINE and CNS DEPRESSANTS may result in increased risk of respiratory and CNS depression.
KETAMINE HYDROCHLORIDE TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of TRAMADOL and CNS DEPRESSANTS may result in increased risk of respiratory and CNS depression.
MEROPENEM VALPROIC ACID	S <u>Major</u>	Excellent	Concurrent use of MEROPENEM and VALPROIC ACID may result in reduced valproic acid exposure and an increased risk of seizures and status epilepticus.
METOCLOPRAMIDE HYDROCHLORIDE MIDAZOLAM	S <u>Major</u>	<u>Fair</u>	Concurrent use of METOCLOPRAMIDE and CNS DEPRESSANTS may result in increased risk of CNS depression.
METOCLOPRAMIDE HYDROCHLORIDE PROPOFOL	S <u>Major</u>	<u>Fair</u>	Concurrent use of METOCLOPRAMIDE and CNS DEPRESSANTS may result in increased risk of CNS depression.
METOCLOPRAMIDE HYDROCHLORIDE TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of METOCLOPRAMIDE and CNS DEPRESSANTS may result in increased risk of CNS depression.
MIDAZOLAM PROPOFOL	S <u>Major</u>	<u>Fair</u>	Concurrent use of FOSPROPOFOL and BENZODIAZEPINES may result in additive cardiorespiratory effects.
MIDAZOLAM TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of TRAMADOL and CNS DEPRESSANTS may result in increased risk of respiratory and CNS depression.
NIFEDIPINE PROPOFOL	S <u>Major</u>	Good	Concurrent use of NIFEDIPINE and CYP3A4 INHIBITORS may result in increased NIFEdipine exposure.
NIFEDIPINE VALPROIC ACID	S <u>Major</u>	<u>Fair</u>	Concurrent use of NIFEDIPINE and VALPROIC ACID may result in increased NIFEdipine exposure.
NIMODIPINE PHENYTOIN	S <u>Major</u>	Good	Concurrent use of FOSPHENYTOIN OR PHENYTOIN and NIMODIPINE may result in reduced niMODipine exposure and reduced efficacy niMODipine efficacy.
OMEPRAZOLE PHENYTOIN	S <u>Major</u>	Fair	Concurrent use of FOSPHENYTOIN OR PHENYTOIN and OMEPRAZOLE may result in increased phenytoin exposure or fosphenytoin (prodrug of phenytoin), an increased risk of phenytoin or fosphenytoin toxicity (ataxia, hyperreflexia,

nystagmus, tremor) and reduced omeprazole exposure.

			exposure.
PHENYTOIN VALPROIC ACID	S <u>Major</u>	Good	Concurrent use of FOSPHENYTOIN OR PHENYTOIN and VALPROIC ACID may result in increased valproate exposure, altered phenytoin or fosphenytoin (prodrug of phenytoin) exposure and an increased risk of valproate-associated hyperammonemia.
PHENYTOIN VECURONIUM BROMIDE	S <u>Major</u>	Good	Concurrent use of FOSPHENYTOIN OR PHENYTOIN and VECURONIUM may result in reduced vecuronium exposure, reduced efficacy of vecuronium and an increased risk of resistance to the neuromuscular blocking action of the nondepolarizing neuromuscular blocking agents.
PHENYTOIN TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of TRAMADOL and CYP3A4 INDUCERS may result in reduced traMADol exposure and reduced efficacy of traMADol.
PROPOFOL VALPROIC ACID	S <u>Major</u>	<u>Fair</u>	Concurrent use of PROPOFOL and VALPROIC ACID may result in increased propofol exposure and risk of sedation or cardiorespiratory depression.
PROPOFOL TRAMADOL HYDROCHLORIDE	S <u>Major</u>	<u>Fair</u>	Concurrent use of TRAMADOL and CNS DEPRESSANTS may result in increased risk of respiratory and CNS depression.
ACETAMINOPHEN PHENYTOIN	Moderate	<u>Good</u>	Concurrent use of ACETAMINOPHEN and PHENYTOIN may result in decreased acetaminophen effectiveness and an increased risk of hepatotoxicity.
ACETYLCYSTEINE CARBAMAZEPINE	Moderate	Good	Concurrent use of ACETYLCYSTEINE and CARBAMAZEPINE may result in subtherapeutic carbamazepine levels.
CARBAMAZEPINE DIPYRONE	<u>Moderate</u>	<u>Fair</u>	Concurrent use of CARBAMAZEPINE and CYP3A INDUCERS may result in reduced carBAMazepine exposure and reduced efficacy of carBAMazepine.
CARBAMAZEPINE FUROSEMIDE	<u>Moderate</u>	Good	Concurrent use of CARBAMAZEPINE and SELECTED DIURETICS may result in hyponatremia.
CARBAMAZEPINE HYDROCHLOROTHIAZIDE	Moderate	Good	Concurrent use of CARBAMAZEPINE and SELECTED DIURETICS may result in hyponatremia.
CARBAMAZEPINE LEVETIRACETAM	<u>Moderate</u>	<u>Good</u>	Concurrent use of CARBAMAZEPINE and LEVETIRACETAM may result in symptoms of carbamazepine toxicity (nystagmus, ataxia, dizziness, double vision).

CLINDAMYCIN HYDROCHLORIDE VECURONIUM BROMIDE	Moderate	<u>Fair</u>	Concurrent use of CLINDAMYCIN and VECURONIUM may result in vecuronium toxicity (respiratory depression).
DEXAMETHASONE HYDROCORTISONE	Moderate	<u>Fair</u>	Concurrent use of HYDROCORTISONE and CYP3A4 INDUCERS may result in reduced hydrocortisone exposure.
DEXAMETHASONE VECURONIUM BROMIDE	Moderate	Good	Concurrent use of VECURONIUM and DEXAMETHASONE may result in decreased vecuronium effectiveness; prolonged muscle weakness and myopathy.
DIPYRONE IRBESARTAN	Moderate	<u>Excellent</u>	Concurrent use of ACE INHIBITORS AND ANGIOTENSIN RECEPTOR BLOCKERS and NSAIDS may result in reduced antihypertensive effect and renal dysfunction and/ or increased blood pressure.
DIPYRONE LABETALOL HYDROCHLORIDE	Moderate	Good	Concurrent use of BETA- ADRENERGIC BLOCKERS and NSAIDS may result in reduced antihypertensive effect.
FLUDROCORTISONE ACETATE FUROSEMIDE	Moderate	Good	Concurrent use of FLUDROCORTISONE and FUROSEMIDE may result in hypokalemia.
FLUDROCORTISONE ACETATE HYDROCHLOROTHIAZIDE	Moderate	<u>Fair</u>	Concurrent use of HYDROCHLOROTHIAZIDE and FLUDROCORTISONE may result in hypokalemia and subsequent cardiac arrhythmias.
FUROSEMIDE HYDROCORTISONE	Moderate	<u>Fair</u>	Concurrent use of FUROSEMIDE and HYDROCORTISONE may result in hypokalemia.
FUROSEMIDE VECURONIUM BROMIDE	Moderate	Good	Concurrent use of FUROSEMIDE and VECURONIUM may result in increased or decreased neuromuscular blockade.
HYDROCHLOROTHIAZIDE HYDROCORTISONE	Moderate	<u>Fair</u>	Concurrent use of HYDROCHLOROTHIAZIDE and HYDROCORTISONE may result in hypokalemia and subsequent cardiac arrhythmias.
HYDROCORTISONE NEOSTIGMINE METHYLSULFATE	Moderate	<u>Fair</u>	Concurrent use of HYDROCORTISONE and NEOSTIGMINE may result in decreased neostigmine effectiveness.
HYDROCORTISONE PROPOFOL	Moderate	<u>Fair</u>	Concurrent use of HYDROCORTISONE and CYP3A4 INHIBITORS may result in increased hydrocortisone exposure and an increased risk of hydrocortisone-related adverse reactions.
HYDROCORTISONE VECURONIUM BROMIDE	Moderate	Fair	Concurrent use of VECURONIUM and HYDROCORTISONE may result in decreased vecuronium effectiveness; prolonged muscle weakness and myopathy.
LABETALOL HYDROCHLORIDE METHYLDOPA	Moderate	<u>Fair</u>	Concurrent use of LABETALOL and METHYLDOPA may result in exaggerated hypertensive response, tachycardia, or

exposure to exogenous catecholamines. LABETALOL HYDROCHLORIDE --**Fair** Concurrent use of NIFEDIPINE and BETA **NIFEDIPINE** BLOCKERS may result in an increased risk of **Moderate** congestive heart failure, severe hypotension or exacerbation of angina. MIDAZOLAM -- OMEPRAZOLE **Fair** Concurrent use of MIDAZOLAM and OMEPRAZOLE may result **Moderate** in benzodiazepine toxicity (CNS depression, ataxia, lethargy). MIDAZOLAM -- PHENYTOIN Good Concurrent use of MIDAZOI AM and PHENYTOIN may result in Moderate decreased efficacy of midazolam. NIMODIPINE -- VALPROIC ACID Good Concurrent use of NIMODIPINE and VALPROIC ACID may result **Moderate** in nimodipine toxicity (dizziness, headache,

arrhythmias during physiologic stress or

flushing, peripheral edema).

Definitions

Severity:

The drugs are contraindicated for concurrent use.

Contraindicated

The interaction may be life-threatening and/or require medical intervention to minimize or prevent serious adverse effects.

Moderate
The interaction may result in exacerbation of the patient's condition and/or require an alteration in therapy.

Minor

The interaction would have limited clinical effects. Manifestations may include an increase in the frequency or severity of the side effects but generally would not require a Major alteration in therapy.

Unavailable Severity ratings are not assigned for the content.

Documentation:

Excellent Controlled studies have clearly established the existence of the

interaction.

Good Documentation strongly suggests the interaction exists,

but well-controlled studies are lacking.

Fair Available documentation is poor, but pharmacologic considerations lead

clinicians to suspect the interaction exists; or, documentation is good for a

pharmacologically similar drug.

Unavailable Severity ratings are not assigned for the content.