


New to Travel Course – Day 1

Thursday 4th July 2019
 Written and taught by
 Jane Chiodini MSc RGM RM FFTM RCPS(Glasg)
 Queen's Nurse



Jane Chiodini
 Travel Health Specialist Nurse

1

About you



Where from?


Your job?

Your travel health experience?

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What do you hope to achieve from this course?



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What are my goals




- 1. Today** - I want you to understand the basic principles of a risk assessment, the diseases and vaccines and the resources to help put safe service together
- 2. By the end of the two days** I'd like you to go away feeling more 'in control' for travel health, hopefully enthused about the subject and to potentially enjoy it in the future!

Competence comes with time and experience!

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4

www.janechiodini.co.uk/education/new-to-travel/july2019/



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Agenda

- Introduction to travel medicine
- Travel risk assessment
- Travel vaccines and related issues
- Travel medicine operational issues
- Recap on resources

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Recap of the subject onlinelearning

<http://www.janechiodini.co.uk/education/online-learning/>

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An introduction to travel medicine – the key issues

- The travelling public needs to be well informed not only about their destinations and all of the cultural richness, but also **aware of the potential risks during their journey**
- Equally there needs to be a nucleus of GPs, practice nurses and other trained health professionals who are **knowledgeable about the risks on a country by country basis and who are confident about advising their patients** about each of the measures necessary to keep them healthy while travelling

Field VE, Ford L, Hill DR, eds. Health Information for Overseas Travel, National Travel Health Network and Centre, London, UK, 2010.

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Who provides travel health advice?

- In other countries its traditionally the doctor who sees the traveller and performs the risk assessment, passing them on to the nurse to administer the vaccines and give some advice
- More recently pharmacists have become involved in travel medicine, especially in Canada and the UK
- In the UK, nurses have been undertaking all aspects of travel health since the early 1990s, from risk assessment to administration of vaccines and providing risk management advice. In some cases, nurses who have obtained a non medical prescribing qualification are not only prescribing but in some circumstances setting up and owning their own travel clinics

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Why is the practice of travel medicine different in the UK?

- National Health Service provides some travel vaccines free of charge – service provided in the majority of primary care settings as GPs are financially rewarded for the service
- Pressure on GPs with their workload so historically, they passed travel health on to the nurses, but now pharmacists are getting very involved as well, with private clinics are growing dramatically
- Some surgeries are ceasing the provision of a travel service – however, they are NOT allowed to do this unless they surrender the provision of the global sum which they receive for immunisation services

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www.janechiodini.co.uk/news/faqs/faq-2/

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Charging for travel vaccines in an NHS setting – covered in day 2

- Vaccines that must always be given as part of NHS provision** (hepatitis A all doses, combination A+B all doses, typhoid, combination typhoid and hep A, polio and cholera)
- Vaccines that cannot be given as an NHS service** (yellow fever, Japanese encephalitis, tick borne encephalitis and rabies for travel and more recently ACWY for travel – but see FAQ page)
- Vaccines that can be given as NHS or private service** (hepatitis B)

Note: Cholera and oral typhoid vaccines are now only NHS vaccines in an NHS setting

see <http://www.janechiodini.co.uk/news/faqs/faq-no-2/>

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12

General Practice in the UK



Bedfordshire

UK GP Statistics 2014 ¹

9,800 practices

- 7,962 in England
- 988 in Scotland
- 470 in Wales
- 363 in Northern Ireland

43,000+ GPs

General Practice Nursing in 2013 ²

14,943 full-time equivalent GP practice nurses




1. BMA Press Briefing – General Practice in the UK July 2014 (accessed January 2016)
<http://www.bma.org.uk/search?query=press%20briefing%20general%20practice%20in%20the%20uk>
 2. http://www.pulsetoday.co.uk/hot-topics/practice-news/gp-practice-nurse-numbers-grow-by-17/2009656_fullarticle

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Governance for health care professionals – working within ‘our codes’




http://www.gmc-uk.org/Good_medical_practice_English_1215.pdf_51527435.pdf
<https://www.pharmacyregulation.org/page>
http://www.nmc-uk.org/Documents/Standards/The_code-M-20100006.pdf

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A key document for those undertaking travel medicine



Includes

- History of travel medicine
- Details about the provision of a travel service
- Risk assessment
- Competencies
- Forms
- Resources

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New inclusions - Page 9

A statement is included for those who run Yellow Fever Vaccination Centres in the UK acknowledging that whilst YF training is not mandatory for all individuals administering the vaccine, both NaTHNaC and Health Protection Scotland (HPS) recommend:

all those responsible for administering YF vaccine complete the training for their own accountability and good practice

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Three levels of nurses


Competent nurse	Experienced / proficient nurse	Senior practitioner / expert nurse
See slides to follow outlining expectations	Fulfils points of competent nurse as well	Fulfils points of competent and experienced nurse as well

While there is a strong focus on the work of a registered nurse, the field of travel medicine is truly multidisciplinary and much of the information provided in this publication is equally applicable to other registered health care professionals including **doctors** and **pharmacists** who provide travel health

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For doctors, pharmacists and nurses specialising in travel medicine

Go to
<http://www.janechiodini.co.uk/about/publications/>

<http://download.journals.elsevierhealth.com/pdfs/journals/1477-8939/PIIS1477893912000671.pdf>

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Core competence for the *Competent Nurse* (or practitioner) in a travel health consultation (pages 21/23)

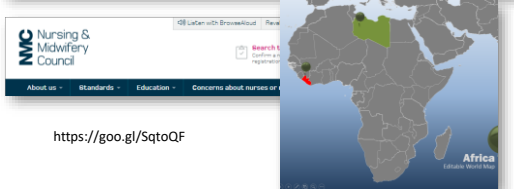
- Demonstrates good geographical knowledge
- Able to perform risk assessment effectively and understands how to interpret potential risk within a trip
- Knows where to 'go' for recommendations for travel advice, immunisations, malaria chemoprophylaxis
- Recognises limit of knowledge and knows when to refer appropriately
- Has good knowledge of common travel related illnesses e.g. TD, hepatitis, typhoid, malaria

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Search on 'travel health' at www.nmc.org.uk/

i) Documented in the patient record that the patient was going to 'Lybia' (sic.) (a reference to Libya, a low risk travel destination for contracting malaria) when in fact the patient was travelling to Liberia (a high risk travel destination for contracting malaria).



<https://goo.gl/SqtoQF>


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Core competence continued

Able to provide individual advice to the traveller

- ✓ Accident prevention
- ✓ Safe food, water and personal hygiene
- ✓ Prevention of blood-borne infections and sexually transmitted diseases
- ✓ General insect bite prevention
- ✓ Prevention of animal bites, particularly rabies including wound management
- ✓ Prevention of sun and heat complications
- ✓ Personal safety and security
- ✓ Malaria awareness, ABCD advice



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Core competence continued

- Communicates information effectively
- Prioritises in a situation when traveller is on a limited budget
- Assesses anxieties and acts appropriately
- Demonstrates an excellent vaccine administration technique
- Completes patient and administrative records after vaccination

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Education and Training – page 23

- Demonstrates evidence of learning to apply skills and knowledge in the field of travel medicine. For example, minimum of 15 hours of relevant learning plus mentorship in clinical skills before undertaking a travel consultation alone
- Ensures travel health knowledge is always up to date
- Attends an annual travel health update study session/conference at a local, national or international event

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Agenda

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Risk Assessment & Management in Travel Health



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Aims and Objectives of this session

- To understand what risk assessment is
- To appreciate the elements of the risk assessment process
- To have a good understanding of the required knowledge and resources needed to perform a risk assessment
- To be able to apply these skills at the end of the course

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Risk assessment

No travel health consultation should take place without conducting a travel risk assessment and documenting the information.

The assessment forms the basis of all subsequent decisions, advice given, vaccines administered and the malaria prophylaxis advice that is offered.

This takes time to perform correctly, and for best practice practitioners should leave sufficient time.



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Time is the major constraint

From the RCN Guidance – page 18

The main consideration is to allocate sufficient time to perform the risk assessment. It would be **unsafe to only allow 10 - 15 minutes** for a new travel appointment.

A 20-minute consultation appointment per person should be allowed to exercise best practice. Travellers with more complex needs such as backpackers or individuals requiring malaria prevention advice relevant to their destination - **may need even longer** consultation time.

The Nursing and Midwifery Council 'Code' is about being professional, about being accountable and about being able to justify your decisions; employers need to respect the complexity of a travel consultation and appreciate that sufficient time must be allowed for nurses to abide by the Code.

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What is risk?



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Would you enjoy this?



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The best way to manage/conduct a risk assessment – one option

Travel risk assessment form completed prior to appointment by traveller

Travel risk assessment form reviewed by travel health adviser

Management of the travel risks discussed with the traveller by the travel health adviser and conclusions reached

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Example of risk assessment form for information recording

Available to download from my 'Tools' page – item no. 1 <http://www.janechiodini.co.uk/tools/>

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Information to be gathered

Traveller information	Traveller's itinerary
<ul style="list-style-type: none"> Age and sex Medical history, past and present Current health status Medication Allergies to drugs and food Previous experience travelling Current interest and knowledge of health risks Previous vaccine history Any special needs 	<ul style="list-style-type: none"> Destinations (s) Date of departure Duration of stay Mode of transport Purpose of trip and planned activities Quality of accommodation Financial budget Healthcare standards at destination Relevant comprehensive insurance provision

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Risk assessment exercise

- Beckham is 10 years old and is travelling to Angola in the summer holidays to stay with his grandparents for 8 weeks
- What are the issues and risks when assessing this traveller?

What if Beckham had been a girl, is there anything else you might consider?

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Risk management

Having performed a risk assessment the risks identified are managed by individualised advice

<ul style="list-style-type: none"> Medical preparation Journey risks Safety risks Environmental risks Food and water borne risks 	<ul style="list-style-type: none"> Vector borne risks Air borne risks Sexual health and blood borne viral risks Skin health Psychological health
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Field VF, Ford L, Hill DR, eds. Health Information for Overseas Travel, National Travel Health Network and Centre, London, UK, 2010.

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What does performing a risk assessment achieve?

It enables you to give:

<p>Appropriate travel health risk advice</p>	<p>Appropriate travel vaccines for travel plans</p>	<p>Appropriate malaria prevention advice</p>
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To perform and provide evidence of best practice

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But what is risk assessment all about?

A very individual process also influenced by the traveller's personal perception of risk

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Booking process and patient expectations

How is the trip booked?

- Travel agent
- Online travel site
- Self organised trips

Patient issues

- Visiting the travel clinic for advice in good time!
- Often annoyance at the risk assessment process
- Focus on the injections with limited understanding of other risks

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Vaccine preventable health risks to travellers abroad ?

In reality, the diseases below are uncommon in travellers, usually occurring less than 1 case per 1,000 overseas visits

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The role of vaccination

Nevertheless, vaccination is one of the most important public health interventions for global infectious disease control and offers protection for travellers at risk of exposure

Field VF, Ford L, Hill DR, eds. Health Information for Overseas Travel, National Travel Health Network and Centre, London, UK, 2010.

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Estimated incidence per month of vaccine preventable diseases in lower-income countries among non-immune Western travellers

Epidemiology: Morbidity and Mortality in Travellers in Travel Medicine 4th Edition, Eds. Keystone et al. Elsevier 2019

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Cause of mortality in travellers

Hargarten SW et al. Ann Emergency Med 20:622-626, 1991
This slide was adapted from the ISTM slide set – Introduction to travel medicine 2nd Ed. www.istm.org

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Implications of the questions we ask

Many sources to increase your knowledge and understanding of pre-travel risk assessment in more detail – including on national databases and international resources



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The following slides provide some examples
but please refer to the resources on previous slide for more information.

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Age and Sex



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Babies and small children


- Increased risk of other hazards e.g. accidents, encounters with animals – need for rabies post exposure
- Small, mobile, inquisitive toddlers, limited hygiene awareness
- Risk of illness more severe – e.g travellers' diarrhoea, malaria – requiring medical treatment abroad
- Restrictions on some choices of vaccines and malaria chemoprophylaxis

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Older travellers

- Immune systems reduced – infection risk increased
- Senses reduced
- PMH more common
- Immunisation status
- Specific problems e.g. yellow fever vaccine




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Female travellers

- Security risk
- Travelling during pregnancy / breast feeding
- Managing contraception
- Coping with menstruation



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www.mooncup.co.uk

mooncup® an innovative alternative to tampons

home
what is it?
why is it better?
is it easy to use?
which size?
price
get convinced?
how to use
faq
history
testimonials
order online
wh stockists
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newsletter
tell a friend
wholesale info
sell the mooncup
resources

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www.whizproducts.co/uk

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Male travellers

20 – 29 year old age group
at greater risk
of accidents

Ref:McInnes R, Williamson L, Morrison A. (2002) Unintentional injury during foreign travel: a review. *Journal of Travel Medicine*, 9:297-307.

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Medical History

"Cat Scan"

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Medical History

1. Past and present medical history and current health status
2. Medication
3. Allergies to drugs or food/reaction to vaccination

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Why consider the medical history ?

For example:

- Specialist advice may be needed e.g. those with severe renal or liver disease & malaria chemoprophylaxis
- Recent surgery or long term medical problems such as respiratory disease may impact of travel and fitness to fly
- Immunosuppression – some live vaccines contra-indicated, other vaccines may be less effective
- Impact on travel insurance with many medical problems
- Elderly people on regular medication need to be aware of continuing regular administration
- Establishing true anaphylaxis

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Implications of the questions we ask

Many sources to increase your knowledge and understanding of pre-travel risk assessment in more detail

The Yellow Book has a lot of information about medical history as do the National databases

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Fitness to fly

The International Air Transport Association

- Is a trade association of the world's airlines. IATA supports airline activity and helps formulate policy and standards
- Its key priority is one of safety

<http://www.iata.org/publications/Documents/medical-manual.pdf>
<https://www.iata.org/publications/Pages/medical-manual.aspx>

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Aircraft Operations

Diagnosis	Assessment by a doctor with previous medicine experience	Accept	Comments
CNS disorders (Central Nervous System)			
TIA (Stroke)	4 days or less	24 hours if stable or improving, with a normal ECG. Passenger travelling in the field & under your care should receive appropriate oxygen.	If an uncomplicated recovery has been made, further escort is not required.
Cranial trauma	28 hrs or less	24 hours if generally well	
Cranial surgery	9 days or less	10 days, clinical test of an and adequate general health.	
Cognitive impairment, Dementia	History of abnormal, unprovoked, aggressive or disturbed behaviour, discrimination, agitation or similar neurocognitive disorder, significant anxiety	100% impairment, independent function and living in the community, no significant paranoid, aggressive behaviour, hallucinations, or agitation, no change in deterioration since recent flight.	Consider support of travel companion
Gastro-intestinal			
GI bleed	24 hours or less following a bleed	2-10 days	10 days can travel if endoscopic or other clear evidence is felt has contributed to risk to indicate bleeding has ceased of healing. See also anemia.
Major abdominal surgery	9 days or less	2-10 days if uncomplicated recovery	e.g. bowel resection, open hysterectomy, gall surgery etc.
Acute abdominal pain	4 days or less	2-8 days if uncomplicated recovery	
Laparoscopic surgery (cholecystectomy)	4 days or less	2-8 days if uncomplicated recovery	e.g. cholecystectomy (gall bladder removal), gall surgery.
Intensive respiratory	28 hours or less	24 hours if gas absorbed	

<http://www.iata.org/publications/Documents/medical-manual.pdf>

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Travellers visiting friends and relatives (VFRs)

- Less likely to observe malaria chemoprophylaxis compliance
- Values and beliefs need to be explored

Chiodini PJ, Patel D, Whitty GM and Laloo DG. Guidelines for malaria prevention in travellers from the United Kingdom. London: Public Health England; November 2018

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Previous vaccine history

- In the absence of documentation, don't **assume**
- Ensure primary immunisations are up to date
- Give traveller a record of vaccines given

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ETHIOPIA

Travel Inspiration

ETHIOPIA

- Highly diverse climate
- Home to over 100 ethnic groups
- Home to over 100 ethnic groups
- Home to over 100 ethnic groups

Other features

- Home to over 100 ethnic groups
- Home to over 100 ethnic groups
- Home to over 100 ethnic groups

Destination location, altitude, climate

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Destination - political and economic situation

GOV.UK

Home > Research, Travel and Living Advice > Travel advice

March 2016

Foreign travel advice

Which country or territory are you look

Egypt: Travel Advice

<https://www.gov.uk/foreign-travel-advice>

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Departure date – season and timing

- Wet season – increases malaria risk
- Dry season – increases meningitis risk
- Last minute, still consider some vaccines e.g. hepatitis A

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Length of stay

A 3 month visit carries a malaria risk around 6 times greater than a 2 week visit*

FIGURE 4 CUMULATIVE RISK OF ADVERSE EVENTS AND OF MALARIA

Compacency can creep in during a longer visit

* Chiodini PL, Patel D, Whitty CIM and Laloo DG. Guidelines for malaria prevention in travellers from the United Kingdom. London: Public Health England; November 2018.

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Mode of transport

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Risk of accidents

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Purpose of trip and planned activities



People often seek adventure and take risks abroad they wouldn't consider when 'back home'.
People vary in their perception of risk

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Quality of accommodation

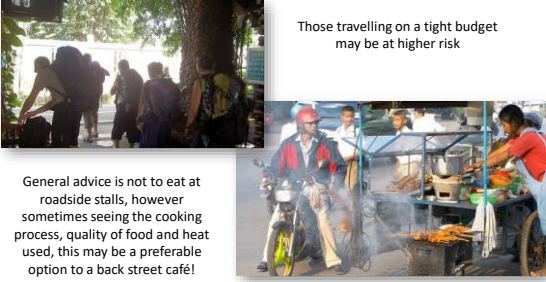


Top quality accommodation is not absolute assurance that there is no risk

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Financial budget



Those travelling on a tight budget may be at higher risk

General advice is not to eat at roadside stalls, however sometimes seeing the cooking process, quality of food and heat used, this may be a preferable option to a back street café!

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Health standards at the destination




- What are the healthcare standards like?
- The reuse of needles and syringes can be a common practice in some resource poor countries
- Carry a sterile medical kit
- Has adequate insurance been purchased?

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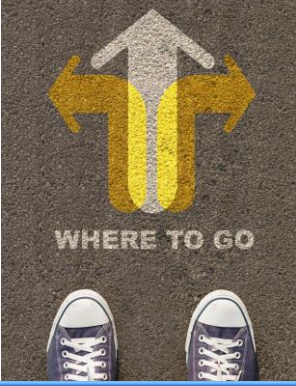
Collected the risk assessment information – then what?



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Do you know which resources we would use to make decisions?



WHERE TO GO

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Be aware of key UK resources for guidance

Immunisation against infectious disease

Guidelines for malaria prevention in travellers from the UK

Vaccine update

The Value of Vaccines

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UK National databases - www.travax.nhs.uk and for the public - www.fitfortravel.nhs.uk

TRAVAX

fitfortravel

Information on how to stay safe and healthy abroad. About Us.

Travel Health Advice

News

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75

<https://travelhealthpro.org.uk/>

TRAVEL HEALTH PRO

COUNTRY INFORMATION LATEST NEWS OUTBREAK SURVEILLANCE IN BRIEF DISEASES FROM A-Z FACTSHEETS WORLD OVERVIEW

Latest News: 28 Sep 2018
New interactive malaria maps

Searchable malaria maps have been added on to the country pages for health, travel and media.

Read more

From NaTHNaC for healthcare professionals and the general public

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76

Access via your page and the 'New to Travel' page on my website

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Home About Education Tools Help Links Contact Us

New to Travel

4th Sep and 19th August 2019 - Sunday 1st and 17th September 2018

Below are the malaria maps that are being displayed for the New to Travel page. This is provided for limited time for members of the other members of the NaTHNaC who are attending the event.

NEW TO TRAVEL - HEALTH SEARCH tool - these are shown on the 'New to Travel' page at the bottom of the home page and on the 'New to Travel' page.

<http://www.janechiodini.co.uk/tools/new-to-travel/>

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Please note, if using TRAVAX, you will still need a user name and password

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- Taking into consideration any patient specific factors (e.g. medical history, how high risk the destination is etc.) review the vaccines advised and decide what is needed – based also on previous vaccine history
- If a malarious area, also decide risk and identify appropriate chemoprophylaxis
- Consider advice required to manage the risks identified

Review the country specific information on a national database e.g. TravelHealthPro or TRAVAX

Travel health case studies

Putting theory into practice

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78


Communicating the risk and providing advice

- Providing information about vaccines sufficient to provide adequate information to obtain informed consent
- Discussion of what is necessary and desirable – taking time and cost into the equation
- Advising on malaria prevention advice and deciding with patient the most suitable chemoprophylaxis
- Delivering other appropriate travel health advice – some will need to be in written format

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Advice leaflet that can be adapted for your use –



See item no. 4 at <http://www.janechiodini.co.uk/tools/> - written in Word format for you to adapt

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Food, water and personal hygiene advice

- Always wash hands before eating or preparing food
- Boiled water, bottled water - this includes ice cubes in drinks and water for cleaning your teeth
- Only eat well cooked fresh food
- Avoid leftovers and reheated food
- Ensure meat is thoroughly cooked
- Eat cooked vegetables, avoid salads
- Only eat fruit you can peel
- Never drink unpasteurised milk and avoid ice cream
- Shellfish is a high risk food

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- Studies have shown that the “Cook it, peel it, boil it or forget it” directive is not followed by many travellers and that conflicting results have been shown in the value of such strict advice
- New thinking in travel medicine is that food and drink can be placed into three categories
 - Safe
 - Probably safe
 - Unsafe
- There is no vaccine available for travellers’ diarrhoea

Ericsson CD. Prevention of Travelers Diarrhea in: Keystone J, Freedman D, Kazarsky P, Connor B and Northrup H. Eds. Travel Medicine 3rd Edition. Saunders, an imprint of Elsevier Inc; 2013. p. 191-196

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82

Food and beverage recommendations for travellers (this poster is on your page)

Category	SAFE	PROBABLY SAFE	UNSAFE
Beverages	<ul style="list-style-type: none"> • Carbonated soft drinks • Carbonated water • Boiled water • Purified water (iodine or chlorine) 	<ul style="list-style-type: none"> • Fresh citrus juices • Bottled water • Packaged (machine-made) ice 	<ul style="list-style-type: none"> • Tap water • Chipped ice • Unpasteurized milk
Food	<ul style="list-style-type: none"> • Hot, thoroughly grilled, boiled • Processed and packaged • Cooked vegetables and peeled fruits 	<ul style="list-style-type: none"> • Dry items • Hyperosmolar items (such as jam and syrup) • Washed vegetables and fruits 	<ul style="list-style-type: none"> • Salads • Sauces and ‘salsa’ • Uncooked seafood • Raw or poorly cooked meats • Unpeeled fruits • Unpasteurized dairy products • Cold desserts
Setting	Recommended restaurants	Local homes	Street vendors

Ericsson CD. Prevention of Travelers Diarrhea in: Travel Medicine 4th Edition, Eds. Keystone et al. Elsevier 2019

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83

Food water and personal hygiene advice...
(this poster is on your page)



84

Travellers’ diarrhoea advice

- **High risk areas** include North Africa, sub-Saharan Africa, the Indian Subcontinent, S.E. Asia, South America, Mexico and the Middle East
- **Medium risk areas** include the northern Mediterranean, Canary Islands and the Caribbean Islands
- **Low risk areas** include North America, Western Europe and Australia

Management

- Rehydration
- Anti diarrhoeal tablets
- Standby emergency treatment could be an option for some

Contact medical help if the affected person has:-

- A temperature
- Blood in the diarrhoea
- Diarrhoea for more than 48 hours (or 24 hours in children)
- Becomes confused

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85

Prevention advice for hepatitis B, C and HIV infection

- Only accept a blood transfusion when essential
- If travelling to a resource poor country, take a sterile medical kit
- Avoid high risk procedures e.g. ear and body piercing, tattooing & acupuncture
- Avoid casual sex, especially without using condoms

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Malaria prevention advice - the ABCD rules !




Photo credit: James Gathany

More information on malaria on day 2 of this course

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Rabies advice

1. Do not touch any animal, even dogs and cats
2. If you are licked on broken skin, scratched or bitten in a country which has rabies, wash the wound thoroughly with soap and running water for 15 minutes then apply antiseptic.
3. Seek medical advice IMMEDIATELY, even if you have been previously immunised.

More information on rabies on day 2 of this course

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88


Prevention of accidents advice

- Avoid alcohol and food before swimming
- Never dive into water where the depth is uncertain
- Only swim in safe water, check currents, sharks, jellyfish etc.
- Avoid alcohol when driving, especially at night
- Avoid hiring motorcycles and mopeds
- If hiring a car, rent a large one if possible, ensure the tyres, brakes and seat belts are in good condition
- Use reliable taxi firms, know where emergency facilities are

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89

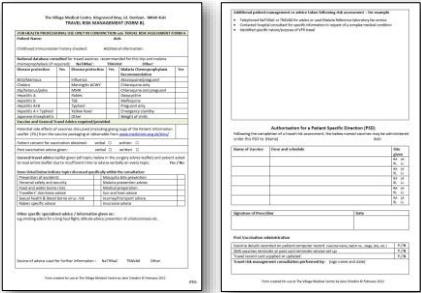
Risk management and the importance of documentation



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Risk management form helps to 'record' best practice within the travel consultation



Form can be found in "Tools" – item no. 2 <http://www.janechiodini.co.uk/tools/>

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FOR HEALTH PROFESSIONAL USE ONLY IN CONJUNCTION with TRAVEL RISK ASSESSMENT FORM A

Patient Name: _____ dob: _____

Childhood immunisation history checked: _____ Additional information: _____

National database consulted for travel vaccines recommended for this trip and malaria chemoprophylaxis (if required): NaTHNaC: _____ TRAVAX: _____ Other: _____

Disease protection	Yes	Disease protection	Yes	Malaria Chemoprophylaxis Recommendation	Yes
BCG/Mantoux		Influenza		Atovaquone/proguanil	
Cholera		Meningitis ACWY		Chloroquine only	
Dip/tetanus/polio		MMR		Chloroquine and proguanil	
Hepatitis A		Rabies		Doxycycline	
Hepatitis B		TBE		Mefloquine	
Hepatitis A+B		Typhoid		Proguanil only	
Hepatitis A + Typhoid		Yellow fever		Emergency standby	
Japanese Encephalitis		Other		Weight of child:	

Vaccine and General Travel Advice required/provided

Potential side effects of vaccines discussed (including giving copy of the Patient Information Leaflet (PIL) from the vaccine packaging or obtainable from www.medicines.org.uk/emc/)

Patient consent for vaccination obtained: verbal written

Post vaccination advice given: verbal written

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Advice leaflet with additional resources – sample leaflet available on my website

See item no. 4 at <http://www.janechiodini.co.uk/tools/> - written in Word format for you to adapt

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General travel advice leaflet given (all topics below in the surgery advice leaflet) and patient asked to read entire leaflet due to insufficient time to advise verbally on every topic: Yes / No

Items ticked below indicate topics discussed specifically within the consultation:

Prevention of accidents	Mosquito bite prevention	
Personal safety and security	Malaria prevention advice	
Food and water borne risks	Medical preparation	
Travellers' diarrhoea advice	Sun and heat advice	
Sexual health & blood borne virus risk	Journey/transport advice	
Rabies specific advice	Insurance advice	

Other specific specialised advice / information given on:
e.g. smoking advice for a long haul flight; altitude advice; prevention of schistosomiasis etc.

Source of advice used for further information: NaTHNaC TRAVAX Other

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Additional patient management or advice taken following risk assessment – for example

- Vaccine(s) patient declined following recommendation, and reason why
- Telephoned NaTHNaC or TRAVAX for advice or used Malaria Reference laboratory fax service
- Contacted hospital consultant for specific information in respect of a complex medical condition
- Identified specific nature/purpose of VFR travel

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Authorisation for a Patient Specific Direction (PSD)

Following the completion of a travel risk assessment, the below named vaccines may be administered under this PSD to:

Name: _____ dob: _____

Name, form & strength of medicine (generic/brand name as appropriate)	Dose, schedule and route of administration	Start and finish dates

Signature of Prescriber _____ Date _____

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- The template of this form could be adapted to use within a computer system, e.g. EMIS or System One
- If using paper copy of the form, then scan in after completion

Post Vaccination administration	
Vaccine details recorded on patient computer record (vaccine name, batch no., stage, site, etc.)	Y / N
SMS vaccines reminder or post card reminder service set up	Y / N
Travel record card supplied or updated:	Y / N
Travel risk management consultation performed by: (sign name and date)	


Form devised and created by Jane Chiodini © Updated May 2013

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Performing vaccination

Preparation of equipment and vaccines




Preparation of the patient

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Documenting the vaccinations

- Record of vaccines used must include the name of the drug, batch number, expiry date, site of administration and names of the administrator
- Ideally provide a written record of vaccinations given to the traveller



Vaccine Record

Variety of options now available, e.g. online, app format

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Conclusion

- No travel health consultation should take place without conducting a travel risk assessment and documenting all the information
- The assessment forms the basis of all subsequent decisions, advice given, vaccines administered and malaria prophylaxis advice that is offered
- Risk assessment and management takes time to perform correctly, and for best practice practitioners should leave sufficient time
- Good documentation is essential

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Case Study Practice



Travel health case studies

Putting theory into practice

PREV NEXT

Access via 'your page' to practice the case studies

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Agenda

- ✓ Introduction to travel medicine
- ✓ Travel risk assessment
- Travel vaccines and related issues
- Travel medicine operational issues
- Recap on resources

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Vaccine preventable diseases and related issues



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Agenda

- ✓ Travel vaccines and related issues
 - Key resources
 - Principles of vaccination and the rules
 - Range of vaccine preventable diseases and the specifics of these vaccines

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Be aware of key UK resources for guidance

Yellow Book not online – may be in your workplace and some information is being placed on the NaTHNaC website

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Key UK resources for guidance

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UK National databases - www.travax.nhs.uk and for the public - www.fitfortravel.nhs.uk

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UK National databases - www.nathnac.org

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108

Working within National Guidelines and knowing one's limitations

Telephone helplines

NaTHNaC
0845 602 6712
Monday – Friday 9am to 11.00 am and 1pm to 2pm x 2
Closed Monday and Friday at 2pm and other days at 3.30pm

TRAVAX
0141 300 1130
Mon. & Wed. 2 to 4pm
Friday 9.30 to 11.30am

MRL e mail service
Download risk assessment form from www.malaria-reference.co.uk, complete and e mail phe.malproph@nhs.net

E mail service – see malaria page, to be discussed next time

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Available from 'TOOLS' item no. 8

110

Vaccines currently available to protect our travellers

111

Diseases for consideration in this section?

PART 1
NHS vaccines (mostly) and provided in an NHS setting (hepatitis B and meningitis can be private)

- Tetanus, diphtheria and polio
- Hepatitis A
- Typhoid
- Cholera
- Hepatitis B
- Meningitis

PART 2
Always private, more specialist vaccines given by those more experienced

Covered on day 2

- Just touching on Yellow fever but separate training is required by NaTHNaC
- Rabies, Japanese B, tick-borne encephalitis

For more details regarding the charging of vaccines see FAQ no. 2 on my website <http://www.janechiodini.co.uk/news/faqs/faq-no-2/>

112

Immunisation timeline

Historical vaccine developments and introduction of vaccines in the UK

Historical information on NHS Choices – an interesting read see: <http://www.nhs.uk/conditions/vaccinations/pages/the-history-of-vaccination.aspx>

Remember – this is item above accessed no. 10 to download from the 'Help' page at <http://www.janechiodini.co.uk/news/help/>

113

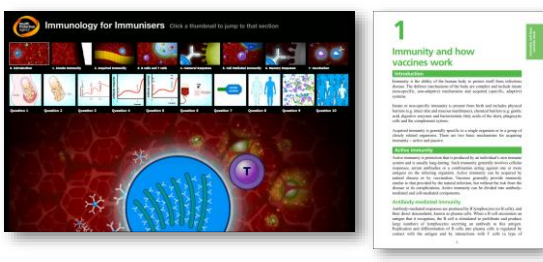
Important to understand the principles of immunology

114

Active Immunity

115

Helpful videos for immunology and FAQs
<http://immunologyanimation.hpa.org.uk>
 and chapter 1 of the 'Green Book'
 access via the 'your dedicated page' on my website



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Another useful video published in May 2018



<https://www.ovg.ox.ac.uk/news/how-do-vaccines-work>

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Do you know what factors might effect the immune response to a vaccine?


Age **Medical history**

- Very young children (especially under 2 years) have difficulty developing an immune response to polysaccharide only vaccines, and conjugated vaccines are used where possible
- Immunocompromised individuals usually cannot receive live attenuated vaccines. Inactivated vaccines are usually safe, but their immune response may be inadequate

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When might there be a contraindication to vaccinate?




- In general, a vaccine is absolutely contraindicated if a person has a confirmed anaphylactic reaction to a previous dose of the vaccine or product contained in the vaccine
- Pregnant women present a special risk group where, if the disease exposure is considered high during travel, most vaccine can be offered, although caution should be used with live vaccines
- All centres administering vaccines must be adequately prepared to deal with anaphylaxis

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Knowledge of the route we give vaccines and how soon they start to work is needed

- Most vaccines given by IM or SC route except BCG and oral vaccines (cholera and live typhoid)
- An active immune response to vaccines begins within a few days of administration and peaks in approximately 10-14 days
- Primary vaccine courses need 2 or 3 doses to complete the series



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Time for Vaccines to become effective

Table adapted from TRAVAX


Vaccine	Time until effective
BCG	6 weeks
Diphtheria	1-2 weeks after 3 rd dose
Hepatitis A (active)	2 weeks for optimum protection (the average incubation period for the disease is 28 days so it is often still useful to give the vaccine even at short notice prior to travel)
Hepatitis A immunoglobulin (passive)	Immediate
Hepatitis B	1 month after the 3 rd dose
Japanese encephalitis	7 days after the 2 nd dose, IXIARO® (Novartis) 1-2 weeks after the 2 nd dose, Green Cross vaccine (MASTA)
Measles/Mumps/Rubella (MMR)	2 weeks
Meningococcal vaccines (including ACW135Y)	2 weeks
Poliovirus (inactivated)	1-2 weeks after 3 doses
Rabies	1-2 weeks (after the 3 rd dose)
Tetanus	1-2 weeks after the 3 rd dose
Tick-borne encephalitis	2 weeks after the 2 nd dose
Typhoid injectable	1-2 weeks
Yellow fever	10 days

<http://www.travax.nhs.uk/vaccination-practice/arranging-schedules/time-for-vaccines-to-become-effective/>

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121

The rules of vaccination



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Travel Health Specialist Nurse

General rules on the vaccination

The Health Advice are compiled from sources of the most frequently asked questions about immunisation. The content has been reviewed and approved on 17/06/2019 and is subject to change. Any updates will be posted on the website.

- Recommendations for immunisation are based on currently available evidence and are subject to change as new evidence emerges. The advice is based on the best available evidence at the time of writing. The advice is based on the best available evidence at the time of writing. The advice is based on the best available evidence at the time of writing.
- The purpose of immunisation is to prevent disease and protect the health of the individual. It is not a guarantee that the individual will not get the disease. It is a way of reducing the risk of getting the disease and the severity of the disease if they do get it.
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on 'your dedicated' page


122

Peter thinks he has had 2 doses of hepatitis A vaccine in the past, but nothing is documented. He's going to travel to do some research work in a hospital in India – what would you do?

In the absence of documentation you cannot

ASSUME

the patient has been vaccinated, therefore further vaccines for protection should be given



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Travel Health Specialist Nurse


123

The evidence

For a variety of reasons, some individuals may not have been immunised or their immunisation history may be unknown.

If children and adults coming to the UK are not known to have been completely immunised, they should be assumed to be unimmunised and a full course of required immunisations should be planned.

Where a child born in the UK presents with an inadequate immunisation history, every effort should be made to clarify what immunisations they may have had. A child who has not completed the routine childhood programme should have the outstanding doses as described in the relevant chapters of the Green Book.



Page 6
REALLY IMPORTANT
to read this chapter


<https://www.gov.uk/government/publications/immunisation-schedule-the-green-book-chapter-11>
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124

Michelle had a first Havrix Junior Monodose at the age of one but never returned to complete the course, how would you proceed?

The evidence – chapter 11 again

Immunological memory from priming dose(s) are likely to be maintained in healthy individuals, increasing that interval will usually lead to a more pronounced response to the later dose. **Therefore, where any course of immunisation is interrupted, there is normally no need to start the course again - it should simply be resumed and completed as soon as possible.** Where vaccination was commenced some time previously however, the product received may have changed and the relevant chapter should therefore be consulted.



<https://www.gov.uk/government/publications/immunisation-schedule-the-green-book-chapter-11>
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125

The rules of vaccines

If a course goes off schedule and there is quite a long time interval, there is no need to re start the course, just pick up where it was left off and continue the course

January

First dose

6-12 months
time interval

Between July and the following January

Booster dose (or completing dose in the course)

January

Forgets!

January 6 years later

Just boost when they turn up

ALWAYS USE AN AGE APPROPRIATE VACCINE

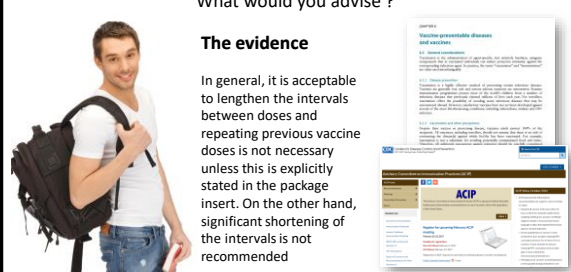
Department of Health, Immunisation against infectious disease (3rd Edition) London: TSO, 2006 Ch 17, p154
https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/147994/Green_Book_Chapter_17.pdf
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Jon is backpacking around SE Asia for 6 months and is having a course of rabies vaccine. He had his day 0 dose today but won't be around for the day 7 dose and asks if he can attend in 5 days instead for his second dose. What would you advise?

The evidence

In general, it is acceptable to lengthen the intervals between doses and repeating previous vaccine doses is not necessary unless this is explicitly stated in the package insert. On the other hand, significant shortening of the intervals is not recommended



Department of Health, Immunisation against infectious disease (3rd Edition) London: TSO, 2006 Ch 17, p154
https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/147994/Green_Book_Chapter_17.pdf
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Site of injection and number that can be given at one time?

- If two or more injections need to be administered at the same time, they should be given in separate sites, preferably in a different limb. If more than one injection is to be given in the same limb, they should be administered at least 2.5cm apart
- Immunisations should not be given into the buttock, due to the risk of sciatic nerve damage and the possibility of injecting the vaccine into fat rather than muscle?

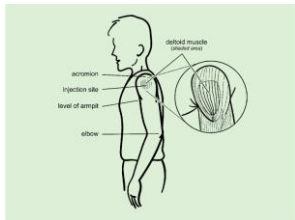


Figure 4.1 Preferred site for intramuscular and deep subcutaneous injections in older children and adults

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/417315/Green_Book_Chapter_4.pdf
 2. Department of Health. Immunisation against infectious disease (2nd Edition) London: HMSO, 2006. Ch.4, p.29
 https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/417315/Green_Book_Chapter_4.pdf

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 Trustee, British Society for Immunology

128

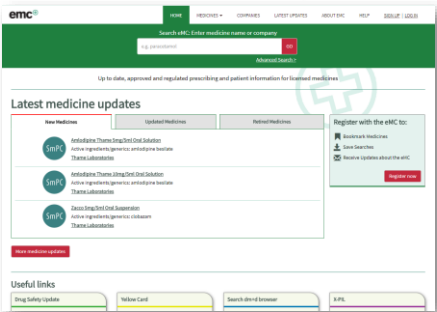
Where do you obtain relevant vaccine information?

- Green Book – for diseases and vaccines, online
- The National Databases (NaTHNaC and TRAVAX)
- Patient Group Directions - in your workplace
- Electronic Medicines Compendium - online
- British National Formulary – book or online
- Pharmaceutical companies - online

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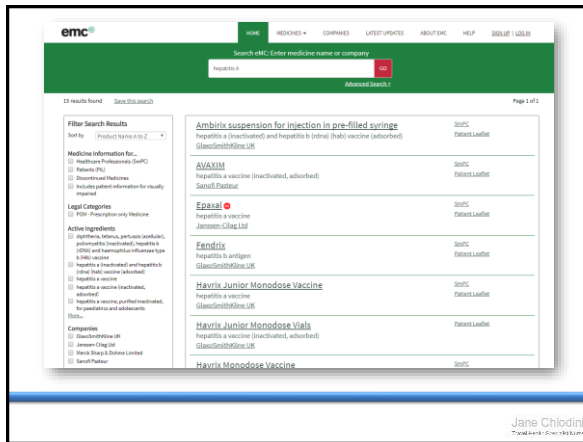
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Electronic Medicines Compendium www.medicines.org.uk



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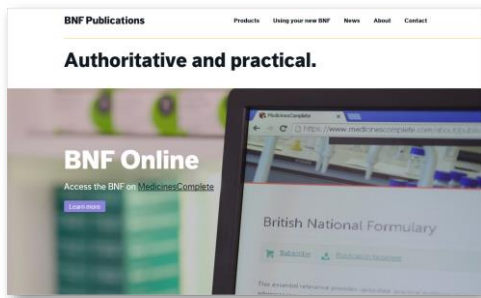
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131

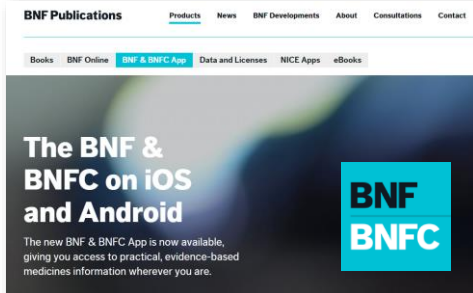
www.bnf.org



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132

BNF and BNFC apps are great!



https://www.bnf.org/products/bnfbnfcapp/

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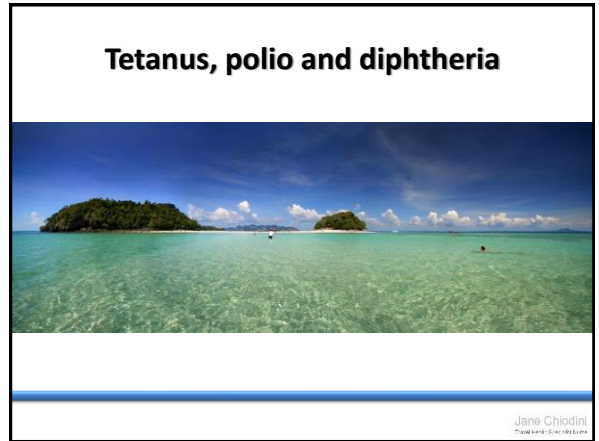
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Document your learnings

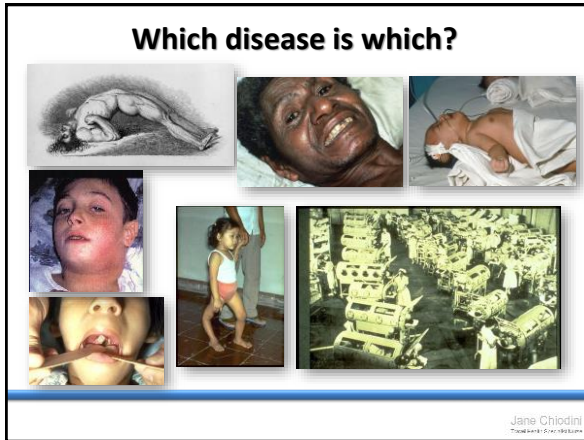
Disease protection	Causative organism	Mode of transmission..... prevention advice required	No. of vaccines in the course	Length of protection
Tetanus, polio and diphtheria				
Measles A				
Typhoid				
Cholera				
Measles B				
Meningococcal meningitis				
Yellow fever				
Rabies				
Japanese encephalitis				
Tick borne encephalitis				

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Thank you to the NHS

134





135



136

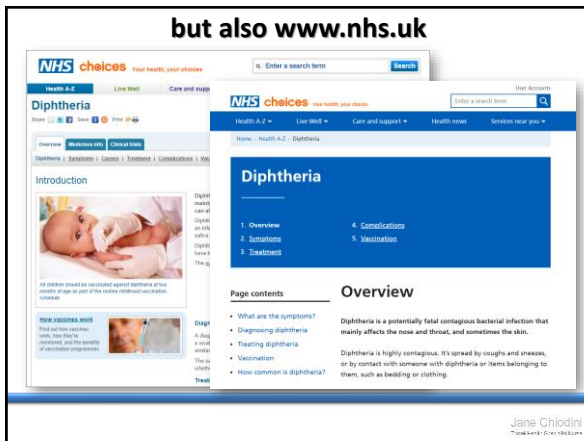
Learning more from Green Book chapters, but also www.nhs.uk

Disease	Organism	Mode of transmission
Tetanus	toxoid	Spores in the environment
Diphtheria	bacterium	Droplet infection
Polio	virus	Faecal oral and saliva

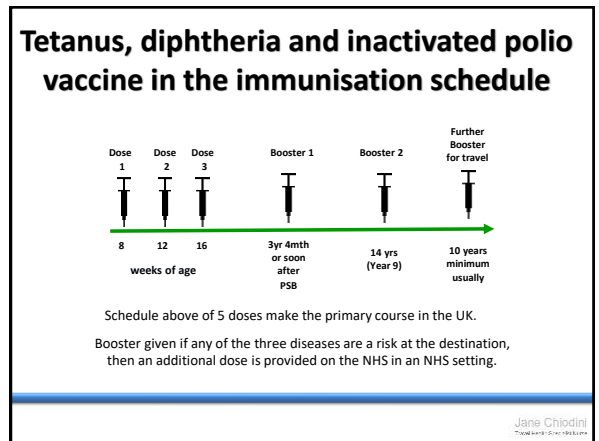



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137



138



139

The routine immunisation schedule from Autumn 2018

Tetanus, polio and diphtheria disease protection is not available in monovalent vaccines, only as one combined vaccine but in different products within the routine immunisation schedule

Infanrix hexa	DTaP/IPV/Hib/Hep B
Infanrix IPV	(DTaP/IPV)
Repevax	(DTaP/IPV)
Revaxis	(Td/IPV)

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Travel Health Clinician

140

Acronyms for vaccines and values of diphtheria content – D and d

Available vaccines^[6]

Diphtheria vaccines are available in two strengths according to dose of toxoid:

- High-dose - vaccines contain ≥30 IU of diphtheria toxoid and are used to achieve satisfactory primary immunisation of children - as in diphtheria/tetanus/acellular pertussis (DTaP) vaccine (capital D = high-dose).
- Low-dose - vaccines contain approximately 2 IU of toxoid and are used for primary immunisation of those aged over 10 years and for subsequent boosters (lower case d signifies low-dose as in dTaP).

<https://www.uks.gov/vaccines/ukca/comm/tra/children/acc/ukca/ukca.html>
<https://www.gov.uk/government/consultations/diphtheria-and-diphtheria-vaccines>

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Travel Health Clinician

141

Would this traveller need Td/IPV?

- Lucy is 19 years old and is going on a two week holiday in the Galapagos Islands
- She is up to date on all her scheduled national programme immunisations
- She hasn't travelled abroad before
- No PMH, she is on the OCP only

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Travel Health Clinician

142

Galapagos Islands off coast of Ecuador

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Travel Health Clinician

143

Would this traveller need Td/IPV?

- James is 26 years old
- he's taking a one year career break back packing around the world
- He last had a tetanus vaccine as a school booster 9½ years ago at the age of 16

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Travel Health Clinician

144

Keeping an eye on other groups

dTaP/IPV vaccine given between gestational weeks 20* and 32 rather than from week 28

*Can be given from 16 weeks but usually offered after the anomaly scan

Examples ?

Boostrix-IPV or Repevax

To find out more about whole cell pertussis and acellular pertussis see <http://www.who.int/biologicals/vaccines/pertussis/en/>

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Travel Health Clinician

145

FAQ on tetanus

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Travel Health Specialist Nurse

146

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Travel Health Specialist Nurse

147

Read the guidance to see definitions

6.2 Management of Tetanus Prone Wounds

Tetanus-prone wounds include:

- puncture-type injuries acquired in a contaminated environment and likely therefore to contain tetanus spores* e.g. gardening injuries
- wounds containing foreign bodies such as wound splinters*
- compound fractures
- wounds or burns with systemic sepsis
- certain animal bites and scratches**

*Note: individual risk assessment is required and this list is not exhaustive e.g. a puncture-wound from discarded needle found in a park may be a tetanus-prone injury but a needledstick injury in a medical environment is not. **Similarly, although smaller bites from domestic pets are generally puncture injuries, animal saliva should not contain tetanus spores unless the animal has been rooting in soil or lives in an agricultural setting.*

High-risk tetanus-prone wounds include:

- heavy contamination with material likely to contain tetanus spores e.g. soil, manure
- wounds or burns that show extensive devitalised tissue
- wounds or burns that require surgical intervention that hours are high risk even if the contamination was not

And on page 9 of the Green Book tetanus chapter

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Travel Health Specialist Nurse

148

Jane Chiodini's Blog

Tuesday, 30 April 2019

Tetanus vaccine: 5 doses?

This isn't travel but because we have to also ensure patients are fully protected with the national immunisation schedule for tetanus I still think it's important.

A little history

For many years the said once you have had 5 doses of tetanus containing vaccine then you are protected for life within the UK. Information written in the Green Book chapter on tetanus published in 2005 and 2007 informed that if you then sustained a tetanus prone you would require treatment which would be tetanus specific immunoglobulin but such documents are not available online any more.

The Patient info website on their page here states: *The primary course of three injections gives good protection for a number of years. The fourth and fifth doses (boosters) maintain protection. After the fifth dose, immunity remains for life and you do not need any further boosters (apart from some travel situations).*

New guidance

new guidance published by the on 3 November 2018 covers Tetanus advice for health professionals. Guidance on the treatment of tetanus cases and management of tetanus prone wounds (and the list of 2019 5 doses) can be accessed here and may be longer. The Green Book chapter 30 for tetanus was also updated on 30 November 2018 and provides a more comprehensive perspective of what is a tetanus prone wound?

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Travel Health Specialist Nurse

149

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150

Hepatitis A

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Travel Health Specialist Nurse

151

Hepatitis A

http://gamapserver.who.int/maplibrary/Files/Maps/Global_HepA_IHTRiskMap.png?ua=1

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Travellers' Clinician

152

Hepatitis A disease

- Viral infection
- Transmitted via contaminated food and water
- Those at higher risk – VFRs, long term travellers, those exposed to conditions of poor sanitation
- Incubation averages 28 - 30 days (range 15 to 50 days)
- Often asymptomatic in young children
- Abrupt onset of malaise, anorexia, nausea, fever followed by jaundice
- Fulminant hepatitis is more likely in those with pre-existing liver disease and in older individuals
- The overall case fatality ratio is low but is greater in older patients and those with pre-existing liver disease

<http://phil.cdc.gov/phil/home.asp>

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/147954/Green-Book-Chapter-17.pdf

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Travellers' Clinician

153

Hepatitis A schedule – what the Green Book says

- The duration of protection from a completed course of vaccine can be expected to be at least 25 years and probably indefinite.
- However, PHE recommend that until further evidence is available on persistence of protective immunity, a booster dose at 25 years is indicated for those at ongoing risk of hepatitis A.

Hepatitis A vaccines can be used interchangeably: Chapter 4, page 145 of the Green Book

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/147954/Green-Book-Chapter-17.pdf

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Travellers' Clinician

154

Five hepatitis A monovalent vaccines

VACCINE NAME	AGE GROUPS	WHEN TO BOOST- general principles and comments
HEPATITIS A vaccine (and schedules)		
• Hep A vaccine SPCs have different timings but note Ch. 4, 1 st paragraph in GB. Ideally, follow the summary of product characteristics but in late-presenting travellers, a course does not need to be restarted (DH 2013). Protection is expected for 25 years from the second dose – also see NaTHNaC info on Hepatitis A ** and detail below within the 'Key' section regarding GSK 'Havrix' vaccines***.		
VAQTA® Paediatric	2 dose schedule of hepatitis A vaccine should be given at day 0 and then 6 to 12 months after the initial dose as recommended in Green Book for hep A vaccines.	1 - 17 years
VAQTA® Adult		18 years and over
Avaxim®		16 years and over
Havrix Junior Monodose®		1 - 15 years
Havrix Monodose®	Regimes may vary in SPCs, see above • & key	16 years and over

KEY
 * Within the Summary of Product Characteristics (SPC)
 ** The Green Book (2013) refers to all hep A products, so the 25 year protection also applies to the combined products and paediatric hepatitis A vaccines. Until further evidence is available on persistence of protective immunity, a further booster at 25 years is indicated for those at ongoing risk. See the Green Book chapter (page 154) and NaTHNaC document at www.nathnac.org/pro/factsheets/hep_a.htm
 *** SPC for Havrix Monodose & Havrix Junior Monodose April 2012 states 'Current recommendations do not support the need for further booster vaccination among immunocompetent subjects after 2 dose course'

See item no. 3 at <http://www.janechiodini.co.uk/tools/>

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Travellers' Clinician

155

Over to you!

- 1 What is the ideal schedule for a course of hepatitis A vaccine?
- 2 What do you do if the patient doesn't return for their booster on time?
- 3 How long does a completed course of hepatitis A vaccine last?
- 4 If a patient had a past medical history of confirmed hepatitis A infection, would you need to vaccinate them?
- 5 Are hepatitis A vaccines interchangeable?
- 6 If a patient had HNIG recorded in their notes would how would you proceed on hep A protection?
- 7 Is the time of the protection taken from the first dose or booster dose of hepatitis A vaccine?
- 8 If you gave a child a first hepatitis A vaccine and they return as an adult and a booster is required – which vaccine is best?
- 9 Could you give hepatitis A vaccine on the day of departure of a trip?

Jane Chiodini
Travellers' Clinician

156

Over to you!

- 1 What is the ideal schedule for a course of hepatitis A vaccine? **0 and 6-12 months**
- 2 What do you do if the patient doesn't return for their booster on time? **Just boost when they turn up**
- 3 How long does a completed course of hepatitis A vaccine last? **25 years**
- 4 If a patient had a past medical history of confirmed hepatitis A infection, would you need to vaccinate them? **No**
- 5 Are hepatitis A vaccines interchangeable? **Yes**
- 6 If a patient had HNIG recorded in their notes would how would you proceed on hep A protection? **Start a course of hep A vaccine**
- 7 Is the time of the protection taken from the first dose or booster dose of hepatitis A vaccine? **25 years from the booster dose**
- 8 If you gave a child a first hepatitis A vaccine and they return as an adult and a booster is required – which vaccine is best? **Boosting with an adult dose**
- 9 Could you give hepatitis A vaccine on the day of departure of a trip? **Yes**



<https://www.gov.uk/government/publications/hepatitis-a-the-green-book-chapter-17>

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157

Some historical hepatitis A protection/vaccine information

Immunoglobulin given in 1980s and early 1990s but discontinued due risk of CJD from UK sourced blood products

Jane Chiodini
Travel Health Specialist Nurse

158

Some historical hepatitis A protection and vaccine information


- Immunoglobulin given in 1980s and early 1990s but discontinued due risk of CJD from UK sourced blood products
- Hepatitis A vaccine introduced in 1992 – Havrix – had to give two doses prior to travel as it had 720 ELISA units of hepatitis A (three doses in total course)
- Havrix Monodose available from 1994 which had 1440 ELISA units of hepatitis A and only one dose required prior to travel (two doses in total course)
- Vaqta Adult – problem in 1990s when some batches thought not to give protection – instructed at that time to disregard doses given previously and re-vaccinate. Vaqta Adult now available again
- See **Nuggets of Knowledge – hepatitis A**

Jane Chiodini
Travel Health Specialist Nurse

159

Would this traveller need hep A vaccine?

- Lucy is 19 years old and is going on a two week holiday in the Galapagos Islands
- She is up to date on all her scheduled national programme immunisations
- She hasn't travelled abroad before
- No PMH, she is on the OCP only
- Which vaccine schedule would you give?




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Would this traveller need hep A vaccine ?


- James is 26 years old
- he's taking a one year career break back packing around the world
- He last had a tetanus vaccine as a school booster 9½ years ago at the age of 16
- He tells you he had one dose of hep A vaccine when he was 12 years old but there is no record of it in the notes
- How would you proceed?



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Travel Health Specialist Nurse

161

Tip



Hepatitis A vaccine provides some of the most frequently asked questions therefore NaTHNaC and TRAVAX both have very helpful documents to help – it's a good idea to be aware of them

<http://travelhealthpro.org.uk/hepatitis-a/>

<http://www.travax.nhs.uk/diseases/vaccine-preventable/hepatitis-a/hepatitis-a-faqs.aspx>

FAQ on Hep A from TRAVAX

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Travel Health Specialist Nurse

162



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Travel Health Specialist Nurse

Home About Education Tools News Links Contact Us

Hepatitis


The basic principles of a hepatitis A vaccine schedule

FAQ

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Travel Health Specialist Nurse

163


Recommend you do this to consolidate knowledge – see on your page



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Thank you to the CDC

164

Hepatitis B



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165

WHO Factsheet – Hepatitis B updated July 2018

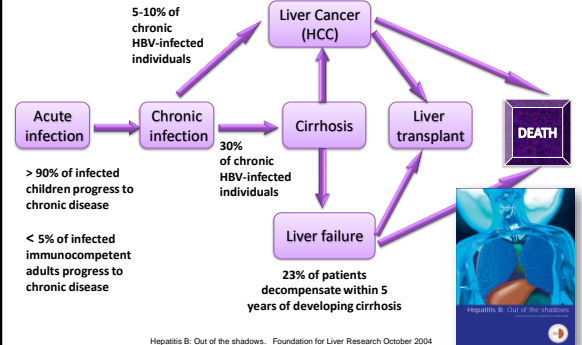
- Hepatitis B is a viral infection that attacks the liver and can cause both acute and chronic disease.
- The virus is transmitted through contact with the blood or other body fluids of an infected person.
- An estimated 257 million people are living with hepatitis B virus infection (defined as hepatitis B surface antigen positive).
- In 2015, hepatitis B resulted in 887 000 deaths, mostly from complications (including cirrhosis and hepatocellular carcinoma).
- Hepatitis B is an important occupational hazard for health workers.
- However, it can be prevented by currently available safe and effective vaccine.

<http://www.who.int/en/news-room/fact-sheets/detail/hepatitis-b>

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Thank you to the CDC

166

Natural history of hep B infection



5-10% of chronic HBV-infected individuals progress to Liver Cancer (HCC)

30% of chronic HBV-infected individuals progress to Cirrhosis

23% of patients decompensate within 5 years of developing cirrhosis

Acute infection leads to Chronic infection


> 90% of infected children progress to chronic disease

< 5% of infected immunocompetent adults progress to chronic disease

Outcomes from Cirrhosis: Liver Cancer (HCC), Liver failure, Liver transplant, DEATH

Hepatitis B: Out of the shadows. Foundation for Liver Research October 2004
<http://www.liverresearch.org.uk/liver-research-files/Hepatitis-B-Out-of-the-Shadows.pdf>

168



This Khmer woman died of hepatoma, four months after arriving in a refugee camp in Thailand

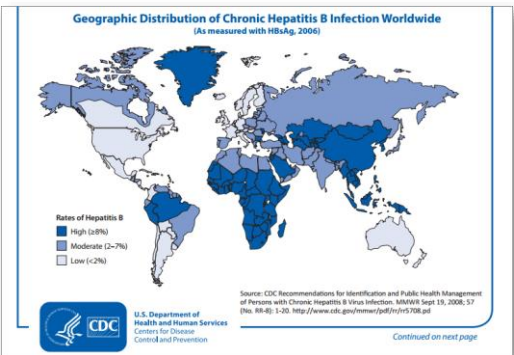
<http://www.vaccineinformation.org/hepb/photos.asp> or
<http://www.immunize.org/photos/hepatitis-b-photos.asp>

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169

Geographic Distribution of Chronic HBV

Geographic Distribution of Chronic Hepatitis B Infection Worldwide (As measured with HBsAg, 2006)



Rates of Hepatitis B

- High (>8%)
- Moderate (2-7%)
- Low (<2%)

Source: CDC Recommendations for Identification and Public Health Management of Persons with Chronic Hepatitis B Virus Infection. MMWR Sept 19, 2008; 57 (No. 38-4): 1-10. <http://www.cdc.gov/mmwr/pdf/w97108a.pdf>

U.S. Department of Health and Human Services
Centers for Disease Control and Prevention

Continued on next page

170

USA and European resources: see 'immunisation resources' and 'hepatitis B' in 'HELP'

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Thank you to the NHS

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Transmission of hepatitis B

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More recent craze – corset piercing

<http://unusual-things.blogspot.com/2011/05/surgeon-blasts-latest-craze-in-body.html>

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The commonest transmission route of hepatitis B?

http://unicef.org.blogspot.co.uk/2010/10/unicef-executive-director-launches_31.html

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Not travel related but important to understand

See page 14, chapter 18 of the Green Book

Age	Routine childhood programme	Babies born to hepatitis B infected mothers
Birth	**	✓ Monovalent HepB
4 weeks	✓	✓
8 weeks	✓	✓
12 weeks	✓	✓
16 weeks	✓	✓
1 year of age	**	✓
3 years and 4 months	**	✓

* Newborn infants born to a hepatitis B negative woman but known to be going home to a household with another hepatitis B infected person may be at immediate risk of hepatitis B infection. In these situations, a monovalent dose of hepatitis B vaccine should be offered before discharge from hospital. They should then continue on the routine childhood schedule commencing at eight weeks.
** Give the recommended non-hepatitis B containing vaccine as per the routine schedule.

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/132637.pdf
<https://www.gov.uk/government/publications/hepatitis-b-antenatal-screening-and-newborn-immunisation-programme-best-practice-guidance>

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175

Green Book recommendations for hepatitis B vaccine

- Injecting drug users
- Individuals who change sexual partners frequently, particularly MSM and commercial sex workers
- Close family contacts of a case or carrier
- Families adopting children from countries with a high or intermediate prevalence of hepatitis B
- Foster carers
- Individuals receiving regular blood or blood products and their carers
- Patients with chronic renal failure
- Patients with chronic liver disease
- Inmates of custodial institutions
- Individuals in residential accommodation for those with learning difficulties

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/148308/Green-Book-Chapter-18.pdf

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Thank you to the NHS

176

The Green Book information regarding travellers

People travelling to or going to reside in areas of high or intermediate prevalence

Travellers to areas of high or intermediate prevalence who place themselves at risk when abroad should be offered immunisation. The behaviours that place them at risk will include sexual activity, injecting drug use, undertaking relief aid work and/or participating in contact sports. Travellers are also at risk of acquiring infection as a result of medical or dental procedures carried out in countries where unsafe therapeutic injections (e.g. the re-use of contaminated needles and syringes without sterilisation) are a risk factor for hepatitis B (Kane *et al.*, 1999; Simonsen *et al.*, 1999). Individuals at high risk of requiring medical or dental procedures in such countries should therefore be immunised, including:

- those who plan to remain in areas of high or intermediate prevalence for lengthy periods
- children and others who may require medical care while travelling to visit families or relatives in high or moderate-endemicity countries
- people with chronic medical conditions who may require hospitalisation while overseas e.g. dialysis
- those travelling for medical care

NB. The Green Book is nothing to do with whether the traveller should pay for vaccine or not

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/148308/Green-Book-Chapter-18.pdf

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177

Schedules for hepatitis B vaccine

HEPATITIS B vaccine (and schedules) Important – Hep B now in the childhood programme not included here		
Engerix B® - 0, 1 and 6 months	Over 16 years	Note: 0, 1, 2 month schedule Green Book doesn't advise 4 th dose at 12 months unless they remain at continued high risk, see Ch.18 Page 16. Note SmpCs do advise a 4 th dose but GB should be followed. Update to policy in the Green Book in June 2017 for hepatitis B for all (which would include travellers) states those who have received a primary course do not require a reinforcing dose of hep B containing vaccine except health care workers (boost once at 5 years), patients with renal failure and at time of significant exposure. Please read Ch. 18 page 13 of Green Book for detail. Testing for evidence of immunity post immunisation is not routinely recommended. See GB. Ch.18. Page 18
Engerix B® - 0, 1, 2 months	Over 16 years	
Engerix B® - 0, 7, 21 days & 12 months	Over 18 years in SmpC But also 16 -18 years in Green Book	
Engerix B® Paediatric 0, 1, 6 months	0 to 15 years	
Engerix B® Paediatric 0, 1, 2 months	0 to 15 years	
Engerix B® Option of two doses of 1 ml (20mcg) for low-compliance adolescents given 6 months apart when the risk of hepatitis B is low and completion of course can be assured before risk is high	11 – 15 years	
HBVaxPRO® 0, 1, and 6 months	16 years and over	
HBVaxPRO® 0, 1, 2 months	16 years and over	
HBVaxPRO® Paediatric 0, 1 & 6 months	0 – 15 years	
HBVaxPRO® Paediatric 0, 1, 2 months	0 – 15 years	

Two products, four presentations

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178

Which schedule?

Hepatitis B Green Book chapter page 12

Pre-exposure immunisation schedule for **high risk individuals**

- For pre-exposure prophylaxis in most adult and childhood risk groups, an accelerated schedule should be used, with vaccine given at zero, one and two months.
- Higher completion rates are achieved with the accelerated schedule (at zero, one and two months) in groups where compliance is difficult (e.g. in people who inject drugs [PWID] and genitourinary medicine clinic attenders) (Asboe *et al.*, 1996).
- This improved compliance is likely to offset the slightly reduced immunogenicity when compared with the zero-, one- and six-month schedule, and similar response rates can be achieved by the opportunistic use of a fourth dose after 12 months.
- An alternative schedule at zero, one and six months should only be used where rapid protection is not required and there is a high likelihood of compliance.
- If the primary course is interrupted it should be resumed but not repeated.)

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179

What about hepatitis B boosters?

Hepatitis B Green Book chapter page 13

Reinforcing doses for those who have received pre-exposure immunisation

The current UK recommendation is that those who have received a primary course of immunisation, including children vaccinated according to the routine childhood schedule and individuals at high risk of exposure, **do not require a reinforcing dose of Hep B-containing vaccine**, except in the following categories:

- healthcare workers (including students and trainees), who should be offered a single booster dose of vaccine, once only, around five years after primary immunisation
- patients with renal failure
- at the time of a significant exposure (see the chapter for more detail)

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/132602/Greenbook_chapter_18.pdf

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180

Statement in this guidance Feb 2018

– not yet in the Green Book

Booster doses in healthcare workers

On the advice of the Joint Committee on Vaccination and Immunisation (JCVI), boosters (priority group 5) will no longer be routinely required in healthy, immunocompetent adults who have completed a primary course of vaccine, including healthcare workers who are known responders.

See page 8

https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/683830/Plan_for_phased_re-introduction_of_hepatitis_B_vaccine_for_lower_priority_groups_2018.pdf

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181

In summary for hepatitis B vaccine given for travel purposes

- Use 0, 1 and 2 month schedule in preference to 0, 1 and 6 month when more rapid protection is needed
- If insufficient time before travel, use a 0, 7, 21 day and then reinforce at 12 months
- No longer boost at 5 years for travel
- Blood test not routinely performed for seroprotection in travellers

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182

Where would you find information about the levels of protection?

Pharmacodynamic properties

Pharmacodynamic group: Hepatitis A vaccine, ATC code: J07CA01

Effectiveness: 100% (12 months)

Population: Healthy subjects aged 16 and including 10 years of age

Schedule: 0, 1, 6 months; 0, 1, 2 - 12 months

Seroprotection rate: At month 7: 95.5%; At month 1: 91.5%; At month 3: 89.5%; At month 13: 85.8%

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Understanding Twinrix

Hepatitis A content

- Havrix Monodose** – content of **1440** elisa units of hepatitis A
- Havrix Junior Monodose** – content of **720** Elisa units of hepatitis A
- Twinrix Adult** – content of **720** Elisa units of hep A plus an adult dose of hep B
- Twinrix paediatric** – content of **360** elisa units of hepatitis A plus paediatric dose of hep B

Red = full dose Green = half dose Blue = quarter of a dose

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Trainer and Content Developer

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Hepatitis A vaccine antigen content within vaccines

Adapted from table 1 of the Public Health England Hepatitis A vaccination in adults temporary recommendations *

Hep A vaccine formulation	Trade name	Hep A vaccine antigen content	Adult dose Hep A antigen equivalent	Made by
Adult monovalent hep A	AVAXIM	160 U	Full dose	SP
	HAVRIX MONODOSE	1440 EU	Full dose	GSK
	VAQTA ADULT	50 U	Full dose	MSD
Paediatric monovalent hep A	HAVRIX JUNIOR MONODOSE	720 EU	Half dose	GSK
	VAQTA PEDIATRIC	25 U	Half dose	MSD
Adult combination hepatitis A/B	TWINRIX ADULT	720 EU	Half dose	GSK
Paediatric combination hepatitis A/B	TWINRIX PEDIATRIC	360 EU	Quarter dose	GSK
	AMBIRIX	720 EU	Half dose	GSK
Combination hepatitis A/typhoid	VIATIM	160 EU	Full dose	SP

Made into a poster and now on your webpage

KEY
SP – Sanofi Pasteur; GSK – GlaxoSmithKline; MSD – Merck Sharpe Dohme Ltd.
<https://www.gov.uk/government/publications/hepatitis-a-infection-prevention-and-control-guidance>
https://assets.publishing.service.gov.uk/government/uploads/system/uploads/attachment_data/file/616228/hepatitis_a_vaccination_recommendations.pdf

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Explaining composition of Twinrix

Two doses of Twinrix (adult or paediatric) must be given to achieve protection against hepatitis A prior to departure

RULE – if you start with Twinrix you can complete with Twinrix. If you start with monovalent vaccines DON'T try to complete with Twinrix

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Trainer and Content Developer

186

Combination A and B vaccines schedules

- Twinrix adult**
 - 0, 1 & 6 months from 16 years
 - 0, 7, 21 days & 12 mths (18yrs)
- Twinrix paediatric**
 - 0, 1 & 6 months
 - Use in 1 – 15 year age group
- Ambirix**
 - 0 and 6 - 12 months
 - Use in 1 to 15 years

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Understanding content of Twinrix and Ambirix

- Havrix Monodose** – content of **1440** elisa units of hepatitis A
- Havrix Junior Monodose** – content of **720** Elisa units of hepatitis A
- Twinrix Adult** – content of **720** Elisa units of hep A plus an adult dose of hep B
- Twinrix paediatric** – content of **360** elisa units of hepatitis A plus paediatric dose of hep B

Ambirix – content of **720** elisa units of hepatitis A plus a full dose of hep B

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Trainer and Content Developer

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Combination hepatitis B vaccines used in the UK

Table 18.2 Dosage of combined hepatitis A and hepatitis B vaccines by age (from the Green Book)


Vaccine product	Ages	Dose HAV	Dose HBV	Volume
Twinrix Adult® Hepatitis A (inactivated) and hepatitis B (rDNA) (HAB) vaccine (adsorbed)	16 years or over	720 ELISA units	20µg	1.0ml
Twinrix Paediatric® Hepatitis A (inactivated) and hepatitis B (rDNA) (HAB) vaccine (adsorbed)	1 – 15 years	360 ELISA units	10µg	0.5ml
Ambirix® Hepatitis A (inactivated) and hepatitis B (rDNA) (HAB) vaccine (adsorbed)	1 – 15 years	720 ELISA units	20µg	1.0ml

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/148308/Green-Book-Chapter-18.pdf

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Travel and Public Health

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Tip



In practice we give hepatitis B for lifestyle risks and travel, but not usually occupational risk. Hepatitis B is a large topic – it would be useful to read the Green Book chapter on this topic at https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/148308/Green-Book-Chapter-18.pdf


And the BMA guidance document at <http://bma.org.uk/practical-support-at-work/doctors-as-managers/managing-your-practice/focus-hepatitis-b-immunisations>

See the hepatitis B document on **your dedicated page !**

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190

Typhoid




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Travel and Public Health

191

Enteric fevers – typhoid & paratyphoid

- Gram-negative bacterial infection
- Transmission by faecal-oral route, water borne and human to human
- Those at higher risk include VFRs, young children, long term travellers and those exposed to conditions of poor sanitation – mainly in Asia
- Incubation 7 to 14 days
- Fever, chills, headache, malaise, weakness, anorexia, abdominal pain, diarrhoea
- Complications in 10% -15%: intestinal perforation, bacteraemia, meningitis
- Chronic carrier status in <3% infected persons



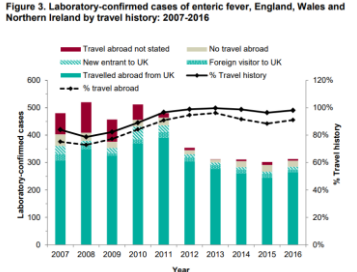
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192

Data from PHE

Enteric fever (typhoid and paratyphoid) England, Wales and Northern Ireland: 2016

Figure 3. Laboratory-confirmed cases of enteric fever, England, Wales and Northern Ireland by travel history: 2007-2016

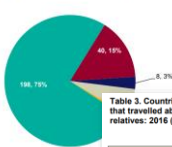


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193

Reason for travel and destination

Figure 4. Reason for travel for laboratory-confirmed cases of enteric fever that travelled abroad from England, Wales and Northern Ireland: 2016 (N=265)



India, Pakistan and Bangladesh were the highest risk country for travellers returning with enteric fever

Table 3. Countries of travel and ethnicity for laboratory-confirmed cases of enteric fever that travelled abroad from England, Wales and Northern Ireland to visit friends and relatives: 2016 (N=198*)

Presumed country of infection	Ethnicity							Total
	Pakistani	Indian	Bangladeshi	Asian other	Black African	Other/mixed	Not stated	
Pakistan	75	2	-	1	-	-	4	82
India	-	87	-	-	-	-	6	93
Bangladesh	-	-	12	-	-	-	-	12
Sub-Saharan Africa	-	-	-	-	2	1	1	4
South America	-	-	-	-	-	1	-	1
Europe	-	-	-	-	-	-	1	1
Other Asia	-	3	-	3	-	-	-	6
Total	75	92	12	4	2	3	12	200*

* Note that some cases travelled to more than one country; all countries are included in this table so the totals in the table will be higher than the actual number of cases.

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Travel and Public Health

194

Two vaccines now for typhoid protection

TYPHOID vaccine		
Typhim Vi® Single dose	2 years and over	3 years
Vivotif® (Ty21a) Oral vaccine on days 0, 2 & 4	5 years and over	3 years (Take with cold or luke warm drink 1 hr before meal, swallow capsule whole)

Injectable typhoid protection is a polysaccharide vaccine and so just one dose makes up 'the course'. After this time period if further typhoid protection is needed a new dose is given.

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Travel Nurse & Clinician

195

Details from the Green Book

Dosage of injectable monovalent typhoid vaccines				
Vaccine product	Ages	Dose	Volume	
Typhim Vi	Two years and older*	25µg	0.5ml	0.5ml
Typhera (discontinued in 2018)	Two years and older*	25µg		

Note
Typhrix and Hepatyrix have now been discontinued – as now highlighted in the Green Book

Dosage of oral monovalent typhoid vaccine		
Vaccine product	Ages	Dose
Vivotif	Five years and older	Three capsules on days 0, 2 and 4

Dosage of combined typhoid and hepatitis A vaccines**				
Vaccine product	Ages	Dose typhoid	Dose HAV†	Volume
Hepatyrix (discontinued in 2018)	15 years and older	25µg	1440 ELISA units	1ml
VIATIM	16 years and older	25µg	160 antigen units	1ml

* Children between the ages of 12 months and two years should be immunised off-license if following a detailed risk assessment the risk of typhoid fever is considered high.

** For booster doses of either typhoid or HAV, single antigen vaccines can be used

† HAV – hepatitis A vaccine

Children between the ages of 12 months and two years should be immunised off-license if following a detailed risk assessment the risk of typhoid fever is considered high

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/248512/Green_Book_Chapter_33_0h_125148.pdf

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Travel Nurse & Clinician

196

Oral typhoid vaccine - Vivotif

- LIVE vaccine*
- Use from 5 years of age
- Three doses on days 0, 2 & 4 (Green Book)

Administration

- The capsule should be taken approximately one hour before a meal with a cold or lukewarm drink (temperature not to exceed body temperature, e.g. 37°C)
- The vaccine capsule should not be chewed and should be swallowed as soon as possible after placing in the mouth

* can be administered at any time before or after other live vaccines.

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Travel Nurse & Clinician

197

Moved to Chapter 11 of the Green Book – in April 2019

Table 11.2 Recommended time intervals when giving more than one live attenuated vaccine

Vaccine combinations	Recommendations
Yellow fever and MMR	A four week minimum interval period should be observed between the administration of these two vaccines. Yellow fever and MMR should not be administered on the same day.
Varicella (and zoster) vaccine and MMR	If these vaccines are not administered on the same day, then a four week minimum interval should be observed between vaccines.
Tuberculin skin testing (Mantoux) and MMR	MMR vaccination and tuberculin skin testing can be performed on the same day. However, if a tuberculin skin test has already been initiated, then MMR should be delayed until the skin test has been read. Strongly recommend separate tuberculin testing, if a child has had a recent MMR, and requires a tuberculin test, then a four week interval should be observed.
All currently used live vaccines (BCG, rotavirus, live attenuated influenza vaccine (LAIV), oral typhoid vaccine, yellow fever vaccine, zoster and MMR)	Apart from those combinations listed above, these vaccines can be administered at any time before or after each other. This includes tuberculin (Mantoux) skin testing.

Co-administration of these two vaccines can lead to sub-optimal antibody responses to yellow fever, mumps and rubella antigens (Nascimento et al, 2011). Where protection is required rapidly then the vaccines should be given at any interval; an additional dose of MMR should be considered.

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198

Oral typhoid - Vivotif®

www.medicines.org.uk

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Travel Nurse & Clinician

199

Interactions with other medication from the SmPC

- Antibiotics – Vivotif may not work if it is taken while you are also taking antibiotics. Take Vivotif no earlier than 3 days after the last dose of an antibiotic
- Medicines to prevent malaria – do not start these until 3 days after the last dose of Vivotif
- Yellow fever vaccine can be given while taking Vivotif

Vivotif® Patient Information Leaflet <http://www.medicines.org.uk/EMC/medicines/24328/PL/Vivotif/>

Jane Chiodini
Travel Nurse & Clinician

200

Charging and administration

Oral typhoid vaccine is an NHS provision in an NHS setting

1. You could buy this vaccine in and bring the patient in for all three doses
2. You could administer the vaccine to the patient for the first dose and given them the other two doses to take home to self administer but the vaccine must be stored at 2 – 8°C
3. You could supply the vaccine on an FP10 and allow the patient to self administer

You need to ensure that your traveller understands the importance of, and can assure the cold chain in points 2 and 3 above.

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Travel Health Consultant

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Who would need typhoid vaccine if there is a shortage?

- Family of four going to an all inclusive break for a 10 days Cancun
- 40 year old couple travelling to stay in 4 star hotel in Bangkok for two weeks
- 26 year old man going to stay in Bangkok in a guest house/hostel type accommodation and he has type 1 diabetes
- Parents and their children of 7, 5 and 2 years travelling to Pakistan for 3 weeks to see family

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Travel Health Consultant

202

Tip



NaTHNaC and TRAVAX have both written information documents on typhoid as well

<http://travelhealthpro.org.uk/typhoid-and-paratyphoid/>
<http://www.travax.nhs.uk/diseases/vaccine-preventable/typhoid/typhoid-faqs.aspx>

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Travel Health Consultant

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Hepatitis A + Typhoid available as a combined vaccine

Why give combination vaccines?

COMBINED vaccines (and schedules)		
VIATIM® (Hepatitis A and typhoid) Single dose	16 years and over	6-12 months for hepatitis A component (then hepatitis A booster given as a monovalent vaccine) and 9 years for the typhoid component

The combination hepatitis A and typhoid vaccine can be given with the hep A protection provided as either the first dose or reinforcing or booster dose of hep A vaccine as long as there is the three year interval to fulfill the typhoid requirement of the vaccine.


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Travel Health Consultant

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Does this traveller need typhoid vaccine ?


Anu is travelling to Mumbai to see relatives for a 4 week stay – she is 22 years old. She had a 1st dose hepatitis A vaccine at the age of 14 years. Would she need a typhoid vaccine and which one would you give if so?

Anu needs a booster dose of hepatitis A vaccine and she needs a typhoid vaccine so it would be very appropriate in this situation to give her a combined hepatitis A and typhoid vaccine.



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Travel Health Consultant

205

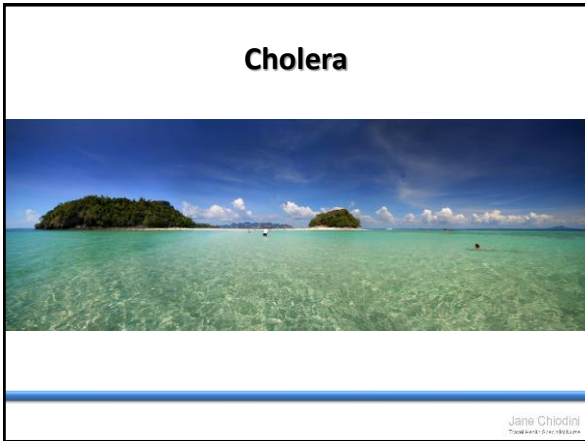


Food and water hygiene advice remain paramount

<https://travelhealthpro.org.uk/news/349/extensively-drug-resistant-typhoid-fever-in-pakistan>

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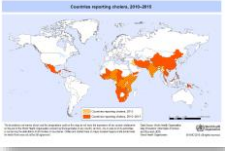


206



207

Cholera

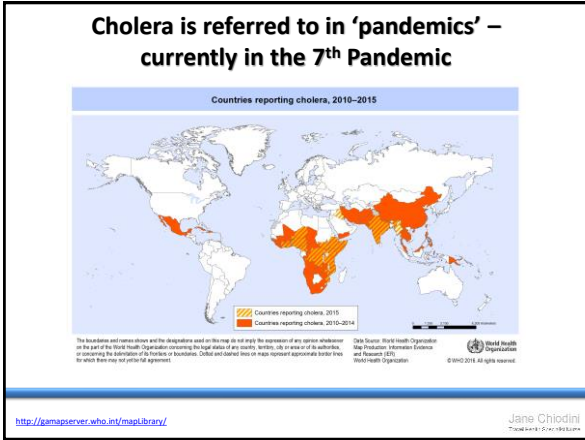
Disease	Organism	Mode of transmission
Cholera	Bacterial infection	Mainly water-borne through ingestion of faecally contaminated water or shellfish and other foods. Person-to-person spread may occur through the faecal-oral route

<http://gamapserver.who.int/maptlibrary/>

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Thailand 2019 © Jane Chiodini

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209




210



211

Cholera

- Acute intestinal infection
- Causal bacterium - *Vibrio cholerae*
- Transmitted faecal orally
- 90% cases are mild to moderate
- 10% cases very severe – leading to profuse diarrhoea, vomiting, circulatory collapse and shock
- Mortality rate can be over 50% in untreated cases, unless rapid rehydration therapy is given promptly
- Chronic carriage is rare
- Organism survives for up to 2 weeks in fresh water and 8 weeks in salt water
- Transmission normally through infected drinking water




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Management of cholera

- Fluid replacement
- Prompt action improves outcome
- IV fluids in severe cases or when vomiting
- Rapid rehydration until signs improve
- NG tube used if IV not possible
- Antibiotic therapy in severe cases



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Cholera recommendations for administration

Advice from the Green Book

- Immunisation against cholera can be considered, following a full risk assessment, for the following categories of traveller:
 - relief or disaster aid workers
 - persons with remote itineraries in areas where cholera epidemics are occurring and there is limited access to medical care
 - **travellers to potential cholera risk areas, for whom vaccination is considered potentially beneficial.**

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Cholera vaccine fact finding


- What is the youngest age at which give cholera vaccine can be prescribed?
- How many doses would you give a child?
- How many doses would you give an adult?
- What is the minimum and maximum time interval between doses?
- How long does cholera vaccine last?

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Cholera Vaccine

CHOLERA vaccine	2 years and over	2 yrs in age 6 to adult
Dukoral® Oral vaccine. 2 doses, minimum 1 wk. apart and maximum 6 weeks apart, from 6yrs of age. 3 doses, in 2 – 6 year olds		6 months in 2 – 6 year olds NBM 1 hr before & after vaccine



Food and drink should be avoided 1 hour before and 1 hour after vaccination. Oral administration of other medicinal products should be avoided within 1 hour before and 1 hour after administration of Dukoral.

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From the PiL

Instructions:

1. To prepare buffer solution dissolve the effervescent granules in a glass of cool water (approx. 150 ml). Do not use any other liquid.
Children 2-6 years: pour away half of the buffer solution.
2. Shake the vaccine bottle (1 bottle = 1 dose).
3. Add the vaccine to the buffer solution. Mix well and drink the mixture. Drink the vaccine within 2 hours after mixing with the buffer solution. Avoid food and drink starting 1 hour before until 1 hour after the vaccination.

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Charging and administration

Cholera vaccine is an NHS provision in an NHS setting


1. You could buy this vaccine in and bring the patient in for subsequent doses
2. You could administer the vaccine to the patient for the first dose and given them the 2nd dose to take home to self administer but the vaccine must be stored at 2 – 8°C
3. You could supply the vaccine on an FP10 and allow the patient to self administer

You need to ensure that your traveller understands the importance of, and can assure the cold chain in points 2 and 3 above.





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For further information to learn more about these diseases, look at the Green Book (online) NaTHNaC and TRAVAX




Tip

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Meningococcal meningitis



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Meningococcal Meningitis

- Meningococcal disease is a rare, but potentially devastating infection
- Caused by the bacteria *Neisseria meningitidis* of which there are 6 disease-causing strains called serogroups (A, B, C, W, Y and X)
- Approximately 10 percent of the general population of the UK are thought to carry *N. meningitidis* in the lining of the nose and throat
- Spread between individuals occurs through coughing, sneezing, kissing or during close contact with a carrier
- Carriers do not have symptoms, but can develop disease when bacteria invade the bloodstream from the nasopharynx
- Invasive disease is a rare but serious outcome usually presenting as septicaemia or meningitis


<http://travelhealthpro.org.uk/diseases/meningococcalmeningitis/>

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Meningococcal Meningitis

- Less commonly, individuals may present with pneumonia, myocarditis, endocarditis, pericarditis, arthritis, conjunctivitis, urethritis, pharyngitis and cervicitis



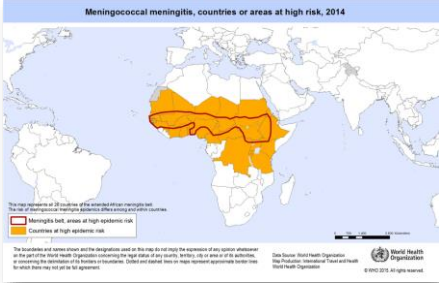
- The incubation period is from two to seven days

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/223749/Green_Book_Chapter_22_v2_3.pdf

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Meningococcal meningitis vaccine given to travellers going to meningitis belt in Africa



Meningococcal meningitis, countries or areas at high risk, 2014

The map indicates all 28 countries of the meningitis belt. The risk of meningococcal meningitis is highest in the meningitis belt. Countries at high epidemic risk are highlighted in orange.

The boundaries and names shown and the designations used on this map do not imply the expression of any opinion whatsoever on the part of the World Health Organization concerning the legal status of any country, territory, city or area or of its authority, or concerning the boundaries and boundaries. ©WHO and NaTHNaC. All rights reserved.

http://pamapserver.who.int/mapi/library/Files/Maps/Global_MeningitisRisk_ITI@RiskMap.png

<http://www.ncdc.gov/travel/yellowbook/2016/infectious-diseases-related-to-travel/meningococcal-disease>

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Vaccine recommendation for travellers to the meningitis belt

Meningococcal disease vaccination

Vaccination is recommended for those whose activities or medical condition put them at increased risk including:

- healthcare workers
- those visiting friends and relatives
- those who live or travel 'rough' such as backpackers
- long-stay travellers who have close contact with the local population
- those with certain rare immune system problems (complement disorders) and those who do not have a functioning spleen

From NaTHNaC ←

Who Should I Vaccinate for Meningococcal Meningitis?

Consider vaccinating:

- Travellers who are likely to have close, prolonged contact with the local population.
- Long stay travellers.
- Those visiting friends and relatives
- Those who will be exposed to crowded areas (e.g. stadia, schools, dormitories, hospitals)
- Travellers visiting an area affected by an ongoing outbreak or epidemic.
- Immunocompromised travellers (including asplenia) visiting endemic areas.

From TRAVAX →

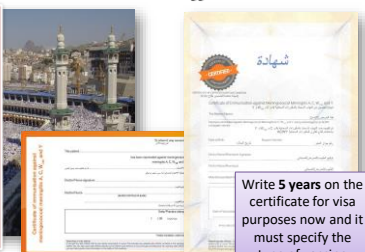
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and travellers going on a pilgrimage to Umrah and Hajj

The KSA Ministry of Health (MoH) currently recommends that, as a precautionary measure, pregnant women and young children, should postpone the performance of the Hajj and Umrah.

Those with severe medical conditions such as terminal cancers, advanced cardiac, respiratory, liver, kidney diseases or senility are exempt from these religious duties



Write **5 years** on the certificate for visa purposes now and it must specify the type of vaccine

Note – information given that vaccine must be given minimum of 10 days prior to entry into the country

Certificates available at:
https://hsa.gov.uk/content/dam/global/hcpportal/en_GB/theravareas/vaccines/pdf/meningococcal-acwy-certificate.pdf
 and <https://globevaccines.media.com/media/7wicket/interfacel3...>

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The ACWY vaccines

Menveo

- Conjugate vaccine
- Use from two years of age
- Available from GSK
- GSK data gives 5 years protection from administration

Nimenrix

- Conjugate vaccine
- Use from 6 weeks now*
- Just had black triangle removed
- Available from Pfizer
- Pfizer studies up to 60 months – refer to pharmacodynamics properties in the SPC


* If needing to give, please check the Green Book, the SPC and TravelHealthPro

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/302904/Green_Book_CBPpgr_22_v2_5.pdf, Jane Chiodini
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When would you boost - it's been very confusing!

- The Joint Committee on Vaccination and Immunisation (JCVI) Committee reviewed information on length of protection following ACWY conjugate vaccination. Antibody against serogroup A disease was the first to wane, and this meant boosting was important for travel, but less important for the routine Men ACWY programme in the UK.
- For travellers at continued risk, the Committee agreed that boosting **every five years** would be a sensible approach until data became available.

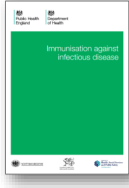


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Tip


For further information to learn more about these diseases, look at the Green Book (online) NaTHNaC and TRAVAX



TRAVAX
Est. 1985

TRAVEL HEALTH PRO

NaTHNaC



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Agenda

- ✓ Introduction to travel medicine
- ✓ Travel risk assessment
- ✓ Travel vaccines and related issues
 - Travel medicine operational issues
 - Recap on resources

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Immunisation Training

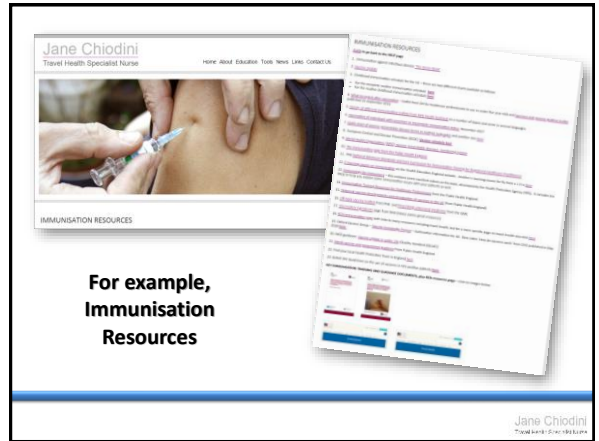


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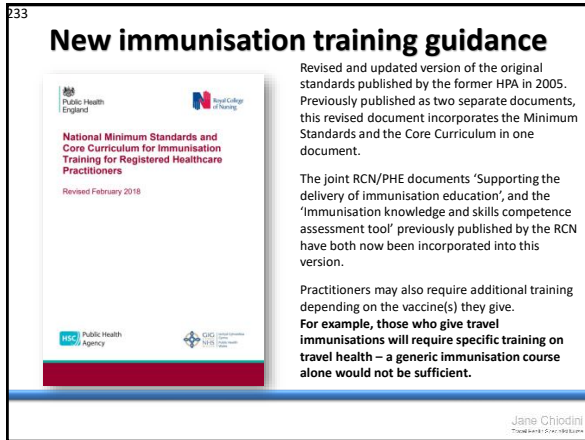
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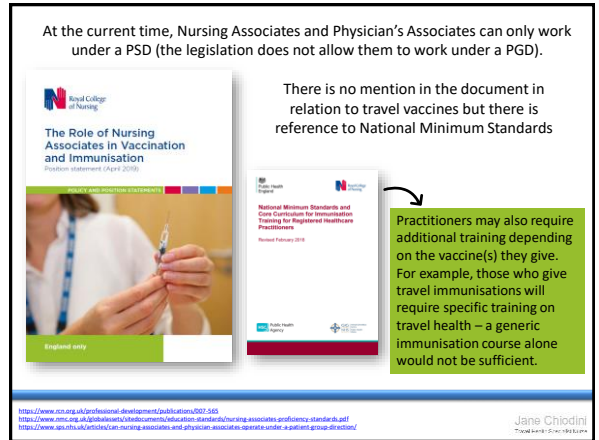
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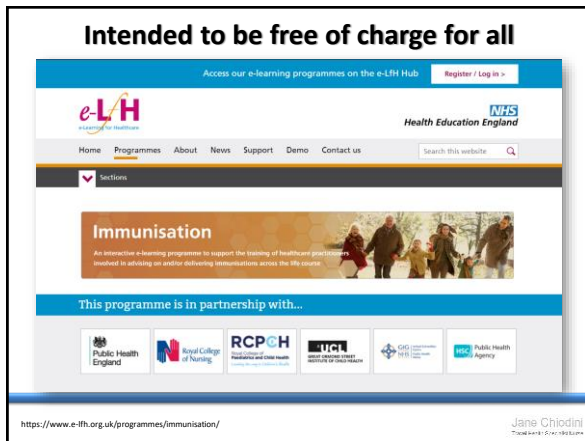
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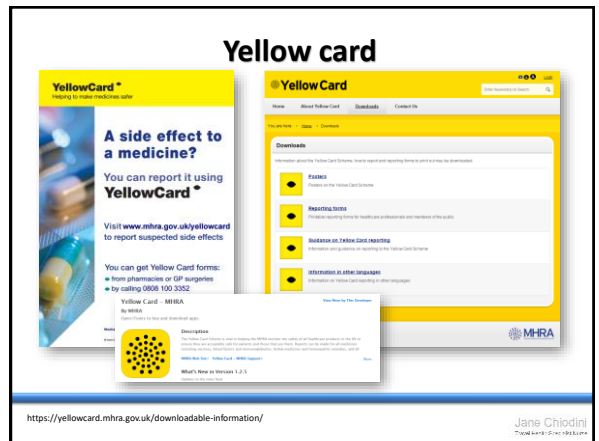
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234



235



236

Licensed, Unlicensed and Off-label

https://www.gov.uk/government/uploads/system/uploads/attachment_data/file/399317/2HE_9173_VU_223_Doc_2014_11.pdf

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Immunisation training includes...

- CPR and anaphylaxis
- Consent
- Prescribing
- Administration
- Documentation
- Vaccine storage – protocol
- Finance

Not part of the National Standards, but topics covered briefly on day 2

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Travel Health Specialist Nurse

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www.resus.org.uk

Annual training for CPR and anaphylaxis should be undertaken

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Travel Health Specialist Nurse

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Vaccine ordering, storage and handling

The perfect fridge!

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Travel Health Specialist Nurse

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Prescribing for travel medicine

Legal requirement – covered on day 2

<https://www.nice.org.uk/guidance/mpg2>

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FAQs (under News) for information on prescribing

<http://www.janechiodini.co.uk/news/faqs/faq-no-1/>

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Travel Health Specialist Nurse

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Consent




Chapter 2 in the 'Green Book'

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Thank you to my students

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Administration

- Cleansing the skin
- Size of needles
- Preparing the vaccine
- Post vaccination waiting time ?



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Equipment



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Importance of Documentation – working within your code



<http://www.nmc-uk.org/>

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Resources for a travel service



247

247

Agenda

- ✓ Introduction to travel medicine
- ✓ Travel risk assessment
- ✓ Travel vaccines and related issues
- ✓ Travel medicine operational issues
- Recap on resources

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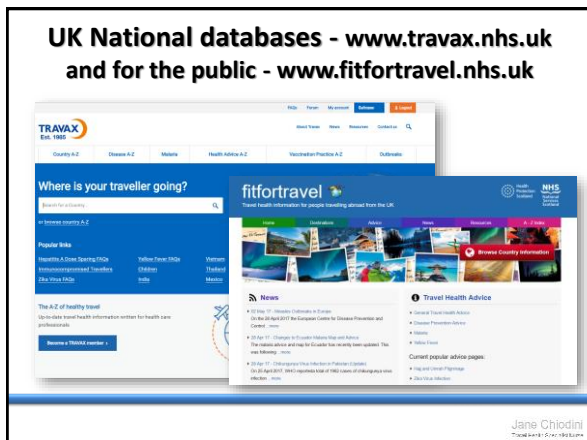
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249



250



251



252



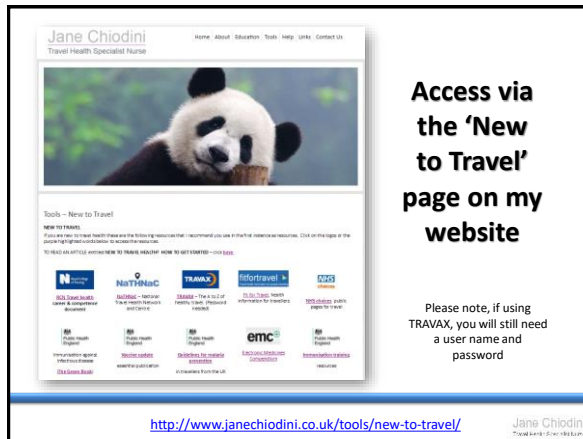
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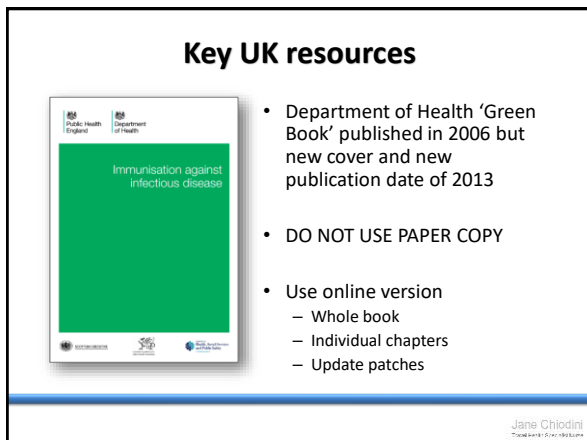
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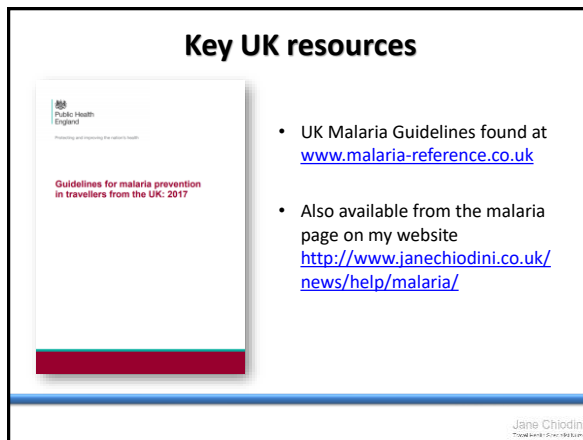
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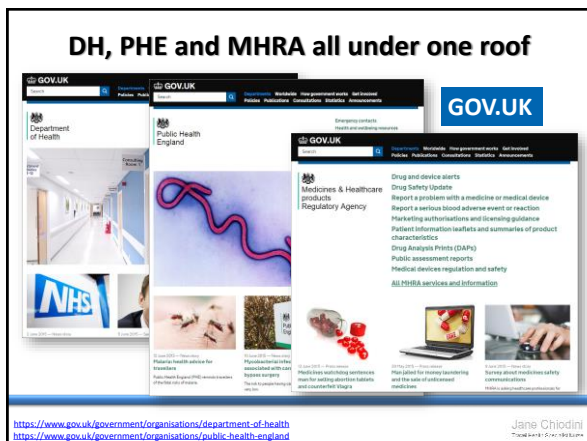
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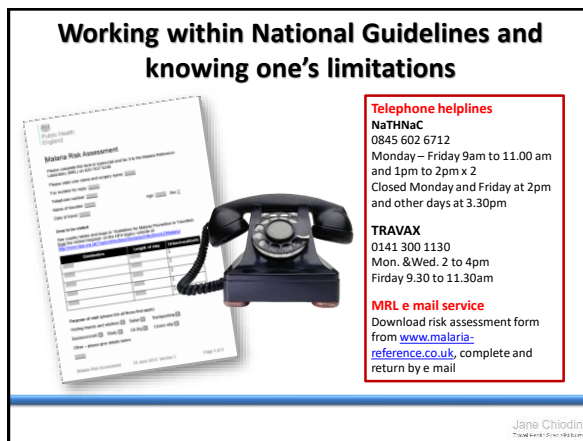
257



258



259



260

Electronic Medicines Compendium

www.medicines.org.uk – provides SmPCs and PILs

Don't forget the protected login area to store your own choices

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Travellers' Clinician

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Immunisation training

National Minimum Standards and Core Curriculum for Immunisation Training for Registered Healthcare Practitioners

Revised February 2018

<https://www.janechiodini.co.uk/help/immunisation-resources/>

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Travellers' Clinician

262

Finding additional destinations

Google search www.google.co.uk
and/or google maps
<http://maps.google.co.uk/>

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Travellers' Clinician

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Worldwide resources

TRAVAX from Shoreland is not the same as UK TRAVAX

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Travellers' Clinician

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Other useful maps also at www.who.int/ith

http://gamapserver.who.int/maplibrary/Files/Maps/Global_CholeraCases_ITHRiskMap.png

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265

World Health Organization

www.who.int

or Google title of the WHO Factsheet required

an extremely useful website with so much information e.g.

Substandard and falsified medical products

- Substandard and falsified medical products may cause harm to patients and fail to treat the diseases for which they were intended.
- They lead to loss of confidence in medicines, healthcare providers and health systems.
- They affect every region of the world.
- Substandard and falsified medical products from all main therapeutic categories have been reported to WHO including medicines, vaccines and in vitro diagnostics.
- Anti-infectives and antibiotics are amongst the most commonly reported substandard and falsified medical products.
- Both generic and brand name medicines can be falsified, ranging from very expensive products for cancer to very inexpensive products for treatment of pain.
- They can be found in illegal street markets, via unregulated websites through to pharmacies, clinics and hospitals.
- An estimated 1 in 10 medical products in low and middle-income countries is substandard or falsified.
- Substandard and falsified medical products contribute to antimicrobial resistance and drug-resistant infections.

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Travellers' Clinician

266

Finding travel clinics abroad www.istm.org

International Society of Travel Medicine
Promoting healthy travel worldwide

The 16th Conference of the International Society of Travel Medicine
5-9 June 2019
Washington, DC
United States of America

Global Travel Clinic Directory

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Additional websites and resources on my website

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Home About Education Tools News Links Contact Us

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Interim work between the two study days listed on your webpage

- Play the vaccine videos to recap on this aspect
- Look around my website
- Find out what travel PGDs you have at work
- See if you have a vaccine storage protocol at work
- Do the practice case study e learning on your page if you have time left
- Remember you can go into the e learning to reflect on today's presentation if you wish – but only available for a limited time.

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Travel Health Specialist Nurse

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Don't forget to view the tasks suggested if you can for the interim period. See you on Friday 9th August 2019

Jane Chiodini
Travel Health Specialist Nurse

270