Effect of Application of Endotracheal Suction Guidelines on Cardiorespiratory Parameters of Mechanically Ventilated Patients

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Abstract

Background One of the highly vital duties done by nurses in the intensive care units (ICU) is endotracheal suctioning (ES). It is done for maintaining airway and improving ventilation and oxygenation; however, it must be done in a way so that, in addition to the mentioned advantages, it can have the least complications for the patient. Kelher et al. ⁽⁵²⁾ showed that some nurses performed ES without consideration to recommended steps so excellence of this nursing intervention was less than predicted. Therefore practicing ES with adherence to recommended guidelines is an inquiry, so it is essential to examine routine ICU method of ES applied by nurses compared with standard one that is based on the guidelines.

Objective To evaluate Effect of application of endotracheal suction guidelines on cardiorespiratory parameters of mechanically ventilated patients.

Method 60 adult patients who admitted to ICU were included in this study. The tool "Cardiorespiratory parameters of mechanically ventilated patient record" was used to collect data. The period of study was from October 2012 to March 2013.

Results Regarding comparing between the standard versus routine ICU methods of endotracheal suction using open suction system, findings revealed that after the standard ES, there was significant improvement in respiratory tidal volume, partial pressure of arterial oxygen, and arterial oxygen saturation.

Conclusions the standard ES method had fewer side effects such as hemodynamic instability, hypoxemia and impaired ventilation when compared with routine ES one.

Key words : Cardiorespiratory, guidelines, nurse, open, routine, suction.

I. Introduction

Mechanical ventilation (MV) and tracheal intubation significantly impair airway secretion clearing causing endotracheal suctioning (ES) of secretions is necessary. Endotracheal suction is defined as: "A component of bronchial hygiene therapy which involves mechanical aspiration of pulmonary secretions from a patient with an artificial airway in place". It is achieved by nurses to obtain effective oxygenation. ⁽¹⁻⁴⁾

Despite the necessity of ES plus some patients reported that the procedure eases their breathing, this approach can lead to complications such as damage to the trachea, bleeding, infection, cardiovascular and hemodynamic disturbances, impaired blood gas exchange and hypoxemia, bronchoconstriction, atelectasis and increased intracranial pressure. Moreover, ES is a source of pain and discomfort for MV patients. This discomfort does not diminish during a course of MV; on the contrary, it may worsen. These complications lead to an increase in the patient's stay in hospital and intensive care unit (ICU). (5, 6)

Technique of ES has a consequence on its harms and usefulness. Nowadays, two systems are available to perform ES: open suction system (OSS) and closed suction system (CSS). OSS requires disconnecting between patient and ventilator when performing suction, which is not necessary to do this disconnection with CSS. Moreover, CSS needs special designed catheter which is more expensive and not easy to be available. Clinical studies have demonstrated that OSS can cause alveolar decruitment and transient hypoxemia. ^(7, 8) In Egypt, OSS is the most common used technique because its suction catheter is less cost and more available.

Studies done by Day et al. ⁽⁹⁾ and Pedersen et al. ⁽¹⁰⁾ provided data on when and how to perform ES reflecting that some patients may receive several times of ES daily according to patient's need. ^(11, 12) In 2010 the American Association of Respiratory Care (AARC) published the AARC Clinical Practice Guidelines on ES of mechanically ventilated patients with artificial airway as ES is done if needed only; providing hyperoxygenation before, during and after ES; ES should be done without disconnection between patient and ventilator; avoiding routine normal saline instillation before ES; use CSS for adults with high FiO2, or PEEP; Suction tube dimeter is less than 50% the lumen of the endotracheal tube in adults, maximum suction duration is

from 10 to 15 seconds. Failure to meet the guidelines in implementation of this procedure can result in numerous side effects. $^{(13, 14)}$

A few studies ⁽¹⁵⁻¹⁸⁾ examined if guidelines for ES were tracked by nurses in ICU. The results demonstrated that nurses were often not alert of these guidelines. Moreover, they found that there was a considerable difference reflected in nurses' performance between ideal standard ES where nurses followed the recommended guidelines and actual routine technique when nurses didn't.⁽¹⁷⁻²⁰⁾ Additionally, Day et al. ⁽¹⁷⁾ and Sole et al. ⁽¹⁹⁾ stated that practice of ES was not always based on current research recommendations, which may lead to inconsistent practice among nurses, affecting patients' experience and increasing health risks associated with ES so it is necessary to examine the actual routine method of ES practiced by nurses in ICU and its effect on patient's condition in comparison with the standard ES that follows evidenced based recommendation. Based on this, the aim of this study was to evaluate effect of application of endotracheal suction guidelines on cardiorespiratory parameters of mechanically ventilated patients.

Research questions

• Is there a difference between the effect of standard and routine ICU method of endotracheal suction on cardiorespiratory parameters of MV patients?

Operational definition

- Endotracheal suction in this study refers to endotracheal suctioning using open suction system.
- Standard endotracheal suction refers to guidelines based endotracheal suction.
- Routine endotracheal suction refers to endotracheal suction according to usual ICU care.

II. Materials and Method

Design was quasi experimental. **Setting**

The study was achieved at Causality Care Unit namely unit one and General Intensive Care Unit namely unit three of Main university Hospital, University of Alexandria, Egypt.

Patients

A convenience sample of 60 adult critically ill patients of both genders was involved in the study based on the power analysis (Epi-Info program) with the following information (population size = 90 over 3 months, expected frequency = 50%, acceptable error = 5%, confidence coefficient = 95%.

Patients were selected according to inclusion criteria; hospitalized < 48 hrs, diagnosed with respiratory failure, on mechanical ventilation with endotracheal tube& pressure controlled mode, attached with arterial line, free of infection were included in this study. Exclusion criteria; V/S unstable, refractory hypoxemia (low oxygen saturation {SpO2<90%} in spite of providing high oxygen concentration {FiO2 \ge 60%}) and diagnosed with chronic obstructive pulmonary disease.

Tool: The used tool was "**Cardiorespiratory parameters of mechanically ventilated patient record**". It was a developed tool by the researcher after reviewing the related literature ^(2, 3, 10, 21-23). It was used to gather data of cardiorespiratory parameters to compare between actual routine method practiced by nurses in ICU and guidelines based method that are restricted to evidenced based recommendations. It consisted of two parts: **Part 1**: it includes data of patient's demographics, past health history, medical diagnosis and smoking.

Part 2: cardiorespiratory parameters record including cardiac parameters (HR and MAP), respiratory parameters (RR and V_t), oxygenation parameters (PaCO₂, PaO₂, and SaO₂). All these parameters used to compare between the routine and standard technique of ES procedure.

Tool development, validity and reliability:

The tool was developed by the researcher based on reviewing the relevant literature ^(2, 3, 10, 21-23). The tool was reviewed for content validity, by professors in the following fields: critical care nursing, critical care medicine, anaesthesia and medical biostatistics. Based on their judgement, the required necessary modifications were done. The tool reliability was tested using internal consistency methods (Alpha Cronbach test) by using SPSS version 20 as following: in the tope toolbar select: Analyze > Scale > Reliability Analysis. Its result was $\alpha = 0.714$, which indicates an accepted reliability of the tool.

Pilot study:

A pilot study was carried out on five (12%) patients who were recruited from study sample to test feasibility and applicability of the tool and necessary modification was done to prepare the tool for final form.

Method

An official approval was obtained from "head of Critical Care & Emergency Nursing Department", and "Nursing Ethical Committee" of Nursing Faculty- Alexandria University and hospital authority "Medical Ethical Committee" after explaining the aim of study. An explanation about the aim and nature of present study

was introduced to critical care unit administrators and nurses. Each nurse was informed that sharing in that study was voluntary.

Patients were selected based on the required criteria. Patients were randomly divided into two groups as following; each selected ICU setting had two halls, first hall patients were assigned to the control group and the second hall patients were assigned to the study group.

Thirty patients (n=30) in each group where:

The control group received routine ES according to usual ICU routine care.

The study group received standard ES based on the recommended guidelines

Data collection

Patient's related characteristics using part 1 of the developed tool were obtained for both groups. Collecting data process occurred once for each patient through three forms; **assessing routine ES method** for the control group, **practicing standard ES method** for the study group and **recording cardiorespiratory parameters** for both groups.

For the **control group** (routine ES method group):

Assessing routine ES method: the researcher assessing routine ES method applied by the assigned nurses as suctioning was done with its usual method with open suction system as following: selecting the suction catheter size did not follow specific principles as. Patient's hyper-oxygenation with 100% oxygen was not always done before, during or after ES. Sometimes suction catheter was entered into the tracheal tube when the suctioning device was switching on or off. There was no control on suction machine pressure level. Duration of suctioning did not have specific time and it was stopped when the nurse diagnosed that there was no remaining secretion. For the study group (standard ES method group):

Practicing standard ES method: After assessing patient's need for ES as increased copious retained secretions in lung, $SpO_2 < 90\%$ and $V_t < 300$ ml. The researcher applied guidelines based ES method using open suction system based on the confirmed protocol of AARC ⁽¹³⁾ as following: choosing suction tube size (for male & female patients) by the formula: endotracheal tube size - 2 x 2 cm. Suction pressure was controlled from 120 to 150 mmHg. The patient was hyper-oxygenated for one minute with 100% oxygen before ES. Peripheral oxygen salutation SpO_2 and heart rate were checked. The suction catheter was brought down to patient's trachea until resistance (max: 6 inches =15 cm) was met then the catheter was drawn about 2 cm. The suctioning machine was then switched on and suctioning was done for a maximum of 15 seconds then suction tube was removed and patient was attached to MV and again the patient was hyper-ventilated for one minute with 100% oxygen in between suction passes and after suction. SpO₂ and HR were rechecked. ^(10, 13)

For **both groups**:

Recording cardiorespiratory parameters: The researcher recorded cardiorespiratory parameters (MAP, HR, RR, V_t , PaCO₂, PaO₂ and SaO₂) using part 2 of the tool. HR was obtained from radial pulse. Blood pressure was measured manually then MAP was calculated by formula of (systolic blood pressure + 2 diastolic blood pressure)/3. An arterial blood sample was taken from arterial line four times; immediately before, after 1, 10, and 30 min after suctioning to determine PaCO₂, PaO₂, and SaO₂.

To evaluate effect of the standard against routine ES method on cardiorespiratory parameters of patients attached with MV, comparison was done between the two groups regarding readings of cardiorespiratory parameters before and after ES.

Administrative and ethical considerations

The study was done at about seven months (March 2012 - September 2013). Informed written consent was obtained from conscious patient or family member (if unconscious patient), after explaining the potential benefits and hazards from participation. The anonymity, privacy, confidentiality, and the right to refuse to participate in the study were assured.

Statistical analysis of the data ⁽²⁴⁾

Data were fed to the computer and analyzed using IBM SPSS software package version $20^{(25)}$ Analysis and interpretation of data were done using the followings; Frequency, mean and standard deviation and t-test were used to test the significance of results of quantitative variables. P is significant if ≤ 0.05 .

III. RESULTS

Table 1 presents the sample distribution according to patients' characteristics. The mean age value was 46.20 ± 12.81 in the control group and 44.73 ± 13.88 in the study group. Most patients were non-smoker in both of the control and study groups (83.3 and 73.3% correspondingly). On admission, about half of patients of the control and study groups had diagnosis related to respiratory and other disorders (46.67% and 56.7% correspondingly). The difference between the control and study groups as regards sex, age, presence of history of respiratory disease, smoking and current diagnosis were statistically non-significant (p = 0.795, 0.672, 0.744, 0.347 and 0.740 correspondingly).

Patients' characteristic	Control (N=30)	Study (N=30)	Significance test	
Sex [N (%)]			$\chi^2 = 0.067$	
Male	14 (46.67%)	13 (43.3%)	$\mathbf{P} = 0.795$	
Female	16 (53.33%)	17 (56.7%)	1 = 0.795	
Age [X ±SD]	46.20 ± 12.81	44.73 ± 13.88	t = 0.425 P = 0.672	
Presence of history of respiratory disease [N (%)]				
Yes	22 (73.33%)	21(70%)	χ2 = 0.082	
No	8(26.67%)	9 (30%)	P = 0.744	
Smoking [N (%)]			0 0.004	
Smoker	5 (16.67%)	8 (26.7%)	$\chi 2 = 0.884$ P = 0.347	
Non smoker	25 (83.33%)	22 (73.3%)	r = 0.347	
Current diagnosis [N (%)]				
Respiratory	6 (20%)	5 (16.67%)	χ2 = 0.603	
Respiratory and other	14 (46.67%)	17 (56.7%)	$\mathbf{P} = 0.740$	
Other	10 (33.33%)	8 (26.67%)		

Table 1: Sample distribution according to patients' characteristics

* P is significant if ≤ 0.05

Table 2 reveals a comparison between the control and study groups as related to cardiac parameters including HR and MAP and respiratory parameters including RR and Vt at immediate before and 1, 10 and 30 min after ES. The results displayed that there was no significant statistical variance between the two groups immediately before suction regarding HR, MAP, RR and Vt (P = 0.547, 0.588, 0.573 and 0.558 correspondingly). While after suctioning, HR increased in both groups with non-significant difference at 1, 10 and 30 min (P = 0.370, 0.344 and 0.072 correspondingly), MAP decreased in both groups with non-significant difference at 1, 10 and 30 (0.562, 0.635 and 0.860 respectively), RR decreased significantly in control group at 1 and 10 min (P = 0.001 and 0.040 correspondingly) and Vt increased significantly in study group at 1, 10 and 30 min (P = 0.000, 0.008 and 0.010 correspondingly).

Table 2: Comparison between the control and study groups as regards cardiac and respiratory parameters							
at immediate before and after endotracheal suction at different intervals.							

Param	eter [X ±SD]		Immediately	after suction						
Group			before suction	1min	10 min	30 min				
a Control			86.43 ± 17.56	96.43 ± 20.29	95.20 ± 18.64	96.67 ± 15.30				
Image: matrix for the standard strength of the strengend of the strength of the strength of the strength of t			88.7 ± 10.59	92.10 ± 16.73	90.90 ± 16.20	88.57 ± 15.16				
HR (Significance	t	0.605	0.903	0.954	1.833				
Η	test	р	0.547	0.370	0.344	0.072				
Control			96.37 ± 16.58	93.59 ± 19.91	92.41 ± 20.65	91.47± 16.82				
APH Study Significance text			96.69 ± 21.26	95.46 ± 20.01	94.95 ± 20.49	92.20 ± 16.82				
Mu	Significance	t	0.065	0.362	0.477	0.177				
I)	test	р	0.948	0.719	0.635	0.860				
Control Mail Study		18.17 ± 4.57	24.30 ± 7.70	20.80 ± 7.54	18.27 ± 4.95					
			19.10 ± 7.761	18.67 ± 3.96	17.60 ± 3.53	17.03 ± 3.02				
RR (Significance	t	0.568	3.564	2.106	1.165				
de test			0.573	0.001*	0.040*	0.249				
(1	Control		486.87 ± 56.22	412.60 ± 54.23	464.10 ± 82.12	479.50 ± 65.20				
(ml)	Study		475.5 ± 89.57	526.10 ± 116.54	545.63 ± 141.17	546.80 ± 122.78				
V,	Significance	t	0.589	4.837	2.734	2.652				
	test	р	0.558	0.000*	0.008*	0.010 *				

HR: Heart rate. MAP: Mean arterial blood pressure. RR: respiratory rate V_t : Tidal volume. * P is significant if ≤ 0.05 **Table 3** reveals comparison between the control and study groups as regards gas exchange parameters as PaO2, SaO2 and PaCO2 at immediate before and 1, 10 and 30 min after ES. In relation to immediate presuction PaO2, SaO2 and PaCO2, there was no a significant statistical variance between the two groups (P = 0.865, 0.991 and 0.586 correspondingly). On the other hand in the study group, after suction PaO2 increased significantly at 1 and 10 min (P = 0.001 and 0.052 correspondingly) and SaO2 increased significantly at 1, 10 and 30 min (0.000, 0.000 and 0.019 correspondingly). Regarding after suction PaCO2, there was no a statistical significant variance between the two groups at 1, 10 and 30 min (P = 0.449, 0.814 and 0.906 correspondingly).

Gas excl	hange parameter		Immediately	After suction						
[X ± SD] Group			before suction	1min	10 min	30 min				
6	Control		97.09 ± 37.22	87.57 ± 34.27	97.15 ± 38.33	100.62 ± 36.41				
PaO ₂ (mmHg)	Study		95.25 ± 40.89	130.32 ± 58.88	117.96 ± 42.78	111.30 ± 42.09				
PaO nmH	Significance Test	Т	0.171	3.437	1.984	1.051				
I)	Significance Test	Р	0.865	0.001*	0.052*	0.297				
	Control		94.31 ± 3.50	93.93 ± 4.76	93.92 ± 4.20	95.28 ± 3.11				
SaO ₂ (%)	Study		94.81 ± 3.61	97.59 ± 2.08	97.35 ± 2.39	97.09 ± 2.67				
	Significance Test	Т	0.548	3.860	3.880	2.422				
	Significance Test	Р	0.586	0.000*	0.000*	0.019*				
PaCO ₂ (mmHg)	Control		36.71 ± 14.50	39.07 ± 11.98	37.12 ± 15.02	36.11 ± 6.90				
	Study		36.75 ± 15.14	36.57 ± 13.37	36.19 ± 15.51	35.72 ± 6.52				
PaCo mmł	Significance Test	Т	0.011	0.762	0.237	0.118				
I I	Significance Test	P	0.991	0.449	0.814	0.906				

Table 3:	Comparison	between	the	control	and	study	groups	as	regards	gas	exchange	parameters	at
immediate before and after endotracheal suction at different intervals.													

PaO_{2:} partial pressure of arterial oxygen SaO_{2:} Arterial oxygen saturation

PaCO₂: partial pressure of arterial carbon dioxide * P is significant if ≤ 0.05

IV. Discussion

Results of this study showed that insignificant increase in HR in the control group more than in the study group and returned to the base status after 30 min in study group compared with before ES. This finding could be due to irritation by suction tube movements plus accompanied fear, pain and stress which are caused by ES technique itself. This was accepted by Keykha *et al.* ⁽²⁶⁾ who compared effect of standard ES and routine methods on vital signs, arterial blood oxygen saturation at ICU as they found that ES also caused increased HR in the two groups, but in study group, HR returned to the base status after five minutes. Also, Iranmanesh *et al.* ⁽²⁷⁾ and Ackerman ⁽²⁸⁾ indicated that HR elevated in the studied groups receiving ES with or without normal saline. Additionally, Lee *et al.* ⁽²⁹⁾ and Clark *et al.* ⁽³⁰⁾ approved that HR after ES elevated from its baseline. Contrary to this finding, Akbaryan Deheki *et al.* ⁽³¹⁾ who compared three methods of suctioning with normal saline, N-acetyl cysteine and without them reported that the level of HR and BP decreased after ES.

The current study illustrated that no significant decrease in MAP after ES in the two groups. Although MAP decrease was slightly lower in study group. This can be accepted as standard ES technique including: short time for MV disconnection and providing hyperoxygenation to avoid occurrence of hypoxemia as hypoxemia can result in hemodynamic instability. This result is in line with other studies ^(32, 33) which showed that there was no statistical significant difference in BP after ES with or without saline. Preusser *et al.* ⁽³⁴⁾ found through studying patients with coronary artery bypass grafting operation that oxygenation before ES resulted in low fluctuations in MAP readings.

Fisher *et al.* ⁽³⁵⁾ described in their study that elevation in BP next to ES. Parsons *et al.* ⁽³⁶⁾ evaluated the effect of hyperventilation at 100% O₂ followed by ES and found that MAP was significantly elevated above baseline. Ozden *et al.* 's ⁽³⁷⁾ study as well as Cereda *et al.* 's ⁽³⁸⁾ study illustrated that MAP increased following open ES. Moreover, Johnson *et al.* 's ⁽³⁹⁾ findings describing effect of open and closed endotracheal suctioning, MAP increased after ES in both groups.

Concerning respiratory parameters, results of this study showed that RR decreased in the study group and increased in the control one with significant difference between them at immediate 1, and 10 min after suction. The result can be related to restlessness and suffering produced by ES technique. Futter *et al.* ⁽⁴⁰⁾ illustrated that there was a significant elevation in RR after ES resulting from suction producing suffering pain and restless. While, Alavi *et al.* ⁽³³⁾ found that the effect of suction with as well as without normal saline on RR was negligible and changes were not significant.

Regarding V_t, it increased significantly in the study group in comparison with the control one after suction. It can be related to that in the study group, there was increased probabilities of improving inspired volume of air as evidenced by decreased time of patient's disconnection from ventilator, suctioning was done for 10 to 15 seconds, external diameter of the suction catheter was less than half of the internal diameter of the patient's tracheal tube, and suction pressure was controlled from 120 to 150 mmHg. Copnell B. *et al.* ⁽²²⁾ reported that the effect of suction method, force, and tube size on lung volume changes during ES in piglets and found that closed suction preserved more lung volume than open one.

In the current study, PaO_2 and SaO_2 increased significantly in the study group when compared with control group after ES. It can be understood through that the standard ES technique comprised MV disconnection short time and hyperoxygenation which helps to avoid hypoxemia. Kelleher *et al.* ⁽⁴¹⁾ and Day *et al.* ⁽⁴²⁾ found that main deficiencies were related to failure to provide hyperoxygenation before as well as after ES procedure plus keeping suitable suction force. Bourgault *et al.* ⁽⁴⁷⁾ found that PaO₂ elevated significantly post suctioning. Also, Sole *et al.* ⁽¹⁹⁾ stated that increased SaO₂ occurred after suctioning with the standard method. Moreover, other studies ^(38, 43- 46) showed that longer time with ventilation interruption, loss of positive airway pressure, and reduction in lung volume lead to hypoxemia during open suctioning system.

Akgul ⁽⁴⁸⁾ and Najaf Yarandi ⁽⁴⁹⁾ and Akbaryan Deheki *et al* ⁽³¹⁾ showed that SaO_2 reduced after ES in both groups with and without normal saline. Also, Iranmanesh *et al*. ⁽²⁷⁾ reported that negative change in SaO_2 was observed in the group who received ES with normal saline.

In the current study, there was nonsignificant increase in $PaCO_2$ in the control group at 1 and 10 min after suction when compared with the study group. This may be related to that in control group, there was partial limitation in oxygen and carbon dioxide exchange during longer period of patient's disconnection from MV, non-constant administering 100% oxygen before, between and after ES and decreased exhaled tidal volume. Zahran *et al.* ⁽⁵⁰⁾ found significant increase in PaCO₂ after ES with normal saline instillation. However Caramez *et al.* ⁽⁸⁾ found PaCO₂ elevations after ES with open and closed method, the Özden *et al.* ⁽³⁷⁾ and U⁻gras *et al.* ⁽⁵¹⁾ didn't

In addition, Kelher *et al.* $^{(52)}$ showed that some nurses performed ES without consideration to recommended steps so excellence of this nursing intervention was less than predicted. Additionally, Jongerden *et al.* $^{(3)}$ performed a practical study evaluating ES in which the suctioning procedure was done during standard way as needed based on advices. They found that ES wasn't linked with practically disorders in physiologic indicators.

V. Conclusion

From the present results, it can be concluded that MV patients who received ES with standard method based on the recommended guidelines had more stable heart rate, blood pressure, respiratory rate, improved tidal volume and arterial blood oxygenation.

Recommendations

Therefore this study recommends that regarding

Clinical practice:

- Nurses in ICU should closely observe HR, RR, V_{t} and SaO $_{2}$ during ES.

Administration:

- Modern apparatus for practicing and monitoring patients during ES correctly should be available in ICUs.

Education and training:

- Conducting in-service education to nurses about standard ES raising their awareness on its advantage other than routine method
- Conducting in-service training about consideration for adhering strictly to steps of standard ES method such as providing 100% oxygen before-during-after the procedure, maximum of three ES pass, a maximum of 15seconds suction for each suction pass and sufficient resting times to be given to patients between ES passes.

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