

Neurological Society of Thailand

Surat Thani Hospital

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Approach to Weakness

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- Clinical data gathering
 History
 Physical examination:
 motor system: tone/power; reflex
 cranial nerves
 sensory system
- ➤ Interpretation, localization
 Upper versus lower motor neuron
 Motor pathway: cortex to muscle





Clinical data gathering

History

Chief complaint:

Present illness:

clearly defined as much as possible

- Quality of symptoms
 - Location
- **-** Time course

distinctive features e.g. character, severity, etc.

onset: abrupt or insidious subsequent course: improving, worsening, exacerbation-remission



Neurologic Evaluation



Progressive:

- tumor
- degenerative

Paroxysmal:

- epilepsy
- syncope
- periodic paralysis
- transiert ischemic attack

Sudden:

- stroke
- trauma

Relapsing-remitting:

- multiple sclerosis
- neuromyelitis optica





Clinical data gathering

History

e.g.: exertion (fatigue)

Present illness:

Associated symptoms

systemic neurologic





Clinical data gathering

History

Past medical history:

- illnesses

- operations

- medication

✓- immunization

Diet, tobacco, alcohol, substance abuse





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Die

Vaccine: protective against organisms causing neurologic disorders (e.g. poliovirus, tetanus, rabies virus, Japanese encephalitis virus, meningococcal, pneumococcal, Hemophilus influenzae Post vaccination: immunological reactions i.e. ADEM (Acute **Disseminated EncephaloMyelitis**





Clinical data gathering

Family history:

First degree relatives: parents, siblings, children

Second degree relatives: grandparents, grandchildren

Social history: education, occupation





- ➤ Physical examination Mental status
- Cranial nerves
- Motor function
- ✓ Reflexes
 Sensory function
 Coordination
 Gait and stance





- > Inspection
- ➤ Muscle tone: tension of a muscle at rest
 - examination
 - interpretation
- Muscle power
 - examination
 - grading system
 - interpretation



asymmetry

- Compare both sides

- Look for muscle atrophy,



- **▶** Inspection
- **Muscle tone**
 - examinatio

 - interpretation
- > Muscle power
 - examination
 - grading system
 - interpretation





- > Inspection
- > Muscle tone: tension of a muscle at rest
 - examination
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- Inspect
- **➤ Muscle**
 - exami
 - interpr
- > Muscle p
 - examir
 - grading
 - interpres

- Compare both sides
- Always check for tone i.e. resistance to passive movement (muscles relaxed)

Tone: upper limbs

- hold one hand in the "shake hands" position
- support at the elbow elbow flex/extend forearm supinate/pronate
- hold forearm above the wrist wrist flex/extend



- Inspect
- **➤ Muscle**
 - exami
 - interpression
- > Muscle p
 - examir
 - grading
 - interpretation

- Compare both sides
- Always check for tone, i.e. resistance to passive movement (muscles relaxed)

Tone: lower limbs
hand on knee, "roll" each
knee from side to side
hands under knee, "snap"
lifting, observe the heel





- > Inspection
- Muscle tone: tension of a muscle at rest
 - examin Tone:
 - interpr Hypotonia >> flaccid
- ➤ Muscle p Normotonia
- - examin Hypertonia: spasticity, rigidity
 - grading system
 - interpretation





- > Inspection
- > Muscle tone: tension of a muscle at rest
 - examination
 - interpretation
- Muscle power
 - examination
 - grading system
 - interpretation

Upper limb Shoulder Elbow

Shoulder Deltoid (abduction)

Biceps (flexion)

Brachioradialis (flexion)

Triceps (extension)

Radio- Pronator

ulnar Supinator

Wrist Flexor

Extensor

Finger Handgrip (flexion)

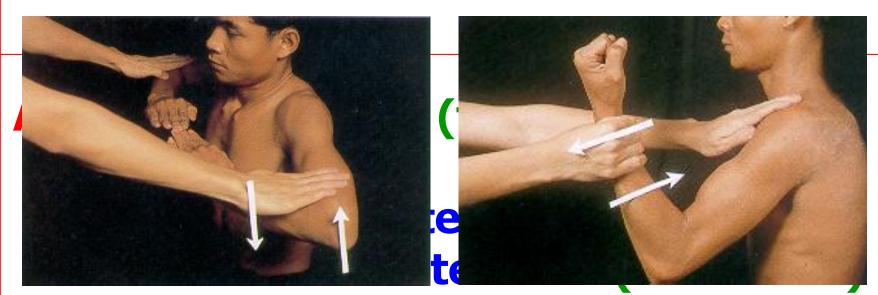
Extensor

Dorsal interossei (adduction)

Palmar interossei (abduction)

Upper limb

- Deltoid: Exert pressure close to elbow
- Biceps: Place one hand at patient's shoulder to stabilize
- Triceps: Test with arm extended is less painful than with arm flexed



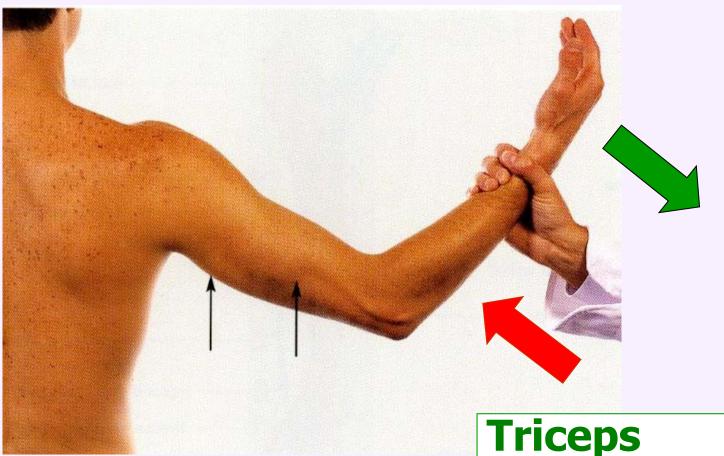


Neurologic Evaluation



MRC

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Elbow extension

Extend elbow against resistance

Upper limb

- Deltoid: Exert pressure close to elbow
- Biceps: Place one hand at patient's shoulder to stabilize
- Triceps: Test with arm extended is less painful than with arm flexed
- Pronator sign to detect mild weakness

Upper limb

Pronator sign

Position:

- raise both arms forward
- fully extend and supinate both arms
- fully extend and abduct fingers
- close the eyes

Interpretation: test positive if

- arm drift down
- elbow flex
- forearm pronate
- wrist and fingers flex

Minimal arm weakness

Lower limb

Hip Gluteus maximus (extension)

Iliopsoas (flexion)

Adductor

Abductor

Knee Quadriceps (extension)

Hamstring (flexion)

Ankle Gastrosolieus (extension)

Tibialis anterior (flexion)

Tibialis posterior (inversion)

Peroneii (eversion)

Toe Flexor

Extensor

Lower limb

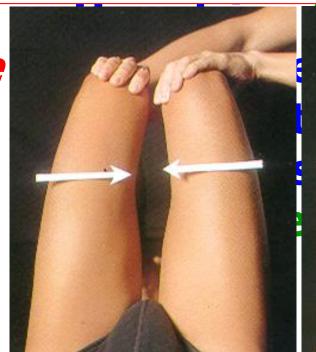
Hip

Gluteus maximus (extension) Iliopsoas (flexion)

Hints:

- Avoid pressing on the thigh
- Use the knee as pressure point

Ankle



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ion)

Toe

	Grade	Character
>	0	No movement
	1	Minimal movement
	2	Horizontal movement
\	3	Against gravity, unsustain
	4	Against resistance, partial
	5	Against resistance, strong (normal)
/	gradin	g system

Grading system: Medical Research Council (MRC); UK





- > Inspection
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Patterns of weakness

- pyramidal weakness

Pyramidal weakness:

upper limb: flexor group stronger "decorticate posture"

lower limb: extensor group stronger

: upper motor neuron (UMN)

: corticospinal tract lesion







Patterns of weakness

- pyramidal weakness
- proximal weakness

Proximal weakness:

more weakness of proximal group (arms and legs)

: muscle disease e.g. metabolic myopathy myositis

: neuromuscular junction





Patterns of weakness

- pyramidal weakness
- proximal weakness
- distal weakness

Distal weakness:

- more weakness of distal group, muscle wasting may be present
 - : peripheral nerve e.g.
 - distal symmetrical polyneuropathy



Neurologic Evaluation



Weakness

Patterns of weakness

- pyramidal weakness
- proximal weakness
- distal weakness

Specific pattern:

- weakness/wasting according to
 - : peripheral nerve distribution
 - : brachial, lumbosacral plexus
- : spinal nerve roots
 - : cauda equina
 - : muscular dystrophy

> Physical examination

Procedure

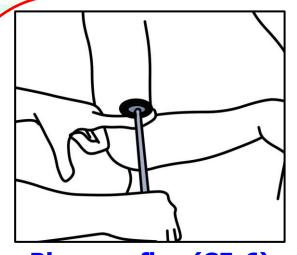
- optimum posture
- symmetry for comparison
- relaxed posture
- brisk and strong percussion
- observe the responses



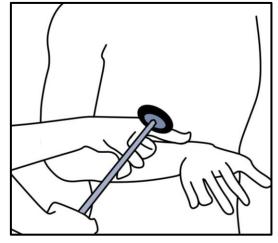


Neurological Examination

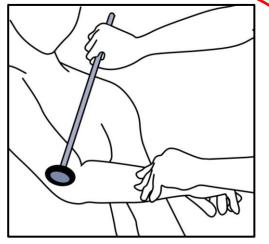
Muscle Stretch Reflex



Biceps reflex (C5-6) Musculocutaneous



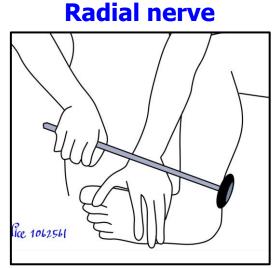
Brachioradialis (C5-6)



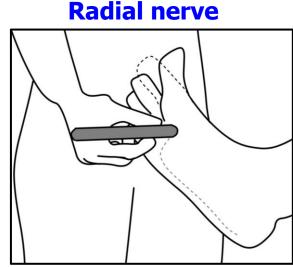
Triceps reflex (C7-8)

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Quadriceps reflex (L2-3)
Femoral nerve



Ankle reflex (S1-2)
Tibial nerve



Plantar reflex
Tibial nerve





Neurological Examination Muscle Stretch Reflex



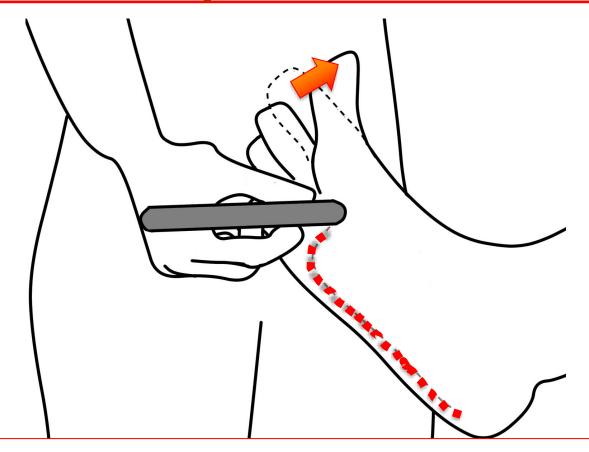
Reflex spreading:
UMN lesion above level
of the muscle to which
reflex spreads, e.g. hip
adduction with knee
jerk

Quadriceps reflex (L2-3) Femoral nerve



Neurological Examination Superficial Reflex





Normal: Plantar flexion of big toe Abnormal: Dorsiflexion of big toe

Tibial nerve





Neurological Evaluation Muscle Stretch Reflex



Interpretation (grading)

Grading	Interpretation
0	absent
1+ or +	diminished
2+ or ++	normal
3+ or +++	brisk
4+ or ++++	brisk with clonus



Neurological Examination Muscle Stretch Reflex



- Signs associated with corticospinal tract lesion
- Clonus
- Grasp reflex
- Tromner sign
- Babinski sign

Tromner sign Flick the hyperextended middle finger results in flexion of other fingers





- Muscle power
 - examination
 - grading system
 - interpretation: localize the lesion

UMN versus LMN versus non neurogenic

UMN: upper motor neuron from motor cortex > corona radiata (corticobulbar/corticospinal tracts) > posterior limb of internal capsule > brainstem > spinal cord (corticospinal tract)
LMN: lower motor neuron from anterior horn cell > ventral rami of spinal nerve > brachial/lumbosacral plexus > peripheral nerve
NMJ: neuromuscular junction



Neurological Evaluation
Motor Syste

Cortical

LMN

Anterior horn cell

Spinal nerve

Nerve plexus

Peripheral nerve

Neuromuscular junction

Sub cortical

> Brain stem

Spinal cord

UMN

Non neurogenic

Muscle









- Muscle power
 - examination
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 - interpretation: localize the lesion

UMN versus LMN versus non neurogenic

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NMJ: neuromuscular junction





Muscle power

Type/Parameters	Upper motor neuron	Lower motor neuron
Tone	Increase*	Decrease
Atrophy	Uncommon	Prominent
Fasciculation	Absent	Present
Deep tendon reflex	Hyperreflexia*	Hyporeflexia/ Areflexia
Plantar reflex	Extensor	Flexor

*Acute phase: hypotonia; hyporeflexia/areflexia





Localization Parameter	Cerebral hemisphere	Brainstem	Spinal cord
Distribution of muscles involved	Contralateral leg and arm, cranial muscles	Contralateral leg and arm, cranial nerve involvement	<i>Ipsilateral</i> leg and arm, specific group (segmental)
Туре	UMN	UMN	LMN and UMN*
Atrophy	Absent or mild	Absent or mild	When involve anterior horn cell
Tendon reflex	^	<u>^</u>	^
Plantar reflex	Extensor	Extensor	Extensor
Sensory symptoms	When involve sensory cortex	When involve sensory tract	When involve STT, posterior column
Example	Tumor, motor neuron disease, stroke	Stroke	Myelitis, Syringomyelia







al hemisphere

muscles and leg; cortical lobe signs **UMN Type**

Atrophy Absent or mild

Tendon reflex Increased

Plantar reflex Extensor

Sensory symptoms Present if involve sensory cortex

Additional deficits Gaze deviation, motor aphasia (in dominant hemisphere)

Tumor, hemispheric infarction, Example motor neuron disease





Localization Cerebr	ام
Parameter	Subcortical
Distribution of muscles	Contralateral cranial muscles, arm and leg (severity leg = arm)
Туре	UMN
Atrophy	Absent or mild
Tendon reflex	Increased
Plantar reflex	Extensor
Sensory symptoms	Present if involve thalamus
Additional deficits	
Example	Lacunar infarction

Cranial nerve

Mid brain: CN III, IV

Pons: CN V, VI, VII, VIII

Medulla: CN IX, X, XI, XII

Evaluation kness



Brain stem

Distribution of Contralateral leg and arm, cranial muscles nerve involvement

Type UMN

Atrophy Absent or mild

Tendon reflex Increased

Plantar reflex Extensor

Sensory symptoms Present if involve sensory tract

Additional deficits CN involvement

Example Brainstem infarction

Sensory

- vibratory, joint position ipsilateral
- pinprick, touch contralateral

ANS: bowel, bladder, sweating





cord

Distribution of	<i>ipsilateral</i> leg and arm, specific
muscles	muscle group (segmental)

Type LMN (at the level)+ UMN (below)*

Muscle pain/ absent tenderness

Atrophy present if involve anterior horn cell

Tendon reflex increased (below)

Plantar reflex extensor

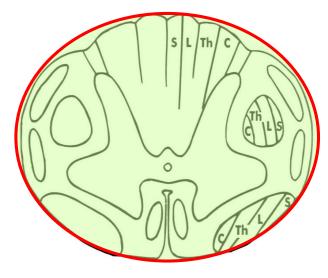
Sensory symptoms present if involve spinothalamic tract, posterior column

Example myelitis, syringomyelia

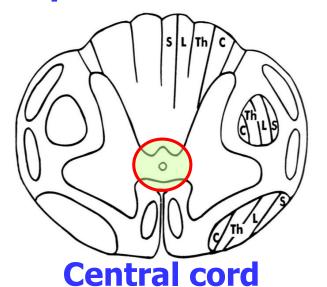
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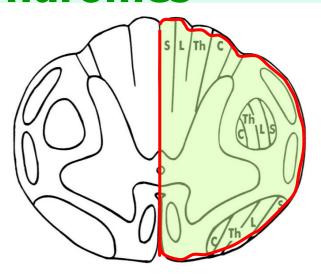




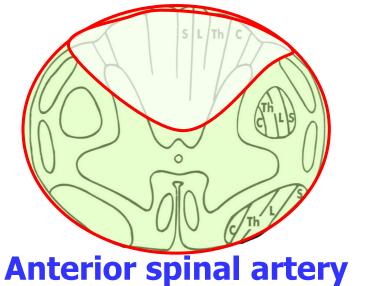


Complete transection



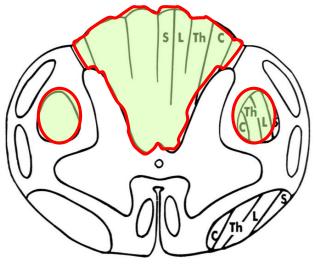


Brown Sequard (hemicord)

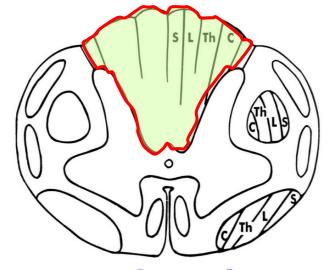




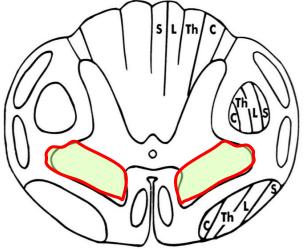




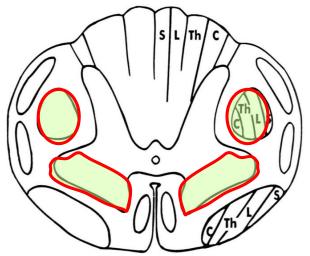
Postero-lateral column



Posterior column



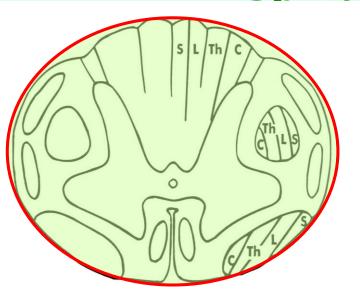
Anterior horn



Anterior horn-pyramidal tract







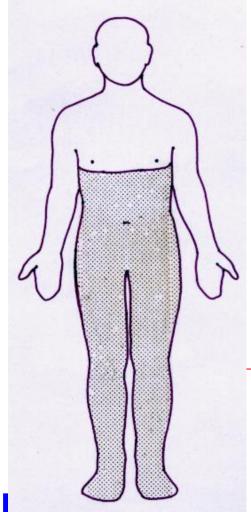
Complete transection

Motor: para-, tetraplegia (*spinal shock); LMN at the level

Sensory: all modalities below

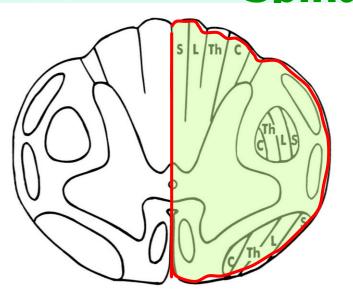
the level

ANS: urgency, retention, incontine









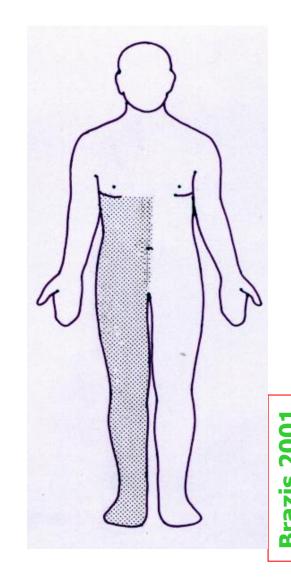
Brown Sequard (hemicord)

Motor: ipsilateral hemiplegia

Sensory: contralateral pain,

temperature; ipsilateral jps,

vibratory (2 segments below)







Localization Parameter	Spinal nerve	Nerve plexus	Peripheral nerve
Distribution of muscles involved	Segmental muscle group	Arms or legs (brachial or lumbo sacral plexus)	Specific groups
Туре	LMN	LMN	LMN
Atrophy	Prominent	Prominent	Prominent
Tendon reflex	4	V	↓
Plantar reflex	Flexor	Flexor	Flexor
Sensory symptoms	Present	Present	Absent in pure motor nerve
Example	Cervical spondylosis	Brachial plexus injury	Carpal tunnel syndrome

Cervical C1-8 Thoracic T1-12 Lumbar L1-5 Sacral S1-5

ologic Evaluation Weakness



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Distribution of Segmental muscle group muscles

Type LMN

Atrophy Prominent

Plantar reflex Flexor

Sensory symptoms Present (segmental)

Additional deficits

Example Cervical spondylosis



Brachial plexus C5-T1

Trunk: upper/middle/lower

Cord: lateral/medial/posterior

Lumbosacral plexus L1-S2

aluation

SS



lerve plexus

Distribution of Arms or legs (brachial or lumbomuscles sacral plexus)

Type LMN

Atrophy Prominent

Plantar reflex Flexor

Sensory symptoms Present to nerve sistribution

Additional deficits

Example Brachial plexus injury





Localization	
Parameter	Peripheral nerve
Distribution of muscles	Specific group (mononeuropathy)
Туре	LMN
Atrophy	Prominent
Tendon reflex	Decreased or normal
Plantar reflex	Flexor
Sensory symptoms	Specific to nerve distribution Absent in pure motor nerve
Additional deficits	
Example	Carpal tunnel syndrome, diabetic mononeuropathy





Parameter	Peripheral nerve
Distribution of muscles	Distal limb (polyneuropathy)
Туре	LMN
Atrophy	Prominent
Tendon reflex	Decreased
Plantar reflex	Flexor
Sensory symptoms	Present (distal limb)
Additional deficits	
Example	Guillain Barré syndrome, diabetic polyneuropathy





Localization Parameter	Neuromuscular junction Muscle		
Distribution of muscles involved	Symmetrical, proximal limb, ocular, bulbar muscle	- : Symmatrical nrovimal limi	
Туре	Non neurogenic	Non neurogenic	
Atrophy	Absent	In muscular dystrophies	
Tendon reflex	Normal or ↓	Normal or ↓	
Plantar reflex	Flexor	Flexor	
Sensory symptoms	Absent	Absent	
Example	Myasthenia gravis Lambert Eaton syndrome	Myositis, hypokalemia, thyrotoxic	





Localization	
Parameter	Neuromuscular junction
Distribution of muscles	Symmetric, usually proximal limbs
Type	Non neurogenic
Atrophy	Absent
Tendon reflex	Normal
Plantar reflex	Flexor
Sensory symptoms	Absent
Additional deficits	Ocular, facial, bulbar muscles
Example	Myasthenia gravis, Lambert Eaton syndrome





Parameter	Muscle	
Distribution of muscles	Symmetric, usually proximal limbs	
Туре	Non neurogenic	
Atrophy	Present in muscular dystrophies	
Tendon reflex	Normal or decreased	
Plantar reflex	Flexor	
Sensory symptoms	Absent	
Additional deficits	Muscle pain/tenderness in inflammatory disease	
Example	Myositis, hypok+, thyrotoxic	





Pattern of Weakness	Localization
Upper limb: extensor muscles Lower limb: flexors muscles	Upper motor neuron (corticospinal tract: CS)
Hemiparesis	Upper motor neuron CS
Multifocal, asymmetric weakness without sensory involvement	Motor neuron disease Multifocal motor neuropathy Myasthenia gravis (uncommon)
Multifocal, asymmetric weakness with sensory involvement	Polyradiculopathy Multifocal neuropathy





Pattern of Weakness	Localization
Symmetric weakness, proximal or generalized without sensory involvement	Myopathy Motor neuron disease NMJ disorder
Generalized motor > sensory	Polyradiculoneuropathy
Asymmetric cranial muscles with or without limbs	Myasthenia gravis
Distal symmetric motor only	Distal myopathies Distal spinal muscular atrophy
Distal symmetric sensory > motor	Length-dependent polyneuropathy





Pattern of Weakness	Localization
Multiple nerves, asymmetric	Multifocal polyneuropathy
Multiple roots, asymmetric	Polyradiculopathy
Multiple nerves and roots single extremity	Plexopathy
Single root	Monoradiculopathy
Single nerve	Mononeuropathy





➤ Clinical data gathering History

Physical examination:

motor system: tone/power; reflex

cranial nerves

sensory system

➤ Interpretation, localization

UMN versus LMN versus non neurogenic

Motor pathway: cortex to muscle