

Acute Ischaemic Stroke

CT or MR SCAN READING FORM

SCAN ID:

DATE OF READING:

SCAN QUALITY: Good Moderate Poor

Comment:

READER ID:

TYPE OF SCAN: CT: Without contrast: With contrast:
 MR: Diffusion: Perfusion

Note – examples and definitions will be on a separate sheet or in drop down boxes if electronic. Please tick Yes or No. Please do not leave blanks. Thank you.

1. Is the scan completely normal? **Y** **N**
 If YES stop here

ISCHAEMIC CHANGES

2. Is there any sign of acute ischaemic change? If in doubt as to whether acute or old, code as acute. **Y** **N**
 If No go to Q.10

3. Which side of the brain shows ischaemic change? **R** **L**
 Tick R and L if both

4. Classify signs of ischaemic change in the main lesion (if more than one recent lesion).
(see examples)

a) loss of grey/white matter cortex definition	Y	N	
	<input type="checkbox"/>	<input type="checkbox"/>	
b) loss of basal ganglia outline	<input type="checkbox"/>	<input type="checkbox"/>	N/A
	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
c) hypodensity present (i.e. more than in a or b so that the lesion appears less dense than white matter)	<input type="checkbox"/>	<input type="checkbox"/>	

d) mass effect (swelling) present

Y	N
<input type="checkbox"/>	<input type="checkbox"/>

If yes –

sulcal effacement

<input type="checkbox"/>	<input type="checkbox"/>
--------------------------	--------------------------

ventricular effacement

<input type="checkbox"/>	<input type="checkbox"/>
--------------------------	--------------------------

midline shift

<input type="checkbox"/>	<input type="checkbox"/>
--------------------------	--------------------------

uncal herniation

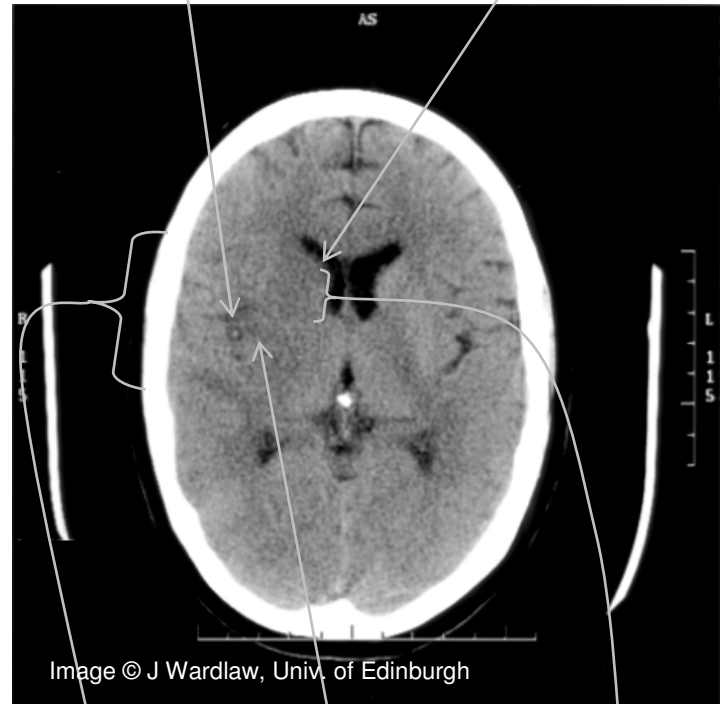
<input type="checkbox"/>	<input type="checkbox"/>	N/A
		<input type="checkbox"/>

hyperdense artery
(see Q.10)

mass effect:
ventricular effacement

hyperdense artery
(see Q.10)

mass effect:
ventricular effacement



loss of grey/white
matter definition

hypodensity

loss of basal
ganglia outline

mass effect:
sulcal effacement

hypodensity

loss of basal
ganglia outline

5. Classify site and size of ischaemic lesion
(see examples)

a) site



- M =MCA*
- AS =Infarct of up to half of ACA territory
- AL =Infarct of more than half of ACA territory
- PS =Infarct of up to half of PCA territory
- PL =Infarct of more than half of PCA territory
- MAS=M+AS*
- MAL=M+AL*
- MPS=M+PS*
- MPL=M+PL*
- MAP=Infarct of whole MCA, ACA and PCA territories
- L =Lacune*
- B =Borderzone*
- C =Cerebellum*
- S =Brainstem*
- CS =Cerebellum and brainstem

* code sub-territory sites in b

b) sub-territory sites



MCA sub-territory codes

- 1=small cortical infarct
- 2=basal ganglia infarct (>2x2x2cm)
- 3=infarct of white matter lateral to the lateral ventricle (>2x2x2cm)
- 4=infarct of anterior half of peripheral MCA territory
- 5=infarct of the posterior half of peripheral MCA territory
- 6=infarct of the whole of peripheral MCA territory
- 7=6+infarct of lateral part of basal ganglia
- 8=infarct of whole of MCA territory

Lacunar/Borderzone sub-territory codes

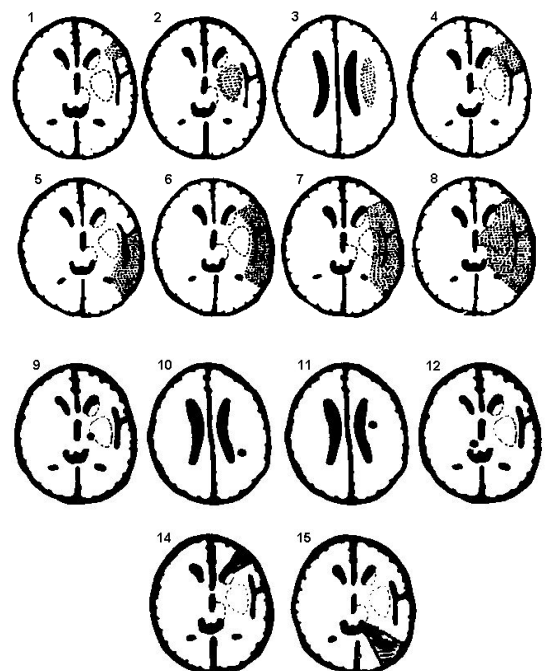
- 9=lacune in internal capsule/lentiform
- 10=lacune in internal border zone
- 11=lacune in centrum semiovale
- 12=lacune in thalamus
- 13=lacune in brainstem, inc. pons (not shown)
- 14=anterior (mainly) border zone
- 15=posterior (mainly) border zone

Cerebellum sub-territory codes

- 16=small cortical (not shown)
- 17=<1/2 hemisphere (medium) (not shown)
- 18=>1/2 hemisphere (not shown)

Brainstem sub-territory codes

- 11=small, i.e.<1/2 medulla (not shown)
- 12=extensive, i.e. pons + medulla (not shown)

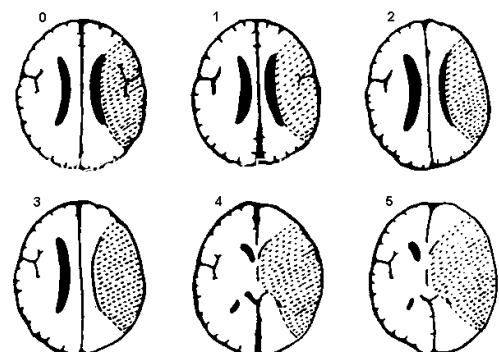


c) degree of mass effect



Mass effect grading

- 0=no swelling
- 1=effacement of the sulci overlying the infarct
- 2=1+minor effacement of adjacent lateral ventricle
- 3=1+complete effacement of lateral ventricle
- 4=1+effacement of the lateral and third ventricle
- 5=4+shift of the midline away from the side of the ventricle
- 6=5+effacement of the basal cisterns



Diagrams © J Wardlaw, Univ of Edinburgh

IF INFARCT IS IN THE MCA TERRITORY ANSWER Q.6 & Q.7 (if not go to Q.8):

6. In your opinion does the new acute ischaemic change involve more than 1/3 of the MCA territory? Y N
-

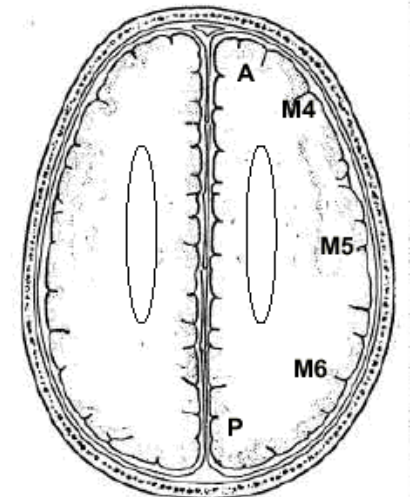
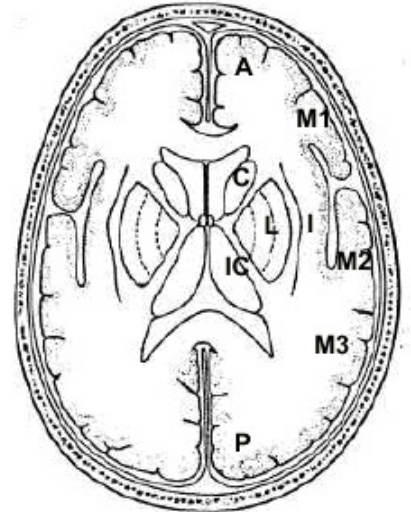
7. ASPECTS for the MCA territory (see examples)

ASPECT Score:

Please indicate if each of the MCA areas shown opposite, in the hemisphere that you think is ischaemic, are normal or show some signs of an infarct (abnormal).

(NB: Does not include areas A or P)

	Norm	Abnorm
Caudate (C)	<input type="checkbox"/>	<input type="checkbox"/>
Lentiform (L)	<input type="checkbox"/>	<input type="checkbox"/>
Insula (I)	<input type="checkbox"/>	<input type="checkbox"/>
Internal Capsule (IC)	<input type="checkbox"/>	<input type="checkbox"/>
MCA1 (M1)	<input type="checkbox"/>	<input type="checkbox"/>
MCA2 (M2)	<input type="checkbox"/>	<input type="checkbox"/>
MCA3 (M3)	<input type="checkbox"/>	<input type="checkbox"/>
MCA4 (M4)	<input type="checkbox"/>	<input type="checkbox"/>
MCA5 (M5)	<input type="checkbox"/>	<input type="checkbox"/>
MCA6 (M6)	<input type="checkbox"/>	<input type="checkbox"/>



Diagrams and score taken from Lancet 2000;355:1670-1674

8. Is there a second (discrete) recent ischaemic lesion? Y N
- If No go to Q.10*

9. Describe second ischaemic lesion:

HYPERDENSE VESSEL SIGN

10. Is there a hyperdense artery? (see example in question 4) Y N
- If No go to Q.12*

11. Name hyperdense artery:

HAEMORRHAGIC CHANGES

12. Is there any haemorrhage anywhere?

Y N

If No go to Q.14

13. Classify haemorrhage (if more than one haemorrhage, tick all present – indicate order of significance) :

a) petechial haemorrhage (example 1 or 2 below)

Y

N

Order
(insert 1, 2, 3 to indicate your estimate of the order of importance)

Size of Haematoma
(max diam.):

<3cm 3-5cm 5-8cm >8cm

b) significant haemorrhagic transformation of infarct (i.e. underlying infarct still visible) (example 3 below)

c) parenchymal haematoma (i.e. no infarct visible)

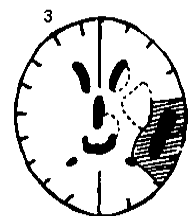
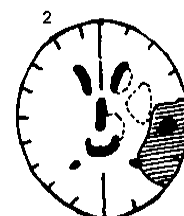
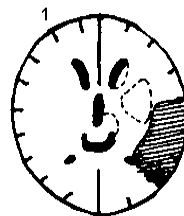
d) parenchymal haematoma clearly remote from infarct

e) subdural haematoma

f) subarachnoid haemorrhage

g) extradural haemorrhage

h) In your opinion, is the haemorrhage a major component of the infarct which is likely to have worsened mass effect or involved more brain in the damage present and so worsened symptoms, or if remote from the infarct, likely to have contributed significantly to the burden of brain damage?



Haematoma with no or only slight mass effect

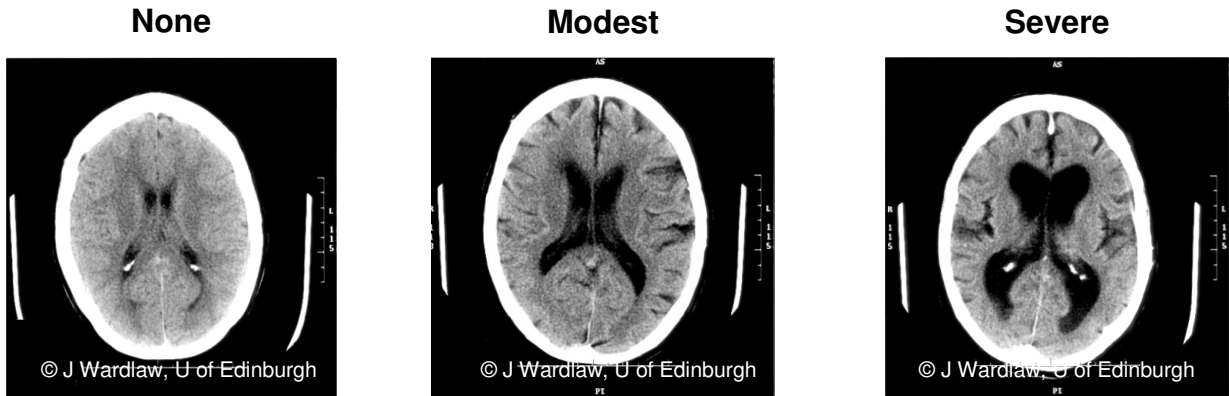
Haematoma with definite mass effect compressing surrounding tissue

Diagrams © J Wardlaw, Univ of Edinburgh

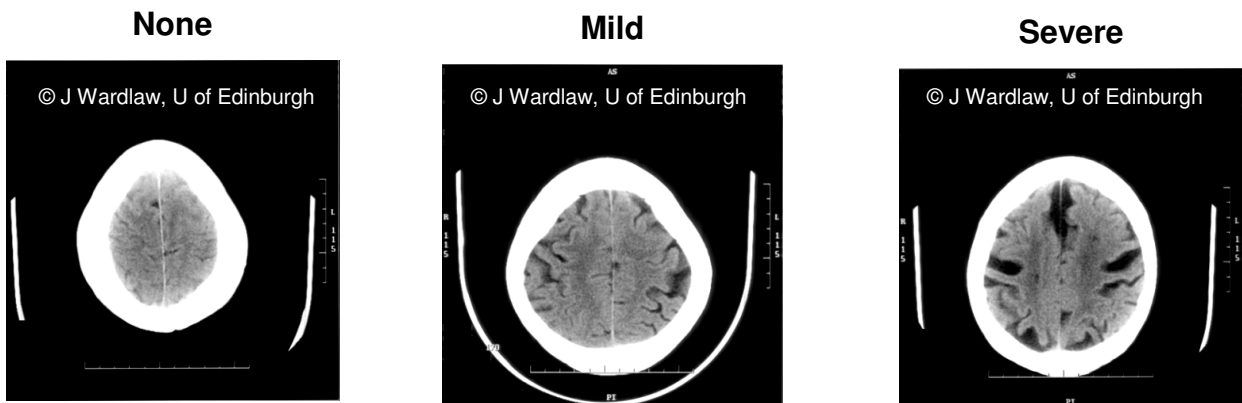
Reduction in brain tissue volume

14. Is there any Reduction in brain tissue volume? **Y** **N**
 If No go to Q.16
15. Classify atrophy (see examples and pick nearest likeness):
- | | | | |
|----------|--------------------------|--------------------------|--------------------------|
| central | None | Mod | Severe |
| | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> |
| cortical | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> |

CENTRAL reduction in brain tissue



CORTICAL reduction in brain tissue



Approach validated in Eur Radiol 2008;19:177-183

PERIVENTRICULAR LUCENCIES

16. Are there any periventricular lucencies?

Y

N

If No go to Q.18

17. Classify extent of white matter lucency

a. Anterior white matter

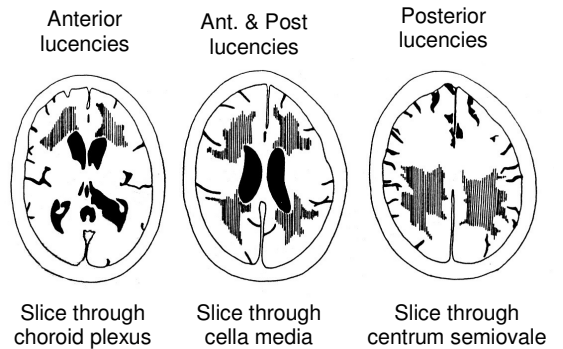
- 0= no lucency
- 1= lucency restricted to region adjoining ventricles
- 2= lucency covering entire region from lateral ventricle to cortex

0,1,2

b. Posterior white matter

- 0= no lucency
- 1= lucency restricted to region adjoining ventricles
- 2= lucency covering entire region from lateral ventricle to cortex

0,1,2

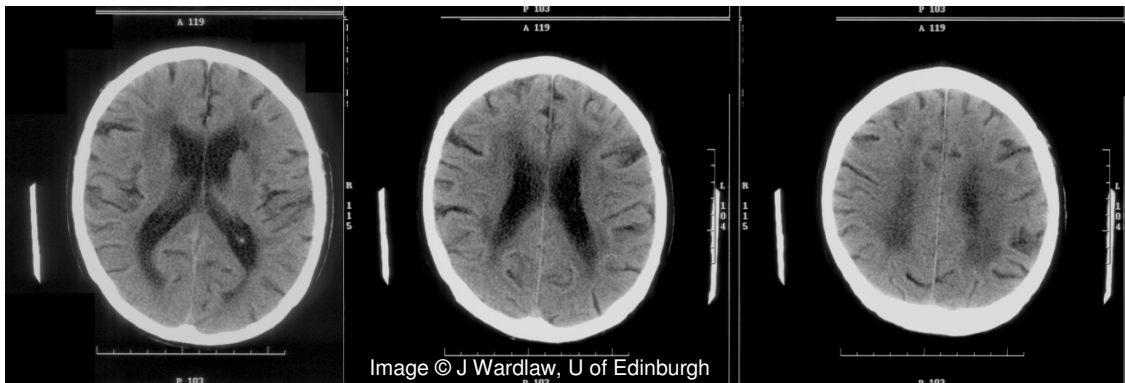


(diagrams from van Swieten et al. JNNP 1990;53:1080-1083)

AWM = 1 PWM = 0



AWM = 2 PWM = 1



OLD VASCULAR LESIONS

18. Are there any old vascular lesions? **Y** **N** *If No go to Q.20*

19. Classify old vascular lesion(s):

- | | Y | N |
|--|--------------------------|--------------------------|
| a) old cortical infarct(s) | <input type="checkbox"/> | <input type="checkbox"/> |
| b) old striatocapsular infarct(s) | <input type="checkbox"/> | <input type="checkbox"/> |
| c) old borderzone infarct(s) | <input type="checkbox"/> | <input type="checkbox"/> |
| d) old lacunar infarct(s) | <input type="checkbox"/> | <input type="checkbox"/> |
| e) old brainstem/cerebellar infarct(s) | <input type="checkbox"/> | <input type="checkbox"/> |
| f) probable old haemorrhage | <input type="checkbox"/> | <input type="checkbox"/> |

NON-STROKE LESIONS

20. Is there a non-stroke lesion, which could have accounted for the patient's stroke syndrome? **Y** **N** *If No go to Q.22*

21. Classify non-stroke lesion:

- | | Y | N |
|---------------------------|--------------------------|--------------------------|
| a) cerebral tumour | <input type="checkbox"/> | <input type="checkbox"/> |
| b) encephalitis | <input type="checkbox"/> | <input type="checkbox"/> |
| c) cerebral abscess | <input type="checkbox"/> | <input type="checkbox"/> |
| d) other (e.g. contusion) | <input type="checkbox"/> | <input type="checkbox"/> |

Specify Other:

22. **COMMENT:**

For further information, contact joanna.wardlaw@ed.ac.uk. This form is freely available for academic use but please acknowledge the source and any specific references therein.