



# Safety Notice 012/17

## Intravenous piperacillin-tazobactam – Disruption to supply

**20 October 2017**

**Distributed to:**

- Chief Executives
- Directors of Clinical Governance
- Director Regulation and Compliance Unit

**Action required by:**

- Chief Executives
- Directors of Clinical Governance

**We recommend you also inform:**

- Emergency Departments
- Intensive Care Units
- Infectious Diseases Physicians
- Directors of Surgery
- Directors of Medical Services
- Directors of Nursing
- Directors of Pharmacy
- Drugs and Therapeutics Committees and subcommittees
- AMS Committees

**Expert Reference Group**

Content reviewed by:

- Office of the Chief Health Officer
- Chief Pharmacist Unit
- Clinical Excellence Commission
- HealthShare

**Clinical Excellence Commission**

Tel. 02 9269 5500  
Fax. 02 9269 5599

Email:  
[CEC-Quality@health.nsw.gov.au](mailto:CEC-Quality@health.nsw.gov.au)

Internet Website:  
<http://www.health.nsw.gov.au/sabs>

Intranet Website  
<http://internal.health.nsw.gov.au/quality/sabs/>

**Review date**

June 2018

**Background**

There is a disruption to the supply of piperacillin-tazobactam 4 g/500 mg intravenous preparation. This disruption has resulted from a global shortage of the active pharmaceutical ingredient (API) of piperacillin-tazobactam. This shortage affects Alphapharm and Sandoz who are the primary suppliers to the Australian market.

A previous communication regarding the shortage was sent on 13 July 2017. Sandoz has provided updated advice and anticipate that based on the utilisation rates of piperacillin-tazobactam during the previous shortage notification period:

- they will be able to continue supply for the remainder of 2017
- they will **not** be able to continue supply in the first quarter of 2018.

Therefore, hospitals must conserve stock of piperacillin-tazobactam for January, February and March 2018 by implementing or tightening measures to control use of piperacillin-tazobactam.

**Further information**

For details of supply options, including Special Access Scheme (SAS) products, email: [HSNSW-contract902@health.nsw.gov.au](mailto:HSNSW-contract902@health.nsw.gov.au)

There is advice on alternative therapy to piperacillin-tazobactam from the National Centre for Antimicrobial Stewardship (NCAS) <https://www.ncas-australia.org/education> and the Australasian Society for Infectious Diseases (ASID) <https://www.asid.net.au/documents/item/1439>. Refer to the Australian Medicines Handbook and Therapeutic Guidelines (eTG complete) for further advice on precautions associated with alternative antibiotic choices. Both can be accessed from CIAP.

**Suggested actions by Local Health Districts/Networks**

1. Distribute this notice to all stakeholders and all clinical departments.
2. Assess the current status of piperacillin-tazobactam intravenous preparations available in each facility, ensuring all locations of stock are identified.
3. Strategies to manage the shortage of piperacillin-tazobactam should be planned and implemented at a local level by the facility/LHD Antimicrobial Stewardship (AMS) or Drug and Therapeutics Committee. These measures should include (after risk assessment at the local level):
  - Remove and quarantine stock from clinical areas where piperacillin-tazobactam intravenous preparations are not routinely used.
  - Reduce and minimise stock levels from clinical areas where piperacillin-tazobactam intravenous preparations are routinely used.
  - Reserve stock of piperacillin-tazobactam intravenous preparations for infective conditions where IV piperacillin-tazobactam is the only available option.
  - Review prescriptions for piperacillin-tazobactam and assess whether antibiotic therapy can be de-escalated.
  - Use alternative antibiotic options; consult your infectious diseases (ID), clinical microbiology or AMS services for advice. NCAS (<https://www.ncas-australia.org/education>) and ASID (<https://www.asid.net.au/documents/item/1439>) have released advice on alternatives.
  - Consider restricting piperacillin-tazobactam intravenous preparations to ID, clinical microbiology or AMS approval only.
4. Ensure a system is in place to document actions taken.