

# 1/Key Recommendations

These **Key Recommendations** are current as of July 2014. There are a number of websites of reputable public health organisations that also provide information on infectious diseases and infectious disease assessments in migrants including:

Public Health England. Migrant Health Guide: <http://webarchive.nationalarchives.gov.uk/20140714084352/http://www.hpa.org.uk/migranthealthguide>

TravelHealthPro: <http://travelhealthpro.org.uk/>

Centers for Communicable Disease Control and Prevention: [www.cdc.gov](http://www.cdc.gov)

DISEASE	KEY RECOMMENDATIONS
<p><b>Chickenpox/Varicella</b></p>	<p><b>Offer test to:</b></p> <ul style="list-style-type: none"> <li>• All healthcare workers (HCWs), unless known to be immune</li> <li>• Migrant women of childbearing age</li> <li>• Immunocompromised individuals and their household contacts</li> </ul> <p><b>Vaccinate non-immune:</b></p> <ul style="list-style-type: none"> <li>• HCWs</li> <li>• Non-pregnant women of childbearing age</li> <li>• Healthy close household contacts of immunocompromised individuals</li> <li>• Some immunocompromised people may be vaccinated, e.g. those with lymphocytic leukaemia in remission, transplant recipients and some children and adults with HIV infection.</li> </ul> <p><b>Offer varicella-zoster immunoglobulin (VZIG) to:</b></p> <ul style="list-style-type: none"> <li>• Non-immune women who have been exposed to varicella or zoster during pregnancy as soon as possible after exposure and ideally within 96 hours</li> <li>• Specific neonate groups (see section 5.1)</li> <li>• Specific immunocompromised individuals (see section 5.1)</li> </ul>
<p><b>Hepatitis B</b></p>	<p><b>Offer test (HBsAg and anti-HBc) to:</b></p> <ul style="list-style-type: none"> <li>• All new migrants originating from countries with a HBsAg prevalence of <math>\geq 2\%</math></li> <li>• Household and sexual contacts of identified acute or chronic cases</li> <li>• All women attending antenatal services</li> <li>• Sex workers and those who have been trafficked</li> <li>• People who inject drugs (PWID)</li> <li>• Men who have sex with men (MSM)</li> </ul> <p><b>Vaccinate:</b></p> <ul style="list-style-type: none"> <li>• All infants according to the routine childhood immunisation schedule, 6 in 1 at 2, 4 and 6 months</li> <li>• All children aged 12 months to &lt;10 years according to the “late entrants catch-up schedule”</li> </ul> <p><b>Vaccinate if non-immune (vaccination not required in anti-HBc positive):</b></p> <ul style="list-style-type: none"> <li>• All migrants originating from countries with a prevalence of <math>\geq 2\%</math></li> <li>• Children born to parents from countries with a prevalence of <math>\geq 2\%</math></li> <li>• Persons at risk of occupational exposure to blood or blood contaminated environments</li> <li>• Household and sexual contacts of persons with acute or chronic infection</li> <li>• Families adopting/fostering children from countries with a prevalence of <math>\geq 2\%</math></li> <li>• Babies born to mothers who have HBV infection (they should receive a complete course of vaccine at 0, 2, 4 and 6 months and also HBIG within 24 hours of birth and have serological testing 2 months after vaccination completed)</li> <li>• HIV exposed and HIV infected infants should be given hepatitis B vaccine at birth and then continue with the routine childhood schedule</li> <li>• Sex workers and those who have been trafficked</li> <li>• Those deemed at risk following an assessment of their health needs</li> <li>• PWID and their contacts</li> <li>• MSM</li> </ul> <p>Refer all HBsAg positive cases to specialist services for review. People who are HBsAg negative, anti-HBc positive should be referred for specialist care if they become immunosuppressed (including that due to chemotherapy or transplantation).</p>

DISEASE	KEY RECOMMENDATIONS
<p><b>Hepatitis C</b></p>	<p><b>Offer test for anti-HCV to:</b></p> <ul style="list-style-type: none"> <li>• All migrants who originate from countries with a prevalence of chronic hepatitis C of <math>\geq 3\%</math></li> <li>• Those with a history of hepatitis C risk exposure/behaviour including people who inject drugs (PWID) and men who have sex with men (MSM)</li> </ul> <p><b>Offer test for HCV RNA:</b></p> <ul style="list-style-type: none"> <li>• All those who have a positive anti-HCV result</li> </ul> <p>Refer all positive cases to specialist services for review. Vaccinate those who are non-immune to hepatitis A and/or hepatitis B with hepatitis A and/or hepatitis B vaccine.</p>
<p><b>HIV</b></p>	<p><b>Offer test for HIV Ag/Ab to:</b></p> <ul style="list-style-type: none"> <li>• All women attending antenatal services</li> <li>• All those with risk factors for HIV including but not limited to                             <ul style="list-style-type: none"> <li>◦ From high HIV prevalence countries (<math>&gt;1\%</math>)</li> <li>◦ Concurrent sexually transmitted infection</li> <li>◦ People who inject drugs (PWID)</li> <li>◦ Sex workers and those who have been trafficked</li> <li>◦ Men who have sex with men (MSM)</li> </ul> </li> <li>• Concurrent TB infection</li> </ul> <p>Refer all positive cases to specialist services for review.</p>
<p><b>Intestinal parasites/helminths</b></p>	<p><b>Offer test (ova, cysts and parasites) to:</b> Symptomatic migrants only, particularly those who have:</p> <ul style="list-style-type: none"> <li>• Lived or travelled in endemic regions</li> <li>• Migrated from Southeast Asia or Sub-Saharan Africa</li> <li>• Eosinophilia</li> </ul> <p><b>Note:</b> Healthcare professionals should also be aware that those with concurrent immunosuppression are at increased risk of developing disseminated parasitic infections, especially strongyloides, as this auto-infects and disseminates widely in those who are immunosuppressed. Note: a raised eosinophil count (<math>&gt;0.4 \times 10^9/l</math>) may be the only indication of a parasitic infection</p>
<p><b>Malaria</b></p>	<p><b>Offer test (thick and thin malaria films) to:</b> <i>Symptomatic migrants only</i>, particularly those who have:</p> <ul style="list-style-type: none"> <li>• Fever</li> <li>• Lived or travelled in malaria-endemic regions within the previous 12 months, particularly in Sub-Saharan Africa</li> </ul> <p>Refer all positive cases to specialist hospital services for review.</p>
<p><b>Measles</b></p>	<p><b>Assess all migrants for previous measles vaccination</b> <b>Vaccinate (MMR):</b></p> <ul style="list-style-type: none"> <li>• All children according to the routine childhood immunisation schedule at 12 months and 4-5 years of age (2 doses)</li> <li>• All others according to the "late entrants catch-up schedule" for children and adults, as follows:                             <ul style="list-style-type: none"> <li>◦ 12 months to 4 years, 1 dose MMR, 2<sup>nd</sup> dose at 4-5 years old</li> <li>◦ 4 years to &lt;18 years old, 2 doses MMR at one month interval</li> <li>◦ Adults aged 18 years and older, 2 doses MMR at one month interval</li> </ul> </li> </ul>
<p><b>Polio</b></p>	<p><b>Assess all migrants for previous polio vaccination</b></p> <ul style="list-style-type: none"> <li>• Be aware that acute cases of polio can present from countries where polio is endemic</li> <li>• Consider post-polio syndrome in patients who may have been infected in childhood</li> </ul> <p><b>Vaccinate:</b></p> <ul style="list-style-type: none"> <li>• All children according to the routine childhood immunisation schedule, 6 in 1* at 2, 4 and 6 months with a booster dose at 4-5 years old</li> <li>• All others according to the "late entrants catch-up schedule" for children and adults as follows:                             <ul style="list-style-type: none"> <li>◦ 12 months to &lt;4 years, three doses of 6 in 1* at two month intervals with booster at 4-5 years old</li> <li>◦ 4 to &lt;10 years, three doses of 6 in 1* at two month intervals with booster dose at least 6 months and preferably 3 years after the primary course</li> <li>◦ 10 to &lt;18 years, three doses of Tdap/IPV^ at one month intervals with booster dose 5 years after primary course</li> <li>◦ 18 years and older, one dose of Tdap/IPV^, followed by two doses of Td/IPV# at one month intervals</li> </ul> </li> </ul> <p>*6 in 1: DTaP/IPV/HiB/HepB ^Tdap/IPV: Tetanus, reduced dose diphtheria vaccine, reduced dose pertussis vaccine/IPV #Td/IPV: Tetanus, reduced dose diphtheria vaccine/IPV</p>

<p><b>Rubella</b></p>	<p><b>Offer test for rubella immunity to:</b></p> <ul style="list-style-type: none"> <li>• All women of childbearing age</li> </ul> <p><b>Vaccinate (MMR):</b></p> <ul style="list-style-type: none"> <li>• All children (two doses)</li> <li>• Non-pregnant seronegative women of childbearing age (one dose)</li> <li>• Non-immune healthcare workers (one dose)</li> <li>• All children and non-pregnant adults from low income countries, without documented evidence of rubella vaccination, should be offered one dose of MMR; two doses may be required to fully protect against measles and mumps</li> </ul>
<p><b>Sexually Transmitted Infections (STI)</b></p>	<p><b>Offer testing:</b></p> <ul style="list-style-type: none"> <li>• All sexually active people who are from countries with a HIV rate of &gt;1% (available from: <a href="http://apps.who.int/gho/data/node.main.622?lang=en">http://apps.who.int/gho/data/node.main.622?lang=en</a>) should be offered a full sexual health assessment. A high HIV rate in a country can be taken as an indicator of likely high rates of other STIs as well.</li> </ul> <p><b>The following sexually transmitted infections should be screened for at a minimum in sexually active asymptomatic individuals from these countries:</b></p> <ul style="list-style-type: none"> <li>○ Serology for HIV</li> <li>○ Syphilis serology</li> <li>○ Urinary nucleic acid amplification test (NAAT) for <i>Chlamydia trachomatis</i> and <i>Neisseria gonorrhoeae</i></li> </ul> <ul style="list-style-type: none"> <li>• Sexually active people who are from countries with an HIV rate of ≤1% should be offered sexual health screening as appropriate for their sexual history.</li> <li>• All people with symptoms of an STI should be offered a clinical assessment, STI testing and treatment</li> </ul> <p>Further information on what is involved in a sexual health assessment can be found at the British Society for Sexual Health and HIV website (<a href="http://www.bashh.org/">http://www.bashh.org/</a>).</p> <p><b>Offer Vaccine (Human Papilloma Vaccine – HPV):</b></p> <ul style="list-style-type: none"> <li>• Females at 12-13 years of age as part of the national vaccination programme</li> <li>• HPV vaccine may be given to females aged 9 to 26 years.</li> <li>• Vaccination with the quadrivalent HPV vaccine should be considered for HIV positive males and females from 9 to 26 years</li> <li>• Hepatitis B should be considered as per section 5.2</li> </ul> <p><b>Health Promotion</b></p> <p>All sexually active people. This should include safer sex and contraceptive advice for both males and females and information for women about cervical screening.</p> <p>Refer to STI services if more specialist services are required.</p>
<p><b>Tuberculosis</b></p>	<p><b>Risk assess:</b></p> <p>All migrants from countries where prevalence of TB disease is known to be change ≥40 cases per 100,000 population as per the national TB guidelines 2010 (see Appendix F for a list of these countries)</p> <p><b>Follow the TB disease algorithm</b></p>