Baseline investigations for a child with suspected HIV and newly diagnosed HIV

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If history and examination lead to a differential diagnosis, which includes HIV, pay special attention to:

History:	
Birth:	Mode of delivery; duration of rupture of membranes; other infections eg. chorioamnionitis; birth weight; infant feeding eg. breast feeding and duration; maternal health including other STI's; and antenatal HIV test result.
Past medical history:	Previous infections (eg. oral candida); swollen lymph nodes; chronic diarrhoea; failure to thrive and nutritional history; recurrent URTIs; childhood exanthems (e.g. chickenpox, rubella); skin infections (eg warts, molluscum); severe infections; hospitalisations; transfusions and IM injections; TB risk factors; immunisations; developmental history; and sexual history (if appropriate).
Social history:	Name and relationship of adult accompanying child; who has parental responsibility; deaths of parents or siblings; significant previous caregivers; travel history; housing; and social circumstances; school attendance and performance.
Drug history:	Previous antiretroviral exposure: in-utero / peripartum / as treatment in another country; other current drugs. Children newly arrived from abroad may be on combination ARV tablets not available in this country; check with a specialist HIV pharmacist.
Examination:	Full examination including: mouth; lymph nodes; parotids; chest; liver; spleen; skin; neurology; developmental assessment, growth (ht, wt, OFC, BMI); pubertal stage (if indicated from screening in history); and BCG scar. Any signs of lipodystrophy if on treatment.

First line HIV diagnostic tests:

Infant < 18 months of age: HIV antibody test and HIV RNA PCR (preferred to HIV DNA PCR in local lab with faster results available).

* NB in the first weeks after delivery an infant at risk of HIV may have a negative RNA PCR. **Child > 18 months of age:** HIV antibody test

Second line confirmatory HIV tests:

HIV RNA PCR viral load and assessment of severity of HIV disease. (If HIV known or clinically very likely then consider doing both first and second line tests together.)

See CHIVA HIV Testing guidelines: https://www.chiva.org.uk/infoprofessionals/guidelines/testing/

HIV parameters	CD4 count and percentage	
	HIV RNA PCR (viral load)	
	Baseline HIV resistance including integrase resistance (and maternal	
	resistance if an infant)	
	HLA-B*5/01	
Haematology	FBC + film	
	Sickle cell and G6PD deficiency screen (if appropriate racial group)	
	Ferritin	
	Consider malaria film if recently arrived from endemic area	
Biochemistry	U+E, Creat Glucose TSH Vitamin D	
	Ca, PO ₄ Amylase Albumin	
	LFT's Lipids Total protein (globulin)	
	Urine dip (mid-stream) – if 1+ or more protein send urine protein/Cr and	
	albumin/Cr ratio (ideally early morning sample)	
Serology	Hepatitis A IgG, HBsAg, anti-HBsAb, anti-HBcAb, HCV IgG, Syphilis	
	serology, IgG for EBV, CMV, HSV, VZV, Toxoplasmosis and SARS-	
	In children over 1 year consider vaccine serology as per CHIVA	
	vaccination guideline: Measles, Mumps, Rubella, Hib, MenC, Tetanus,	
	serotype specific pneumococcal serology and if appropriate SARS-CoV-	
	2 serology.	
1/2 1 202	NB. Low CD4 count could affect serology results	
Viral PCRs	Plasma CMV PCR should be undertaken in infants & children with	
	advanced disease	
	HCV PCR – should be undertaken in infants at risk of exposure and	
	those with advanced disease (this can be positive even if the child is	
Oulture	HCV antibody negative)	
Cultures	According to symptoms / travel history:	
	Stools (including ova, cysts and parasites) / unine / throat swabs / blood	
TD correction	Cultures / malaria films / sexual nealth screen if appropriate	
IB screening	CXR, mantoux, IGRA	
Oliviaal	If active TB suspected – consider gastric aspirate, induced sputum, BAL	
	BP, urinalysis, neight / weight / nead circumference	
Investigations	Formal ophtnalmological examination	
Radiology	Baseline CXR	
	Consider bone age (if small for age) with advice of endocrine specialist	
	infants / children with neurological signs, evidence of congenital	
Development	Intections of severe co-intection: MKI of brain	
Development	Full formal baseline neurodevelopment/neuropsychology assessment if	
Assessment	available of clinically indicated	

PCP Prophylaxis

Infants < 12 months of age	Children > 1 year of age
If > 6 weeks and under 12 months of age,	Start Co-trimoxazole
start Co-trimoxazole irrespective of CD4	1-4 yrs: CD4 count <15% or <500 x 10 ⁶ /L
count	5 yrs or older: CD4 count <15% or <200 x 10 ⁶ /L

Important: Any child that is diagnosed with HIV and was born in the UK, should be investigated as an incident and reported back to the obstetric unit where they were born, as all infant infection is potentially preventable.

Assess the child's clinical stage according to WHO and/or CDC criteria. More information on treatment of HIV infected children in the PENTA guidelines: https://penta-id.org/hiv/treatment-guidelines/

For newly diagnosed, see PENTA risk calculator, where the child's 12 month's risk of progression to AIDS and death can be checked: <u>https://penta-id.org/education/educational-tools/</u>