

Guillain Barre Syndrome in Children : Case Report

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Abstract

Background: The incidence of GBS at the age of <15 years is 1.5 cases per 100,000 population and peaks at the age of 70-79 years, reaching 8.6 cases per 100,000 population. The peak of this disease at the age of 15-35 years and 50-75 years. Infants are said to have the lowest risk of developing GBS. Male have 1.5 times greater risk than female

Case: a 7 years old male children was presented with main complaint of weakness at extrimites on both side since March 27th, 2019. Patient felt weakness at lower extremities first and then followed by upper extremities two days later. patient cannot use his hand and leg at all. patient was unable to transfer from lying to sitting independently and unable to slide his buttocks. patient felt numbness on his both arms and leg. patient got dyspepsia one week before weakness. Based on physical and diagnostic examination patient diagnosed with Tetraparesis LMN type due to Guillan Barre Syndrome (GBS). Patient got rehabilitation program consist of NMES, ROM, active breathing, endurance, strengthening, transfer and mobilization exercise, and occupational therapy. Patient got improvement in muscle power of upper and lower extremities after five (5) month rehabilitation medicines. Barthel index improved from 25 to 95 and activity daily living independently.

Conclusion: NMES, ROM, active breathing, endurance, strengthening, transfer and mobilization exercise, and occupational therapy can improve muscle power of upper and lower extremities for GBS of male children in five month rehabilitation program.

Keywords: Guillain Barre Syndrome; children; physical exercise.

Introduction

Guillain Barre Syndrome (GBS) is a demyelinating neuropathy characterized by ascending weakness and reduced/loss of reflexes and can affect cranial nerves. GBS is the most important cause of acute flaccid paralysis (Andary, 2021). The incidence of GBS in the United States is 1.2–3 out of 100,000 population/year and increases with age.

The incidence of GBS at the age of <15 years is 1.5 cases per 100,000 population and peaks at the age of 70-79 years, reaching 8.6 cases per 100,000 population. The peak of this disease at the age of 15-35 years and 50-75 years. Infants are said to have the lowest risk of developing GBS. Men have 1.5 times greater risk than women (Van Doorn and Drenthen, 2014; Andary, 2021).

The cause of GBS itself is still unknown for certain, but there is an opinion that one of the triggers for GBS is a viral or bacterial infection such as Campylobacter Jejuni or Citomegalovirus (Craig, Richardson and Ayyangar, 2016). GBS begins with motor weakness and takes time to heal. Although GBS is said to be self-limiting disease, some patients experience sequelae. therefore, it is important for physical medicine and rehabilitation specialists to recognize the signs of GBS and determine the appropriate rehabilitation program for individuals affected by GBS.

Case Report

a 7 years old male children was presented with main complaint of weakness at extrimites on both side since March 27th, 2019. Patient felt weakness at lower extremities first and then followed by upper extremities two days later. patient cannot use his hand and leg at all. patient was unable to transfer from lying to sitting independently and unable to slide his buttocks. patient felt numbress on his both arms and leg. patient got dyspepsia one week before weakness.

History of present illness are felt weakness at both of arm, hand, leg, and foot. Weakness at both of legs since 5 days before hospitalized. Weakness more prominent until he cannot move his foot and cannot walk. Weakness of the arm and hand since two days before hospitalized, but he can move his upper extremities a little. He also felt difficulty in swallowing. Patient also experienced difficult of breath. After his breathing difficulties improved, he was hospitalized for 2 days. His condition when discharge (18 April 2019) was stable in respiration but still bed ridden and cannot use both of hand. Difficulty of swallowing was absence. Slurred speech and asymetrical face was denied, difficulty in urination and defecation was denied, history of cough, fever denied, he only felt dyspepsia one week before weakness.



Prenatal history of Patient is a 2nd child of 2 siblings, her mother was taken antenatal care routinely, aterm, normal delivery labor with 3000 gr weight and 50 cm height. Complete immunization, no history of seizure and lung Tuberculosis.

On April 1st, 2019 General examination shows that patient is bed ridden, Glasgow Coma Points: E4M5V6, BP: 110/60 mmHg, Pulse: 116 bpm, Respiration: 24x/min, T: 36,6°C, SpO₂: 98% room air, weight: 18 Kg, weight: 120 cm, BMI: 16,67 Kg/m², Barthel index is 25 (severely dependent). Head and neck: not anaemic, no jaundice, and no cyanosis. Heart sound is nomal, no murmur and no gallop sound, lung sound is vesicular on both side, no ronchi, no wheezing, abdomen is soft, liver and spleen is unpalpable.

Musculosceletal status on the table 1, neurological and functional examination on the table 2

Table 1. Musculosceletal Status

Part of Body	Movement	ROM	MMT
Neck	Flexion	F	1
	Extension	F	1
	Lateral Bending	F/F	1/1
	Rotation	F/F	1/1
Trunk	Flexion	F	Impressed <3
	Extension	F	Impressed <3
	Lateral Bending	F/F	Impressed <3/ Impressed <3
	Rotation	F/F	Impressed <3/ Impressed <3
Shoulder	Flexion	F/F	2/2
	Extension	F/F	2/2
	Abduction	F/F	2/2
	Adduction	F/F	2/2
	External Rotation	F/F	2/2
	Internal Rotation	F/F	2/2
Elbow	Flexion	F/F	2/2
	Extension	F/F	2/2
	Pronation	F/F	2/2
	Supination	F/F	2/2
Wrist	Flexion	F/F	2/2
	Extension	F/F	2/2
Thumb	Flexion MCP	F/F	1/1
	PIP	F/F	1/1
	Extension	F/F	1/1
	Abduction	F/F	1/1
	Adduction	F/F	1/1
	Opposition	F/F	1/1
Fingers	Flexion MCP	F/F	1/1
	PIP	F/F	1/1
	DIP	F/F	1/1
	Extension	F/F	1/1
	Abduction	F/F	1/1
	Adduction	F/F	1/1
Нір	Flexion	F/F	
	Extension	F/F	1/1
	Abduction	F/F	1/1
	Adduction	F/F	1/1
	External Rotation	F/F E/E	1/1
17		F/F	
Knee	Flexion	F/F E/E	1/1
A -= 1-1 -	Extension Dama flamian	F/F	1/1
Апкіе	Dorsonexion	F/F E/E	1/1
	Inversion	Г/Г Е/Е	1/1
	Evention	Г/Г Е/Е	1/1
Dig Too	Elevien		1/1
Dig 10e	Extension	171' F/F	1/1 1/1
Toos	Elavion		1/1
1005	Extension	1'/F F/F	1/1 1/1
	LAUISIOII	171	1/1

Table 2. Neurological and Functional examination Status



No	Examination	Result
A.	Neurological Status	
1	Cranial Nerve	Normal Limit
2	Physicological Reflexes	
	BPR KPR	+1/+2 +1/+1
	TPR APR	+1/+2 +1/+1
3	Pathological Reflexes	
	Babinski	- -
	Chaddock	- -
	Hoffman	- -
	Tromner	- -
4	Sensory Deficit	Difficult to evaluate
5	Spasticity	
6	Tonus	Decrease
В.	Functional Examination	
1	Balance	
	Sitting	
	Static	Unable
	Dynamic	Unable
	Standing	
	Static	Unable
	Dynamic	Unable
2	Pulmonary Function	
	Count Test	9
	Chest Expansion	T2-T4-T6: 2 cm-2 cm-2,5 cm
3	Hand Function	
	Grasp	NF/NF
	Spherical	NF/NF
	Cylindrical	NF/NF
	Lateral tip	NF/NF
	Pinch	NF/NF
	Hook	NF/NF

Chest X Ray on April 3rd, 2019 shows normal cardiac and suspect pneumoniae (Figure 1), EMG on April 16th, 2019 shows clinical neurophysiological showed demyelinating motor polyradiculoneuropathy.



Figure 1. Chest X-ray AP position (April 3rd , 2019), suspect pneumoniae, normal cardiac.

Based on physical and diagnostic examination patient diagnosed with Tetraparesis LMN type due to Guillan Barre Syndrome (GBS). Functional diagnosis are 1) Impairment: Tetraparesis; 2) Disability: Barthel index was 25 (severely dependent: bathing, dressing, grooming, toilet use, transfer, mobility, stairs); 3) Handicap: patient could not go to school and could not handle normal daily activity.



Table 3. ICF Classification

	No		Classification	
A	Body Functi	on	Clussification	
	1. b1522	His mother worried t	hat her son's condition will last forever	
	2. b4350	Immune response		
	3. b730	Muscle power function	on	
	4. b7500	Stretch motor reflex		
	5. b7603	Supportive functions	of arm or leg	
В.	Body Struct	ure		
	s198	Structure of the nerve	ous system, other specified	
	s730	Structure of upper ex	tremity	
	s750	Structure of lower ex	tremity	
С.	Activity and	Participation Limitating		
	1. d230	Carrying out daily ro	utine	
	2. d420	Transferring oneself		
	3. d430-c	1449 Carrying, moving and	d handling objects	
	4. d450-c	Walking and moving		
	5. d530	Toileting		
	6. d540	Dressing		
	7. d820 8 d020	School education		
D	8. U920	recreation and leisur	e	
<u>D</u> .		Ital Factors	victore.	
	1. E380	Health services and s	ystem	
		Tetra	nparesis LMN Type e.c Guillen Bare]
,	L .		Activity Limitation	Ţ
Body Fun b1522 wor the conditi last foreve b4350 imm response b730 muse function	action rrying that ion will rr nune cle power	Body Structure S198structure of the nervous system S730 structure of upper extremity S750 structure of lower extremity	d230 carrying out daily routine d420 transferring oneself d430-d449 carrying, moving and handling objects d450-d469 walking and moving d530 toileting d540 dressing	Participatio n Limitation d820 school education d920 recreation and leisure
	Ei he	nviromental Factors alth services and systems	Enviro health s	mental Factors services and systems

Figure 2. ICF Guillain-Barre Syndrome

Patient medical problem are Tetraparesis LMN Type ec GBS, rehabilitation medicine are R1: dependent ambulation with wheelchair; R2: severely dependent (Barthel Index 25); R7: weakness of upper and lower extremities bilaterally and transfers lying to sitting disturbance. There is no problem in R3 until R6.

Goal of treatment are: 1) short term: Sitting independently and increase the muscle strength; 2) long term: independently ambulation and improvement of quality of life.

Rehabilitation medicines on the table 4



Rehabili	tation Explanation
R1 Amb	ulation: dependent ambulation with wheelchair
PDx	
PTx	Continue medication from neurology pediatric department:
	Metilcobalamin 3 x 150 mg
	Modality:
	NMES at shoulder abductor D/S, elbow flexor D/S, wrist extensor D/S, finger fleksor D/S vicible muscle contraction,
	20 min everyday
	NMES hip extensor and flexor D/S, knee extensor D/S, ankle dorsiflexor D/S, big toe extensor D/S, visible muscle
	There are to the second s
	PPOM exercise of upper extremities D/S
	PROM exercise of lower extremities D/S
	Weight shifting exercise
	Breathing exercise with blowing trumpet and tissue
	Axial loading exercise helped by therapist
PMx	Clinical sign, ROM, MMT
Pex	Health education/Home Exercise Program
	Explain the patient's condition
	Continue exercise at home
	Continue NMES at home once/day,@20 minutes.
	Explain precaution of exercise (fatigue)
R2 ADL	: severely dependent
PDX DT _W	- Assisted by his family
PMv	Absisted by initial family ADI (barbel index)
PEx	Health education/Home Exercise Program
	Explain the patient's condition
	Practice doing ADL as much as possible
R3 Com	munication: no problem
R4 Psycl	nological:no problem
PDx	-
PTx	-
PMx DE-	Psychologic condition
R5 Socia	
R6 Voca	tional: He couldn't go to school and play with his friend as usual
PDx	
PTx	Give psychological support and give the idea to play something with sitting position and study at home
PMx	Psychologic condition
PEx	Health education
	Explain to the patient and her family about his condition
	Give psychological support (from family, clinician, environment)
	• Doing mediation with his teacher by explain his health condition to teacher, and help to postpone final
P7 Othe	examination until ne can write.
K/ Othe	rs; waaknass of upper and lower extremities D/S
	transfer lving to sitting disturbances
•	decrease of count test and chest expansion
PDx	-
PTx	Continue medication from neurology pediatric department:
	Metilcobalamin 3 x 150 mg
	Modalities:
	NMES at shoulder abductor D/S, elbow flexor D/S, wrist extensor D/S, finger fleksor D/S vicible muscle contraction,
	20 min/everyday
	invites inplextension and nexor D/s, knee extension D/s, ankle dorsinexor D/s, big toe extension D/s, visible industre
	Thera exc -
	PROM exercise of upper extremities D/S
	• PROM exercise of lower extremities D/S
	• Weight shifting exercise
	Breathing exercise with blowing trumpet and tissue
	• Axial loading exercise helped by therapist
PMx	Clinical sign and symptoms and MMT
PEx	Health education/Home Exercise Program
	• Explain the patient's condition
	Continue exercise at home
	• Eating with small portions but often and rich in nutrition
	Explain precaution of exercise (fatigue)



Table 5. Progress note of treatment

Date		Subjective		Objective		Assessment	Planning
May	1.	Both hand	General status :			Tetraparesis	Continue medication from
22 nd		become	• GCS 456, whee	lchair dependent		LMN type ec	neurology pediatric department:
2019		stronger	• BP : 110/70 mm	Hg, HR : 116 x/mn	t, RR : 24 x/mnt, SpO2	Guillain	Metilcobalamin 3 x 150 mg
	2.	Can turn	: 98%			Barre	
		his body to	Physical examination:			Syndrome	Modality:
		the	Th : C : S1-2 sing	le, mur2 -, gallop –		(day-35) with	NMES at shoulder abductor D/S,
	2	right/left	P : Rh -/- Wi	1 -/-		improvement	elbow flexor D/S, wrist extensor
	5.	around 30	Cnest exp : 12	/14/16 2/2/2,5 cm			muscle contraction 20
		minutes	Musculosceletal status				min/everyday
		mutes	Widseuloseeletai status	ROM	MMT		NMES hip extensor and flexor
			Head/ Neck	F	4		D/S, knee extensor D/S, ankle
			Trunk	F	Impressed >3		dorsiflexor D/S, big toe extensor
			Upper extr				D/S, visible muscle contraction,
			- Shoulder	F/F	3/3		20 minutes/everyday.
			- Elbow	F/F	3/3		Thera exc.:
			flexion	F/F	3/3		 AAROM exercise of lower extremities D/S
			extension	F/F	3/3		Iower extremities D/S
			Pronation	F/F	3/3		 Breathing exercise with blowing whistle
			Supination	F/F	3/3		• Avial loading evercise
			- Wrist Finger	F/F E/E	3/3		helped by therapist
			- Fliger	г/г F/F	3/3		Endurance exercise:
			Lower Extr	1/1	5/5		F: 3x/week
			- Hip	F/F	2/2		I: Heart rate rest $+20$
			- Knee	F/F	2/2		T: 5 minutes warm up, 20 minutes
			- Ankle	F/F	2/2		conditioning, 5 minutes cooling
			- Toes	F/F	2/2		down
			- Big Toes	F/F	2/2		T : arm crank
			 Neurologic 	al status			Bridging exercise
			- cranial nerve : No	rmal			• OT:
			 Physiological ref 	lexes :			AROM exercise AGA
			BPR : ++	-/++ KPR:+	+/++		D/S Jactonia strongthening
			TPR : ++	+++ APR:+-	+/++		AGA D/S
			Pathologica	ll reflexes: -			Hand function exercise
			Barthel inc	lex (BI) : 50		for ADL (especially	
			Grooming: 5 B1	Bower: 10			writing)
			Bathing 0 Tr	auder : 10			PMx :
			Dressing : 5 M	obility : 0			 Clinical signs &
			Toileting : 5 S	tairs : 0			symptoms, vital sign
			Hand Funct	ion: weak functiona	al bilateral		• ROM, MMT, count test,
							chest expansion, hand
							function, barthel index
							PEx :
							• Avoid fatigue during
							exercise Control to DMD
							Collifor to PMR outpatient clinic routinely
							Home based NMES
							everyday
							• Prevent hanging foot
							when sitting on
							wheelchair
June	1.	Hands felt	General Status			Tetraparese	PDx :
12 th ,	-	stronger	• GCS 456, wheel	chair independent		ec Guillain	PTx:
2019	2.	Hand	• BP : 110/90 mm	Hg, HR : 88 x/mnt	, RR : 22 x/mnt, SpO2 :	Barre	Rehabilitation program :
		function	99%			Syndrome	Modality:
	2	improved	Physical Examination				NMES 50 Hz on muscle belly hip
	з.	her buttock	Th: $C: SI-2$ sing	ie, mur2 -, gallop –			dorsiflexors knee flexor knee
		ner buttock	r: Kn -/- Wl Chest exp : T2/T4/T4	1 -/- 3/3/3 cm			extensor D/S ankle plantar flexors
			Count test · 22	5/5/5 CIII			D/S everyday. intensity visible
			Musculosceletal status	•			muscle contraction. Faradic type.
			museurosceretar status	ROM	MMT		20 minutes.
			Head/	F	5		
			Neck	-	-		 Therapy Exercise (as
			Trunk	F	Impressed >3		patient's tolerance)
			Upper extr		•		AROM exercise AGA
			·		-		D/S

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Date	Subjective		Objective		Assessment	Planning 512
		- Shoulder	F/F	4/4		AAROM lower
		- Elbow				extremities D/S
		Flexion	F/F	4/4		• Isotonic strengthening
		Extension	F/F	4/4		evercise upper
		Propation	E/E	4/4		avtromity D/S
		Supination	1/1 E/E	4/4		E 1
		Supination	Γ/Γ	4/4		• Endurance exercise:
		- wrist	E/E			F: 3x/week
		Flexion	F/F	4/4		I: Heart rate rest $+ 20$
		Extension	F/F	4/4		T: 5 minutes warm up, 30 minutes
		Ulnar dev	F/F	4/4		conditioning, 5 minutes cooling
		Radial dev	F/F	4/4		down
		- Finger	F/F	4/4		T : arm crank
		- Thumb	F/F	4/4		
		Lowerextr	- , -			• Active breathing
		Lower extr	E/E	2/2		• Active breathing
		- Inp	171° E/E	2/2		exicise (deep
		- Kliee		2/2		breathing)
		- Ankle		2/2		 Bridging exercise
		- Toes	F/F	2/2		
		- Big Toes	F/F	2/2		 OT: Hand function
						exercise for ADL
		 Neurological sta 	tus :			(using clothes,
		- Cranial nerves	: normal			drawing books
		- Physiological	efleves ·			playing card)
			VDD · · · / · ·			DM _w .
		DPK : ++/++	NPR : ++/++			PMX :
		1PK : ++/++	APK : ++/++			 Clinical signs &
		- Pathological	reflexes: -			symptoms, vital sign
		Barthel Index :	60			 ROM, MMT, chest
		Feeding : 10	Bowel : 10			expansion, count test,
		Grooming : 10	Bladder: 10			hand function, barthel
		Bathing: 5 T	ransfer : 5			index
		Dressing : 5	Aobility : 5			DE _v .
		Toileting : 5	toirs : 0			• Avoid fotions during
		Tolleting . 5 5				• Avoid laugue during
		Hand Function:	functional bilatera	1		exercise
						 Control to PMR
						outpatient clinic routinely
						 Prevent hanging food,
						when sitting on the
						wheelchair (use booth
						shoos)
I1	Detient een	Comound Status			T-4	DDa i
July,	Patient can	General Status			Tetraparese	PDX :
16 ^m	stand by hold	• GCS 456, depend	lent wheelchair		(improved)	PIX:
2019	walker/ wall	 BP : 110/70 mmH 	Ig, HR : 110 x/mnt,	RR : 20 x/mnt, SpO2 :	ec. Guillain	<u>Rehabilitation program</u> :
		98% ,temp ax 36.	5 degree celcius		Barre	 Modality:
		Physical examination	•		Syndrome	NMES 50 Hz on muscle belly hip
		Th : C : S1-2 sin	ole, mur2 -, gallon –		(3 months)	extensor D/S, ankle plantarflexor
		$P \cdot Rh _{-}/_{-} W$	h _/_		· · · ·	D/S everyday, 20 minnutes.
		Chast avp : $T2/T4/T6$	$\frac{11}{2}$ /2/2 om			_ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~
		Cliest exp : 12/14/10	5/ 5/ 5 CIII			Therapy everyise: (do as
		Count test : 24				- Therapy exercise. (do as
		Musculosceletal statu	3:			patients tolerate)
			ROM	MMT		 AROM exercise upper
		Head/ Neck	F	5		extremity D/S
		Trunk	F	5		 AROM exercise lower
		Unner extr	=	<u> </u>		extremity D/S
		Shoulder	E/E	1/1		• Active breathing exercise
		- Shoulder	Г/Г Г/Г	4/4		(deep breathing)
		- Elbow	F/F	4/4		(deep bleating)
		- Wrist	F/F	4/4		Isotonic Strengthening
		- Finger	F/F	4/4		exercise upper extremity
		- Thumb	F/F	4/4		D/S
		Lower extr				 Sitting to standing
		- Hip				exercise (by holding
		Flexion	F/F	3/3		walker)
		Extension	F/F	2/2		Endurance exercise •
		Abduction	1/1 F/F	3/2		F. 3v/week
		Adduction	17/1° E/E	3/3 2/2		I. JA/WEEK
		Adduction	F/F	5/5		1. Heart rate rest $+ 20$
		Int. Rot	F/F	3/3		1: 10 minutes warm up, 30
		Ext. Rot	F/F	3/3		minutes conditioning, 10 minutes
		- Knee	F/F	3/3		cooling down
		- Ankle	F/F	2/2		T: arm crank
		- Toes	F/F	3/3		PMx :
		- Rig Toes	F/F	3/3		 Clinical signs &
			1/1	5/5		symptoms vital sign
		Ineurological st	atus :			DOM MAT 1
		Cranial nerves : norm	ai			- KOWI, WIWI I, Dartnel
		Physiological reflexes	:			index
						Precaution fatigue

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Date	Subjective	Obj	ective		Assessment	Planning 512
		BPR:+2/+2 KPR:	+2/+2			PEx : 313
		$TPR:+2/+2 \qquad APR:$	+2/+2			 Explain about patient
		Pathological reflexes: -				condition rehabilitation
		Hand Function: Functional Bilateral				program to the patient
		• Standing balance static:	poor		and his family	
	Standing balance state: poor Standing balance dynamic: poor					 Practice exercise
		BI : 75	lei poor			minimum 2x/day
		Feeding: 10 Bowel: 10				 Avoid fatigue during
		Grooming : 10 Bladder : 10)			exercise
		Bathing : 5 Transfer : 5				 Control to PMR
		Dressing : 10 Mobility : 5	5			outpatient clinic routinely
		Toileting : 10 Stairs : 0				 Using booth shoes
						routinely when go to
						school
						 Avoid hanging foot when
						sitting position
						 Home based NMES
						routinely everyday
August	Patient walk	General Status			Tetraparese	PDx :
15 th ,	independent	GCS 456, ambulation indep	pendent, wadd	lling gait	(improved)	PTx :
2019	with out assitive	• BP : 120/70 mmHg, HR : 80	x/mnt, RR : 20) x/mnt, SpO2 :	ec Guillain	Rehabilitation program :
	device	98%		-	Barre	 Therapy exercise: (do as
		Physical Examination			Syndrome	patients tolerate)
		Th : C : S1-2 single, mur2 -,	gallop –		(5 months)	AROM exercise upper and
		P : Rh -/- Wh -/-				lower extremity D/S
		Chest exp : T2/T4/T6 3/3/3 cm				(except hip extensor D/S)
		Count test : 24				 AAROM exercise hip
		Genu recurvatum +				extensor D/S, ankle
		Musculosceletal status :				plantarflexor D/S
			ROM	MMT		• Active breathing exercise
		Head/Neck	F	5		(deep breathing)
		Trunk	F	5		• Strengthening exercise
		Upper extr				upper and lower extremity
		Shoulder	F/F	4/4		(except hip extensor D/S
		Elbow	F/F	4/4		and ankle plantarflexor
		Wrist	F/F	4/4		D/S)
		Fingers	F/F	4/4		 Endurance exercise with
		Thumb	F/F	4/4		walking:
		Lower extr				F: 5x/week
		Hip				I: Heart rate rest $+20$
		Flexion	F/F	4/4		T: 10 minutes warm up, 20
		Extension	F/F	3/3		minutes conditioning, 10 minutes
		Abduction	F/F	4/4		cooling down
		Adduction	F/F	4/4		T : aerobic
		Int. Rot	F/F	4/4		
		Ext. Rot	F/F	4/4		PMx :
		Knee	F/F	4/4		 Clinical signs &
		Ankle	F/F	2/2		symptoms, vital sign
		Toes	F/F	4/4		 ROM, MMT, barthel
		Bigtoes	F/F	4/4		index
		Neurological status		<u></u>		PEx:
		Cranial nerves : normal				 Explain about patient
		Physiological reflexes :				condition and
		BPR : $+2/+2$ KPR :	+2/+2			rehabilitation program to
		TPR: +2/+2 APR:	+2/+2			the patient and his family
		Pathological Reflexes: -				 Do Home Base NMES
		Barthel Index: 95				everyday
		Feeding: 10 Bowel · 10				 Avoid fatigue during
		Grooming : 10 Bladder : 1	0			exercise
		Bathing: 10 Transfer: 1	5			 Use booth shoes when well-ing
		Dressing : 10 Mobility : 15	5			waiking
		Toileting : 10 Stairs : 0				- Doing nome based
		Standing balance static: good				INIVIES IOUUIIEIY
		Standing balance dynamic: poor				
		Hand Function: Functional bilate	eral			



Discussion

This case explained that 7 years old male children. was presented with main complaint of weakness at extrimites on both side since March 27th, 2019. Diagnosed Tetraparesis LMN type due to GBS. Patient diagnosed by general, physical, and diagnostic examination. Examination shows that patient suffered weakness at lower extrimites gradually followed by upper extrimities, felt numbness on all of extrimity, patient are unable sit and stand, losing all of hand function, difficult to breath suspected of pneumonia, patient experience a stomach discomfort. Barthel index is 25 (severely dependent).

2nd month rehabilitation, patient shows improvement of upper extrimity muscle tone, grasp, turn his body to the right and left, patient also can sit around 30 minutes. BI increased to 50. 3rd month rehabilitation, patient shows improvement of upper extrimity muscle tone, grasp, move his buttocks. BI increased to 60. 4th month rehabilitation, patient shows improvement of lower extrimity muscle tone, patient can stand by hold walker/wall. BI increased to 75. 5th month rehabilitation, patient can walk independent without assistive device. BI increased to 95.

GBS more favorable in children than in adults (Andary, 2021). The average incidence was 0.82 cases per 100,000 children aged <15 years (Landaverde et al., 2010). This patient experiencing decreased tendon reflexes in the affected extremity All children had weakness of bilateral limbs and disappearance or reduction of tendon reflex, and limb weakness reached the highest level of severity within 4 weeks (Ju-Fang et al., 2021) and pneumonia (Katirji, 2016). Some patients have symptoms of cranial nerve disorders, most commonly facial, oculomotor or bulbar weakness and ataxia known as Miller Fisher Syndrome (Thomas and Therattil, 2010). Patient got rehabilitation medicines consist of NMES, ROM. Occupational therapy professionals should be involved early in the rehabilitation program to promote positioning, posture, upper body strengthening (Craig, Richardson and Ayyangar, 2016), range of motion (ROM), and activities that aid functional self care (Andary, 2021).

Patient also undergo active breathing, endurance, strengthening, transfer and mobilization exercise, occupational therapy. Patient got improvement in muscle power of upper and lower extremities after five (5) month rehabilitation medicines. The recovery period often weeks to months, with a median estimated recovery time of 6–12 months. Full recovery within 3–12 months is experienced by 90–95% of pediatric patients with GBS (Andary, 2021). Barthel index also improved to 95. Patient can do activity daily living independently

Family counseling and education is extremely important early in the illness. The family must be prepared for a prolonged and potentially complicated course of illness.

Conclusion

This report established that male children suffering tetraparese due to GBS was able to increase physical strength and Activity Daily living following a five (5) months rehabilitation medicine using the NMES, ROM, active breathing, endurance, strengthening, transfer and mobilization exercise, and occupational therapy.

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