

| DRUG | 'Spread' & Dilute | Localise & Neutralise | Management (see Table) |
|-----------------------------|---|--|------------------------------------|
| Aclarubicin | No | Yes | А |
| Aldesleukin (Interleukin 2) | Yes | No | В |
| Amsacrine | No | Yes | A |
| Arsenic Trioxide | Yes | No | General |
| Bleomycin | Yes | No | В |
| Carboplatin | Yes-after 1st 24hrs | Yes-during initial inflammatory reaction | С |
| Carmustine | No | Yes | D |
| Chlormethine Hydrochloride | No | Yes | E |
| Cisplatin | Yes-for treatment administered within 24hrs | Yes-for treatment commenced 24hrs post extravasation | F (see additional comments) |
| Cladribine | Yes | No | В |
| Crisantaspase | Yes | No | В |
| Cyclophosphamide | Yes | No | В |
| Cytarabine | Yes | No | В |
| Darcarbazine | No | Yes | A |
| Dactinomycin | No | Yes | A |
| Daunorubicin | No | Yes | A |
| Liposomal Daunorubicin | No | Yes G then A | |
| Docetaxel | Yes | No | (see additional comments) H |
| Doxorubicin | No | Yes | A |
| Liposomal Doxorubicin | No | Yes | G then A (see additional comments) |
| Epirubicin | No | Yes | À |
| Etoposide | Yes | No G | |
| Etoposide Phosphate | Yes | No | G |
| Fludarabine | Yes | No | В |
| Fluorouracil | No | Yes | G |
| Gemcitabine | Yes | No | В |
| Idarubicin | No | Yes | A |
| Ifosfamide | Yes | No | В |



| α-Interferon | Yes | No | В |
|--------------|-----|-----|--------------------------------|
| Irinotecan | Yes | No | G |
| Melphalan | Yes | No | В |
| Methotrexate | No | Yes | G |
| Mitomycin | No | Yes | A |
| Mitoxantrone | No | Yes | С |
| Oxaliplatin | Yes | No | I |
| Paclitaxel | Yes | No | H (see additional comments) |
| Pentostatin | Yes | No | В |
| Streptozocin | No | Yes | A |
| Teniposide | No | Yes | G |
| Thiotepa | Yes | No | В |
| Topotecan | No | Yes | D |
| Vinblastine | Yes | No | J |
| Vincristine | Yes | No | J |
| Videsine | Yes | No | J |
| Vinorelbine | Yes | No | J |



| A | Apply topical dimethylsulphoxide at extravasation site. Once area has dried, apply hydrocortisone 1% cream followed by 30 mins cold compression. Repeat 2 hourly for the first 24 hours after extravasation. For the next 7-10 days, apply dimethylsulphoxide 6 hourly alternating with hydrocortisone 1% cream, so treatment is being applied every 3 hours on an alternating basis. Avoid contact with good skin. If blistering occurs, stop applying dimethylsulphoxide and seek further advice |
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| В | If a large volume has extravasated aspirate as much fluid as possible. Where a large volume is present in tissues, causing the patient pain, use the pin cushion technique to infiltrate the site with hyaluronidase (1500units in 2mL water for injection or sodium chloride 0.9%). Apply heat and compression to assist natural dispersal of the drug. |
| С | Aspirate as much fluid as possible. Give 100mg hydrocortisone injection via the cannula. Administer 100mg hydrocortisone by subcutaneous injection, in 0.2mL aliquots, around the circumference of the affected area. Apply hydrocortisone 1% cream and cover the affected area with an ice pack, on an intermittent basis, for first 24 hours. |
| D | Follow general procedure of management of cytotoxic extravasation. Treat with cold compression also. |
| E | Using the pin cushion technique infiltrate the area with 1-3mL sodium thiosulphate 2.98% followed by 100mg hydrocortisone injection to the infiltrated area. Apply cold compression intermittently for 12 hours. |
| F | Using the pin cushion technique infiltrate the affected area with 1-3mL of sodium thiosulphate 2.98%. Aspirate back, then give 1500units of hyaluronidase around the area. Apply heat and compression. |
| G | Give 100mg hydrocortisone injection via the cannula. Administer 100mg hydrocortisone by subcutaneous injection, in 0.2mL aliquots, around the circumference of the affected area. Apply hydrocortisone 1% cream and treat with pulsed cold compression for up to 24 hours. |
| Н | Reconstitute 100mg hydrocortisone injection and mix with 10mg chlorphenamine injection, in a volume of 10mL. Infiltrate the extravasated area with 1-3mL of this mixture as 0.2mL pin cushion subcutaneous injections. Depending on the size of the area it may not be necessary to use the whole 3mL. Large volume extravasations may need as much as 10mL. Follow this with 1500units of hyaluronidase and warm compression. Use topical antihistamine cream for 4 days. In particularly severe cases give 1g sodium cromoglycate orally as soon as possible after injury. This can be followed by oral sodium cromoglycate 200mg QID for the next 3 days. |
| I | Infiltrate the area with hyaluronidase (1500units in 2mL water for injection) using the pin cushion technique. Gently massage the area to facilitate dispersion. Treat with warm compression. Depending on the nature and severity of the extravasation the medical team should consider the following: prescribe high dose oral steroids (dexamethasone 8mg BD for 2-3 days), prescribe oral analgesia (e.g. diclofenac SR 75mg BD) and consider a PPI. Consider referral to Plastic Surgery and/or Physiotherapy. |
| J | Infiltrate the area with hyaluronidase (1500units in 2mL water for injection or sodium chloride 0.9%), in 0.2mL aliquots, over and around the circumference of the affected area. Treat affected area with warm compression for first 24 hours. For the next 7 days apply a non-steroidal anti-inflammatory cream to the affected area, QID. |



| Cisplatin | As an intact molecule, cisplatin causes few problems when extravasated. Problems arise when it is left untreated. Within 4 to 6 weeks of an acute event a subcutaneous deposit of platinum precipitates in the tissues causing pain, inflammation and necrosis. Injuries not treated within 24 hours should be treated with intermittent cold compression and managed symptomatically. |
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| Liposomal Daunorubicin & Liposomal Doxorubicin | Whilst the drug contained within the liposome is a vesicant, the formulation offers some protection. If untreated, liposomes may be degraded in the body over the next 2-3 weeks resulting in a full-blown extravasation within the next 7 to 10 days. |
| Paclitaxel | Inflammation and soft tissue reactions at the injection site have been reported after infusion of paclitaxel. This can progress to serious necrotic injury if not treated promptly. Paclitaxel has a greater risk classification than docetaxel because of the cremophor in its formulation. Prolonged infusions should be avoided. |

Interrupt any systemic anti-cancer therapy including oral drugs until management discussed with the Acute Oncology Team, the patient's treating oncology team or Cancer Centre oncology on-call team