

## Predictors of Tracheostomy in Mechanically Ventilated Guillain-Barré Syndrome (GBS) Patients in a Tertiary care Hospital in Bangladesh

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### Abstract

**Background:** Patients suffering from Guillain-Barré syndrome (GBS) may frequently develop an acute respiratory failure and need ventilatory support with prolonged mechanical ventilation (MV) and tracheostomy. **Objective:** The present study was designed to search for factors that could predict the need for prolonged MV and ultimately need of tracheostomy after completion of immune therapy. **Methodology:** This retrospective study was performed in patients with GBS admitted to the intensive care unit (ICU) of National Institute of Neurosciences and Hospital, Dhaka, Bangladesh from January 2017 to December 2018. The patients were initiated mechanical ventilation within the first week of admission and were received mechanical ventilation for more than 2 weeks. Demographic, clinical, biological and electrophysiological data and times of endotracheal intubation, tracheostomy, and mechanical ventilation weaning were prospectively collected for all patients. Sequential daily neurological testing used standardized data collection by the same investigators all along the study period. **Results:** A total number of 74 patients were recruited for this study of which 50% cases required tracheostomy; 34(45.9%) patients needed prolonged mechanical ventilation more than 14 days. The mean duration of mechanical ventilation was 20 days. The strongest observed predictors of tracheostomy were muscle weakness, high CSF protein and Plasma exchange and prolonged mechanical ventilation (P value <0.05). These patients are more likely to need of tracheostomy. **Conclusion:** In conclusion ventilated GBS patients who have more muscle weakness at presentation, high CSF protein and prolonged mechanical ventilation are high risk of tracheostomy. [*Journal of National Institute of Neurosciences Bangladesh, January 2022;8(1): 23-27*]

**Keywords:** Guillain-Barré syndrome; mechanical ventilation; tracheostomy

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### Introduction

Respiratory failure is a life-threatening manifestation of the Guillain-Barré syndrome (GBS) that occurs in 20.0% to 30.0% of patients with GBS<sup>1-4</sup>. The duration of the required MV varies widely in GBS, ranging from a few days to several months and even longer than 1 year. Immunomodulatory treatment reduces the proportion of

patients who require mechanical ventilation (MV) as well as the duration of mechanical ventilation<sup>5-6</sup>. The patients may need duration of mechanical ventilation for variable days may complicate this decision in tracheostomy in clinical practice. In general, tracheostomy should be considered when the expected ventilation duration is more than 14 days<sup>7-8</sup>.

Delayed tracheostomy in ventilated patients may result in avoidable damage of the vocal cords, laryngeal mucosa, and recurrent laryngeal nerves due to decubitus or local pressure from the endotracheal tube<sup>7</sup>. On the other hand, early tracheostomy may be unnecessary because of clinical improvement and exposes patients to the risk of perioperative bleeding, infection, esophageal perforation, pneumothorax, and tracheal stenosis and, in all cases, leaves a permanent scar<sup>9</sup>. Previous studies showed that the clinical course of GBS in individual patients can be predicted with reasonable accuracy<sup>10-14</sup>. In the current study, it had been described the need of tracheostomy on the basis of duration of mechanical ventilation and characteristics of GBS patients and aimed to identify predictors of tracheostomy. These predictors may support individual clinical decision-making about indication and timing of tracheostomy in patients with GBS patients in the course of their disease.

### Methodology

**Study Setting and Population:** This retrospective study data were collected from patients with GBS in National Institute of Neurosciences and Hospital, Dhaka, Bangladesh from January 2017 to December 2018, who initiated mechanical ventilation within the first week of admission and who received mechanical ventilation for more than 2 weeks. Early tracheostomy was defined as tracheostomy performed within 14 days of mechanical ventilation. Potential predictors for prolonged mechanical ventilation, defined as duration of  $\geq 14$  days, were considered using crosstabs. Consecutive patients with GBS having respiratory involvement were prospectively evaluated. The study has been approved by Institute Ethics Committee. Prospectively collected data were used from GBS patients requiring mechanical ventilation who fulfilled the diagnostic criteria for GBS, were treated with either plasma exchange or intravenous immunoglobulin.

**Evaluation:** Evaluation of detailed medical history including preceding events such as flue like illness, rash, diarrhea, vaccination, trauma, pregnancy, childbirth and surgery were noted. Baseline characteristics were noted. Presence of cranial nerve palsy was recorded. Muscle weakness was graded on a 0 to 5 MRC (Medical Research Council) scale. The patients were closely followed up in ICU. Pulse, blood pressure, respiration and pulse oxymetry were continuously monitored in all. The ABG analysis was done daily or more frequently if needed. Blood

counts, hemoglobin and serum electrolytes were done twice weekly and X-ray chest weekly or earlier if indicated. Blood culture, endotracheal or tracheostomy tube aspirate culture and urine culture were carried out as indicated. Pneumonia, pneumothorax, urinary infection, pressure sore, sepsis, deep vein thrombosis and pulmonary thromboembolism were closely monitored. Tracheostomy was done usually after 2 weeks of intubation based on the clinical condition. The patients admitted within 14 days of illness were prescribed IVIg 400 mg/kg/day for 5 days or plasma exchange every alternative day for consecutive 5 cycles which one they could afford. Enoxaparin prophylaxis was given to the patients who had severe weakness (MRC grade).

**Study Procedure and Data Collection:** Data collected prospectively for all patients were age, gender, preceding infections, number of days from onset of weakness to hospital admission, date of intubation and extubation, and neurological examination (cranial nerve testing, sensory and motor testing; using the Medical Research Council [MRC] grade during admission, the muscle scale grades muscle power on a scale of 0 to 5 in relation to the maximum expected for that muscle. For this study, we recorded neurological examination after admission in ICU requiring MV. Nerve conduction studies were performed in the first 2 weeks after inclusion, and the data were used to classify GBS as acute inflammatory demyelinating polyradiculoneuropathy (AIDP), acute motor axonal neuropathy (AMAN) and AMSAN

**Endpoints:** The primary endpoint in this study was the occurrence of prolonged mechanical ventilation which was defined as mechanical ventilation of more than 14 days, as an established criterion to consider tracheostomy<sup>7-8</sup>. Predictors of tracheostomy were sought at day 14 after start of ventilation, as a clinical decision point for considering tracheostomy.

**Statistical Analysis:** The patients with respiratory impairment requiring mechanical ventilation were compared for demographic details, age, sex, comorbidity, antecedent events, pattern of weakness, MRC grade, type of NCS, mode of treatment, need of tracheostomy, using X<sup>2</sup> for categorical and independent 't' test or Mann-Whitney U-test for continuous variables. The predictors of tracheostomy were evaluated by Chi-square test. The P-values of  $<0.05$  were considered as significant. The statistical analysis was done by SPSS version 16 software.

**Results**

A total number of 74 patients were recruited for this study. The median age was 27 (range 3 to 65) years and 51 were male and 23 were females. The triggering events were patients diarrheal illness 53 (71.8%), RTI illness in 18(24.3%), others in 3 (4.1%) patients. The median peak disability was MRC grade -1 (56.8%). Facial weakness was present in 28(37.8%), bulbar weakness in 36(48.6%). After nerve conduction

studies, the patients were categorized into AIDP in 19 (25.7%), AMAN in 46(62.2%), AMSAN in 3(4.1%). Intravenous immunoglobulin was given to 58 (78.4%) patients and plasma exchange in 16 (21.6%) patients (Table 1).

Out of 74 GBS patients with respiratory compromise, all needed mechanical ventilation based on ABG criteria after a median duration of 20 days of hospitalization. 63.71% patients were prolonged mechanical ventilation (> 14 days). Comparing the clinical data between the tracheostomized and

Table 1: Base line characteristics of BGS patients

Variables	Frequency	Percent
<b>Age Group</b>		
Less Than 20 Years	20	27.0
21 to 40 Years	36	48.6
More than 40 Years	18	24.3
<b>Gender</b>		
Male	51	68.9
Female	23	31.1
<b>Comorbidity</b>		
Diabetes	36	48.6
Brochial Asthma	10	13.5
Ischemic heart disease	18	24.3
Hypertension	10	13.5
<b>Antecedent events</b>		
Respiratory tract infection	18	24.3
Gastrointestinal tract infection	53	71.6
Others	3	4.1
<b>Pattern of weakness</b>		
Ascending	37	50.0
Descending	37	50.0
<b>Cranial nerve involvement</b>		
Facial nerve	28	37.8
Bulbar	36	48.6
No	10	13.5
<b>MRC score on admission</b>		
Grade-0 (Complete paralysis)	8	10.8
Grade-1 (minimal contraction)	42	56.8
Grade-2 (active movement with gravity eliminated)	17	23.0
Grade-3 (muscle movement is possible against gravity)	7	9.5
Grade-4 (weak contraction against gravity)	0	
Grade-5 (normal strength)	0	
<b>Types of Nerve conduction study</b>		
AIDP	19	25.7
AMAN	46	62.2
AMSAN	3	4.1
NCS not done	6	8.1
<b>Immunomodulatory treatment</b>		
Plasma exchange	16	21.6
IV Immunoglobulin	58	78.4
<b>Duration of Mechanical ventilation</b>		
Less Than 14 days	27	36.48
More Than 14 days	47	63.7

Table 2: Predictors of Tracheostomy

Variables	Tracheostomy		P value
	Yes	No	
<b>Age Group</b>			
3 to 20 Years	14	22	0.292
21 to 40 Years	16	34	
More than 41 Years	7	11	
<b>Gender</b>			
Male	29	22	0.079
Female	8	15	
<b>Co-morbidity</b>			
Yes	6	30	0.384
No	9	27	
<b>Pattern of weakness</b>			
Ascending	20	17	0.642
Descending	17	20	
<b>Antecedent events</b>			
RTI	12	6	0.324
GI Infection	24	29	
Chicken pox	1	2	
<b>Cranial nerve involvement</b>			
Facial nerve	20	12	0.131
Bulbar palsy	17	25	
<b>MRC Grade</b>			
Grade: 0-3	36	1	0.035
Grade : >3	1	36	
<b>NCS</b>			
AIDP	8	11	0.213
AMAN	25	25	
AMSAN	4	2	
<b>CSF Protein</b>			
Less Than 100 mg/dl	26	6	0.032
More Than 100 mg/dl	11	28	
<b>CSF Lymphocyte</b>			
0 to 3 per ml	30	7	0.409
4 to 6 per ml	29	8	
<b>Mechanical Ventilation</b>			
Less Than 14 Days	7	30	0.017
More Than 14 Days	34	3	
<b>Immunomodulatory Treatment</b>			
IV Immunoglobulin	23	35	0.001
Plasma exchange	14	2	

non-tracheostomized patients, there was no significant difference in the age ( $p=0.29$ ), sex ( $p=0.07$ ), co-morbidity ( $p=0.38$ ), pattern of weakness ( $p=0.64$ ) NCS variety ( $p=0.21$ ), CSF cell count ( $p=0.40$ ) and antecedent events ( $p=0.32$ ). The tracheostomized patients had significantly more disability ( $p=0.03$ ), high CSF protein ( $p=0.032$ ) and prolong duration of mechanical ventilation ( $p=0.017$ ). The patients who were treated with intravenous immunoglobulin were significantly less need of tracheostomy in comparison to who were treated with plasma exchange ( $p=0.001$ ) (Table 2).

### Discussion

In this study, all patients with GBS had respiratory compromise and all needed mechanical ventilation on the basis of clinical and ABG criteria. Severity of weakness (MRC grade) was an independent predictor of mechanical ventilation. In our study, we have not intubated the patients with bulbar weakness unless they had hypoxia or ABG abnormality. In this study, AIDP constituted 25.7% and AMAN in 62.2% patients. Higher frequency of AMAN although have been reported in the developing countries of Asia and America but on detailed nerve conduction studies, we have also found AMAN as a predominant subtype of GBS<sup>15-20</sup>.

In the current study, ventilated and tracheostomized patients were not significant difference on variety of GBS requiring prolonged mechanical ventilation. One study revealed AMAN variety were associated more need of tracheostomy and ultimately prolonged ventilation<sup>21</sup>. One study indicated that the presence of AIDP was associated with a higher chance of respiratory failure<sup>22</sup> but we were unable to confirm that finding. The electrophysiology results are influenced by the applied classification criteria and the timing of the NCS. In Western countries, the axonal forms of GBS are relatively rare compared to AIDP and are found in 5.0% to 10.0% of GBS patients. In addition, NCS performed at 1 week is less accurate for identifying axonal GBS, as the axonal pattern may appear only after 2 to 4 weeks. At 1 week of admission, patients more frequently show unexcitable nerves in NCS. These patients may have either AIDP or axonal forms, but in all cases this is a sign of severe diffuse neuropathy. As such, unexcitable nerves may be a more frequent indication than AMAN for early tracheostomy. In most studies, older age is a predominant prognostic factor for poor outcome in GBS, including those of our own group.

In the current study, we found no association between age and prolonged MV neither did other previous studies. Also the presence of a preceding *Campylobacter jejuni* infection, which is a general poor prognostic factor in GBS, was not predictive for prolonged mechanical ventilation.

The decision for tracheostomy in patients with GBS depends on the expected duration of respiratory failure, which may range from a few days to more than 6 months. In the current study, 63.71% patients were prolonged mechanical ventilation (more than 14 days). The chance of prolonged MV were MRC grade 0 or 1 bilaterally, high protein in CSF and plasma exchange instead of IVIG. In these patients, it may be considered to perform an early tracheostomy. Several opinions still ongoing about the optimal timing of tracheostomy. A consensus report on MV indicated that the translaryngeal route is preferred when the expected duration is not exceeding 10 days, while tracheostomy is preferred for expected durations longer than 21 days<sup>23</sup>. Prolonged MV via the translaryngeal route carries significant risks, while tracheostomy has its own complications and leaves permanent disfigurement. Nowadays prospective trials show that early tracheostomy was associated with less sedative and analgesic administration, less frequent prescriptions of haloperidol to treat agitation or delirium, earlier oral nutrition, and out-of-bed mobilization. Early tracheostomy does not seem to shorten the duration of MV, length of hospital stay, mortality, or frequency of infectious complications<sup>8, 24-27</sup>.

The current study has several limitations that need to be addressed. First, the group of ventilated patients was too small to be able to develop and validate a prognostic model, as was done previously for predicting respiratory failure in the first week in GBS<sup>11</sup>. Second, the patient population investigated was biased toward adult patients and patients with AMAN, which is the predominant GBS subtype in our study. The observed finding at present cannot be extrapolated to pediatric GBS or countries where AIDP forms predominate. Third, in this multicenter study, differences in the duration of MV may reflect variation in local clinical management; intubation or extubation criteria were not used in our patient group. Also, usage and timing of tracheostomy was not recorded in our cohort, and this probably influenced the duration of MV. Fourth, we cannot exclude that the criteria for extubation and supportive care have changed over time. In the future, larger cohorts of GBS patients, with clear



definitions regarding extubation criteria, will be needed to substantiate our findings.

### Conclusion

Ventilated GBS patients who have more muscle weakness at presentation, high CSF protein, prolonged MV are high risk of tracheostomy. Whereas the patients who were treated with intravenous immunoglobulin were significantly less need of tracheostomy in comparison to who were treated with plasma exchange.

### References

- Dhar R, Stitt L, Hahn AF. The morbidity and outcome of patients with Guillain-Barré syndrome admitted to the intensive care unit. *J Neurol Sci.* 2008; 264(1-2):121-128.
- Fletcher DD, Lawn ND, Wolter TD, Wijdicks EF. Long-term outcome in patients with Guillain-Barré syndrome requiring mechanical ventilation. *Neurology.* 2000;54(12):2311-2315.
- Rees JH, Thompson RD, Smeeton NC, Hughes RA. Epidemiological study of Guillain-Barré syndrome in south east England. *J Neurol Neurosurg Psychiatry.* 1998;64(1):74-77.
- Winer JB, Hughes RA, Osmond C. A prospective study of acute idiopathic neuropathy. I. Clinical features and their prognostic value. *J Neurol Neurosurg Psychiatry.* 1988;51(5):605-612.
- Hughes RA, Swan AV, van Doorn PA. Intravenous immunoglobulin for Guillain-Barré syndrome. *Cochrane Database Syst Rev.* 2014;(9):CD002063.
- Raphaël JC, Chevret S, Hughes RA, Annane D. Plasma exchange for Guillain-Barré syndrome. *Cochrane Database Syst Rev.* 2012;(7):CD001798.
- Durbin CG., Jr Tracheostomy: why, when, and how? *Respir Care.* 2010;55(8):1056-1068.
- Trouillet JL, Luyt CE, Guiguet M, et al. Early percutaneous tracheotomy versus prolonged intubation of mechanically ventilated patients after cardiac surgery: a randomized trial. *Ann Intern Med.* 2011;154(6):373-383.
- Wijdicks EF, Lawn ND, Fletcher DD. Tracheostomy scars in Guillain-Barré syndrome: a reason for concern? *J Neurol.* 2001;248(6):527-528.
- van Koningsveld R, Steyerberg EW, Hughes RA, Swan AV, van Doorn PA, Jacobs BC. A clinical prognostic scoring system for Guillain-Barré syndrome. *Lancet Neurol.* 2007;6(7):589-594.
- Walgaard C, Lingsma HF, Ruts L, et al. Prediction of respiratory insufficiency in Guillain-Barré syndrome. *Ann Neurol.* 2010;67(6):781-787.
- Walgaard C, Lingsma HF, Ruts L, van Doorn PA, Steyerberg EW, Jacobs BC. Early recognition of poor prognosis in Guillain-Barré syndrome. *Neurology.* 2011;76(11):968-975.
- Lawn ND, Wijdicks EF. Post-intubation pulmonary function test in Guillain-Barré syndrome. *Muscle Nerve.* 2000;23(4):613-616.
- Fourrier F, Robriquet L, Hurtevent JF, Spagnolo S. A simple functional marker to predict the need for prolonged mechanical ventilation in patients with Guillain-Barré syndrome. *Crit Care.* 2011;15(1):R65.
- Kalita J, Misra UK, Goyal G, Das M. Guillain-Barré syndrome: subtypes and predictors of outcome from India. *J Peripher Nerv Syst* 2014; 19:36-43
- Islam Z, Jacobs BC, van Belkum A, Mohammad QD, Islam MB, Herbrink P, et al Axonal variant of Guillain-Barré syndrome associated with *Campylobacter* infection in Bangladesh. *Neurology* 2010; 74:581-7
- Kalita J, Misra UK, Das M. Neurophysiological criteria in the diagnosis of different clinical types of Guillain-Barré syndrome. *J Neurol Neurosurg Psychiatry* 2008; 79:289-93
- Nachamkin I, Arzarte Barbosa P, Ung H, Lobato C, Gonzalez Rivera A, Rodriguez P, et al Patterns of Guillain-Barré syndrome in children: results from a Mexican population. *Neurology* 2007; 69:1665-71.
- Misra UK, Kalita J. Patterns of Guillain-Barré syndrome in children: results from a Mexican population. *Neurology* 2008; 71:1203-4.
- Fourrier F, Robriquet L, Hurtevent JF, Spagnolo S. A simple functional marker to predict the need for prolonged mechanical ventilation in patients with Guillain-Barré syndrome. *Crit Care.* 2011;15(1):R65.
- Lawn ND, Wijdicks EF. Tracheostomy in Guillain-Barré syndrome. *Muscle Nerve.* 1999;22(8):1058-1062.
- Durand MC, Porcher R, Orlikowski D, et al. Clinical and electrophysiological predictors of respiratory failure in Guillain-Barré syndrome: a prospective study. *Lancet Neurol.* 2006;5(12):1021-1028. doi: 10.1016/S1474-4422(06)70603-2.
- Plummer AL, Gracey DR. Consensus conference on artificial airways in patients receiving mechanical ventilation. *Chest.* 1989;96(1):178-180.
- Nieszkowska A, Combes A, Luyt CE, et al. Impact of tracheotomy on sedative administration, sedation level, and comfort of mechanically ventilated intensive care unit patients. *Crit Care Med.* 2005;33(11):2527-2533.
- Rumbak MJ, Newton M, Truncate T, Schwartz SW, Adams JW, Hazard PB. A prospective, randomized, study comparing early percutaneous dilational tracheotomy to prolonged translaryngeal intubation (delayed tracheotomy) in critically ill medical patients. *Crit Care Med.* 2004;32(8):1689-1694.
- Terragni PP, Antonelli M, Fumagalli R, et al. Early vs late tracheotomy for prevention of pneumonia in mechanically ventilated adult ICU patients: a randomized controlled trial. *JAMA.* 2010;303(15):1483-1489.
- Wang F, Wu Y, Bo L, et al. The timing of tracheotomy in critically ill patients undergoing mechanical ventilation: a systematic review and meta-analysis of randomized controlled trials. *Chest.* 2011;140(6):1456-1465.