

7 Neurology

STROKE

ABCD2 for TIA

Age > 60	1pt	
BP ≥ 140/90	1pt	0-2 = low risk
Clinical feat:		= 0.2-0.8% risk of stroke next 7d
Unilateral weakness	2pt	
Speech disturbance	1pt	
Other	0pt	
Duration		
≥ 60min	2pt	
10-59min	1pt	
<10min	0pt	
Diabetes	1pt	

Admission criteria for TIA

- 4 in 2 weeks, 2 in 24hrs
- High-grade **carotid stenosis**
- Crescendo TIA
- Likely **cardiac source** (AF)
- Anticoagulated

Stroke Prevention

- **Aspirin** 150mg/day
- Carotid endarterectomy
- **Warfarin**

Stroke Mimics

MISTA

- Migraine
- Infection: Sepsis, Cerebral
- Seizure eg Todd's paresis, SOL
- Trauma, Tox
- **Aortic dissection**

Cerebellar Signs

RAIN

- Rapid alt movt (dysidiokinesis)
- Ataxia (esp truncal)
- Intention tremor, past pointing
- Nystagmus

Stroke Syndromes

ACA: Contralateral LL paralysis, confusion, behaviour, urinary incontinence

MCA: Contralateral hemiplegia UL > LL, hemisensory loss, homonymous hemianopia

Dom MCA = global aphasia; Non-dom MCA = neglect, contractual apraxia

- M1 global aphasia, L side neglect
- M2 expressive aphasia, L side neglect
- M3 receptive aphasia, minimal weakness

PICA: multiple CN syndromes

- LMS CN 7,9,10, CL SL tract
- **Wallenberg's syndrome** (LMS)
 - ipsilateral facial paraesthesia
 - contralateral arms/legs pain/temp
 - seen in vertebral artery dissection

Stroke Thrombolysis

ACEM Independent Review

- NNTB @ 3hrs = 7
- NNTH (ICH) = 42
- NNTH (Death) = 122

NINDs

	tPA	Placebo	
Improved @ 3m	47	39	+8%
90-day mortality	17	21	- 4%
ICH < 36hrs	6.4	0.6	+5.8%
ICH < 3m	7.7	1.3	+6.4%
New stroke	5.8	0.4	+0.4%

Medication

- **Alteplase** 0.9mg/kg max 90mg
- 10% as bolus over 1min then rest over 1hr

Indications

No perfusion scan

- < 1/3 MCA territory
- < 25 **NIHSS score**
- < 90min to lysis
- < 80yo

Perfusion scan

- < 1/3 MCA territory
- < 25 **NIHSS score**
- < 6hr
- Demonstrated perfusion
- NOT ICA or Basilar

Contraindications

Absolute (ST. ABCD)

- Stroke < 3m
- Trauma, sig to face or head
- Anytime **ICH**
- Bleeding, active (not mens)
- Cerebral **mets** or vasc **malformation**
- Dissection, suspected Ao

Relative

- Isch stroke > 3m
- Dementia
- Pregnancy
- Trauma wishes
- CPR > 10min
- Current anticoagulation
- Non-compress vascular puncture
- Major surgery < 3w
- Internal bleeding incl PUD
- Severe HTN SBP > 180
- Intracranial abnormality

General Stroke Mx

- BP < 220/120 (**labetalol** 10-20mg IV)
- BP < 200 if haemorrhagic stroke
- Prevent pressure sores, aspiration

ACUTE VESTIBULAR SYNDROME

severe vertigo, N&V, spont nystagmus, postural instability

Causes

- Peripheral: **vestibular neuronitis, labyrinthitis**
- Central: **posterior infarct**

HINTS Test

Differentiate central vs peripheral AVS

Head Impulse

- move head L&R with eyes fixed on nose
- head turn to effected side = **corrective saccade**
- means **peripheral cause**

Nystagmus

- Peripheral: unidirectional (lesion fast phase), horizontal
- Central: rotational, vertical, or direction-changing

Test of Skew

- Look at nose, cover eye; uncover eye & look at gaze
- Gaze fixed to nose = -ve skew = peripheral
- Gaze corrects to nose = +ve skew = central

Peripheral Vertigo

Vestibular Neuronitis

- Most common AVS
- Related to **viral illness**
- Can try **prednisolone** 125mg for 3d → taper

Labyrinthitis

- Rare
- Essentially same as vest neuritis but with **hearing loss**
- Usually toxic as well

BPPV

- Stuck **otoliths**
- Most common cause of vertigo (not AVS)
- **Episodic**, sudden onset
- Antiemetics, **stemetil, largactil**, Eply's manoeuvre

Meniere's

- Lack of **endolymph** resorption (or excess production)
- **Episodic** but **Tinnitus!** ≠ BPPV
- Restrict salt intake, avoid caffeine, try **thiazides**

HEADACHE

DDx Headache

- Infection: Meningitis, encephalitis, sinusitis
- Ischaemia: CVA, TIA
- Haemorrhage: Trauma, SAH
- Vascular: Dissection, GCA, venous thrombosis
- O&G: Eclampsia
- Cardio: HTN
- Eyes: glaucoma, optic neuritis
- Neuralgia: trigeminal, post-herpetic
- Systemic illness (sinister in 33%)
- Withdrawal
- Sinusitis
- Headache syndromes
- TMJ syndrome
- Intracranial HTN
- Tumour
- Post-LP

Drug Classes in Headache

- Simple analgesia/Opiate
- Serotonin agonist eg **tryptans, ergots**
- Dopamine agonists eg **largactil, stemetil**
- Other: **Dexamethasone**
- **Amtriptyline, Gabapentin**
- **Haloperidol, droperidol**
- **Propofol**

DELIRIUM

- Treat underlying cause
- Familiar faces, less stimulation
- Rx: **respiridone** 0.5mg
- **haloperidol** 0.5mg
- **olanzapine** 2.5mg
- Prevention eg hearing aids, visual aids, prevent sleep deprivation, maintain mobility, low stimulation environment

Delirium vs Dementia vs Psychosis

	Delerium	Dementia	Psychosis
Main	Confusion	Memory loss	Loss of contact with reality
Onset	Acute	Gradual	Either
Course	Fluctuant	Progressive	Chronic with exacerbation
Duration	Hrs-Mths	Mths-Yrs	Mths-Yrs
Conscious State	Clouded	Normal	Normal
Attention	Impaired	Normal (until late)	Normal
Orientation	Fluctuant	Poor	Normal
Speech	Incoherent	Mild errors	Normal
Thought	Disorganised	Impovershed	Disorganised
Illusion/Hallucination	Common visual	Rare	Common
Reversible	Usually	Rare	Rare
EEG	Mod-severe bckgrd slowing	Normal-mild	Normal

ALTERED CONSCIOUSNESS

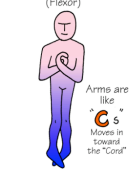
Glasgow Coma Scale

	1	2	3	4	5	6
E	None	Pain	Voice	Open	-	-
V	None	Incomp	Inapp	Conf	Orient	-
M	None	DecEr	DecOr	Wdraw	Local	ObeY

DecErebrate

- extensors
- straight elbows
- pointed toes

DECORTICATE (Flexor)



Problems With Cervical Spinal Tract or Cerebral Hemisphere.

DDx ALOC

AEIOU TIPS

- **A**cidosis, arrhythmia, NH₄, Etoh
- **E**pilepsy
- **I**nfection
- **O**verdose, underdose
- **U**raemia
- **T**rauma
- **I**nsulin
- **P**sychogenic
- **S**troke, SOL

Coma Cocktail

- 50ml 50% **Dextrose**
- 400 mcg **Naloxone**
- 100mg **Thiamine**

Reye's Syndrome

- **Aspirin** associated ∴ only use in Kawasaki in kids now
- Rapidly progressive encephalopathy
- Tx supportive (mostly target cerebral oedema)

AUTOIMMUNE

Guillain Barre Syndrome

- Sensory & motor
- Demyelination, infection (**campylobacter, CMV, EBV**)
- Predominant **symmetrical ascending flaccid weakness**
- Loss of DTRs
- **Ascending glove & stocking paraesthesia**
- **Miller-Fischer** variant = CN 1st
- **↑ CSF protein (> 0.55 g/L)**, all else ok (if WCC ↑ not GBS)
- Don't use **sux**, use **IVIG** or **plasma exchange**

Myasthenia Gravis

- AChR autoantibodies = pure motor
- **Isolated muscle weakness (esp ocular, prox limbs)**
- **Fatigability**, relieved with rest
- Mx **pred**, thymectomy, **IVIG**, plasma exchange
- **Pyridostigmine, neostigmine** regular

Complications

- **Myesthenic crisis:** common, missed dose
- **Cholinergic crisis:** OD on **Pyridostigmine**

Eaten-Lambert

- Proximal weakness
- **Lambert sign** - better with exercise

Multiple Sclerosis

- Demyelination from loss of oligodendrocytes
- LP **oligoclonal bands**
- Mx: **methylpred, ACTH**, plasma exchange

Common manifestations

- Cord: limb weakness & UMN signs
- **Optic neuritis** (esp colour), worse after heat
- Brainstem: diplopia etc
- Cerebral: seizure, ↓IQ

SEIZURES

- **6% population** have a non-febrile seizure
- 50% of the 6% have a 2nd episode
- 75% of the 50% go on to have epilepsy (~2%)

Cause

- Idiopathic
- ARF
- Trauma
- SOL
- Degen
- Infection
- Congenital malformation
- Metabolic: low Na, Glc, Ca
- Vascular (intracerebral or pre)
- Drugs & OD eg sertraline
- Structural brain lesion
- **Non-compliance**

Causes (neonates)

- hypoxia, haemorrhage, hypoglycaemia
- pyridoxine def
- infection
- drug withdrawal
- familial

Seizure Mimics

- TIA
- Syncope
- Arrhythmia
- Movement disorder
- Sleep disorder
- Psychogenic
- Migraine

Risk Factors for Recurrence

Adults

- < 50yo
- FHx
- Hx of neuro insult eg trauma, hypoxia
- Structural lesion
- 2 in 1 week

Paeds

- Abnormal EEG
- Seizure in sleep
- Complex Febrile seizure
- Todd's paresis

Pseudo-seizure Features

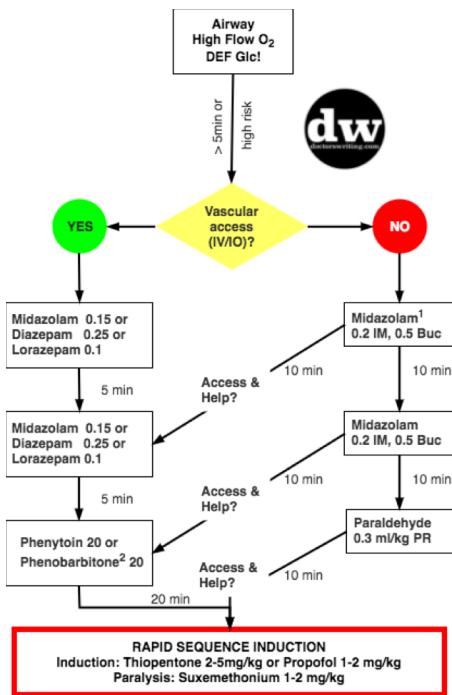
Presence of:

- Side-to-side head movt
- Pelvic thrusting
- Non-sync movt
- Out-of-phase movt
- Eyes look away

Absence of:

- No tongue biting
- No incontinence
- No post-ictal state
- Avoidance manoeuvres
- Normal serum lactate

Paed Mx



All doses are mg/kg unless otherwise stated

¹Diazepam PR 0.5 mg/kg is also available

²Use phenobarbitone if already on phenytoin

Seizure Drugs in Paeds

	Drug	Dose ¹	Route
1st	Midazolam	0.15, 0.20, 0.50	IV/IM/Bc
	Diazepam	0.25, 0.5	IV/PR
	Lorazepam	0.1	IV
2nd	Phenytoin	20	IV
	Phenobarbitone ²	20	IV
	Paraldehyde	0.3	PR
RSI	Thiopentone	2-5	IV
	Propofol	1-2	IV
	Suxemethonium	1-2	IV
PD ³	Pyridoxine	10mg	IV

¹All doses mg/kg except paraldehyde ml/kg

²Use if already on phenytoin

³Pyridoxine def doesn't respond to others - seen from 0-6m

Febrile Seizures

3% incidence

	Simple	Complex
Seizure	Generalised	Focal
Duration	< 15min	≥ 15min
Recovery	< 1 hr	≥ 1hr
Neurology	No	Focal
Episode	1 st	> 1 st for illness

Prognosis

- recurrence **1/(1 + age)** (1yo 50%, 2yo 30%)
- epilepsy:

Risk Factor	Interpretation
- Complex febrile seizure	No RF 01%
- Low fever	1 RF 02%
- Low age	> 1 RF 10%
- Neurodevelopmental	
- FHx	

Seizure Complications

Kids

- 4% mortality if status
- arrhythmia
- HTN
- APO
- Hyperthermia
- DIC

Neonate

- CP
- Spasticity
- Feeding difficulties

Status

- cont seizure or no return to normal > 30min
- start Tx at 5min (most abort at 2min)

Adult Mx

- Glc ± Thiamine
- A: left lateral
- B: O₂ high flow
- C: IV access
- D: Abort seizure
- 1st line: Midazolam 5-10mg IM/IV
- 2nd line: Midazolam infusion
- D: Prevent further seizures
- Clonazepam 1mg BD for 3/7 or
- Phenytoin 2g max 50mg/min
- Look for cause

MENINGITIS

Bacterial

- S pneumoniae or N meningitidis
- < 2m old GBS, e coli, listeria
- > 2m old S. pneumoniae, n meningitidis, HiB

Aseptic

- Viral: Mumps, CMV, HIV, EBV, Coxsackie
- Fungi/Parasites: Cryptococcus, toxoplasma
- Autoimmune: SLE, sarcoid, Wegners

Atypical Bacteria

- TB, Mycoplasma, Borrelia, Treponema

Meningitis Mx

- Dexamethasone 10mg (0.15mg/kg)
- Ceftriaxone 4g (100mg/kg), Vanc/Cipro 400mg if allergic

± BenPen 2.4g to cover listeria if at risk (immunocompromised, > 50, alcohol, pregnant, debilitated)

- ± Vanc if mastoid
- ± Aciclovir 10mg/kg
- < 2m cefotaxime 50mg/kg + BenPen 60mg/kg

LUMBAR PUNCTURE

Indications

- SAH
- Infection
- Autoimmune: GBS, MS, MG
- Benign Intracranial HTN (BIHTN)

Contraindications

- Anticoagulation
- Coagulopathy
- Plt < 50
- ALOC
- Seizure
- ↑ ICP
- Focal neurology
- Cushing's reflex

Intracranial Pressure

- normal adult 5-13 mmHg
- normal adolescent 5-15 mmHg
- normal < 8yo 1-7 mmHg

Causes of Raised

- Infection incl abscess
- Tumour
- ICH incl SAH
- BIHTN

Causes of Lower

- Hyperventilation
- SAH block
- CSF leak
- Diabetic coma
- Dehydration
- Degeneration

Complications

- Herniation/coning
- Post LP headache
- Disc laceration
- Infection
- Poor specimen
- Spinal epidural haematoma
- Worsening paraparesis
- Nerve root puncture/graze
- Intra-spinal epidermoid cyst

Post LP Headache

2-15% incidence in 24-48hrs

Risk

- technical: large needle, bevel tip, orientation of tip, no stylet when withdrawing
- Pt: young, female, Phx same, low BMI, headache prior

Mx

- Fluids
- Rest supine
- Caffeine
- Blood patch (unreliable)

CSF/LP Interpretation

	White cells		Biochem		
	Neut	Lymph	Protein	Glc: ¹	G Stain
Normal	0	5	0.4	0.6	-ve
Bacterial	100-10k	<100	>1	< 0.4	80% +ve
BPT ²	100-10k	100-10k	>1	<0.4	20% +ve
Viral	< 100	100-10k	0.4-1.0	0.6	-ve
TB	< 100	50-1k	1-5	< 0.3	-ve

¹CSF:Blood

²Bacterial - partially treated

NB abscess has HIGH protein

NEUROLOGICAL TOXINS

Botulism

- Ach blocker
- Cause: poor canning, honey, corn syrup
- Ex: anticholinergic, cranial/bulbar early (ie descending)
- Mx: lavage/charcoal, antitoxin
- Adults: equine botulism antitoxin
- Infants: human botulism Ig 100mg/kg
- **DON'T GIVE AMINOGLYCOSIDES** (releases more toxins when bacteria dies)

Diphtheria

- Toxin destroys everything
- Myocarditis or Hepatitis cause death
- Motor paralysis is earliest sx
- Mx: Antitoxin 20K-120K units

Tetanus

- Clostridium → Tetanospasmin **migrates along nerves**
- Inhibits GABA = **sustained paralysis**

Clinical Features

- Triad: **rigidity + muscle spasm + autonomic dysfunction**
- Incubation 3-14d → onset 1-7d → rigid 2w → spasm 2w → auto dysfunction 2w
- Resp: chest wall rigidity, laryngospasm, aspiration
- Autonomic: ↑HR, BP, Salivation, Bronchorrhoea
- MSK: Rhabdo/AKI

Acute Management

- Source control: metronidazole, Ig (300u/kg), Vaccine
- High dose relaxant & sedation
- Rigidity: BZDP, MgSO₄

Prophylaxis

Time ¹	Wound	TTox	Tlg ²
Complete Immunisation (3x TTox)			
< 5y	all	No	No
5-10y	clean/minor	No	No
	all other	Yes	No
> 10y	All	Yes	No
Incomplete Immunisation (< 3 TTox)			
	clean/minor	Yes	No
	all other	Yes	Yes

TTox Tetanus Toxoid vaccine; Tlg Tetanus Immunoglobulin

¹ Time since vaccination

² Provides 4 weeks protection; 250IU IM < 24h, 500IU IM > 24h, 4000IU IV if clinical evidence of tetanus

Tetanus Prone Wound

ABCDEF

- Avulsed tooth re-implanted
- Burns, Bites
- Compound #, Crush
- Deep/penetrating
- Environmental: obvious contamination eg soil, manure
- FB (esp wood splinters)

Indications for Tlg

Tetanus prone wound and:

- Immunocompromised eg HIV
- Evidence of tetanus
- Vaccination incomplete/unknown
- Allergy to TTox

Prognosis (Poor)

RUST

- Route of spread: umbilicus, uterus, burns, #, IM injection
- Unwrap: incubation < 7d; Onset < 48h
- Spasm
- Temp > 38.4, Tachy > 120 (adults) > 150 (neo)

HICCUGH DDX

- Stroke
- Gastric distension
- Lung Ca
- Pre-term baby
- Uraemia
- Low Na or Ca
- Vagal or phrenic n. stimulation

Mx

- **Largactil**, nebulised **lignocaine**, pharyngeal stimulation, Valsalva, CPAP mask

DYSTONIC REACTIONS

- not genetic or dose related
- oculogyric crisis prominent
- can be delayed (50% D1; 95% by D5)

Cause

- anti-emetics
- anti-psychotics
- anti-histamines
- tox: CO

Mx

- **Benztropine** 2mg IV then 2mg PO TDS 48hr

DDx

- Akathisia (anti emetics)
- Tardive dyskinesia (antipsychotics)
- Chorea (antipsychotics)

NEURO SIGNS & SYMPTOMS

Cranial Nerves

- UMN vs LMN: U has sparing of forehead
- Bells' **Pred** 1mg/kg max 100mg OD 5d w/in 48hrs
- Ramsay Hunt: CN VII, EAC, **valaciclovir** 1g TDS

Visual Field Defects

Defect	Anatomy	Causes
Tunnel	Concentric diminution	Glaucoma, papilloedema, syphilis
↑ Blind Spot	ON enlarged	↑ ICP
Central Scotoma	ON → Chiasm	Demyelinating, toxic
Unilateral LOV	Ocular, ON	Vascular, tumour
BT Hemianopia	OC	Pit tumour
HO Hemianopia	OT → OC	Vascular, tumour
UQ Hemianopia	Temporal	Vascular, tumour
LQ Hemianopia	Parietal	Vascular, tumour

ON Optic Nerve; ICP Intracranial Pressure; LOV Loss of Vision; OC Optic Chiasm; OT Optic Tract; BT Bitemporal; HO Homonymous; UQ Upper Quadrant; LQ Lower Quadrant

Parietal Lobe Defects

Dominant

- **AALF**
- **Acalculia**
- **Agraphia**
- **Left-right disorientation**
- **Finger agnosia**

Non-dominant

- Graphaesthesia
- Tactile extinction

Dysphasia

BeW! Brocas = Broken speech

- **Brocas** Expressive dysphasia (motor)
Dom frontal lobe
- **Wernickes** Receptive dysphasia (sensory)
Dom temporal lobe
- Arcuate Fasciculus Conductive dysphasia

WEAKNESS

Causes

Non-neuromuscular

- Anaemia
- Chronic disease
- Malignancy
- Psychological
- Electrolytes eg ↓ K, Ca
- Chronic Fatigue
- Rheum
- Dehydration
- Malnutrition
- Cardiac failure
- Rx eg **statins**
- Sepsis
- ↓ Thyroid

Neuromuscular

- Autoimmune: GBS, MS, Myasthenia
- Cord compression
- Environmental: botulism, tetanus, tick paralysis, marine
- Myopathy: HypoK periodic paralysis, Cushing's, Addison's

Clinical Features of Neuromuscular Weakness

	UMN	LMN	NMJ	Myopathy
Atrophy	-	+++	+	+
Fasc	-	Common	-	-
DTR	++	-	-	-
Tone	Spastic	Flaccid	Flaccid	Reduced
Distribution	Region	Distal	Variable	Prox>Dist
Plantar	↑	↓	↓	↓

UMN Upper Motor Neuron; LMN Lower Motor Neuron; NMJ Neuromuscular Junction; Fasc Fasciculations; DTR Deep Tendon Reflexes

GAIT

Types of gait

- Broad-based = cerebellar
- Hemiplegic = cerebral
- Tabetic: foot slapping (sensory/dorsal column)
- Steppage: due to foot drop
- Duck waddle: myopathy, esp hips
- Scissor gait: UMN eg spastic paraparesis
- Parkinsonian: shuffling

Cerebellar

Midline Cerebellar Vermis

- Postural instability
- Includes alcoholic cerebellar degeneration

Spinocerebellar

- Truncal ataxia (Drunken Sailor)
- +ve Babinski
- Hereditary eg Friedrichs

Cerebellar Hemisphere

- lateral hemispheres
- Intention tremor
- Cerebellar dysarthria (feat explosive variation to voice)
- Dysidiadochokinesia
- Dysmetria (inability to judge distance)

Sensory

- Dorsal column (lack of proprioception)
- Cerebellum, thalamus, parietal lobes are inputs
- +ve rhombegs

Vestibular

See vertigo

Movement

- NM junctions
- LMN

Cranial Nerve	Type	Function	Symptoms/Signs
I Olfactory	S	Smell	Anosmia
II Optic	S	Vision	↓ VA
III Oculomotor	M	Eye movt: elevation, adduction Pupil: constriction	Eye ' down-&-out ' Loss of meiosis/accom (outer layer)
IV Trochlear	M	Eye movt: depression of adducted eye	Lateral eye deviation
V Trigeminal	B	M: Mastication S: Sensation	M: weakness/loss of mastication S: Facial anaesthesia
VI Abducens	M	Eye movt: abduction	Medial eye deviation
VII Facial	B	M: Facial expression, salivation, lacrimation S: Taste	M: facial muscles, dry mouth/no tears S: Loss of taste ant 2/3
VIII Vestibulocochlear	S	Balance & Hearing	Vertigo, nystagmus, hearing loss
IX Glossopharyngeal	B	M: Salivation S: Taste, pharyngeal sensation M: Swallow, phonation	M: dry mouth S: no gag ref, no taste post 1/3 M: dysphagia/dysphonia
X Vagus	B	S: Cardiac, GIT, Resp, Taste	S: no cough ref, no taste (hard palate)
XI Accessory	M	Pharyngeal/laryngeal, neck & shoulder	Weak head turn/shrug
XII Hypoglossal	M	Tongue	Tongue atrophy, deviation, fasciculation