Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	difuent	concentration	duration/rate	RT	REF	requirement)	
Abatacept (Orencia) (non-chemo) REF	Mix 250 mg vial with 10 mL SWFI (only use silicone-free disposable syringes for preparation)	25 mg/mL	NS 100 mL (withdraw a volume of NS equal to volume of abatacept required)	Maximum 10 mg/mL	30 minutes	24 hr.	24 hr.	IV: Administer through 0.2-1.2- micron, low protein-binding filter (e.g. secondary tubing with inline filter) Do not shake SQ: Allow to warm to RT for 30-60 min	Lexicomp
Ado-trastuzumab emtansine (Kadcyla) Sound-alike, look- alike warning: do not confuse with trastuzumab or trastuzumab or trastuzumab S Q REF	Mix 100 mg vial with 5 mL SWFI Or Mix 160 mg vial with 8 mL SWFI	20 mg/mL	NS 250 mL	N/A	First cycle: 90 minutes Subsequent cycles: 30 minutes (if tolerates prior infusions)	Immediate Use	24 hr.	Do not administer IV push or bolus Administer through a 0.2 or 0.22- micron inline non-protein adsorptive polyethersulfone filter in PVC bag (non-PVC bag ok) Do not shake Do not use D5W Do not administer with other medications	Lexicomp
Aldesleukin (Proleukin) REF	Mix 22 Million Unit vial (1.3 mg) with 1.2mL SWFI (PF) (do not use BSWFI or NS for reconstitution or dilution)	18 million units/mL (1.1 mg/mL)	D5W 50 mL	30-70 mcg/mL (If aldesleukin final conc. < 30 mcg/mL, add human albumin to achieve final albumin conc. 0.1% (to ↓ adsorption)	15 minutes (flush line before and after with D5W)	48 hr.	48 hr.	PVC bag recommended Do not filter Do not shake Protect lyophilized power from light during storage Allow to reach RT	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	oility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unucht	concentration	uurauomrate	RT	REF	requirement)	
Alemtuzumab	RTU					8 hr.	8 hr.	Do not administer IV push or bolus	Lexicomp
(Lemtrada) Lemtrada is on REMS Campath is available only through Sanofi	Lemtrada 12 mg /1.2ml vial	Lemtrada 10 mg/mL	NS 100 mL	N/A	Lemtrada: 4 hr.		(use within 8 hrs. of dilution)	Do not shake Ok to infuse in PVC or use low-protein-binding filter	
Campath Distribution Program REF	Campath 30 mg/1 mL vial	Campath 30 mg/mL	NS 100 mL	N/A	Campath: 2 hr.			To order Campath (off the market): 1-877-422-6728 (Campath is not on REMS)	
Amifostine	Mix 500 mg vial with	500 mg/10mL	Straight drug, IV push		Give 15-30 minutes prior to	5 hr.	24 hr.	Ok to use PVC bag for IV bolus	Lexicomp
(Ethyol)	9.7 mL NS		Or in NS if		Radiation:3			-	
(non-chemo) RT	SQ: 500 mg vial with 2.5 mL NS or SWFI		diluted to 5-40 mg/mL		minutes 30 min prior to cisplatin: 15 minutes				
(Rybevant®)	RTU 350 mg / 7 mL vial	Withdraw and then discard the volume of solution from the 250 mL infusion bag equal to the volume of RYBREVANT to be added. Withdraw 7 mL of RYBREVANT from each vial and add it to the infusion bag. The final volume in the infusion bag should be 250 mL.	5% dextrose solution or 0.9% sodium chloride			Diluted solutions should be administered within 10 hours [including infusion time (2-5 hours)] at room temperature 59°F to 77°F	Store intact vials at 2°C to 8°C (36°F to 46°F). Store in original carton to protect from light	Only use infusion bags made of polyvinylchloride (PVC), polypropylene (PP), polyethylene (PE), or polyolefin blend (PP+PE)	Package insert
Arsenic trioxide (Trisenox) RT	RTU New formulation: 12 mg liquid vial (2 mg/ml)	2 mg/mL (prior formulation: 1 mg/ml)	NS 100–250 mL	N/A	1-2 hours (may extend up to 4 hrs. if acute vasomotor reactions are observed)	24 hr.	48 hr.	Irritant Verify concentration prior to admixture	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uurauom/rate	RT	REF	requirement)	
Atezolizumab	RTU	60 mg/mL	NS 250 mL	N/A	First cycle:	6 hr.	24 hr.	Do not administer as an IV push or bolus	Lexicomp
(Tecentriq)	1200 mg/20 mL vial				60 minutes			Use polyvinyl chloride (PVC) bag	
REF	840 mg/14 mL vial				Subsequent cycles: 30 minutes (if tolerated)			May be infused with or without a 0.2- to 0.22-micron sterile, non-pyrogenic, low-protein binding in-line filter	
					(ii tolciated)			Dilute only with NS	
								Do not shake	
								Do not infuse with other medications	
Azacitidine (Vidaza)	Mix 100 mg vial with SWFI:	IV: 10 mg/mL	NS 50-100 mL	N/A	IV: 10 minutes SQ: volume of	IV: 1 hr. (short stability)		Do not filter after reconstitution	Lexicomp
RT	IV: 10 mL SQ: 4mL	SQ: 25 mg/mL	(Limited stability and must be prepared immediately prior to each dose)		injection may vary based on dose		SQ: 8 hr.	SQ: Dose greater than 4 mL may be equally divided into 2 syringes	
BCG (TICE BCG) Intravesical (non-chemo but requires biohazard precaution) REF	Intravesical: Mix 1 vial (50 mg) with 1 mL NS (PF), then dilute with 49 mL NS (PF) to put total volume = 50 mL Percutaneous or intracavitary route: 50 mg in 50 mL NS, then the tube is clamped x 1 hour. Pressure is set up in a way to maintain low intra-renal pressure to ensure infusion	50 mg/mL	NS 50 mL (PF)	N/A	N/A	2 hr. (use within 2 hrs. of reconstitution)		Intravesical administration only Percutaneous or intracavitary has been used Dispense in a catheter tip syringe Biohazard labeling required Do not filter Do not shake Protect from light in storage and reconstitution only	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	• •	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	durunomrute	RT	REF	requirement)	
Belantamab mafodotin (Blenrep) REF	SWFI 100 mg with 2 mL	50 mg/mL	NS 250 mL	0.2- 2 mg/mL	30 min	4 hr.	24 hr (once removed from fridge, administer within 6 hrs (including infusion time)	Ok to use PVC or polyolefin bag May use polyethersulfone-based filter (0.2 micron) Mix by gentle inversion; do not shake. Solutions diluted for infusion should be clear and colorless; discard if particulate matter is observed	Lexicomp
Belimumab (Benlysta) (Non-chemo)	SWFI 120 mg vial with 1.5 mL 400 mg vial with 4.8 mL (Allow to stand 10 to 15 minutes to reach room temperature)	80 mg/mL	NS 250 mL (Before reconstitution, remove and discard diluent volume equal to volume of drug solution)		1 hour	8 hr. (complete within 8 hrs. of drug reconstitution)	drug	Do not administer as an IV push or bolus Protect from light during storage Administer through a dedicated IV line	Lexicomp
Bendamustine HCl (Treanda) Drug manufacturer phased out in March 2017 REF	SWFI 25 mg vial: 5mL 100 mg vial: 20mL	5 mg/mL	NS 500 mL (transfer to 500 mL infusion bag within 30 mins of reconstitution)	Powder for solution: 0.2-0.6 mg/mL	CLL: 30 minutes NHL: 60 minutes	3 hr.	24 hr.	Irritant with vesicant-like property Powder formulation: CSTDs or adaptors containing polycarbonate or acrylonitrile-butadiene-styrene (ABS) are safe to use with the lyophilized powder formulation Protect from light during storage	Lexicomp
Bendamustine HCl (Bendeka) (contains polyethylene glycol) Sound-alike, look- alike warning: do not confuse with Treanda	RTU 100 mg/4mL MDV (Allow to come to RT prior to use)	25 mg/mL	NS 50 mL (Use D5W for pts with restricted Na)	1.85-5.6 mg/mL	CLL and HLL: 10 minutes	NS: 3 hr.	NS: 24 hr.	Irritant with vesicant-like property Protect from light during storage	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe	Standard infusion duration/rate	Sta	bility	• •	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uurauom rate	RT	REF	requirement)	
Bevacizumab (Avastin) Sound-alike, look-alike warning: do not confuse with bevacizumab-awwb (Mvasi) REF	RTU 100 mg/4 mL vial 400 mg/16 mL vial	25 mg/mL	IV: NS 100 mL (do not administer or mix with D5W) Intravitreal: straight drug	1.4-16.5 mg/mL	Flat rate: 30 min Or first cycle: 90 minutes Second cycle: 60 minutes (if the initial infusion is well tolerated) Third and subsequent: 30 minutes (if the 60- minute infusion is well tolerated)	IV: No data Intravitreal: No data	IV: 8 hr. Intravitreal: 3 months	incision is fully healed) after surgery	IV: Lexicomp Intra- vitreal: Paul M et al
Bevacizumab-???? (Alymsys) Sound-alike, look-alike warning Bevacizumab-awwb (Mvasi)	RTU 100 mg/4 mL vial	S	IV: NS 100 mL (do not administer or mix with D5W)	1.4-16.5 mg/mL	Flat rate: 30 min Or 90/60/30 min rate	IV: no data	IV: 8 hr.	Same precaution as in bevacizumab	Lexicomp
Blinatumomab (Blincyto)	Mix 35 mcg single dose vial with SWFI (direct stream toward side of the vial, gently swirl; do not shake)	12.5 mcg/mL	24- or 48-hour infusion: add 270 mL NS to an empty IV bag 7-day infusion (for pts > 22 kg): add 90 mL bacteriostatic NS to an empty IV bag						

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concenti ation	uurauon/rate	RT	REF	requirement)	
Bleomycin (Blenoxane) REF	IV: NS Only 15-unit vial: 5 mL 30-unit vial: 10 mL IM/SQ: 15 units vial: 1 mL NS	IV: 3 units/ml IM/SQ: 15 units/ml	IV: NS 50 mL Intrapleural: 60 units in NS 50-100 mL	N/A	Minimum 10 minutes	24 hr.	No data	Irritant Do not reconstitute with dextrose Test dose (optional): IM, IV, SQ: administer 1 to 2 units of bleomycin in NS 50 mL	Lexicomp
Bortezomib (Velcade) RT Generic formulation: approved for IV route only	Mix 3.5 mg vial with NS: IV: 3.5 mL SQ: 1.4 mL	IV: 1 mg/mL SQ: 2.5 mg/mL	IV push: Straight drug, do not further dilute Administer IV push directly	N/A	Rapid IV push: 3-5 seconds	IV push: 3 d SQ: 8hr	IV push: 5 days SQ: no data	For IV or SQ administration only Irritant Protect from light during storage and reconstitution only	Lexicomp
Brentuximab vedotin (Adcetris)	Mix 50 mg vial with 10.5 mL SWFI	5 mg/mL	Minimum NS 100 mL	0.4-1.8 mg/mL	30 minutes	Immediate Use	hrs. of	Do not administer as IV push or bolus Do not shake Do not mix or infuse with other medications When administered with AVD, begin brentuximab within 1 hr. after completing AVD	Lexicomp
Busulfan (Busulfex) REF	RTU 60 mg/10 ml vial	6 mg/mL	NS or D5W (Dilution volume should be 10 times the volume of busulfan injection)	Minimum of 0.5 mg/mL			12 hr. (infusion must be completed within 12 hrs.)	Irritant Always add drug to the diluent (not diluent to busulfan) Incompatible with many CSTDs (contains N,N-dimethyl acetamide), refer to CSTD manufacturer for compatibility Do not use polycarbonate syringes Filter required (5-micron filter needle provided with ampule)	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	um anon/rate	RT	REF	requirement)	
Cabazitaxel (Jevtana) RT	Mix 60 mg/1.5 mL vial with 5.7 mL of supplied diluent	10 mg/mL	NS 250 mL non-DEHP bag (dilute within 30 mins)	0.1-0.26 mg/mL (Total doses >65 mg will require a larger infusion volume; final concentration should not exceed 0.26 mg/mL)	1 hour	8 hr. (infusion must be completed within 8 hrs.)	he completed	Infuse using a 0.22 micron inline filter Do not prepare or administer in PVC- containing infusion containers or polyurethane infusion sets Do not shake Allow to reach RT prior to infusion	Lexicomp
Carboplatin (Paraplatin) RT	RTU 50 mg per 5 ml vial 150 mg per 15 ml vial 450 mg per 45 ml vial 600 mg per 60 ml vial	10 mg/mL	NS 250 mL	0.5- 10 mg/mL	30-60 minutes (30 min in central line ok)	NS or D5W: 8 hr.	No data	Irritant Protect from light during storage Concentrations used for desensitization vary based on protocol Do not use with aluminum needle/administration sets	Lexicomp
Carfilzomib (Kyprolis) REF	Mix 60 mg vial with 29 mL SWFI	2 mg/mL	D5W 50-100 mL (may give as IV push or further dilute in D5W 50 mL only, use regular secondary tubing)	N/A	10 or 30 minutes (depends on regimen)	4 hr.	24 hr.	Do not administer as an IV bolus Requires hydration 250-500 mL NS prior to/after infusion (continue in subsequent cycle if needed) Do not shake Protect from light Do not administer with other medications	Lexicomp
Carmustine (BCNU) REF	Mix 100 mg vial with 3 mL provided diluent (dehydrated alcohol), then further dilute with 27 mL SWFI	3.3 mg/mL in 10% ethanol	NS or D5W 250 – 500 mL (use non-DEHP bag or glass container)		Minimum 2 hours (infusions <2 hrs. may lead to injection site pain or burning)	8 hr.	24 hr. (followed by additional 6 hr. at RT)	Irritant Prepared in either glass or polyolefin container or non-PVC container Protect from light in storage and dilution (does not require light protection in admin) Do not use if product has oil film on the vials	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	oility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	untent	concentration	duration/rate	RT	REF	requirement)	
Cemiplimab-rwlc (Libtayo) REF	RTU 350 mg/7 mL	50 mg/mL	NS or D5W	1 – 20 mg/mL	30 minutes	8 hr.	24 hr.	Contains polysorbate 80 (monitor for infusion reactions) Do not shake Allow to reach RT prior to infusion Infuse through 0.2-5 micron in-line or add-on filter	Lexicomp
Cetuximab (Erbitux) REF	RTU 100 mg/50 mL vial 200 mg/100 mL vial	2 mg/mL	Straight drug, do not dilute		Loading dose: 2 hours Maintenance dose: 1 hour	8 hr.	12 hours	Do not administer as IV push or bolus Administer through a low protein- binding 0.22 micrometer in-line filter Do not shake	Lexicomp
Cisplatin (Platinol) RT	RTU 50 mg/50 mL vial 100 mg/100 mL vial 200 mg/200 mL vial	1 mg/mL	>1/4 NS 500 mL must have a minimum conc. of 0.2% NaCl if only D5W is used, solution will PPT	1 mg/mL 0.05 – 2 mg/mL	60 min - 4 hours (varies, depends on dose and protocol) 1 mg/min rate has been used Continuous infusion has been used (e.g. ESHAP regimen)	20 hr. (Stability may vary due to pH, light and concentration of chloride)	Do not REF	Do not administer as a rapid IV infusion Vesicant (> 0.4 mg/mL); Irritant (≤ 0.4 mg/mL) Vesicant if > 20mL of 0.5 mg/mL solution infiltrates Do not dilute in D5W only Protect from light in storage. No light protection required during administration Do not use with aluminum administration set	Lexicomp
Cladribine (Leustatin) REF	RTU 10 mg/10 mL vial	1 mg/mL	24-hour infusion: 500 mL NS 7-day infusion: QS to total volume of 100 mL in CADD using bacteriostatic NS	IV: 0.016 mg/mL SQ: 1 mg/mL	Continuous infusion: 24 hours Off-label: 30 mins or 2 hours (infusion rate varies by indication and/or protocol)	IV: dilutions in NS should be used promptly SQ: 7 days	IV 24-hour: up to 8hr IV 7-day: up to 8hr prior to admin Bacteriostatic NS in CADD: 7 days	Irritant 24-hour infusion: filter with a 0.22-micron hydrophilic syringe filter prior to adding to infusion bag 7-day infusion: filter diluent and cladribine with a 0.22-micron hydrophilic filter prior to adding to cassette/reservoir	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter requirement)	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	duranon/rac	RT	REF		
Clofarabine	RTU	1 mg/mL	NS 100 mL	0.15-0.4 mg/mL	2 hours	24 hr.	No data	Filter drug through a 0.2-micron filter prior to dilution	Lexicomp
(Clolar)	20 mg/20 mLvial							Do not administer any other medications through the same intravenous line	
Cyclophosphamide (Cytoxan) RT	SWFI or NS: 500 mg vial: 25 mL 1 gm vial: 50 mL 2 gm vial: 100 mL	20 mg/mL	≤1 gm: 100 mL NS >1 gm: 250 mL NS	0.24 - 20 mg/mL	60 min (250 mL) (infusion rate may vary based on protocol)	24 hr.	6 days	Irritant Do not use vials with signs of melting which may appear as droplets Solutions reconstituted in SWFI are hypotonic and should not be used for direct IV push administration	Lexicomp
Cytarabine (Cytosar) RT	RTU 2 gm solution IV: Mix 1-gram vials with 10 mL BWFI (PF) IT: 100 mg for IT use: 1mL NS (PF) to 100 mg/mL	IV: 100 mg/mL IT: 100 mg/mL	NS 500 mL	32 mg/mL	AML: Continuous infusion 100-200 mg/m²/day High dose: 1-3 hours (Doses ≥ 1gm/m² should be infused over > 2 hours) (other rates have been used, refer to specific reference)	Powder for reconstitution 48 hrs. IV solution: 8 days IT: 8 hours		Irritant IT: use PF drug for IT doses or when cytarabine dose > 1 gm/m ²	Lexicomp
Cytarabine liposomal (Depocyte) REF	RTU 50 mg/5 mL vial (Allow vial to warm to RT)	10 mg/mL	Straight drug; Do not dilute	N/A	N/A	4 hr. (use within 4 hrs. of withdrawal from the vial)	No data (use within 4 hrs. of withdrawal from the vial)	Intrathecal route only Avoid aggressive agitation Do not filter Injection should be made slowly (over 1 to 5 minutes) Do not mix with any other medications	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uurauon/rac	RT	REF	requirement)	
Dacarbazine (DTIC-Dome) REF	SWFI: 100 mg with 10 mL 200 mg with 19.7 mL 500 mg with 49.25 mL	10 mg/mL	NS or D5W 250 mL	N/A	30-60 minutes Continuous infusion: up to 72 hrs.	8 hr. (stability of 72 hr. has been reported) (extended stability is important for continuous infusion)	24 hr. (stability of 4 days has been reported)	Irritant Protect from light for continuous infusion 7-day refrigerated stability in conc. up to 1.4 mg/mL (in D5W)	Lexicomp
Dactinomycin (Cosmegan, Actinomycin)	Mix 0.5 mg vial with 1.1 mL PF SWFI	500 mcg/mL	D5W or NS 50 mL	≥ 10 mcg/mL	15 minutes	4 hr. (varies based on protocol)	No data	Do not administer IM or SQ Vesicant Dilute in D5W or NS in glass or polyvinyl chloride (PVC) containers Protect from light in storage (does not require light protection in administration) Do not filter with cellulose ester membrane filters	Lexicomp
Daratumumab (Darzalex) REF	RTU 100 mg/5mL vial 400 mg/20 mL vial	20 mg/mL	NS 500-1,000 mL (remove the volume of NS from the bag that is equal to the required volume of the daratumumab dose)	N/A	Variable, max rate (typically 7-10 hrs.) If expected to infuse > 5 hrs., may split dose into 2 bags, to give on days 1, 2 sequentially	15 hr. (complete infusion within 15 hrs.)	24 hr.	Do not administer IV push or as a bolus Protect from light Administer with an inline filter (0.22 or 0.2 micron) Ok to use PVC bag Do not mix with or infuse with other medications	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concenti ation	um anon/rate	RT	REF	requirement)	
Daratumumab hyaluronidase, subcutaneous (Darzalex Faspro)	RTU 1,800 mg daratumumab and 30,000 units hyaluronidase per 15 mL in single dose vial	N/A	N/A		3-5 min subcutaneously	Immediate use	N/A	Do not substitute daratumumab/hyaluronidase for daratumumab IV and vice versa Administer SQ over ~3 to 5 min Compatible with PVC syringe	Lexicomp
Daunorubicin (Cerubidine, Daunomycin) RT	RTU 20 mg/4mL (4 mL)	5 mg/mL	NS or D5W 50 mL	5 mg/mL	15 -30 minutes	24 hr.	No data	Do not administer IM or SQ Vesicant Protect from light in storage (does not require light protection in administration)	Lexicomp
Daunorubicin liposomal (Daunoxome) REF Sound-alike, look alike warning: do not confuse with Daunorubicin and cytarabine liposomal	RTU 50 mg/25mL vial	2 mg/mL	1:1 concentration in D5W	1 mg/mL	60 minutes	Immediate use	6 hr.	Irritant Dilute with D5W only Protect from light Do not filter Do not mix with other drugs	Lexicomp
Daunorubicin and cytarabine liposomal (Vyxeos, CPX-351) REF Sound-alike, look alike warning: do not confuse with Daunorubicin liposomal	Mix each vial (44/100 mg) with 19 mL SWFI	2.2 mg/mL	NS or D5W 500 mL	N/A	90 minutes	Immediate use	4 hr.	Do not administer IM or SQ Irritant Flush line with NS or D5W after infusion Do not mix with other medications	Lexicomp
Decitabine (Dacogen) RT	Mix 50 mg vial with SWFI: IV: 10mL SQ: 5mL	IV: 5 mg/mL SQ: 10 mg/mL	NS 50 mL (use cold NS bags to prepare IV doses)	IV: 0.1-1 mg/mL	1-3 hours (AML) (Rate of administration depends on protocol)	No data	IV: 4 hr. (if prepared in cold NS)	Short stability	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	* *	Ref
condition/special warning	RTU information	Concentration	dituent	concentration	duration/rate	RT	REF	requirement)	
Denileukin diftitox (Ontak) (in freezer ≤ 10 °C)	RTU 300 mcg/2ml vial	15 mcg/mL	NS (PF) (thaw frozen vials for 1- 2 hrs. at RT or 24 hrs. refrigerated)	>/= 15 mcg/mL	30-60 min (can administer over up to 80 min, depending on severity of infusion reaction)	6 hr.		Do not administer by bolus IV injection Irritant Do not shake, filter or re-freeze vials Do not mix with other drugs	Lexicomp
Dexrazoxane (Zinecard) Do not use Totect for extravasation (non-chemo) RT	Brand (Zinecard): mix 250 mg with 25 mL SWFI Or mix 500 mg with 50 mL SWFI Generic (dexrazoxane): Reconstitute with supplied diluent (1/6 M sodium lactate Continuous infusion data available in Lexicomp		Cardioprotection Zinecard: LR only Dexrazoxane: NS 50-100mL Anthracycline extravasation Dexrazoxane: NS 1,000 mL	Zinecard: 1.3-3 mg/ml Dexrazoxane: 1.3-5 mg/ml Anthracycline extravasation: not to exceed 10 mg/mL	Cardioprotection Zinecard: 15 minutes Dexrazoxane: 15 minutes Anthracycline extravasation Dexrazoxane: 1-2 hours Cardioprotection: Infuse over 10-15 min before anthracycline extravasation: infuse over 2-3 hr.	Zinecard: 1 hr. (in LR) Dexrazoxane: 6 hr. (in NS)	(in LR) Dexrazoxane: 6 hr. (in NS)	Cardioprotectant Do not administer by IV push Administer doxorubicin within 30 minutes after completion of the dexrazoxane infusion Anthracycline extravasation: IV administration only Infusion solution should be at RT prior to administration Must initiate within 6 hr. of extravasation Do not order Totect Kit when anthracycline induced extravasation is suspected, use dexrazoxane	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	oility	• •	Ref
condition/special warning	RTU information	Concentration	unuent	Concenti ation	dui attom race	RT	REF	requirement)	
Docetaxel (Taxotere) REF Sound-alike, look alike warning: verify strength prior to admixture	RTU 20 mg 80 mg 160 mg 200 mg PF and non-alcohol forms are available (docetaxel is available in multiple concentrations and multiple dosage forms)	20 mg/mL (do not use 10 mg/mL)	IV: NS or D5W 250 mL (use non-DEHP bag) Intravesical: in 50 mL syringe (preferred, to avoid spasm), do not flush	j	60 minutes	Taxotere: 6 hr. Generic: 6 hr. Non-alcohol formulation: no data	No data	Irritant with vesicant-like property Mix in Non-DEHP bag and administer through non-DEHP or non- PVC tubing Do not shake Use of filter is not necessary (although has been commonly practiced) Protect from light	Lexicomp
Doxorubicin (Adriamycin) RT Sound-alike, look alike warning: do not confuse with doxorubicin liposomal (Doxil)	RTU 2 mg/mL (5 mL, 10 mL, 25 mL) (multiple vial sizes are available) (doxorubicin solid powder is reserved for trans-arterial chemoembolization or TACE)	2 mg/mL			Intermittent infusion: 15 minutes Continuous infusion: Rate of admin varies by protocol (e.g. infusional EPOCH, AIM)	6 hr. (per USP 797) (Extended stability has been reported)	(Extended stability has been reported)	Do not administer IM or SQ Vesicant Avoid contact with alkaline solutions For TACE: loading time with doxorubicin liquid form < < than that of solid (needs overnight preparation). See Appendix B	Lexicomp
Doxorubicin liposomal (Doxil) REF Sound-alike, look alike warning: do not confuse with doxorubicin (Adriamycin)	RTU 20 mg/10mL vial 50 mg/25mL vial	2 mg/mL	Dose ≤ 90 mg: D5W 250 mL Dose >90 mg: D5W 500 mL	0.36 mg/mL	60 minutes (initial rate of 1 mg/minute recommended to minimize risk of infusion reactions)	No data	24 hr.	Do not administer IV push Irritant Do NOT administer undiluted Dilute ONLY in D5W Do not mix with bacteriostatic agents Do not filter Do not mix with other medications	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuciit	concentration	durunomrute	RT	REF	requirement)	
Durvalumab (Imfinzi) REF	RTU 120 mg/2.4 mL 500 mg/10 mL	50 mg/mL	NS or D5W	1 - 15 mg/mL	60 minutes	Immediate use	24 hr.	Contains polysorbate 80 (monitor for infusion reactions) Protect from light Do not shake Administer with low-protein binding 0.2-0.22 micron in-line filter Do not mix with other medications	Lexicomp
Eculizumab (Soliris) (non-chemo) REF	RTU 300 mg (30 mL)	10 mg/mL	Add drug to an infusion bag (e.g. viaflex bag), dilute with equal volume of NS to final conc of 5 mg/mL (e.g. 300 mg dose to a total volume of 60 mL and etc.)	5 mg/mL	Adults: 35 minutes Pediatrics: 1-4 hrs. (Do not exceed a max 2-hr duration of infusion in adults)	24 hr. (use within this time frame)	24 hr. (use within this time frame)	Do not administer as an IV push or bolus Do not shake Protect from light in storage and dilution (no light protection in administration) Allow to reach RT prior to administration Prescriber must be enrolled in REMS program	Lexicomp
Efgartugnid Alfa (Vyvgart©)	400 mg (20 mL)	20 mg/mL	Dilute the dose in NS to make a total volume 125 mg	Varies	Infuse over 1 hour via a 0.2 micron in-line filter	4 hr	8 hr	Only administer with PE, PVC, EVA, or polyurethane/polypropylene infusion lines. Do not mix efgartigimod alfa with other medications or inject other medications into infusion side ports. Flush the entire line with 0.9% Sodium Chloride Injection, USP following infusion.	Lexicomp
Efgartigimod Alfa- hyaluronidase (Vyvgart© Hytrulo)	RTU 1,008 mg efgartigimod alfa and 11,200 units hyaluronidase /5.6 mL	RTU 1,008 mg efgartigimod alfa and 11,200 units hyaluronidase /5.6 mL	n/a - RTU	RTU 1,008 mg efgartigimod alfa and 11,200 units hyaluronidase /5.6 mL	Over 30-90 seconds	PF - use immediately once withdrawn from vial	PF - use immediately once withdrawn from vial	Use a winged infusion set made of polyvinyl chloride (PVC), 25G, 12 inches tubing, maximum priming volume of 0.4 mL	Package insert

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	oility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	Concentration	uui auoii/1 ate	RT	REF	requirement)	
Elotuzumab (Empliciti) REF	300 mg: 13 mL SWFI 400 mg: 17 mL SWFI	25 mg/mL	NS 230 mL NS (PVC bag or polyolefin bag) (may adjust volume not to exceed 5 mL/kg of body (pt. weight)	Volume of diluent < 5 mL/kg (patient weight in kg)	First infusion: start at 0.5 mL/min, escalate by 1 mL/min to max 2 mL/min Second infusion: start at 3 mL/min for 1st 30 min; if tolerated, escalate to max 4 mL/min Subsequent infusion: if no prior reaction, infuse at 5 mL/min	8 hr.	24 hr. (including 8hr in RT)	Do not administer IV push or as a bolus Administer with a sterile, non-pyrogenic, low protein-binding filter (0.2 to 1.2 micron) Do not shake After dissolution, allow the reconstituted vials to stand for 5 to 10 minutes Protect from light in storage Do not mix with or infuse with other medications	Lexicomp
Emapalumab (Gamifant) REF	RTU 10 mg/2 mL 50 mg/10 mL	5 mg/mL	NS	0.25 – 2.5 mg/mL	1 hour	Immediate Use	4 hr.	Infuse through IV line containing a sterile, non-pyrogenic, low-protein binding 0.2-micron inline filter Store in original carton to protect from light Bring solution to RT prior to infusion Do not mix with other drugs	Lexicomp
Enfortumuamb vedotin (Padcev) REF	20 mg: mix with 2.3 mL SWFI 30 mg: mix with 3.3 mL SWFI	10 mg/mL	NS 250 mL	0.3-4 mg/mL	30 min	Immediate use	8 hr.	Do not administer IV push or bolus Maximum dose is 125 mg	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	dituent	concentration	duration/rate	RT	REF	requirement)	
Epcoritamab-bysp (Epkinly) REF	dilutions): Add 4.2 mL of NS to 0.8 mL (4 mg) of epcoritamab for an initial concentration of 0.8 mg/mL. Transfer 2 mL of this dilution to empty vial and add 8 mL		NS		N/A (subcutaneous injection)	12 hrs	24 hrs	Allow epcoritamab solution to equilibrate to RT for no more than 1 hr before administration Do not freeze Do not shake intact vials Store in carton to protect from light Filtering of diluted solutions is not required	Lexicomp
Epirubicin (Ellence) REF Sound-alike, look-alike warning: do not confuse with eribulin (Halaven)	RTU 50 mg 200 mg	2 mg/mL	NS or D5W 100 mL	N/A	15 min	Immediate use	24 hr.	Vesicant Protect from light	Lexicomp
Eptinezumab (Vyepti) REF	RTU 100 mg 300 mg	100 mg/mL	NS 100 mL	100 mg/mL	30 min	8 hrs.	No data	Do not administer as IV push or bolus injection Do not mix or infuse other medications in same infusion set Following infusion, flush line with 20 mL NS	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion	Stal	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uurauon/rate	RT	REF	requirement)	
Eribulin (Halaven) RT Sound-alike, look-alike warning do not confuse with epirubicin (Ellence)	RTU 1 mg/2 mL (2 mL)		NS 100 mL or undiluted in syringe	N/A	2-5 min	4 hr.	24 hr.	Do not administer other medications through the same IV line, or through a line containing dextrose	Lexicomp
Etoposide (Toposar) RT Sound alike, look alike warning: do not confuse with etoposide phosphate (Etopophos)	RTU 100 mg 500 mg 1 g	20 mg/mL	NS 500 mL (Use a non-DEHP bag) Continuous infusion is available (e.g. infusional EPOCH)	Max concentration not to exceed 0.4 mg/mL, adjust volume of diluent and infusion rate if necessary	(90 min or longer if not tolerated) (Higher dose may be infused over longer time periods depending on the	0.2 mg/mL: 96 hr. 0.4 mg/mL: 24 hr. (extended stability has been reported for continuous EPOCH)	ino data	Do not infuse by rapid infusion: may cause hypotension Irritant Concentration > 0.4 mg/mL is unstable and may PPT Administer infusion with filter per HM standard May administer through a non-PVC (low sorbing) tubing Do not infuse by rapid infusion, may cause hypotension	Lexicomp
Etoposide phosphate (Etopophos) Sound alike, look alike warning: do not confuse with etoposide (Toposar) REF	Mix 100 mg vial with 5 mL SWFI	20 mg/mL	Straight drug Or NS (use a non- DEHP bag)	0.1mg/mL - 20 mg/mL	5 minutes - 3.5 hours	24 hr.	24 hr.	Do not interchange with etoposide (unless in drug shortage per policy) (113.5 mg etoposide phosphate = 100 mg etoposide) IV bolus over at least 5 minutes Mix in non-DEHP bag, non-DEHP tubing with 0.2-micron filter	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion	Stal	oility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent			RT	REF	requirement)	
Fam-trastuzumab deruxtecan (Enhertu) Sound alike, look alike warning: do not confuse with trastuzunab (Herceptin) or adotrastuzumab emtansine (Kadcyla)	Mix 100 mg vial with 5 mL SWFI	20 mg/mL	D5W 100 mL (do not dilute in NS)	N/A	1 st dose: 90 min Subsequent dose over 30 min (if tolerated)	4 hr.	24 hr.	Administer in bag made of polyolefin or polybutadiene (ok for PVC bag) and a 0.2- or 0.22-micron in-line polyethersulfone or polysulfone filter Allow solution to reach RT prior to administration Do not shake the reconstituted or diluted solution. Cover infusion bag to protect from light. Do not mix fam-trastuzumab deruxtecan with other medications.	Lexicomp
Degarelix (Firmagon)	Reconstitute with provided prefilled syringe containing preservative free sterile water for injection	Loading dose 40 mg/mL each syringe x 2 syringes Maintenance: 20 mg/mL x1 syringe		Loading dose 40 mg/mL each syringe x 2 syringes Maintenance: 20 mg/mL x1 syringe	Subcutaneous injection	Administer within 1 hour of reconstitution	Administer within 1 hour of reconstitution	N/A	Lexicomp
Fludarabine (Fludara) REF	RTU Or mix 50 mg vial with 2mL SWFI	25 mg/mL	NS or D5W 100 – 125 mL	1 mg/mL	30 minutes	8 hr.	No data	Protect from light	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uuruuomraee	RT	REF	requirement)	
	RTU 500 mg 1 g 2.5 g (multiple vial sizes are available)	50mg/ml	NS Outpatient: use CADD pump (use only NS as diluent), volume depends on dose (> 5,000 mg requires 250 mL cassette; < 5,000 mg: use 100 mL cassette) Inpatient: add daily dose to 500 mL NS, hang bag daily	50 mg/mL	15 minutes- variable (based on protocol)	Varies 7 days (in CADD pump if diluted) (follow USP 797 standard for beyond use date)	reported)	Irritant May be administered by IV push, IV bolus, or continuous infusion	Lexicomp
Gemcitabine (Gemzar) RT	RTU 200 mg vial: 5 mL 1 gm vial: 25 mL 2 gm vial: 50 mL 200 mg vial: 5 mL	IV: 38 mg/mL Intravesical: 40 mg/mL	IV: NS 250 mL Intravesical: 50 mL NS (preferred, to avoid urinary spasm), do not flush	IV: 0.1 mg/mL - 25 mg/mL Intravesical: 20 mg/mL or 40 mg/mL	30 min Fixed dose rate at 10 mg/m ² /min has been reported for sarcoma, pancreatic, biliary and urothelial ca)	IV: 24 hr. Intravesical: 24 hr.	Do not REF (may PPT)	Intravesical: admix in 50- or 100-mL NS (usual dwelling time: 1 hr.) Protect from light	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uurauon/rate	RT	REF	requirement)	
Gemtuzumab Ozogamicin (Mylotarg)	4.5 mg vial: 5 mL SWFI Allow to warm to room temperature prior to reconstitution	Dose < 3.9 mg should be prepared in a syringe to reduce the impact of drug adsorption	0.3 mg up to 0.45 mg: use 4 mL syringe 0.45 mg up to 0.75 mg: use 6 mL syringe 0.75 mg up to 1.89 mg: use 10 mL syringe 1.89 mg up to 3.9 mg: use 25 mL syringe 3.9 mg up to 11.7 mg: use 50 mL IV bag 11.7 mg up to 21.6 mg: use 100 mL IV	Final cone: 0.075-0.234 mg/mL	2 hours	6 hr.	12 hr.	Premedicate appropriately Do not administer as IV push or bolus Protect bag from light using a light- blocking cover; infusion line does not need protection from light Administer through a low protein- binding (0.2 to 1.2 micron) in-line filter Do not co-administer other drugs through the same infusion line	Lexicomp
Glucarpidase (Voraxaze) REF	IV: mix each vial (1,000 units) with 1 mL NS IT: mix 2,000 units with 12 mL preservative-free NS	IV: 1000 units/mL	bag IV: 1 mL NS IT: 12 mL PF NS		IV: 5 minutes IT: 5 minutes via lumbar route, ventriculostomy, Ommaya reservoir (within 3-9 hr. of IT methotrexate overdose)	Immediate Use	4 hr.	IV: flush line before and after administration Administer leucovorin at least 2 hr. before and 2 hr. after glucarpidase dose Mix gently by rolling/tilting vial; do not shake Clinical monitoring guideline for methotrexate level is available	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	* *	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uuruuomraee	RT	REF	requirement)	
Hemin (Panhematin)	Mix 350 mg vial with 48 mL SWFI	7 mg/mL	N/A	7 mg/mL	30 min (flush with NS 100 mL after	Immediate Use	No data available	Reconstitute immediately prior to use	Lexicomp
(ramemaun)	46 IIIL SWFI				infusion) With albumin added:			Shake vial well for 2-3 minutes to dissolve	
(non-chemo)	Albumin 25% has been added to decrease risk of phlebitis in some protocols				max rate ≤ 1 mg/min			Maximum dose of 6 mg/kg/24 hours	
RT	(mixing instruction with albumin embedded in DoseEdge)							Use ≤ 0.45-micron filter Do not mix with other	
	DoseBage)							medications	
Idarubicin	RTU	1 mg/mL	Straight drug	N/A	15 minutes	No data	No data	Do not administer IM or SQ	Lexicomp
(Idamycin)	5 mg		or dilute in NS 50			(follow USP	(follow USP	Vesicant	
REF	10 mg 20 mg		mL			797 for beyond use date)	797 for beyond use date)	Protect from light during storage (does not require light protection during administration)	
Ifosfamide	1 g/20 mL (20 mL)	50 mg/mL	NS 500 mL	0.6-20 mg/mL	1-3 hours	No data	24 hr.	Irritant	Lexicomp
(Ifex)	3 g/60 mL (60 mL)	·		, and the second	(Infusion duration may vary, refer to			Should be given with mesna (concurrent, before or after	
RT					specific protocol for duration)			ifosfamide infusion, check protocol) and hydration	
								Requires hydration	
Imiglucerase	400-unit vial: mix with	40 units/mL	NS 100-200 mL		1-2 hours	12 hr.	24 hr.	NOT cytotoxic	Lexicomp
(Cerezyme)	10.2 mL SWFI		(4 1 4 CT					Diluted solution may be filtered through an in-line low protein-	
(non-chemo)			(translucent fiber may appear after dilution, do not use if discolored or					binding 0.2-micron filter NOT cytotoxic	
REF			opaque particles appear)						

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unucit	concentration	duration/race	RT	REF	requirement)	
inebilizumab-cdon (Uplizna)	100 mg/10 mL (10 mg/mL) solution in a single-dose vial must be diluted d in 250 mL of 0.9% Sodium Chloride Injection, USP prior to administration	10 mg/mL	250 mL of 0.9% Sodium Chloride	1 mg/mL	0-30 minutes = 42 mL/hr 31-60 minutes = 125 mL/hr 61-completion = 333 mL/hr Total infusion time ~90 minutes	4 hours	24 hours	Administer through an intravenous line containing a sterile, low-protein binding 0.2 or 0.22 micron in-line filter.	Package insert
Infliximab (Remicade) (non-chemo) REF	Mix 100 mg vial with 10 mL SWFI (use 21-guage or smaller needle; direct SWFI to the wall of the vial. Allow to stand for 5 minutes)	10 mg/mL	NS to total volume of 250 mL (add reconstituted infliximab slowly)	0.4- 4 mg/mL	Minimum of 2 hours (for first dose: may follow titration rate per HM standard)	3 hr. (Infusion should begin within 3hr of reconstitution & dilution.)	24 hr. (14 days have been documented in one study)	Risk of hypersensitivity reactions, premedicate appropriately Infuse using in-line low protein binding filter (≤1.2 micron) Do not shake Do not dilute reconstituted infliximab solution with any other diluent Contraindicated in moderate to severe heart failure Do not infuse with other agents	Lexicomp
Interferon alfa-2b (Intron A) REF	RTU Mix 10 million units vial, 18 million units vial or 50 million units vial with 1 mL of SWFI	10 million units/mL 18 million units/mL 50 million units/mL	NS 100 mL	Must be ≥ 10 million units/100 mL	20 minutes	IV: Immediate Use	IV: 24 hr.	Refer to package insert for specific route (IM or SQ) if needed Not all routes are available for all dosage forms	Lexicomp
Ipilimumab (Yervoy) REF	RTU 50 mg 100 mg	5 mg/mL	QS with NS	1-2 mg/mL	90 minutes (melanoma) 30 min (colorectal, hepatocellular, NSCLC or renal ca)	24 hr.	24 hr.	Drug must sit at RT for 5 minutes prior to admixing Do not shake Administer with low protein binding, 0.2-micron filter	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uurauon/acc	RT	REF	requirement)	
Irinotecan	RTU	20 mg/mL	D5W 500 mL	0.12-2.8 mg/mL	90 minutes	24 hr.	48 hr.	Irritant	Lexicomp
(Camptosar)	40 mg 100 mg					(in D5W)	(in D%W)	Protect from light (does not require light protection in admin)	
RT	300 mg							Do not REF dose diluted in NS, drug may PPT	
Irinotecan liposomal	RTU 43 mg/10 mL (10 mL)	4.3 mg/mL	D5W or NS 500 mL	N/A	90 min	4 hr.	24 hr.	Mix by gentle inversion, protect diluted solution from light	
(Onivyde) Sound-alike, look-alike warning: do not confuse with irinotecan (Camptosar) REF								Allow diluted solution to come to RT prior to administration	
Ixabepilone	Use supplied diluent:		QS with LR to ~250 mL in non-DEHP	0.2-0.6 mg/mL	3 hours	6 hr.	No data	Irritant	Lexicomp
(Ixempra)	15 mg vial: 8 mL		mL in non-DEHP bag			(if pH is maintained 6-		Use non-DEHP administration set (e.g. polyethylene); filter with a 0.2	
REF	45 mg vial: 23.5 mL (reconstitute only with supplied diluent) (vials must be allowed to warm to RT for 30 min prior to constitution)				hrs.)	9)		to 1.2-micron inline filter Protect from light (does not require light protection in admin)	
Lecanemab (Leqembi®)	Withdraw the required volume of LEQEMBI from the 100 mg/mL vial(s) and add to an infusion bag containing 250 mL of 0.9% Sodium Chloride Injection, USP	Each vial contains a LEQEMBI concentration of 100 mg/mL	0.9% Sodium Chloride Injection, USP	variable	1 hour	4 hours	4 hours	terminal low-protein binding 0.2 micron in-line filter	Package insert

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	bility	• •	Ref
condition/special warning	RTU information	Concentration	unuent	Concenti ation	duration/rate	RT	REF	requirement)	
Leucovorin calcium (folinic acid) (non-chemo) RT Sound-alike, lookalike warning: do not confuse with levoleucovorin (Fusilev)	100 mg/10 mL (10 mL) 500 mg/50 mL (50 mL) Or use solid powder: 100 mg vial: mix with 10 mL SWFI 200 mg vial: mix with 20 mL SWFI	10 mg/mL 10 mg/mL 10 mg/mL 10 mg/mL 20 mg/mL	D5W or NS 50-500 mL (for methanol toxicity, dilute in D5W)	2-10 mg/mL	May give as IM, IV push IV infusion: 15 minutes - 2 hours (maximum rate of 160 mg/min due to calcium content)	No data	prepared in BSWFI in one study)	Not intended for intrathecal use Store refrigerated Protect from light Leucovorin should not be mixed in same infusion as fluorouracil Leucovorin should not be administered concurrently with methotrexate Commonly initiated 24 hours after start of methotrexate (toxicity to normal tissues may be irreversible if leucovorin is not initiated by ~40 hours after the start of methotrexate	Lexicomp
(non-chemo) Sound-alike, lookalike warning:	RTU 175 mg/17.5 mL (17.5 mL) 250 mg/25 mL (25 mL) Or use solid powder: Mix 50 mg vial with 5.3 mL PFNS Mix 175 mg with 3.6 mL PF NS Mix 300 mg with 6.2 mL PF		NS	0.5-5 mg/mL	hours (maximum rate of 160 mg/min due to calcium content)	NS: 4 hr. D5W: 4 hr. Lyophilize d powder: 12 hr. in NS (4-12 hrs. in D5W)	No data available	Do not administer intrathecally NOT cytotoxic Do not prepare with other products in the same admixture; may cause PTT	Lexicomp
Loncastuximab tesirine (Zylonta) REF	10 mg vial; mix with 2.2 mL SWFI	5 mg/mL (gentle swirl, do not shake)	D5W 50 mL	N/A	30 min	8 hr		Infuse through a dedicated line with low-protein binding 0.2 or 0.22 micron in-line or add-on filter and catheter. Do not infuse with other medications Loncastuximab tesirine does not have incompatibilities with PVC, polyolefin, or copolymer of ethylene and propylene	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	* *	Ref
condition/special warning	RTU information	Concentration	dituent	concentration	duradon/rate	RT	REF	requirement)	
Lurbinectedin (Zepzelca)	4 mg vial: 8 mL of SWFI	0.5 mg /mL	NS or D5W	100 mL (central line) or ≥ 250 mL (peripheral line)	60 minutes	24 hr.	24 hr.		Lexicomp
RT									
Luspatercept (Reblozyl) REF	25 mg vial with 0.68 mL SWFI	50 mg/mL	N/A	50 mg/mL	N/A (subcutaneous injection)	8 hr.	24 hr.	Doses requiring larger reconstituted volumes (>1.2 mL) should be divided into separate syringes	Lexicomp
	75 mg with 1.6 mL SWFI							Direct stream onto powder and allow to stand for 1 minute. Gently swirl for 30 seconds, let vial sit for additional 30 seconds. repeat until powder is completely dissolved	
Margetuximab (Margenza) REF	RTU 250 mg/10 mL (10 mL)	25 mg/mL	NS 100 mL or NS 250 mL (based on final	0.5 – 7.2 mg/mL	Initial dose: 120 minutes; infuse subsequent doses over at least 30 minutes.	4 hr	24hr	Infuse via a sterile, nonpyrogenic, low- protein binding polyethersulfone 0.2 micron (in-line or add-on) filter. May use non-PVC bag made with	Lexicomp
			conc.)					polyolefins (polyethylene and polypropylene) and polyamide or polyolefins	
								Do not administer as an IV push or bolus	
Mechlorethamine	10 mg vial with	1 mg/mL	Straight drug;	1 mg/mL syringe	IV push: 3-5 minutes	Immediate	Immediate	Vesicant	Lexicomp
(Mustargen)	10 mL SWFI		Do not dilute- due to short stability,	Give iv push (free flowing IV at gravity)	in free flowing IV fluid	use	use	Protect from light	
RT	Use immediately (IV		give iv push	in ing it at gravity)				Use immediately (IV solution decomposes rapidly)	
	solution decomposes rapidly)							Infusion of the drug is not recommended (phlebitis, venous irritation)	

Generic name	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	bility	Special precautions/filter	Ref
(brand)/storage condition/special warning	RTU information	Concentration	dituent	concentration	duration/rate	RT	REF	requirement)	
preparation, storage, and dosing may	Mix 50 mg vial with 10 mL of supplied diluent Time between reconstitution/dilution and administration must be kept to a minimum (reconstituted and diluted solutions are unstable)	5 mg/mL	NS 250 mL	≤ 0.45 mg/mL	15-20 minutes	4 hrs.	Do not REF (may PPT)	Irritant Do not mix or combine formulations Protect from light Shake vigorously	Lexicomp
Mesna (Mesnex) (non-chemo) RT	RTU 100 mg	100 mg/mL		(if given together with ifosfamide in same bag, ifosfamide final conc. ≤ 50 mg/mL)	as an IV bolus (5-15 min), short infusion (30 min) or as continuous infusion (12-2 4 hrs.)	(24 hr. in one study) (follows USP 797 beyond- use-date rule)	Varies (48 hr. in one study) (follows USP 797 beyond-use- date rule)	IV bolus dose is equal to 20% of the ifosfamide dose at the time of ifosfamide administration (may give after ifosfamide infusion and 4 and 8 hours after each dose) Total dose of mesna is 60% of ifosfamide dose (as intermittent infusion) May also be given continuous infusion	Lexicomp

Generic name (brand)/storage	Reconstitution and/or RTU information	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	bility	Special precautions/filter requirement)	Ref
condition/special warning	K1 U illioi mauon	Concentration				RT	REF	requirement)	
Methotrexate (Trexate) RT	RTU 50 mg 250 mg 1 g Must use PF NS or SWFI if using for intrathecal or intraventricular chemotherapy and if dose used is ≥ 1 g/m2	IV: 25 mg/mL IV: 100 mg/mL (high dose)	Bolus: NS 100 mL(bolus) Continuous infusion: 1,000 mL IT: QS with PF NS to a final volume of up to 6 mL Intra- ventricular: QS with NS or SWFI to a final volume of up to 3-6 mL	25 mg/mL IT: 8 hours		IV: 24 hr. IT: immediate use (no data per USP 797 rule, 4 hrs. has been suggested in one facility)	No data	High dose should be followed by leucovorin calcium rescue to prevent toxicity Protect from light Light (does not require light protection in administration)	Lexicomp
Mirvetuximab Soravtansine-gynx (Elahere TM) RT or REF	RTU 100 mg/20 mL single-use vial	5 mg/mL	IV: D5W 500 mL	1-2 mg/mL	First dose: 1 mg/min If well tolerated after 30 min, increase to 3 mg/min If well tolerated after 30 min, increase to 5 mg/min (Max rate) (Subsequent infusions should be started at the maximally tolerated rate)	IV: 8 hrs	IV: 12 hrs	Hazardous – Handle with care 0.2 or 0.22 µm polyethersulfone (PES) in-line filter Do not shake (If refrigerated, allow the infusion bag to reach room temperature prior to administration)	Lexicomp
Mitomycin (Mutamycin)	5 mg vial: SWFI 10mL 20 mg vial: SWFI 40 mL 40 mg vial: SWFI 80mL Intravesical:	IV: 0.5mg/mL Intravesical: 2 mg/mL (1 mg/mL has been used)	also dilute in sodium lactate) Intravesical: mitomycin 40 mg in	IV: 0.02 - 0.04 mg/mL Intravesical: 2 or 1 mg/mL Intraocular: 0.2mg/mL	IV: 15 minutes		Intravesical: no	Vesicant Shake to dissolve Protect from light	Lexicomp Au J et al
	40 mg in 20 mL NS or SWFI (40 mg in 40 mL NS or SWFI has been used) Intraocular: see appendix A	Intraocular: 0.2mg/mL (0.02%) 0.4 mg/mL (0.04%)	20 mL NS/SWFI 40 mg in 40 mL NS/SWFI has been used Intraocular conc.: see appendix A	(0.02%) and 0.4 mg/mL (0.04%)		Immediate use Intraocular: Immediate use	Intraocular: no	Vesicant	

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	Concentration	uurauon/rate	RT	REF	requirement)	
Mitoxantrone (Novantrone) RT	RTU 20 mg MDV 25 mg MDV 30 MDV	2 mg/mL	NS or D5W 100 mL	0.5 mg/mL	15 minutes	per drug manufacturer)		Do not administer intrathecally, subcutaneously, intramuscularly or intraarterially Vesicant Protect from light Do not mix with other drugs	
Mosunetuzumab- axgb (Lunsumio)	1 mg/ 1 mL single-use vial 30 mg/ 30 mL single-use vial RTU	1 mg / mL	0.9% sodium chloride or 0.45% sodium chloride	Variable See package insert for chart	Cycle 2+: administer over 2 hours if cycle 1	per drug manufacturer) Solutions may be stored at 9°C to 30°C (48°F to 86°F) for up to 16 hours	per drug manufacturer) Store intact vials at 2°C to 8°C (36°F to 46°F). Do not	Do not use an in-line filter to administer LUNSUMIO Only use infusion bags made of polyvinyl chloride (PVC) or polyolefin (PO) such as polyethylene (PE) and polypropylene. Allow solution to reach room temperature prior to infusion	PI
Moxetumomab Pasudotox (Lumoxiti) REF	Mix each vial (1 mg) with 1.1 mL SWFI Add 1 mL IV solution stabilizer to 50 mL NS infusion bag (before adding reconstituted drug)	1 mg/mL	NS 50 mL	N/A	30 minutes	4 hr.	24 hr.	Do not reconstitute with other solutions Flush line with NS (same rate as infusion) to ensure entire dose delivered Protect from light during storage Do not shake Do not mix with other drugs	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uurauomrate	RT	REF	requirement)	
Natalizumab	RTU	20 mg/mL	NS 100 mL	2.6 mg/mL	60 minutes	Immediate use	8 hr.	Do not administer by IV bolus or push	Lexicomp
(Tysabri)	300 mg/15 mL							Do not shake	
(non-chemo)							No data	Protect from light	
REMS program (TOUCH)								Patient, prescriber, and facility must be	
REF								enrolled in REMS program (TOUCH)	
								Allow solution to warm to RT prior to administration	
Nelarabine	RTU	5 mg/mL	Straight drug	N/A	hours (adults)	8 hr.	No data	Irritant	Lexicomp
(Arranon)	250 mg		Do NOT further dilute		1 hr. (pediatrics)			Do not dilutee	
RT			Transfer dose to an empty PVC or glass infusion container						
Nivolumab	RTU	10 mg/mL	NS 100 mL NS	1-10 mg/mL	30 min	8 hr.	24 hr.	Administer with 0.1 to 1.2-micron low protein binding in-line filter	Lexicomp
(Opdivo)	40 mg 100 mg 240 mg							Do not shake	
REF	2-40 mg							Protect from light	
								When administered in combination with ipilimumab, infuse nivolumab first followed by ipilimumab on the same day	
								Discard if cloudy or has color change (should be clear to pale yellow)	
								Do not administer other medications through the same IV line	

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	* *	Ref
condition/special warning	RTU information	Concentration	dituent	concentration	duration/rate	RT	REF	requirement)	
Nivolumab/Relatli mab-rmbw (Opdualag)	Nivolumab 240 mg and relatlimab rmbw 80 mg per 20 mL (20 mL)	(per mL)	Dilute with NS or D5W. The concentration for administration should range from 3 to 12 mg/mL (nivolumab component) and from 1 to 4 mg/mL (relatlimab component); the maximum infusion volume should be 160 mL	The concentration for administration should range from 3 to 12 mg/mL (nivolumab component) and from 1 to 4 mg/mL (relatlimab component	Infuse over 30 minutes via an IV line containing a sterile, nonpyrogenic, low protein binding polyethersulfone, nylon, or polyvinylidene fluoride 0.2 to 1.2 micrometer inline filter. Flush IV line at the end of the infusion	8 hrs	24 hrs	Infuse over 30 minutes via an IV line containing a sterile, nonpyrogenic, low protein binding polyethersulfone, nylon, or polyvinylidene fluoride 0.2 to 1.2 micrometer in-line filter. Flush IV line at the end of the infusion. Do not administer other medications through the same IV line. Monitor for signs/symptoms of infusion-related reactions.	
Obinutuzumab (Gazyva)	RTU 1,000 mg/40 mL		100 mg dose (cycle 1, day 1 dose): 4 mL of drug added to		CLL: starts at 25 mg/hr. (1st dose), up to max 400 mg/hr.	Immediate Use	24 hr.	Do not administer IV push or as a bolus May use PVC or non-PVC tubing/bag	Lexicomp
REF			NS 100 mL 900 mg dose (96 mL of drug) in cycle 1, day 2 dose) or 1,000 mg dose (cycle 1 day 8 and 15) and all subsequent cycles): NS 250 mL		NHL: starts at 50 mg/hr. (1st dose), up to max 400 mg/hr. (Rate of administration depends on indication and/or protocol; refer to specific references)			Do not shake, do not Freeze Administer through dedicated iv line (do not mix with other medication) Do not use D5W	
Ocrelizumab (Ocrevus)	RTU 300 mg/10 mL	withdraw 10 mL from	300 mg: 250 mL NS 600 mg: 500 mL NS		Initial 2 doses (300 mg): 30 mL/hr., increase by 30 mL/hr. q30 min to maximum		≤ 24 hrs. + additional 8 hr. at RT	Administer diluted solution through a separate IV line using a 0.2- or 0.22-mcm inline filter	Lexicomp
REF		600 mg vial: withdraw 20 mL from vial into 250 mL NS			180 mL/hr. Subsequent doses (600 mg): 40 mL/hr., increase by 40 mL/hr. q30 min to maximum 200 mL/hr.				

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	diucit	concentration	uurauon/rate	RT	REF	requirement)	
Ofatumumab (Arzerra) REF	RTU 100 mg 1,000 mg	20 mg/mL	1,000 mL	2 mg/mL	Incremental (rate of administration depends on indication and/or protocol; refer to specific regimen)	Immediate use	24 hr.	Do not administer IV push, IV bolus, or as a subcutaneous injection Protect from light Filter during administration with manufacturer provided filter Administer with 0.2-micron low protein binding in-line filter Do not shake Do not mix with or infuse with other medications	Lexicomp
Olaratumab (Lartruvo) REF	RTU 190 mg/19 mL 500 mg/50 mL	10 mg/mL	NS 250 mL (NS only)	N/A	60 minutes	No data	24 hr. (+ additional 4 hrs. at RT)	Do not infuse as an IV push or bolus Do not shake Allow infusion solution to reach RT prior to administration Do not co-administer with electrolytes or other medication	Lexicomp
Oxaliplatin (Eloxatin) RT	RTU 50 mg 100 mg	5 mg/mL	D5W 500 mL (may use 250 mL)	0.2-0.7 1.3 mg/mL	2 hours (Extend to 6 hrs. for acute toxicities)	6 hr.	24 hr.	Irritant with vesicant like properties Do not use needles or administration sets containing aluminum When used in combination with a fluoropyrimidine (e.g. fluorouracil), infuse oxaliplatin first	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uurauon/rate	RT	REF	requirement)	
Paclitaxel (Taxol) RT	RTU 30 mg 100 mg 150 mg 300 mg	6 mg/mL	NS or D5W 250-500 mL (Use non- DEHP bag)	0.3 – 1.2 mg/mL	1-3 hours (depending on indication/protocol; 24-hr rate has been used, associated with erratic PPT)	27 hrs.	No data	Irritant with vesicant like properties Infuse through a 0.22-micron inline filter and non-PVC administration set Sequence of administration may vary by regimen; refer to specific regimen for sequence recommendation Non-DEHP bag and tubing use Taxol primary tubing with inline filter 0.2 micron	Lexicomp
Paclitaxel nanoparticle albumin bound (nab) (Abraxane) Also known as solvent-free or surfactant-free paclitaxel	Mix 100 mg vial with 20 mL NS	5 mg/mL	Straight drug; Do not dilute (Use of non-DEHP containers or administration sets is not necessary)	N/A	30 minutes	4 hr.	24 hr.	Irritant Protect from light Do not filter Sequence of administration may vary by regimen; refer to specific protocol for sequence of administration Paclitaxel nab should be given first, followed immediately by carboplatin (NSCLC) or gemcitabine (pancreatic cancer) Inject 20 mL air into empty container (to minimize waste) is advised	Lexicomp
Pamidronate (Aredia) (non-chemo) RT	RTU 30 mg 60 mg 90 mg Or mix 30 mg vial and 90 mg vial with 10 mL SWFI	3 mg/mL 9 mg/mL	NS 500-1,000 mL	0.06-0.18 mg/mL	2- 4 hrs. (minimum 2 hrs.) Up to 24 hrs has been given	24 hr.	No data	Irritant Corrected Calcium levels should be calculated before determining dose Caution with renal impairment	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	duration/rate	RT	REF	requirement)	
Panitumumab (Vectibix) REF	RTU 100 mg 400 mg	20 mg/mL	Doses ≤ 1,000 mg: NS 100 mL Doses >1000 mg: NS 150 mL	≤ 10 mg/mL	Doses ≤ 1,000 mg: 1 st dose: 60 min Subsequent: 30 -60 min Dose > 1,000 mg: 90 min	6 hr.	24 hr.	Do not administer IV push or as a bolus Administer through a 0.2 or 0.22 micron in-line filter Doses > 1,000 mg over 90 min in NS 150 mL Do not shake Protect from light	Lexicomp
Pegaspargase (Oncospar) Asparaginase- no longer used in HHS	RTU 3750 units vial (750 units/mL, 5 mL)	750 units/mL	NS 100 mL		2 hours	48 hr.	48 hr.	Administer through 0.22-micron filter Do not administer IV push Do not filter Protect from light IM: max volume per syringe = 2 mL Do not give as IV Push	Ikesue H et al
Pemetrexed (Alimta) RT	NS (PF): 100 mg vial: 4.2 mL 500 mg vial: 20 mL	25 mg/mL	QS with NS to 100 mL	2 – 20 mg/mL	10 min	Immediate use	24 hr.	Use in-line filter	Lexicomp
Pembrolizumab (Keytruda) REF	RTU 100 mg/4 mL (4 mL)	25 mg/mL	NS 100 mL	1-10 mg/mL	30 min	6 hr.	24 hr.	Infuse through a 0.2 to 5 micron inline or add-on filter Do not shake, allow up to 5 min for bubbles to dissipate Protect from light during storage (no light protection in admin) Allow to reach RT prior to administration Do not infuse other medications through the same infusion line	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	undent	Concentration	uurauon/rac	RT	REF	requirement)	
Pemivibart (Pemgarda)									See FACT SHEET FOR HEALTHC ARE PROVIDE RS: EMERGEN CY USE AUTHORI ZATION OF PEMGARD A
Pentostatin (Nipent)	Mix 10 mg vial with 5mL SWFI	2 mg/mL	NS 50 mL	0.18-0.33 mg/mL	30 minutes	8 hr.	No data	Stable for 24 hrs. in D5W	Lexicomp
REF									
Pertuzumab (Perjeta)	RTU 420 mg/14 mL vial	420 mg/14 mL (30 mg/mL)	NS 250 mL		Loading dose (840 mg): 60 minutes	Immediate Use	24 hr.	Do not administer IV push or as a rapid bolus Incompatible with D5W	Lexicomp
REF					Maintenance dose (420 mg): 30-60 minutes			Do not shake Do not mix with other medications	
								May use PVC or non-PVC bag (regular secondary tubing) Pertuzumab and trastuzumab may be administered in any order; however, docetaxel should be given after pertuzumab and trastuzumab Protect from light (only for storage)	

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuciit	concentration	uurauom/rate	RT	REF	requirement)	
Pertuzumab, trastuzumab and hyaluronidase (Phesgo)	RTU Loading: pertuzumab 1,200 mg, trastuzumab 600 mg, hyaluronidase 30,000 units Maintenance: pertuzumab 600 mg, trastuzumab 600 mg, hyaluronidase 20,000 units	Loading: Pertuzumab 80 mg, trastuzumab 40 mg, and hyaluronidase 2,000 units per mL (15 mL) Maintenance: Pertuzumab 60 mg, trastuzumab 60 mg, and hyaluronidase 2,000 units per mL (10 mL)	N/A	15 ml (loading) 10 ml (maintenance)		4 hrs. (if not used immediately)	24 hrs. (if not used immediately)	Do not administer by another route, SQ only Do not freeze Avoid unnecessary storage Do not interchange with Famtrastuzumab deruxtecan	Lexicomp
Polatuzumab vedotin (Polivy) REF	Mix each vial (140 mg) with 7.2 mL SWFI	20 mg/mL	NS, minimum 50 mL	0.72 – 2.7 mg/mL	Initial: 90 minutes (monitor for infusion reaction for at least 90 min after completion) If tolerated initial: 30 minutes for subsequent doses (monitor for at least 30 min after completion) After infusion reaction resolution: Resume at 50% of prior rate	4 hr. (in NS)	24 hr. (in NS)	Admin antihistamine and antipyretic 30-60 min prior to each infusion If infusion reaction occurs, interrupt infusion and administer supportive treatment Use line with sterile, non-pyrogenic, low-protein binding inline or addon 0.2-0.22-micron filter Gently invert bag to mix; do not shake	Lexicomp
Porfimer (Photofrin)	Mix 75 mg vial with 31.8 mL NS	2.5 mg/mL	Straight drug; Do not dilute	2.5 mg/mL	3-5 min	Immediate Use	No data	Protect from light Shake well until dissolved	Lexicomp
Pralatrexate (Folotyn) REF	RTU 20 mg 40 mg	20 mg/mL	Straight drug, do not dilute	20 mg/mL	3-5 min	Immediate Use	No data	Protect from light	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	um anom rate	RT	REF	requirement)	
Ramucirumab	RTU	10 mg/mL	NS QS to 250 mL	N/A	60 minutes	4 hr.	24 hr.	Do not administer as an IV push or bolus	Lexicomp
(Cyramza) REF	500 mg 1,000 mg				(30 min in subsequent if tolerated)			Administration via 0.22micron or less protein sparing filter (e.g. primary non-DEHP tubing with 0.2-micron filter) is recommended	
								Do not shake	
								Do not use D5W	
								Do not infuse in the same IV line with electrolytes or other medications	
								Administer prior to docetaxel, paclitaxel, or FOLFIRI if administering in combination	
Rasburicase (Elitek) (non-chemo)	Supplied diluent: 1.5 mg vial: 1 mL 7.5 mg vial: 5 mL (for cost saving, dose is	•	QS with NS to 50 mL	N/A	30 minutes	No data available	24 hr.	Do not administer as a bolus infusion Do not shake or vortex Not considered cytotoxic chemotherapy Do not use filter	Lexicomp
REF	ordered in flat dose, e.g., 3 mg, 6 mg)							Administer through a separate line if possible	
Ravulizumab-cwvz (Ultomiris)	RTU 300 mg	300 mg/30 mL	NS	5 mg/mL	Infusion rate is weight-based (refer to Lexicomp)	6 hr.	24 hr.	Allow solution to adjust to RT prior to infusion Do not shake Protect diluted solution from light	Lexicomp
Risankizumab-rzaa (Skyrizi)	Withdraw 10 mL Risankizumab from vial	600 mg/10mL in 100, 250, or 500 mL	D5W	1.2-6 mg/mL	Infuse over at least 1 hour	8 hr.	20 hr.	Allow solution to reach RT prior to infusion	Lexicomp
	and add to infusion bag or glass bottle of D5W							Do not shake	
	5							Protect from light	
								Do not freeze	

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	bility	• •	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	um auom ate	RT	REF	requirement)	
Rituximab	RTU	10 mg/mL	QS with NS to 1	1-4 mg/mL;	First Infusion:	24 hr.	24 hr.	Do not administer IV push or bolus	Lexicomp
(Rituxan)	100 mg 500 mg		mg/mL final conc. Rapid rate: mix dose in 250 mL NS	(standard: to admix as 1 mg/ mL infusion)	50 mg/hr. If tolerated, increase by 50 mg/hr. q30 min to max of 400 mg/hr.			Refrigerate	
					Subsequent Infusions: 100 mg/hr. If tolerated, increase by 100 mg/hr. q30-min to max of 400 mg/hr.				
					Rapid rate: 90 min				
Rituximab and hyaluronidase (Rituxan Hycela) REF	RTU Rituximab 1,600 mg/hyaluronidase 26,800 units (CLL) Rituximab 1,400 mg/hyaluronidase 23,400 units (DLBCL)	Rituximab 1,600 mg/hyaluronidase 26,800 units (CLL) Rituximab 1,400 mg/hyaluronidase 23,400 units (DLBCL)	N/A	N/A	SQ 5-7 min	Immediate use (within 8 hrs.)	48 hr. (+ additional 8 hrs. at RT)	Rituximab/hyaluronidase (for subcutaneous administration) and rituximab (for IV administration) are not interchangeable To avoid clogging the needle, change the needle to a 1/2-inch to 5/8-inch needle	Lexicomp
Romidepsin	Mix 10 mg vial with	5 mg/mL	NS 500 mL		4hr	24 hr.	No data		Lexicomp
(Istodax)	2mL of supplied diluent	Ū					available		
RT									
Romiplostim (Nplate)	SWFI: 250 mcg (0.72 mL) 500 mcg (1.2 mL)	500 mcg/mL	N/A	N/A	N/A	24 hr.	24 hr.	SQ: Administration volume may be small; use appropriate syringe (with graduations to 0.01 mL) for administration	Lexicomp
(non-chemo)	oss meg (1.2 m2)							Do not use BSWI	
								Do not shake	
REF								Protect reconstituted solution from light	

Generic name	Reconstitution and/or	Reconstitution and/or RTU	Standard	Final bag/syringe	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
(brand)/storage condition/special warning	RTU information	Concentration	diluent	concentration	duration/rate	RT	REF	requirement)	
Rozanolixizumab (Rystiggo)									See package insert or Lexicomp
Sacituzumab govitican (Trodelvy) REF	Mix 180 mg vial with 20 mL NS	10 mg/mL	NS 500 mL	1.1-3.4mg /mL	3 hrs. (1- 2 hrs. if well tolerated)		4 hrs. (including infusion time)	Do not administer IV push or bolus Do not administer with other medications If the required dose is >1.7 g, divide the dose equally between two 500 mL infusion bags and infuse sequentially Observe patients during for at least 30 min after each infusion	Lexicomp
Secukinumab IV(Cosentyx)									See package insert or Lexicomp
Streptozocin (Zanosar) REF	Mix 1 gm vial with 9.5 mL NS	100 mg/mL	NS 50-250 mL	100 mg/mL	Over at least 15 min to 6hrs (may give as raid IV push)	No data (6 hrs. per USP 797 beyond use date data)	No data	Refrigerate Vesicant	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	oility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	Concentration	uui auoii/1 ate	RT	REF	requirement)	
Tafasitamab (Monjuvi) REF	Mix 200 mg with 5 mL SWFI	40 mg/mL	NS 250 mL	2-8 mg/mL	Infuse initial infusion at 70 mL/hr for the first 30 min, then increase rate so infusion is administered within 1.5- 2.5 hrs Infuse subsequent infusions over 1.5-2 hrs	12 hrs.	18 hrs.	Protect from light during storage Gently swirl vial until completely dissolved; do not shake or swirl vigorously Complete dissolution may take ~5 minutes Determine the volume of reconstituted solution needed for the dose; remove a volume equal to the required dose from a 250 mL NS infusion bag and discard it Do not administer other medications through the same infusion line No incompatibilities have been observed between tafasitamab and infusion sets made of polyurethane (PUR) or PVC	Lexicomp

Frozen Stored in ultra-low freezer at -90°C to -70°C (-130°F to -					N/A (intralesional	RT After thawing,	REF Thawed vials	requirement) Protect from light	
Aherparepvec Imlygic) Frozen Stored in ultra-low freezer at -90°C to -70°C (-130°F to -		PFU/mL 10 ⁸ (100 million)					Thawed vials	Protect from light	
94°F)						light and store in original vials no longer than the specified times below:	are stable REF , when protected from light: 10 ⁶ (1 million) PFU/mL: 24 hrs 10 ⁸ (100 million) PFU/mL: 7 days		Lexicomp
ntrathecal; IV= intrave	ostatic water for injection; C. mously; N/A= not applicable U = ready to use; SWFI = ste	; NHL= non Hodgkin	lymphoma; $NS = 0$. $SQ = $ subcutaneously	9% sodium chloride; PF	= preservative free; PP	ethylhexyl phth F = precipitate;	alate; D5W = De Pt(s) = patient(s)	xtrose 5% in water; IM = intramuscular; PVC = polyvinyl chloride; REF = refri	g

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unten	concentration	uur auton/rate	RT	REF	requirement)	
Teclistamab (Tecvayli)	30 mg/3mL vial	30 mg/3mL 153 mg/1.7mL	N/A	N/A	SQ 5 min	20 hr	20hr	Doses requiring >2 mL total volume should be equally divided into multiple syringes	Lexicomp
								Do not shake vials	
								SQ: Allow to warm to RT for 15 min	
Temozolomide (Temodar)	Mix 100 mg vial with 41 mL of SWFI	2.5 mg/mL	Straight drug; Do not dilute	2.5 mg/mL	90 minutes	14 hr. (must be completed	No data	Bring vial to room temperature RT before prior to reconstitution	Lexicomp
REF						within 14 hrs. of reconstitution)		Do NOT shake Do not administer other medications through the same IV line	
Temsirolimus (Torisel)	Mix 1.8 mL of supplied diluent to 30 mg/1.2 mL vial	10 mg/mL	NS 250 mL (Use non-DEHP bag)	N/A	30 - 60 minutes	6 hr.	No data	Protect from light during storage, preparation, and handling Protect from light	Lexicomp
REF	(Do not add undiluted temsirolimus to aqueous solution)							Must use non-PVC bags and tubing, use primary Taxol tubing with inline filter (0.2 2 micron) Avoid excessive shaking	
Teniposide (Vumon)	RTU 10 mg/mL (5 mL)	10 mg/mL	NS 250 – 500 mL (Use non-DEHP bag)	0.1 mg/mL 0.2 mg/mL 0.4 mg/mL 1 mg/mL	30 - 60 minutes	≤ 0.4 mg/mL- 24hr > 0.4 mg/mL-	Do not REF	Do not administer by rapid IV injection	Lexicomp
REF			QS with NS to final concentration of 0.1 mg/mL, 0.2 mg/mL, 0.4 mg/mL, or 1 mg/mL			4hr		Irritant	

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uurauom/rate	RT	REF	requirement)	
Thiotepa	SWFI:	~10 mg/mL	IV: NS only 50 mL	IV: 0.5 -1 mg/mL	IV: 15 minutes	IV: 4 hr.	24 hr.	Irritant	Lexicomp
(Thioplex)	15 mg vial: 1.5 mL 30 mg vial: 3mL	(actual concentration = 10.4 mg/mL due to overfill)	(if dose < 250 mg)	(consult protocol if high dose is used)	Intravesical: 1-2 hrs.	Intravesical: immediate use, no data		Use 0.22-micron filter during preparation	
REF	100 mg: 10 mL	overm)	IT: PF NS 10 mL	IT: 1 mg/mL		IT: 4 hrs. (per USP 797)		Intravesical: Instill directly into the bladder and retain for 2 hours	
			Intravesical:	Intravesical: 1 – 2				For IT doses use PFNS	
			NS 30 - 60 mL	mg/mL				Contains N, N-dimethylacetamide, which may be incompatible with some closed system transfer devices (CSTDs), consult CSTD manufacturer	
				Irrigation: 1 mg/mL					
Tocilizumab (Actemra)	RTU 80 mg/4 mL 200 mg/ 10 mL	20 mg/mL	Children <30 kg: NS 50 mL total or children ≥30 kg and adults: NS 100 mL total	N/A	60 minutes	24 hr.	24 hr.	Do not administer IV push or IV bolus Allow diluted solution for infusion to reach RT prior to administration	Lexicomp
(non-chemo)			(must withdraw equal volume of NS					Protect from light storage Infuse through dedicated IV line	
REF			for the volume of drug required)					and through country in the	
Tofersen (Qalsody	RTU 100 mg/15 mL	6.7 mg/mL	Do not dilute		injection over 1 to 3 minutes	immediately (within 4 hours of removal from the vial)	n/a	External filters are not required	Package insert
Topotecan	RTU	1 mg/mL	NS 50 mL	N/A		Lyophilized powder:	IV: 7 days	Irritant	Lexicomp
(Hycamtin)	4 mg/4 mL (4 mL)					IV: 24 hrs.		Protect from light	
RT	PF form is available for IT route					IT: 4 hrs. (or per USP		For IT route, use PF NS	
						797 beyond use date guideline)			

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	Concentration	uui auoii/1 atc	RT	REF	requirement)	
Trabectedin (Yondelis) REF	Mix 1 mg vial with 20 mL SWFI	0.05 mg/mL	NS 500 mL (inpatient use) If ordered as CADD pump for outpatient use: must order administration set 21-7394-24 directly from InfusSystem to attach to NS 500 mL bag, no cassette is required		Overien concer	Complete infusion within 30 hrs. of drug reconstitution	No data available	Vesicant Prime in non-PVC, non DEHP bag with polyethersulfonate (e.g. primary paclitaxel tubing with filter) inline 0.22-micron filter Shake until completely dissolved Combination therapy with doxorubicin liposomal: Administer doxorubicin liposomal first (flush line with D5W) then follow with trabectedin infusion Do not mix with other medications	Lexicomp
Trastuzumab (Herceptin) Sound-alike, look-alike warning: do not confuse with ado- trastuzumab emtansine (Kadcyla) REF	Mix 150 mg SDV with 7.4 mL SWFI Or mix 420 mg MDV with 20 mL BSWFI (use SWFI if pt. has hypersensitivity to benzyl alcohol) Trastuzumab intrathecal or Ommaya: do not use manufacturer supplied diluent, use SWFI	21 mg/mL	IV: NS 250 mL IT: SWFI (PF) 6 mL	N/A	Loading dose: 90 minutes Maintenance dose: 30 minutes (if tolerated)	Immediate Use	IV: 24 hr. IT: 8hr	Do not administer IV push or by rapid bolus Do not shake Do not administer with D5W Reconstituted vial stable for 28 days in fridge For IT doses, use PF SWFI Do not mix with any other medications	Lexicomp
Trastuzumab-anns (Kanjinti) Sound-alike, look- alike warning: do not confuse with trastuzumab (Herceptin)	Mix 150 mg SDV with 7.4 mL SWFI	21 mg/mL	IV: NS 250 mL	N/A	Loading dose: 90 minutes Maintenance dose: 30 minutes (if tolerated)	Immediate Use	IV: 24 hr.	Do not administer IV push or by rapid bolus Do not shake Do not administer with D5W Do not mix with any other medications	Lexicomp

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concenti ation	durauon/rate	RT	REF	requirement)	
Trastuzumab-???? (Ogivri)									
Sound-alike, look- alike warning									
Trastuzumab and hyaluronidsae (Herceptin Hylecta)	RTU trastuzumab 600 mg and hyaluronidase 10,000 units per 5 mL (5 mL) Using transfer needle, withdraw solution from vial into syringe	Trastuzumab 600 mg and hyaluronidase 10,000 units per 5 mL (5 mL)	N/A	N/A	Admin SQ over 2-5 min	4 hr.	24 hr.	For SQ use only Alternate injection site between left and right thigh (at least 2.5 cm from previous site) Do not admin other SQ medications at same sites	Lexicomp
								Store in original carton to protect from light Do not shake	
Tremelimumab (Imjudo)	Withdraw appropriate tremelimumab dose volume from the vial (do not use if cloudy, discolored or if visible particulates are present) and transfer to IV bag containing NS or D5W	300 mg/15 mL (15 mL) OR 25 mg/1.25 mL (1.25 mL, 15 mL)	NS or D5W is acceptable	Final concentration of 0.1 to 10 mg/mL The final concentration of the diluted solution should not exceed 10 mg/mL	Infuse IV over 60 min through IV line	24 hr.	24 hr.	For IV use only low-protein binding 0.2- or 0.22-micron filter Do not freeze Do not shake	Lexicomp
Valrubicin (Valstar) REF	RTU 40 mg/mL (5 mL) (vials must be thawed at RT for 30 min prior to	40 mg/mL	800 mg dose (20 mL) added to NS 55 mL(or QS to 75 mL) (Use non-DEHP bag)	14.5 mg/mL	N/A 2 hr.	12 hr.	No data available	For intravesical use only Allow vials to slowly warm to RT prior to use Use non-PVC containers and administration sets	Lexicomp
	admixing)							Retain in the bladder for 2 hours, then void/drain Do not mix with other drugs	

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Sta	bility	• •	Ref
condition/special warning	RTU information	Concentration	unuent	Concenti ation	uurauon/rate	RT	REF	requirement)	
Vedolizumab (Entyvio)	Mix 300 mg vial with 4.8 mL SWFI	60 mg/mL	NS 250 mL	N/A	30 min	12 hr. (in NS)	24 hr. (in NS)	Do not administer by IV push or bolus	Lexicomp
(non-chemo)	(results in 5 mL after dissolving)							Do not vigorously shake or invert Do not use the vial if the drug product is not dissolved within 30 minutes	
REF								Flush with NS 30 mL after infusion	
Vinblastine	RTU	1 mg/mL	NS 50 mL	1 mg/mL	15 min		6 hrs. (per USP	FATAL IF GIVEN INTRATHECALLY	Lexicomp
(Velban)	10 mg					Use	797) (21 days in one	Vesicant	
REF							study)	Dispense in a minibag (NOT in a syringe)	
								Protect from light	
Vincristine (Oncovin)	RTU 1 mg	1 mg/mL	Intermittent infusion:	1 mg/mL	15 min	Immediate use	6 hrs. (per USP 797)	FATAL IF GIVEN INTRATHECALLY Vesicant	Lexicomp
(Oncovin)	2 mg		NS 50 mL					Dispense in a minibag (NOT in a	
REF			Continuous infusion:					Syringe)	
			NS 500 mL NS (as in infusional EPOCH)						
Vincristine liposomal	See 25-step process in drug package insert	0.16 mg/mL	NS or D5W 100	N/A	60 minutes	12 hrs.	No data		Lexicomp
(Marqibo)	5 mg/31 mL		mL					Do not administer IV push or bolus	
***************************************	o mg or min							Irritant	
REF								Do not use with in-line filters	
								Follow package insert for admixing and calibration	

Generic name (brand)/storage	Reconstitution and/or	Reconstitution and/or RTU	Standard diluent	Final bag/syringe concentration	Standard infusion duration/rate	Stal	bility	Special precautions/filter	Ref
condition/special warning	RTU information	Concentration	unuent	concentration	uurauon/rate	RT	REF	requirement)	
Vinorelbine (Navelbine) REF	RTU 10 mg 50 mg	10 mg/mL	NS 50 mL	0.5 – 2 mg/mL	6-10 minutes (followed by NS 75- 125 mL)		No data	FATAL IF GIVEN INTRATHECALLY Vesicant	Lexicomp
Zoledronic Acid (Zometa) (non-chemo) Sound alike look-alike warning: do not confuse with zoledronic 5 mg vial (Reclast) RT	RTU 4 mg/5 mL (5 mL) Premixed bag is available: 4 mg/100 mL bag	0.8 mg/mL	NS 100 mL	N/A	15-30 minutes	Immediate use		Allow solution to reach RT before administration Infuse in a line separate from other medications Caution in patients with renal impairment	Lexicomp

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Appendix A

To prepare a 0.2 mg/mL (0.02%) or 0.4 mg/mL (0.04%) mitomycin eye drop:

- (1) Reconstitute 20 mg vial of mitomycin with 40 mL SWI to give a concentration of 0.5 mg/mL
- (2) For 0.2 mg/mL (0.02%): transfer 6 mL (3 mg) to a sterile 15 mL eye dropper bottle Then add 9 mL SWI to the eye dropper bottle to give final conc. of 0.2 mg/mL (0.02%)
- (3) For 0.4 mg/mL (0.04%): transfer 12 mL (6 mg) to a sterile 15 mL bottle Then add 3 mL SWI to the eye dropper bottle to give final conc. of 0.4 mg/mL (0.02%)

Appendix B

Compounding of TACE with doxorubicin liquid formulation

Dose of doxorubicin	# of doxorubicin liquid vial (2 mg/mL) needed	# of LC (or DC) beads (2 mL each vial)	Loading time required for doxorubicin liquid with LC beads 100-300 µm
50 mg	1	1	5 hours
75 mg	2	1	7 hours
100 mg	2	2	No data (presumed > 10 hours)
150 mg	3	2	No data (presumed > 10 hours)

Disclaimer: Follow most recent Houston Methodist guideline whenever there is discordance