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Case Study

## The Impact of CBAHI Accreditation on Critical Care Unit Outcome Quality Measures: A Case Study

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## ABSTRACT

Quality indicators are the tools to measure the patient safety, effectiveness, equity, patientcenteredness, timeliness, and efficiency as defined by the Institute of Medicine (IOM). These measures are also categorized as structure, process and outcome by Dr. Avedis Donabedian.

**Objectives:** To study the impact of the Central Board for Accreditation of Healthcare Institutions (CBAHI) Accreditation on the outcome measures of critical care units in a tertiary care hospital.

**Methods:** This is a library research methodology, in which the analysis of historical records and data was done before and after the CBAHI Accreditation.

**Significance of Research:** It was observed during pre CBAHI Accreditation (from May 2016 to October 2016) and post CBAHI Accreditation (November 2016 to April 2017) that there was no significant improvement in the outcome measures of Critical Care Units.

**Hypothesis:** Null Hypothesis (Ho) and Alternative Hypothesis (H1) were used and tested to compare the pre CBAHI and post CBHAI impact.

**Study Design:** Outcome Quality Measures as per CBAHI Standards were monitored in pre and post CABHI Accreditation and were compared statistically to study the impact of CBAHI Accreditation. **Study Population**: The Outcome Quality Measures for the Critical Care Units as per the CBAHI Standards third edition were monitored from May 2016 to October 2016 (before CBAHI Accreditation) and from November 2016 to April 2017 (after CBAHI Accreditation)

**Data Collections:** Primary data were collected from all Critical Care Units before and after CBAHI Accreditation. Secondary data were collected from relevant published journals, articles, research papers, academic literature and web portals.

**Conclusion:** There was no statistically significant difference between pre-test and post-test results. Although a number of rates significantly differed across units. Those rates were namely mortality rate, DAMA rate, return to the critical care unit within 48 hours of discharge/transfer rate, average length of stay, rate of initial physical assessment done by nurses with acceptable time, patient identification compliance rate and hospital acquired

pressure ulcer (HAPU) rate. Hence, Null Hypothesis (Ho) is accepted and Alternative Hypothesis (H1) is rejected.

*Key words:* Central Board for Accreditation of Healthcare Institutions (CBAHI), Quality Indicators, Critical Care Units, Joint Commission International (JCI) Accreditation

## **INTRODUCTION**

The Saudi Central Board for Accreditation of Healthcare Institutions (CBAHI) is the official agency authorized to grant accreditation certificates to all governmental private healthcare and facilities operating today in Saudi Arabia. CBAHI has emerged from the Saudi Health Council as a non-profit organization. The principal function of CBAHI is to set the healthcare quality and patient safety standards against which all healthcare facilities are evaluated for evidence of compliance.

The foundation of CBAHI dates back to 2001 as Makkah Region Quality Program (MRQP), an initiative aimed at improving quality of healthcare delivery in the Makkah Region. In 2005, under a Ministerial Order, MRQP was developed and named as Central Board for Accreditation of Healthcare Institutions (CBAHI) and its jurisdiction was expanded to the whole country. In 2006, with the help of healthcare quality experts from the public and private sectors, CBAHI developed the first set of national standards for hospitals. In 2012, CBAHI's 2nd edition of national standards for hospitals was certified by the International Society for Quality in Healthcare (ISQua). In late 2013, when a Cabinet of Ministers Decree called for changing CBAHI's official name to the "Saudi Central Board Accreditation for of Healthcare Institutions", it also mandated the national accreditation by CBAHI on all healthcare facilities. In addition, the Ministry of Health is mandated CBAHI accreditation as a prerequisite for renewal of the operating license – a step towards encouraging more participation in this ambitious national initiative. It is mandatory for all public and private health care delivery facilities (hospitals, polyclinics, blood banks and medical laboratories) in Saudi Arabia to comply with national standards set by CBAHI and obtain its accreditation through a survey process set forth by the Center.

The Essential National Requirements for Patient Safety (ESR) is a list of 20 national standards for hospitals. They are deemed to be basic conditions that must be fully observed to ensure patient safety and protection against health care related errors (CBAHI, 2017).<sup>[1]</sup>

## **REVIEW OF LITERATURE**

The increased international focus on improving patient outcomes, safety and quality of care has led stakeholders, policy makers and health care provider organizations adopt standardized processes for measuring health care systems.

Based on the Institute of Medicine (IOM, 1990) definition of quality of care as "the degree to which health care services for individuals and populations increase the likelihood of desired health outcomes and are consistent with current professional knowledge," <sup>[2]</sup> a quality indicator is a tool that enables the user to quantify the quality of a selected aspect of care by comparing it with a criterion (NQMC, 2013). <sup>[3]</sup>

Intensive-Care Units (ICUs) are the most expensive part of a hospital. It is therefore extremely important that they are used in the most efficient way. As in any other business, high quality and costeffective performance in Intensive-Care Medicine (ICM) can best be achieved when responsibility and management are given to those who have the special expertise.

In the past decade, it has become evident that a greater input of intensivists leads to better outcomes for patients and more efficient resource use. This became obvious from a discussion in the United States of America (USA), where ICU

structures differ greatly from those in Western Europe. In the USA, most ICUs are so-called 'open' units, in which critically ill patients in the ICU are cared for by their primary physicians, who are not specialists in ICM. In contrast, a 'closed' unit is one in which a full-time intensivists (or a team of intensivists) provides ICM. Closed ICUs predominate in Western Europe. Now there seems to be an increasing awareness in the USA that the closed ICU may be more efficient (Hilmar Burchardi, Onnen Moerer, 2001). <sup>[4]</sup> There are statistically significant effects (all improvements) associated with accreditation with reduction in return to the Intensive Care Unit (ICU) within 24 hours of ICU discharge; reduction in staff turnover; and completeness of medical records. The net impact of accreditation was a 1.2 percentage point reduction in patients who returned to the ICU, 12.8% reduction in annual staff turnover and 20.0% improvement in the completeness of medical records. Pooling both hospitals over 3 years, these improvements translated into the total savings of US\$ 593 000 in Jordan's healthcare system (Y.A. Halasa, W. Zeng, E. Chappy and D.S. Shepard, 2015).<sup>[5]</sup>

In the recent studies, the researchers have proved that there is a positive impact of health care accreditation on the health care services. The accreditation has a positive impact on the satisfaction of Physiotherapy Department (Shaikh, 2017), <sup>[6]</sup> Pharmacy Department Service (Shaikh, 2017), <sup>[7]</sup> Dietary Department Services (Shaikh, 2017), <sup>[8]</sup> Laboratory Department Services (Shaikh, 2017), <sup>[9]</sup> Emergency Department Services (Shaikh, 2017), <sup>[10]</sup> Out-Patient Department Services (Shaikh, 2018). <sup>[11]</sup> In-Patient Department Services [12] 2017), (Shaikh. Haemodialysis Department Services (Shaikh, 2017), <sup>[13]</sup> Radiology Department Services (Shaikh, 2017), <sup>[14]</sup> Ambulance Services (Shaikh, 2016), <sup>[15]</sup> and also has positive impact on the Occurrence Variance Reports (Shaikh 2018), <sup>[16]</sup> completeness of personnel files in Human Resource Department (Shaikh

2017). <sup>[17]</sup> A comparative study of laboratory and blood bank performance by using the quality indicators revealed that the mean rating of the second half (after the accreditation) is better than the mean rating of the first half (before accreditation) (Shaikh, 2018). <sup>[18]</sup>

The researchers have compared the healthcare accreditation standards and revealed that there are variations among them despite of being accredited by the International Society for Quality in Health Care (ISQua). The critical analysis of Patient and Family Rights (PFR) standards (Shaikh, 2017), <sup>[19]</sup> Patient and Family Education (PFE) standards (Shaikh, Al-[20] and Towyan & Khan, 2016) International Patient Safety Goals (IPSG) standards (Shaikh, Al-Towyan & Khan, 2016) [21] in the Joint Commission (JCI) Accreditation and International Central Board for Accreditation of Healthcare Institutes (CBAHI) standards for hospitals clearly show that the PFR and PFE standards are very comprehensive than the JCI Accreditation standards whereas the IPSG standards in JCI Accreditation are comprehensive than much CBAHI Standards. The critical analysis of Staff **Oualifications** and Education (SOE) standards in JCI Accreditation and Medical Staff (MS) & Staffing Management (SM) standards in Det Norske Veritas (DNV) Accreditation for hospitals clearly shows that the SQE Standards in JCI Accreditation are very comprehensive than the DNV's National Integrated Accreditation for Organizations Healthcare (NIAHO) Accreditation (Shaikh, Al-Towyan & Khan, 2016). [22]

## DATA ANALYSIS:

# **1.1 Descriptive statistics of the various rates**

The following table represents the pre-test, post-test and overall means and their standard deviation.

C. No	Table1. Descriptive statistics	Pre-test Post-test Total						Ν
Sr.No.	Туре							IN
		М	SD	М	SD	М	SD	
1	Mortality Rate	2.95	3.11	3.70	3.92	3.33	3.52	24
2	Discharge Against Medical Advise (DAMA) Rate	8.87	4.65	6.88	3.43	7.87	4.16	24
3	Re-Intubation within 48 Hours Post Extubation Rate	1.58	2.77	3.89	7.49	2.73	5.71	24
4	Return to the critical care unit within 48 Hours of	1.12	2.23	1.04	1.30	1.08	1.80	24
	Discharge/Transfer Rate							
5	Average Length of Stay	5.72	3.65	5.48	3.82	5.60	3.70	24
6	Rate of Initial Physical Assessment done by Nurses with	96.33	3.95	97.51	2.63	96.92	3.37	24
	Acceptable Time							
7	Patient Falls Rate	0.00	0.00	0.00	0.00	0.00	0.00	18
8	Patient Fall with Injury Rate	0.00	0.00	0.00	0.00	0.00	0.00	18
9	High Alert Medication Compliance Rate	99.63	1.55	100.00	0.00	99.81	1.10	24
10	Patient Identification Compliance Rate	99.69	1.08	99.82	0.53	99.75	0.84	24
11	Ventilator Associated Pneumonia (VAP) Rate	1.44	2.64	1.99	4.74	1.71	3.79	18
12	Catheter Associated Urinary Tract Infection (CAUTI) Rate	2.35	9.03	0.59	1.94	1.47	6.50	18
13	Central Line Associated Blood Stream Infection (CLABSI) Rate	5.80	11.96	3.64	5.85	4.72	9.35	18
14	Hospital Acquired Pressure Ulcer (HAPU) Rate	0.90	1.11	0.74	1.11	0.82	1.09	12
	Total	27.87	40.38	27.94	40.54	27.91	40.43	318

#### Table1. Descriptive statistics of the rates

## **1.2 Pre, post-test and across units differences**

#### **1.2.1 Mortality Rate**

To identify whether the mortality rate differed at pre-test and post-test stages and in various units (ICU, CCU, PICU and NICU)a two-way ANOVA was carried out. The results indicated that mortality rate differed across the units ( $F_{3, 40}$ = 17.455, *p* = 0.000). However there was no significant difference between pre-test and post-test mortality rates ( $F_{1, 40}$  = 1.082, *p* = 0.304).

Table2. Two-way ANOVA results of the Mortality rate

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared
Corrected Model	333.986 <sup>a</sup>	7	47.712	7.685	.000	.574
Intercept	531.735	1	531.735	85.651	.000	.682
Unit	325.094	3	108.365	17.455	.000	.567
Test	6.720	1	6.720	1.082	.304	.026
Unit * Test	2.171	3	.724	.117	.950	.009
Error	248.327	40	6.208			
Total	1114.047	48				
Corrected Total	582.312	47				

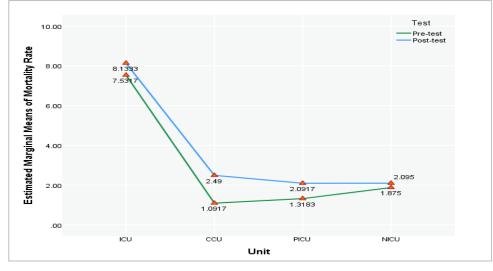


Figure 1. Mean plot of mortality rates in various units and at pre, post-test stages

As in the following figure, ICU had the highest pre-test  $(7.53 \pm 1.77)$  and posttest  $(8.13 \pm 4.61)$  mean mortality rates. Lowest mean was reported by two different

units where at the pre-test stage it was reported by the CCU ( $1.09 \pm .91$ ) and PICU ( $2.09 \pm 1.36$ ) at the post-test stage. These

statistics provide further evidence for the fact that mortality rate differs across units.

Figure 1 also provides graphical interpretation to why there wasn't a significant difference in the pre-test and post-test mortality rate where at each unit pre-test means were closely followed by the post-test mean mortality rates. Hence the pre-test and post-test mortality rates were statistically equal.

## 1.2.2 Discharge against Medical Advise (DAMA) Rate

DAMA rate had the same characteristics when it comes to difference at pre-test, post-test stages and difference between units where DAMA rate differed between units ( $F_{3, 40} = 5.396, p = .003$ ) and not differed between tests ( $F_{1, 40} = 3.638, p = .064$ ). The interaction was not significant either ( $F_{3, 40} = .761, p = .522$ ).

1 abit.	Tables. I we way mano va results of the interaction differences in Dama rate										
Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared					
Corrected Model	290.245 <sup>a</sup>	7	41.464	3.159	.009	.356					
Intercept	2975.490	1	2975.490	226.676	.000	.850					
Unit	212.505	3	70.835	5.396	.003	.288					
Test	47.760	1	47.760	3.638	.064	.083					
Unit * Test	29.980	3	9.993	.761	.522	.054					
Error	525.064	40	13.127								
Total	3790.799	48									
Corrected Total	815.309	47									

Table3. Two-way MANOVA results of the interaction differences in DAMA rate

Two-way ANOVA indicated that pre-test and post-test DAMA rates did not differ significantly. Though the Figure shows some sort of a difference in the pretest and post-test DAMA rates across four units, overall mean of pre-test DAMA rate(M =  $8.8708 \pm 4.64561$ ) and post-test DAMA rate (M =  $6.8758 \pm 3.43366$ ) had no statistically significant difference. Due to that pre-test and post-test DAMA rate means can be considered as statistically equal

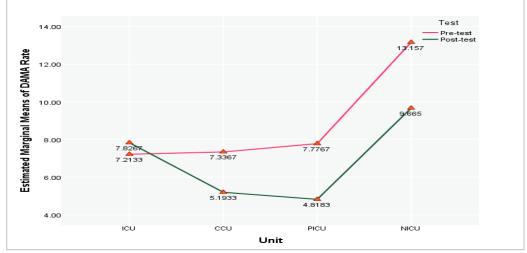


Figure2. Mean plot of DAMA rates in various units and at pre and post-test stages

## **Re-Intubation within 48 Hours Post-Extubation Rate**

Though the Mortality and DAMA rate differed at least across units, the reintubation within 48 hours post-extubation rate didn't differ at least in that aspect ( $F_{3, 40} = .827, p = .487$ ). Also the interaction of unit and the test was not significant either (F<sub>3, 40</sub> = .739, p = .535). Hence the reintubation within 48 hours post-extubation rate can be considered as equal between pre and post-test and across four units. This is mainly due to the fact that in most of the data sample units, rate value was zero, hence the mean was affected by large a standard deviation.

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared
Corrected Model	218.550 <sup>a</sup>	7	31.221	.950	.480	.143
Intercept	358.723	1	358.723	10.920	.002	.214
Test	64.218	1	64.218	1.955	.170	.047
Unit	81.486	3	27.162	.827	.487	.058
Test * Unit	72.846	3	24.282	.739	.535	.053
Error	1313.968	40	32.849			
Total	1891.241	48				
Corrected Total	1532.518	47				

Table4. Two-way ANOVA results of Re-Intubation within 48 Hours Post Extubation Rate

Following figure further proves the fact that there was a considerable number of sample units were their rate was equaled to zero. Hence at the pre-test stage two units

namely CCU and PICU had a mean of zero, which may guide to insignificant test results.

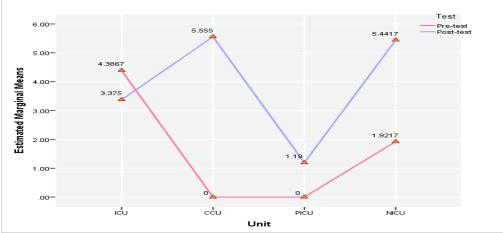


Figure 3. Mean plot of Re-Intubation within 48 Hours Post Extubation Rate

## 1.2.3 Return to the critical care unit within 48 Hours of Discharge/Transfer Rate

Return to the critical care unit within 48 hours of discharge/transfer rate was differed across the four units ( $F_{3, 40} = 4.555 \pm p = .008$ ) here. Yet the difference of the

rate between pre-test and post-test was not significant ( $F_{1, 40} = .028 \pm p = .867$ .)Which indicated that return to the critical care unit within 48 hours of discharge/transfer rate was statistically equal between the pre-test and post-test.

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared
Corrected Model	45.481 <sup>a</sup>	7	6.497	2.416	.037	.297
Intercept	55.492	1	55.492	20.630	.000	.340
Unit	36.760	3	12.253	4.555	.008	.255
Test	.076	1	.076	.028	.867	.001
Unit * Test	8.645	3	2.882	1.071	.372	.074
Error	107.593	40	2.690			
Total	208.566	48				
Corrected Total	153.074	47				

Table5. Two-way ANOVA results of the return to the critical care unit within 48 hours of discharge/transfer rate

#### **1.2.4** Average Length of Stay

As most of the other rates, the average length of stay only differed across units ( $F_{3, 40} = 41.866$ , p = .000). The interaction of the two factors namely unit

and the test was not significant ( $F_{3, 40} = 1.166$ , p = .335). Which indicated that average length of stay was not affected by both the test and unit at once.

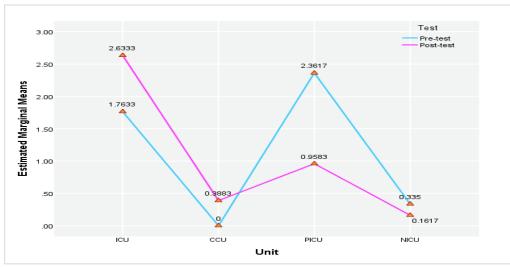


Figure 4. Mean plot of return to the critical care unit within 48 hours of discharge/transfer rate

Table6. Two-way ANOVA results of the average length of stay										
Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared				
Corrected Model	490.727 <sup>a</sup>	7	70.104	18.467	.000	.764				
Intercept	1505.056	1	1505.056	396.461	.000	.908				
Unit	476.794	3	158.931	41.866	.000	.758				
Test	.658	1	.658	.173	.679	.004				
Unit * Test	13.274	3	4.425	1.166	.335	.080				
Error	151.849	40	3.796							
Total	2147.632	48								
Corrected Total	642.576	47								

Both the pre-test and post-test means were close to each other at each unit, for example, the difference between ICU and CCU post-test and pre-test means were

lower than 1. As a result pre-test and posttest means were not significantly differed and can be considered statistically equal.

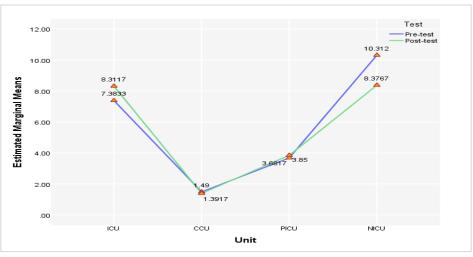


Figure 5. Mean plot of average length of stay

#### **Rate of Initial Physical Assessment** 1.2.5 done by Nurses with Acceptable Time

As most of the other rates, the rate of initial physical assessment done by nurses

with acceptable time differed across units  $(F_{3, 40} = 13.373, p = .000)$ . The interaction of the two factors namely unit and the test was not significant (F<sub>3, 40</sub>= 1.123, p = .351). in other words mean rate of initial physical

assessment done by nurses with acceptable time was different across units but it was

equal between pre-test and post-test.

Table7. Two-way	ANOVA results of	the rate of initial phy	sical assessment don	e by nurses wit	th acceptable time

Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared
Corrected Model	286.483 <sup>a</sup>	7	40.926	6.600	.000	.536
Intercept	450910.608	1	450910.608	72712.232	.000	.999
Unit	248.782	3	82.927	13.373	.000	.501
Test	16.803	1	16.803	2.710	.108	.063
Unit * Test	20.897	3	6.966	1.123	.351	.078
Error	248.052	40	6.201			
Total	451445.143	48				
Corrected Total	534.535	47				

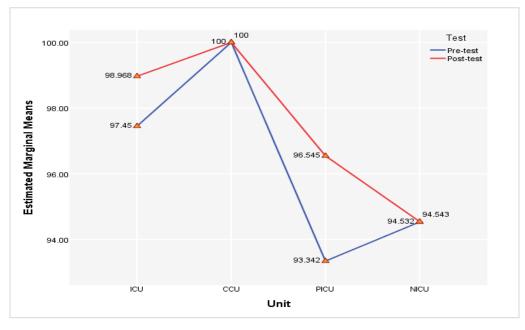


Figure6. Mean plot of the rate of initial physical assessment done by nurses with acceptable time

#### **1.2.6** Patient Falls Rate

Can't compare rates due to the fact that all the values were zero.

#### 1.2.7 Patient Fall with Injury Rate

Can't compare rates due to the fact that all the values were zero.

#### 1.2.8 High Alert Medication Compliance Rate

High alert medication compliance rate wasn't significantly differed between pre-test and post-test ( $F_{1, 40} = 1.470, p =$ .232), across the four units ( $F_{3, 40} = 1.470 p$ = .237). Considering the interaction there was no significance either ( $F_{3, 40} = 1.470 p =$ .237). These statistics indicated that high alert medication compliance rate is completely independents from the tests and four units.

Tables. Two-way ANOVA results of the light alert incucation compliance rate										
Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared				
Corrected Model	11.577 <sup>a</sup>	7	1.654	1.470	.206	.205				
Intercept	478219.654	1	478219.654	425020.384	.000	1.000				
Test	1.654	1	1.654	1.470	.232	.035				
Unit	4.962	3	1.654	1.470	.237	.099				
Test * Unit	4.962	3	1.654	1.470	.237	.099				
Error	45.007	40	1.125							
Total	478276.238	48								
Corrected Total	56.584	47								

Table8. Two-way ANOVA results of the high alert medication compliance rate

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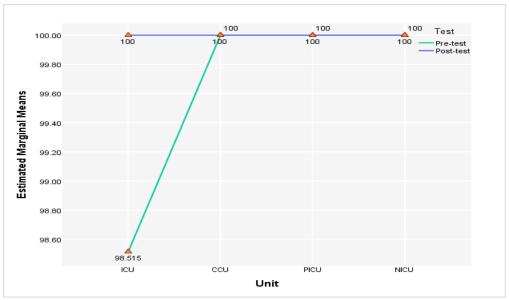


Figure7. Mean plot of the rate of high alert medication compliance rate

### 1.2.9 Patient Identification Compliance Rate

Unsurprisingly the patient identification compliance rate was also differed across four units ( $F_{3, 40} = 5.012, p = .005$ ). As most of the other rates, this rate

was also not differed in the pre-test and post-test stage ( $F_{1, 40}$ = .370, p = .775.). Which indicated that patient identification compliance rate wasn't affected by the program carried out.

Tables. Two-way ANOVA results of the patient identification compliance rate										
Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared				
Corrected Model	9.705 <sup>a</sup>	7	1.386	2.360	.041	.292				
Intercept	477624.945	1	477624.945	812844.309	.000	1.000				
Test	.217	1	.217	.370	.546	.009				
Unit	8.836	3	2.945	5.012	.005	.273				
Test * Unit	.652	3	.217	.370	.775	.027				
Error	23.504	40	.588							
Total	477658.154	48								
Corrected Total	33.209	47								

 Table9. Two-way ANOVA results of the patient identification compliance rate

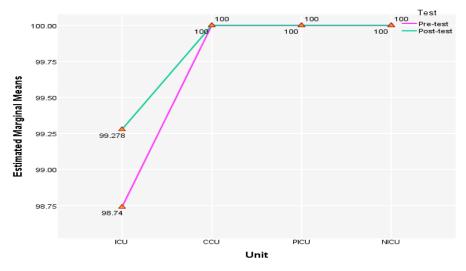


Figure8. Mean plot of patient identification compliance rate

## 1.2.10 Ventilator Associated Pneumonia (VAP) Rate

Surprisingly the interaction of test and unit significantly affected the VAP rate ( $F_{2, 30} = 3.794$ , p = .034). Yet when the unit (F<sub>2, 30</sub>= 2.271, p = .121) and the test (F<sub>2, 30</sub> = .230, p = .635) considered individually they didn't affected the VAP rate significantly.

Table 10. 1 wo-way ANO VA results of the Ventilator Associated 1 neumonia (VAI) Rate								
Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared		
Corrected Model	147.024 <sup>a</sup>	5	29.405	2.472	.055	.292		
Intercept	105.507	1	105.507	8.870	.006	.228		
Test	2.739	1	2.739	.230	.635	.008		
Unit	54.019	2	27.009	2.271	.121	.131		
Test * Unit	90.266	2	45.133	3.794	.034	.202		
Error	356.851	30	11.895					
Total	609.381	36						
Corrected Total	503.874	35						

Table10. Two-way ANOVA results of the Ventilator Associated Pneumonia (VAP) Rate

The main reason behind the insignificant pre-test and post-test VAP rate differences was a large number of sample units in the sample had a value of zero. As in the following figure at the pre-test stage, CCU and PICU had mean VAP rate of zero which means at those stages those units did not record a VAP rate value, thesame characteristic can be seen in CCU at the post-test stage as well. Hence the pre-test and post-test means were not significantly differed from each other.

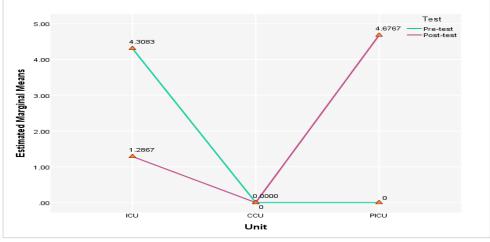


Figure9. Mean plot of VAP rate

## 1.2.11 Catheter Associated Urinary Tract Infection (CAUTI) Rate

From the sample units (36 sample units) collected regarding CAUTI rate, 86.1% (31 sample units) were equaled to

zero. Only 5 sample units had some sort of a positive value. Hence the conducted twoway ANOVA was insignificant; interaction ( $F_{2, 30} = 1.159$ , p = .328), unit ( $F_{2, 30} = .645$ , p = .428), test ( $F_{1, 30} = .645$ , p = .428).

Table11. 1 wo-way ANOVA results of the CAUTI fate									
Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared			
Corrected Model	190.132 <sup>a</sup>	5	38.026	.885	.503	.129			
Intercept	77.704	1	77.704	1.809	.189	.057			
Test	27.685	1	27.685	.645	.428	.021			
Unit	62.912	2	31.456	.732	.489	.047			
Test * Unit	99.535	2	49.768	1.159	.328	.072			
Error	1288.673	30	42.956						
Total	1556.510	36							
Corrected Total	1478.806	35							

Table11. Two-way ANOVA results of the CAUTI rate

A large number of zero values have also affected the estimated marginal means as well. Figure 11shows those effected marginal means where there mean values were lower than zero (negative).

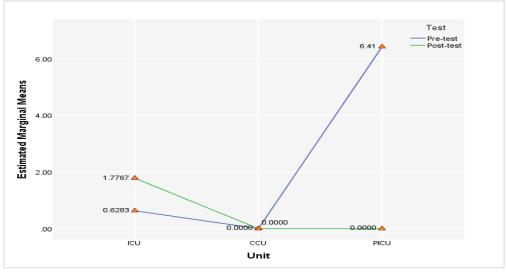


Figure10. Mean plot of CAUTI rate

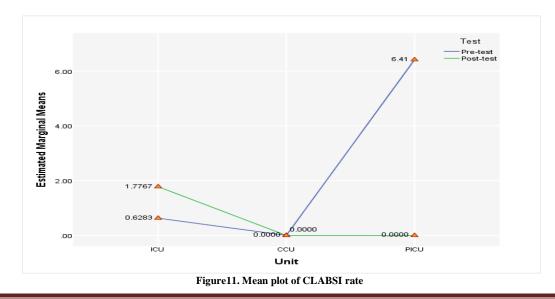
## 1.2.12 Central Line Associated Blood Stream Infection (CLABSI) Rate

CLABSI rate and unit had no significant interaction (F<sub>2, 30</sub> = 3.758, p = .035), or individual effect; unit (F<sub>2, 30</sub> = 3.043, p = .063) and test (F<sub>1, 30</sub> = .609, p =

.441). This is also mainly due to the fact that sample had a large proportion of zero values. In this case, 66.7% (24 units from 36) of the sample units were zeros. As a result, the results were insignificant in every possible way.

rable 12. 1 wo-way 11 to VII results of the CEMBSI fate							
Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	
Corrected Model	982.668 <sup>a</sup>	5	196.534	2.842	.032	.321	
Intercept	802.589	1	802.589	11.607	.002	.279	
Test	42.120	1	42.120	.609	.441	.020	
Unit	420.807	2	210.403	3.043	.063	.169	
Test * Unit	519.742	2	259.871	3.758	.035	.200	
Error	2074.417	30	69.147				
Total	3859.674	36					
Corrected Total	3057.085	35					

Table12. Two-way A	NOVA	results of th	he CLABSI	rate



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#### 1.2.13 Hospital Acquired Pressure Ulcer (HAPU) Rate

Though the interaction was insignificant ( $F_{1, 20} = .010$ , p = .923) individually two units which taken in to consideration significantly differed in HAPU rate ( $F_{1, 20} = 17.301$ , p = .000).

However pre-test and post-test results were not significantly differed from each other  $(F_{1, 20} = .218, p = .646)$ . So the HAPU rate had the characteristic which most of the other rates had where it differed across units but stayed same at the pre-test and post-test stages.

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Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared
Corrected Model	12.737 <sup>a</sup>	3	4.246	5.843	.005	.467
Intercept	16.220	1	16.220	22.322	.000	.527
Test	.158	1	.158	.218	.646	.011
Unit	12.572	1	12.572	17.301	.000	.464
Test * Unit	.007	1	.007	.010	.923	.000
Error	14.532	20	.727			
Total	43.489	24				
Corrected Total	27.269	23				

Table13. Two-way ANOVA results of the Hospital-Acquired Pressure Ulcer (HAPU) Rate

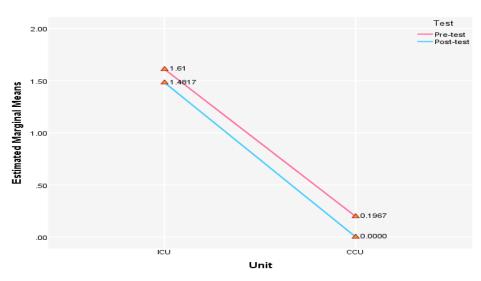


Figure12. Mean plot of HAPU rate

#### **1.3 Overall interaction**

A two-way ANOVA was conducted to identify the overall mean differences between units (ICU, CCU, etc.) and tests (pre-test and post-test).There was statistically insignificant two-way interaction between tests and units; $F_{3, 628} = .013$ , p = .998.Howeverrates were significantly differed across units,  $F_{3, 628} = 4.818$ , p = .003.

Table 14. Two-way Alvo v A results of the overall interaction							
Source	Type III Sum of Squares	df	Mean Square	F	Sig.	Partial Eta Squared	
Corrected Model	23411.066 <sup>a</sup>	7	3344.438	2.070	.045	.023	
Intercept	517400.405	1	517400.405	320.258	.000	.338	
Test	.486	1	.486	.000	.986	.000	
Unit	23349.340	3	7783.113	4.818	.003	.022	
Test * Unit	60.888	3	20.296	.013	.998	.000	
Error	1014581.148	628	1615.575				
Total	1533261.874	636					
Corrected Total	1037992.214	635					

Table14. Two-way ANOVA results of the overall interaction

#### **CONCLUSION**

In conclusion there was no statistically significant difference between

pre-test and post-test results. Although a number of rates significantly differed across units. Those rates were namely mortality

rate, DAMA rate, return to the critical care unit within 48 hours of discharge/transfer rate, occupancy rate, average length of stay, rate of initial physical assessment done by nurses with acceptable time, patient identification compliance rate and hospital acquired pressure ulcer (HAPU) rate. Hence, Null Hypothesis (Ho) is accepted and Alternative Hypothesis (H1) is rejected.

## LIMITATIONS OF THE STUDY:

This study is limited to the study hospital (Joint Commission International and HIMSS-6 accredited) and for a limited period from May 2016 to April 2017 only.

## DIRECTIONS FOR FUTURE RESEARCH:

In future such research should be conducted to study the overall impact of national accreditation on other departments of the hospital.

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The study hospital is a private, tertiary care hospital with 315 in-patient beds and has JCI, HIMSS-6 accreditations.

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