## **Hepatitis B Vaccination**

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## Hepatitis B Vaccination and Serology

- If a higher-risk for infection checking for chronic disease (Page 224 IHB)
- Post-vaccination testing, if required to confirm immunity due to being at higher risk (Table 8.6 IHB)
- Routine pre-vaccination serology not recommended





## **Hepatitis B Vaccination Schedules**

Routine

- 0, 1 and 6mths
- Children aged 11-15 years 2 doses (10mcg) with 4-6 gap

ASAP schedule

0,1 and 2mths (min 4week gap)

If serology require should be done one month post final dose





# Hepatitis B Vaccination Schedules – eligible adults

Dialysis, liver & kidney0, 1 & 6 months (40mcg)

HIV • 0, 1, 2 and 12mths (10 or 20mcg)

Other eligible • 0, 1 and 6 months (10 or 20mcg)





### Hepatitis B non-responder

- Second full course of Hepatitis B vaccine
- Serology 4 weeks later
- If still no seroconversion
- Some evidence that a course double HepA&B vaccine may correct this
- Treat as unprotected, although vaccine failure are rare eg HBIG on exposure





## **Eligibility for funded vaccines**

- All children <18 years of age irrespective of immigration status
- Adults >18 years of age
  - New Zealand residents
  - Registered refugees
  - Migrants holding a visa which provides health services i.e. 2 year working visa





Immune-suppression Hepatitis Tuberculosis Contact Pneumococcal Kidney Immune deficiency Transplant Asplenia Stem Post Pregnancy Chemotherapy Influenza Pre Non-consensual Needle-stick **Decompensation Meningococcal** 





# Additional vaccines funded for special groups

Eligibility criteria is defined in the Pharmaceutical schedule on the PHARMAC website

<u>www.pharmac.govt.nz</u> – 'Health Professionals',

'Pharmaceutical schedule', 'Vaccinations'







### Funded vaccines for special groups from 1st July 2017



Please refer to individual vaccines on the following pages for detailed eligibility criteria and to the Immunisation Handbook 2017 for vaccine administration schedules	
Asplenia — Functional or Pre- or Post-Splenectomy Immunisation Programme • Hib, influenza, meningococcal, pneumococcal, and Tdap vaccines	Influenza Immunisation Programme » Pregnancy,
<ul> <li>Chemotherapy – following</li> <li>Hib, HPV, influenza, pneumococcal, Tdap, and varicella vaccines</li> <li>Also consider immunosuppression for longer than 28 days</li> <li>Hepatitis B and meningococcal vaccines</li> </ul>	» Children aged 6 months to under 5 years who have been hospitalised for respiratory illness or have a history of significant respiratory illness,
	<ul> <li>Individuals aged 6 months to under 65 years with an eligible medical condition,</li> <li>Individuals aged 65 years or older,</li> </ul>
Cochlear implant     Hib, influenza, and pneumococcal vaccines	<ul> <li>Individuals aged under 18 years living in specified earthquake or flood affected areas in Bay of Plenty, Nelson Marlborough or Canterbury District Health Boards.</li> </ul>
Error of metabolism at risk of major metabolic decompensation <ul> <li>Influenza and varicella vaccines</li> </ul>	Influenza vaccine  Kidney disease
Haematopoietic stem cell transplantation (HSCT) – following	Hepatitis B, Hib, influenza, pneumococcal, Tdap, and varicella vaccines
Hib, HPV, influenza, meningococcal, pneumococcal, Tdap, and varicella vaccines     Also consider immunosuppression for longer than 28 days	Liver disease     Hepatitis A and varicella vaccines
Hepatitis B vaccine	Meningococcal disease case – contact with
Hepatitis A case – contact with	Meningococcal vaccine
Hepatitis A vaccine	Needle stick injury – following
Hepatitis B case – contact with	Hepatitis B vaccine
<ul> <li>Infants born to mothers who are hepatitis B surface antigen (HBsAg) positive</li> <li>Hepatitis B vaccine and hepatitis B immunoglobulin (HBIG) at birth</li> </ul>	<ul> <li>Non-consensual sexual intercourse – following</li> <li>Hepatitis B vaccine</li> </ul>
Household and sexual contacts of known acute hepatitis B cases or carriers • Hepatitis B vaccine	Pneumococcal disease – increased risk  Additional pneumococcal vaccines
Hepatitis C positive individual     Hepatitis B vaccine	Pregnancy     Influenza and Tdap vaccines in every pregnancy
HIV positive individual • Hepatitis B, HPV, influenza, meningococcal, pneumococcal, and varicella vaccines	Rubella – women of childbearing age who are not immune to rubella         • MMR vaccine
Immune deficiency/immunosuppression Individuals with an immune deficiency • Influenza, meningococcal, and pneumococcal vaccines	Solid organ transplantation Prior to solid organ transplantation • Hib, meningococcal, pneumococcal, Tdap, and varicella vaccines
Household contacts of children or adults who will be/are immunosuppressed <ul> <li>Varicella vaccine</li> </ul>	<ul> <li>Following solid organ transplantation</li> <li>Hepatitis A, hepatitis B, Hib, HPV, influenza, meningococcal, pneumococcal, and Tdap</li> </ul>
Prior to elective immunosuppression for longer than 28 days <ul> <li>Varicella vaccine</li> </ul>	vaccines
<ul> <li>Varicella vaccine</li> <li>Following immunosuppression for longer than 28 days</li> <li>Hepatitis B, Hib, influenza, meningococcal, and Tdap vaccines</li> </ul>	Tuberculosis – infants and children aged under 5 years at risk of tuberculosis (TB) exposure • BCG vaccine
Vaccine key - PCC: tuberculacie: Hity Ucementilus influenzes ture to HDV: human nanillemeniaus: MMD: messles, mumae xuballe: Tden: totanus, diabtheria, acallular nertusie: varicella; chickenney	

Vaccine key - BCG: tuberculosis; Hib: Haemophilus influenzae type b; HPV: human papillomavirus; MMR: measles, mumps, rubella; Tdap: tetanus, diphtheria, acellular pertussis; varicella: chickenpox.

## Hepatitis B vaccine for babies born to HBsAg-positive mothers

## There is a protocol for managing infants born to hepatitis B positive mothers

### Information for health professionals

on the management of babies born to HBsAg positive mothers

Local arrangements may differ from the information suggested below. Check with your Medical Officer of Health or Immunisation Coordinator for your local arrangements or further information.

Screen all women early in pregnancy for hepatitis B carriage

**HBsAg positive** 

HBsAg negative

No further action required Discuss routine National Immunisation Schedule

**Hepatitis B** 

#### Whoever ordered the test (lead maternity carer or GP) should:

- advise and discuss this result with the woman (and her partner)
- refer the woman for HBeAg and hepatitis B virus DNA testing and discuss the results with a specialist, or refer the woman to an
  infectious diseases specialist early in pregnancy for ongoing oversight. The specialist would normally recommend women who have
  a hepatitis B viral load > 10<sup>8</sup> IU/mL treatment with an antiviral medicine in the weeks before birth to reduce the especially high risk of
  transmitting the hepatitis B virus at or before birth
- record the results in the woman's maternity notes and advise other carers in all referral/consultation/transfer of clinical responsibility documentation
- ask other family members and/or sexual partners to get their immunisation status checked by their GP, and if necessary be immunised
- provide the woman with the information leaflet (HE1402) and consent form (HE1446), and go through this information with her.

#### The lead maternity carer should:

- before the birth, discuss the need for the baby to receive the hepatitis B vaccine and hepatitis B immunoglobulin and for the Medical Officer of Health to be notified following the birth
- after the birth, obtain consent from the woman for the baby to receive the hepatitis B vaccine and immunoglobulin and consent to contact the Medical Officer of Health
- provide information to the woman about the National Immunisation Register and how immunisations are recorded on the Register (HE1327).

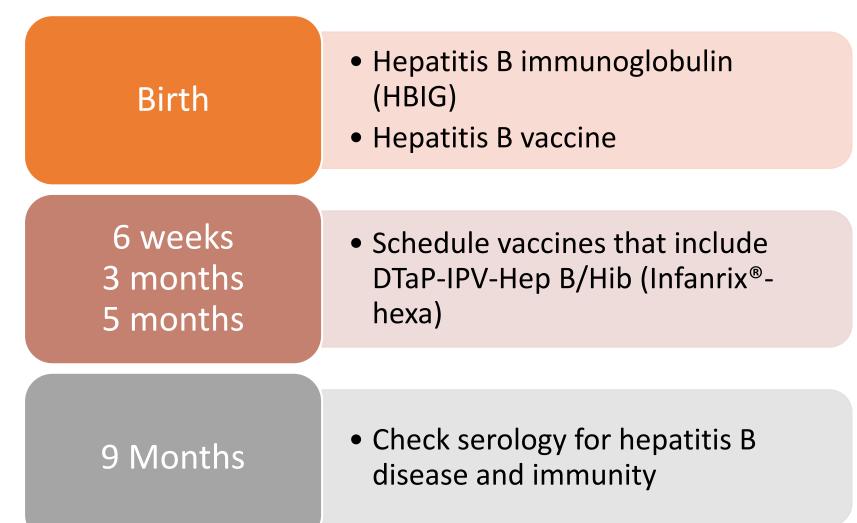
### At birth the lead maternity carer should:







# Management of a baby of a HBsAg-positive woman





Immunisation Handbook 2017 <u>www.health.govt.nz</u>

Ministry of Health resources <u>www.healthed.govt.nz</u>

IMAC website, fact sheets and videos <u>www.immune.org.nz</u>



