

Guideline

Management of newborns born to women with HIV (Human Immunodeficiency Virus) infection

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Applicable to	All Children's Health Queensland staff				
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Purpose

This guideline provides best practice recommendations for obstetricians, paediatricians, neonatologists and Human Immunodeficient Virus (HIV) physicians regarding management of infants born to women with HIV infection.

Scope

This Guideline provides information for all Children's Health Queensland (CHQ) employees (permanent, temporary and casual) and all organisations and individuals acting as its agents (including Visiting Medical Officers and other partners, contractors, consultants and volunteers) caring for paediatric patients.

Related documents

Procedures, Guidelines, Protocols

- [CHQ-PROC-01035 Antimicrobial Restrictions](#)
- [CHQ Antimicrobial Restrictions List](#)
- [CHQ-GDL-01221 Immunisation Guideline for Medically at Risk Children](#)

Abbreviations

AMS	Antimicrobial Stewardship
ART	Anti-retroviral treatment
BCG	Bacille Calmette-Guerin (Mycobacterium tuberculosis vaccine)
CD4	<i>Refers to CD4 or T-helper cells</i>
CHQ	Children's Health Queensland
DNA	Deoxyribonucleic acid
EDTA	Ethylenediaminetetraacetic acid
FBC	Full blood count
HIV	Human Immunodeficiency Virus
ID	Infectious Diseases specialist
IgG	Immunoglobulin G
IMPS	Infection Management Prevention Service
PCR	Polymerase Chain Reaction
PJP	Pneumocystis jiroveci pneumonia
RNA	Ribonucleic acid
QCH	Queensland Children's Hospital
VL	Viral load

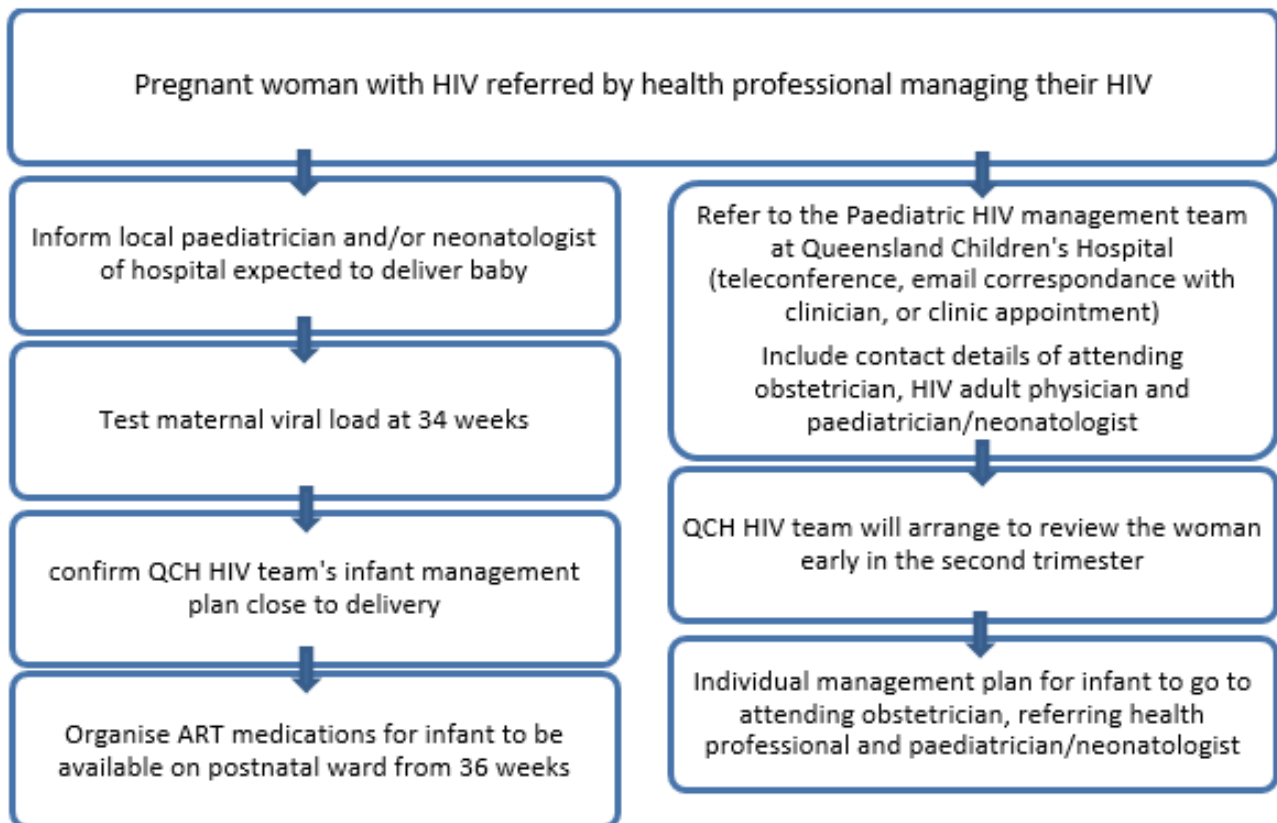
Guideline

1. Communication and assessment prior to birth of infant

- a. When a pregnant woman with HIV infection is identified:
 - Ensure referral to/attendance at Obstetrician and adult HIV physician for management (including delivery planning) for the pregnant woman.
 - Inform local Paediatrician +/-Neonatologist at obstetric unit where infant expected to deliver.
 - Refer woman to CHQ Infection Management and Prevention Service (IMPS) at Queensland Children's Hospital (QCH). Please include details of:
 - woman's antiretroviral treatment (ART);
 - most recent HIV test results (e.g. HIV viral load, HIV Proviral Deoxyribonucleic Acid (DNA) test result, CD4 (T-helper cells) count and HIV resistance tests), if available;
 - contact details for managing Obstetrician, HIV physician and Paediatrician/Neonatologist.

- To refer please complete the [CHQ Specialist Referral form](#).
 CHQ IMPS (team responsible for HIV management) will arrange to review the woman in outpatient clinic early in the second trimester and will make recommendations for neonatal testing, treatment, and follow up.
 For women residing outside of Brisbane, CHQ IMPS (team responsible for HIV management) can offer advice by teleconference or by secure email with patient's consent.
 - Ensure woman receives third trimester pertussis vaccination and seasonal influenza vaccine.
- b. CHQ IMPS will provide an individual written/emailed management plan for the infant. This will be emailed/posted to attending Obstetrician, HIV adult physician and Paediatrician / Neonatologist.
 - c. As the plan will be contingent on maternal viral loads near delivery (e.g. 36 weeks gestation), the local Paediatrician or Neonatologist should review the management plan for the neonate when woman reaches the third trimester and confirm infant management plan with CHQ IMPS. Anticipated ART medications for the infant need to be available on the postnatal ward from 36 weeks gestation onwards.
 - d. In the event of a woman presenting for obstetric management late in pregnancy (e.g. in third trimester or in labour), CHQ IMPS should be contacted urgently before infant is delivered.

Figure 1. Referral pathway



2. Recommended antiretroviral treatment (ART) for newborn and post-natal care

A standard newborn assessment should be performed at birth by the local paediatric / neonatal team. Testing of newborn and ART should be commenced as soon as possible after delivery (ideally within 4 but no later than 12 hours). ART will continue for four (4) weeks.

Discharge infant with enough oral medications to complete four (4) weeks of treatment. Please remind parent/carer to stop infant's ART at four (4) weeks of age which will be before follow up appointment at six (6) weeks of age.

Figure 2. Management of infant born to woman with HIV infection (refer to [Table 1](#) for antiretroviral treatment dosing recommendations for all age groups)

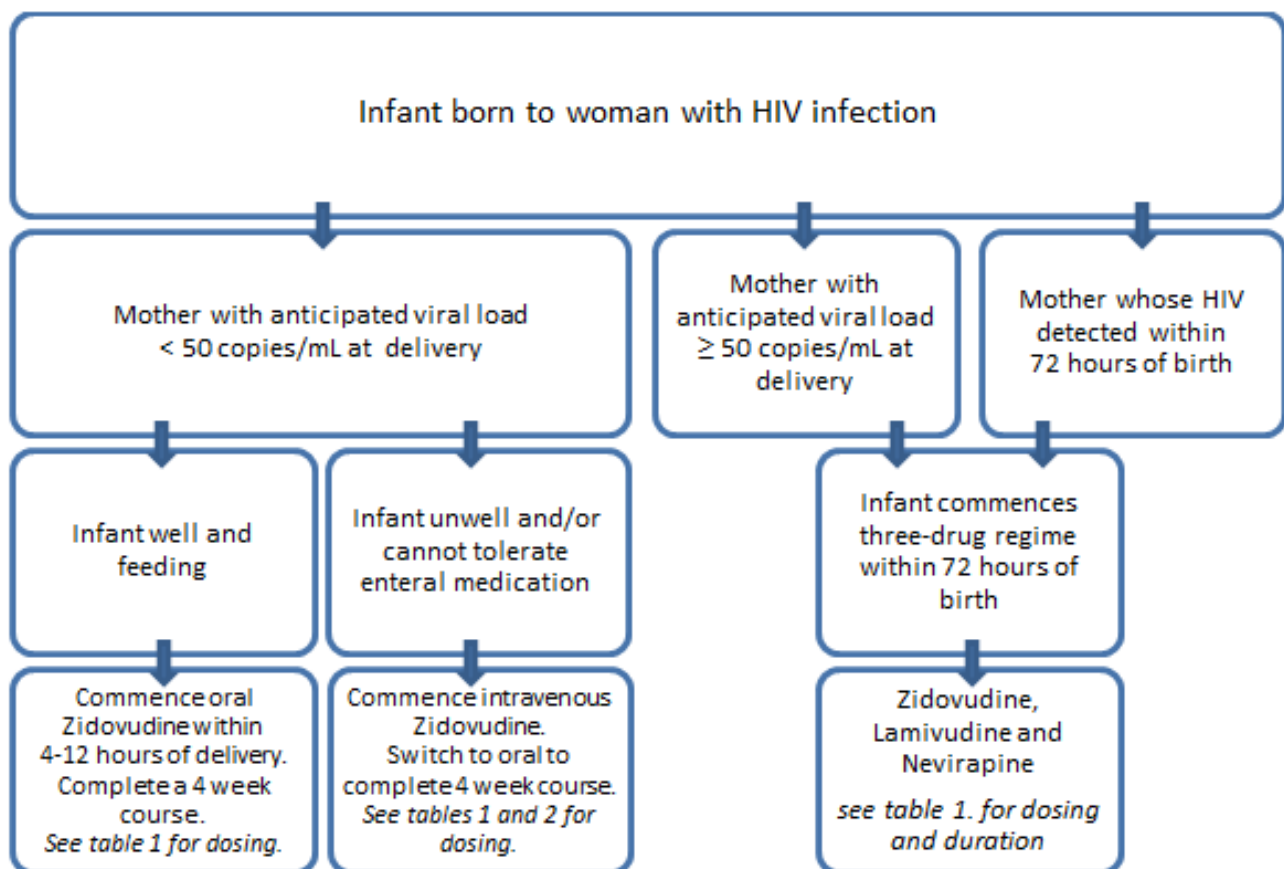


Table 1. Recommended antiretroviral treatment for infants born to HIV infected women

Criteria	Gestation	Treatment
Low risk less than 2% Infants of pregnant women who have an undetectable viral load (less than 40 copies per mL) at delivery	Infants should commence treatment as soon as possible after birth with:	
	More than 35 weeks	Commence Zidovudine 4 mg/kg/dose orally 12 hourly for 4 weeks (total treatment duration of 4 weeks).
	30 to 34 weeks	Commence Zidovudine 2 mg/kg/dose orally 12 hourly for 2 weeks then 2 mg/kg/dose orally 8 hourly for 2 weeks (total treatment duration of 4 weeks).
	Less than 30 weeks	Commence Zidovudine 2 mg/kg/dose orally 12 hourly for 4 weeks (total treatment duration of 4 weeks).
High risk less than 2% Infants born to pregnant women who have detectable viral loads at delivery OR Infants born to pregnant women who are not on therapy / present late or in labour / without testing available OR Women whose HIV infection is detected after birth (infant treatment indicated if within 72 hours of infant's birth)	Infants should commence 3-drug treatment as soon as possible after birth (and no later than 72 hours after birth if maternal HIV is diagnosed late)	
	More than 35 weeks	A. Zidovudine 4 mg/kg/dose orally every 12 hours for 4 weeks (total treatment duration of 4 weeks). B. Lamivudine 2 mg/kg/dose orally every 12 hours for 4 weeks (total treatment duration of 4 weeks). C. Nevirapine 2 mg/kg/dose, once daily for first week then 4 mg/kg/dose once daily for second week, then stop (total treatment duration of 2 weeks). If mother has received nevirapine for 3 or more days just before delivery: Neonatal metabolism of nevirapine has been induced. Hence the full dose of 4mg/kg/dose once daily should be commenced at birth and continued for 2 weeks
	30 to 35 weeks	A. Zidovudine 2 mg/kg/dose orally every 12 hours for 2 weeks then every 8 hours for 2 weeks (total treatment duration of 4 weeks). B. Lamivudine 2 mg/kg/dose orally every 12 hours for 4 weeks. (total treatment duration of 4 weeks). C. Nevirapine 12 mg per dose (or 8 mg per dose if less than 2kg) orally for 3 doses in the first week of life. Give 1 st dose within 48hrs of birth, then the 2 nd dose 48 hours after 1st dose followed by the 3 rd dose 96hrs after 2 nd dose (total treatment duration of 1 week).
	Less than 30 weeks	A. Zidovudine 2 mg/kg/dose orally every 12 hours for 2 weeks then 3 mg/kg/dose orally every 12 hours for 2 weeks (total treatment duration of 4 weeks). B. Lamivudine 2 mg/kg/dose orally every 12 hours for 4 weeks (total treatment duration of 4 weeks). C. Nevirapine 12 mg per dose (or 8 mg per dose if less than 2kg) orally for 3 doses in the first week of life. Give 1 st dose within 48hrs of birth, then the 2 nd dose 48 hours after 1st dose followed by the 3 rd dose 96hrs after 2 nd dose (total treatment duration of 1 week).

Zidovudine monotherapy is recommended if maternal transmission risk is low (woman has an undetectable viral load) even if the pregnant woman has a previous history of Zidovudine resistance.

Newborns who are unwell and/or cannot tolerate oral Zidovudine can be given intravenous Zidovudine (**Please note:** IV dose is 75 % of the oral dose administered at the same interval – refer to [Table 2](#)).

Table 2. Recommended intravenous Zidovudine dosing for pre-term and term neonates/ infants (week 1 to 4 of life) born to HIV infected women

Gestation	Postnatal age	Treatment
More than 35 weeks	Days 1 to 28 of life	Zidovudine IV 1.5 mg/kg/dose every 6 hourly
30 to 35 weeks	Days 1 to 14 of life	Zidovudine IV 1.5 mg/kg/dose every 12 hourly
	Days 15 to 28 of life	Zidovudine IV 2.3 mg/kg/dose every 12 hourly
Less than 30 weeks	Days 1 to 28 of life	Zidovudine IV 1.5 mg/kg/dose every 12 hourly

There is no intravenous formulation for Lamivudine or Nevirapine. These medications should commence as soon as possible after delivery but within 72 hours of birth and only if the newborn is able to tolerate enteral medications.

Zidovudine caution

Results of a French National Trial using Zidovudine and Lamivudine to prevent vertical transmission highlighted a concern in that two (2) out of 200 babies studied developed an extremely rare and fatal neurological disease. Similar clinical findings have not been confirmed at any other centres although the potential for nucleoside analogues to interfere with mitochondrial function is recognised. The possibility of mitochondrial toxicity must be considered in babies exposed to ART who present unwell to your service. Manifestations include lactic acidosis and disturbed neurological findings.

For information about drug interactions with ART: University of Liverpool HIV Drug Interaction Checker (www.hiv-druginteractions.org).

3. Mode of feeding

Formula feeding is recommended for all infants born to women with HIV infection.

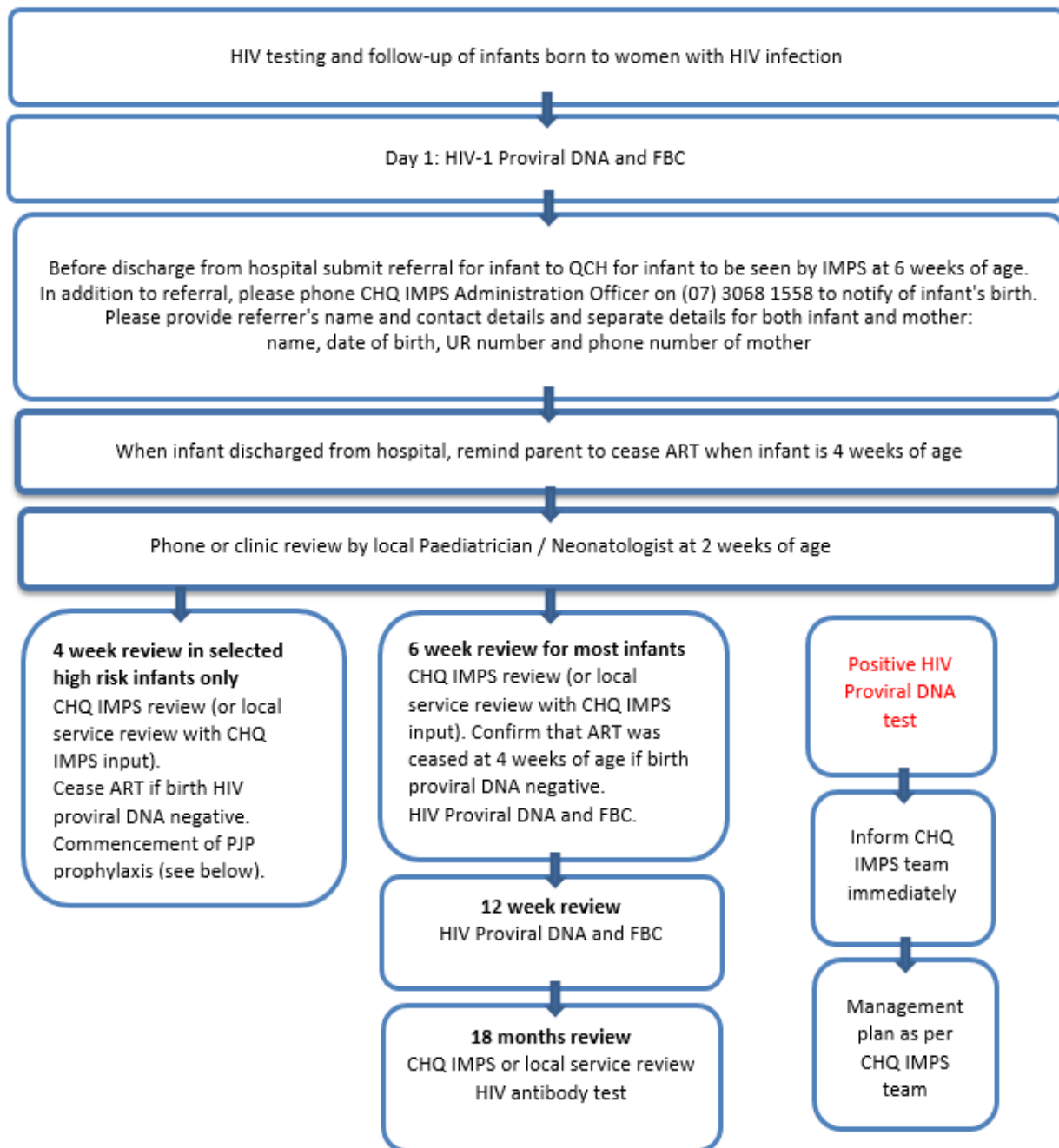
4. HIV testing and follow-up of the infant (see [Figure 3](#))

HIV testing of the infant is required on day one (1) of life. Blood must be taken from the newborn and **NOT** from umbilical cord. HIV ribonucleic acid (RNA) viral load is no longer available for the purposes of HIV diagnosis, hence testing of exposed infants is with HIV Proviral DNA.

Blood tests that are taken on day one (1) and at follow-up appointments are:

- HIV Proviral DNA (1 mL whole blood – EDTA, minimum testing required); and
- Full Blood Count (FBC).

Figure 3. HIV testing and follow-up of infants born to women with HIV infection.



A positive HIV test result within 72 hours of birth is taken as evidence of intrauterine transmission.

Babies who are not breastfed and who have a HIV Proviral DNA Polymerase Chain Reaction (PCR) test that is negative up to and including the test at three (3) months of age are not HIV infected.

All babies born to HIV positive pregnant women will have passively acquired Immunoglobulin G (IgG) antibodies to HIV. The median time to loss of antibody is 10 months, but it may be as long as 18 months.

If birth, six (6) week and three (3) month HIV Proviral DNA tests are negative, then one further test (HIV antibody only) is indicated at 18 months of age to document clearance of maternal HIV antibodies.

Yearly follow-up is recommended for the exposed but not infected child particularly in the first five (5) years of life to assess health and developmental progress.

5. *Pneumocystis jiroveci* pneumonia (PJP) prophylaxis

- In HIV infected children, PJP occurs most frequently at three (3) to six (6) months of age. Trimethoprim/Sulfamethoxazole is recommended from four (4) weeks of age **ONLY** for babies considered to have a high risk of acquiring HIV from their mother. If the HIV proviral DNA PCR at three (3) months of age is negative *Pneumocystis Jiroveci* Pneumonia (PJP) prophylaxis is discontinued.
- Trimethoprim/Sulfamethoxazole dose is 5 mg/kg/day (Trimethoprim component) (orally) in 1 to 2 divided doses daily (equates to approximately 0.63 mL/kg/day of the 8/40 mg/mL oral solution).

6. Immunisation of the infant born to woman with HIV infection

- HIV exposed babies should receive immunisations as per the National Immunisation Program. If Bacille Calmette-Guerin (BCG) vaccination at birth is indicated, it is deferred and can be given usually at three months of age if three Proviral DNA tests are negative.
- For children found to have acquired HIV infection follow QCH HIV service advice.
- HIV infected children will receive additional vaccines including yearly influenza vaccines.

7. Testing for other infections

- Babies at risk of HIV may also be at risk of other infections. If the mother is found to have Hepatitis B Virus, Hepatitis C Virus or Syphilis then the infant will need appropriate evaluation.

Consultation

Key stakeholders who reviewed this version:

- Dr Clare Nourse, Paediatric Infection Management Specialist (QCH)
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- Nicolette Graham, Pharmacist Advanced - Antimicrobial Stewardship (QCH)
- Leanne Philips, Nurse Practitioner (QCH)
- Social Worker, Paediatric HIV service (QCH)

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Guideline revision and approval history

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1.0	Infectious Diseases Consultant Team (IMPS) Infectious Diseases Fellow (IMPS) AMS Pharmacist (QCH)	Medicines Advisory Committee (QCH)	Executive Director Medical Services
2.0	Infectious Diseases Consultant Team (IMPS) AMS Pharmacist (QCH)	Medicines Advisory Committee (QCH)	Executive Director Medical Services
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Accreditation references	NSQHS Standards (1-8): Standard 3 – Preventing and Controlling Healthcare Associated Infections Standard 4 – Medication Safety