Alert
Not be used in infants < 4 weeks of age.
Dose is expressed as trimethoprim (TMP) component.
The Antimicrobial Stewardship Team recommends this drug is listed under the following category: Neonates: Restricted; Infants > 4 weeks of age: Oral — unrestricted and IV — restricted.

Indication
Prophylaxis of urinary tract infections (UTI).
Treatment of mild–severe infections including UTI and acute otitis media.
Prophylaxis in HIV-exposed infants

Action
Sulfamethoxazole is a sulfonamide that prevents the formation of dihydrofolic acid, a bacterial compound necessary for survival. Trimethoprim is a synthetic antibiotic that interferes with the production of folic acid by dihydrofolate reductase.

Drug Type
Sulfonamide with antifolate

Trade Name
Oral: Bactrim Oral Suspension [Roche]; Septrin Sugar Free Suspension [Aspen]
IV: DBL Sulfamethoxazole 400 mg and Trimethoprim 80 mg Concentrate Injection BP [Hospira]

Presentation
Oral liquid: Trimethoprim 8 mg/mL and sulfamethoxazole 40 mg/mL, 100 mL
IV ampoule: Trimethoprim 16 mg/mL and sulfamethoxazole 80 mg/mL 5mL ampoule

Dosage / Interval
Dosage recommendations are based on trimethoprim component.

UTI prophylaxis
PO: 2 mg TMP/kg/dose daily or 5 mg TMP/kg/dose twice weekly.

Prophylaxis in HIV-exposed infants < 6 months of age
To commence from 4–6 weeks of age at a dose of 20 mg trimethoprim once daily (not per kg basis) (equates to 2.5 mL oral liquid daily)

Treatment of mild–severe infections (e.g. UTI, acute otitis media)
Mild to moderate infections
PO: 3–6 mg TMP/kg/dose 12 hourly (AAP Guidelines 2011).
Severe infections
IV: 2–3 mg TMP/kg/dose 6 hourly.

Maximum daily dose
Route PO, IV

Preparation/Dilution
PO: Oral liquid does not require preparation

IV: Draw up 2 mL (32 mg trimethoprim and 80 mg sulfamethoxazole) and add 48 mL of sodium chloride 0.9%, glucose 5% or glucose 10% to make a final volume of 50mL with a concentration of 0.64 mg/mL of TMP.

For severely fluid restricted neonates: Draw up 2 mL (32 mg trimethoprim and 80 mg sulfamethoxazole) and add 18 mL of glucose 5% to make a final volume of 20mL with a final concentration of 3.2 mg/mL of TMP and infuse ONLY VIA A CENTRAL LINE as it is an alkaline solution. Also, follow up with a flush of up to 20 mL.

Administration
PO: Administer with feeds. Shake well before measuring dose.
IV: Infuse over 60–90 minutes. Follow-up with a flush of up to 20 mL.

Monitoring
Watch for skin reactions and blood dyscrasias.
Monitor renal function and full blood count.

Contraindications
Hypersensitivity to sulfonamides or trimethoprim.
Infants < 4 weeks of age (manufacturer says < 8 weeks).

Precautions
Use with caution in renal impairment. Dosage adjustment is required in renal impairment. Suggested adjustment (Product Info) is as follows (MIMS):

<table>
<thead>
<tr>
<th>Renal Impairment Dose Adjustments</th>
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<tbody>
<tr>
<td>CrCl (mL/min)</td>
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<tr>
<td>Above 25</td>
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</table>

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This RHW document is a modification of Neomed version. Dosage schedules remain the same. However, information on the commercial preparations not used at RHW might have been deleted. The risk rating might have been modified as per the local health district policy.
<table>
<thead>
<tr>
<th>Newborn Use Only</th>
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<td><strong>Sulfamethoxazole and Trimethoprim</strong></td>
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<table>
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<tr>
<th>Dosage Schedule</th>
<th>Recommendation</th>
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<tbody>
<tr>
<td>15 to 25</td>
<td>One-half the standard regimen</td>
</tr>
<tr>
<td>Below 15</td>
<td>Not recommended</td>
</tr>
</tbody>
</table>

Concomitant use of potassium sparing diuretics can lead to hyperkalaemia. In individuals with glucose-6-phosphate dehydrogenase deficiency, haemolysis may occur.

**Drug Interactions**
- Risk of prolonged QT interval with concurrent use of chloral hydrate, erythromycin and fluconazole.

**Adverse Reactions**
- Gastrointestinal upset (vomiting, diarrhoea).
- Severe dermatologic reactions, blood dyscrasias, hepatotoxicity.
- Prolonged use may result in fungal or bacterial superinfection.
- Prolonged QT interval, torsades de pointes, ventricular tachycardias have been reported in adults.

**Compatibility**
- Fluids: Glucose 5%, glucose 10%, sodium chloride 0.9%, sodium chloride 0.45%
- Y site: Aciclovir, amino acid solutions, amphotericin B liposomal, lipid emulsions, metronidazole, milrinone, morphine, panceuronium, piperacillin-tazobactam, vecuronium, zidovudine.

**Incompatibility**
- Y site: Amikacin, aminophylline, amiodarone, amphotericin b lipid complex, ampicillin, atropine, calcium chloride, calcium gluconate, cefazolin, cefotaxime, ceftazidime, ceftriaxone, chloramphenicol, clindamycin, dexamethasone, diazepam, diazoxide, digoxin, dobutamine, dopamine, adrenaline (epinephrine), erythromycin, fentanyl, fluconazole, folic acid, furosemide, ganciclovir, gentamicin, heparin, hydralazine, hydrocortisone, indomethacin, insulin, isoflurane, ketamine, lactated ringer’s, lidocaine (lignocaine), methylprednisolone, midazolam, multiple vitamins injection, noradrenaline (norepinephrine), benzylpenicillin, phenobarbitol (phenobarbitone), phenytoin, piperacillin, potassium chloride, propranolol, pyridoxine, ranitidine, sodium bicarbonate, tobramycin, urokinase, vancomycin.

**Stability**
- IV: Start infusion immediately after diluting – infusion must be complete within 2 hours of preparation. Monitor for precipitation, particularly with concentrated infusions.

**Storage**
- Store IV and oral preparations below 30°C. Do not refrigerate. Protect from light.

**Evidence summary**
- **Prophylaxis in vesicoureteric reflux**
  - The proportion of infants with high grade vesicoureteric reflux (VUR) among all infants with febrile UTIs is small. There is no statistically significant benefit of prophylaxis in preventing recurrence of febrile UTI/pyelonephritis in infants without reflux.\(^1\) There is benefit of prophylaxis in reducing the recurrence of infections in children with VUR but no change in renal scarring and concerns regarding multi-resistant strains among treated children.\(^2,3\)

- **Treatment duration of infections**
  - McMullan et al reviewed the evidence for minimum intravenous and total antibiotic duration in children younger than 18 years with bacterial infections, comparing shorter courses with traditionally longer durations. In many infections, especially when clinical improvement is rapid, emerging data suggest that traditional long durations of intravenous antibiotics might be unnecessary and that intravenous to oral switch can occur earlier. In most of the other infections, evidence for routine longer courses is sparse.\(^4\) In a Cochrane review of childhood lower urinary tract infection, no difference in persistent bacteriuria or recurrence was noted between 2–4 days and 7–14 days of oral antibiotics. Results from a subsequent Cochrane review showed that a single-dose antibiotic was associated with more persistent bacteriuria than was 10 days of antibiotics, although there was no difference in symptom duration or recurrence. A large retrospective study of infants younger than 6 months found no difference in treatment.
Prophylaxis in HIV-exposed infants

All HIV-exposed infants born to mothers living with HIV must receive co-trimoxazole prophylaxis, commencing at 4–6 weeks of age (or at first encounter with the healthcare system) and continued until HIV infection can be excluded.7

References