	PICC		CVC			
Variable	No. (N = 1,377)	Adherence, %	No. (N = 2,304)	Adherence, %	Statistics	P Value
Hand hygiene performed before insertion	1,374	99.8	1,880	81.6	277.945	<.001
Appropriate skin prep before insertion	1,377	100.0	2,304	100.0	NA	NA
Skin prep agent completely dried before insertion	1,368	99.4	1,836	79.7	295.318	<.001
Maximal sterile barriers (MSB) used before insertion	1,267	92.0	1,161	50.4	664.978	<.001
Sterile gloves	1,364	99.1	2,278	98.9	0.28	.597
Sterile gown	1,297	94.2	1,267	55.0	626.556	<.001
Cap	1,349	98.0	2,244	97.4	1.203	.273
Mask	1,357	98.7	2,259	98.1	1.246	.264
Large sterile drape	1,360	98.8	1,768	76.7	327.637	<.001

TABLE 1. Adherence to Central Line Insertion Practices (CLIP) Between PICC and CVC

NOTE. CLIP, central-line insertion practices; PICC, peripherally inserted central catheter; CVC, central venous catheter; NA, not available; Statistics, Pearson Chi-square test.

placed by nurses. Our results suggest better adherence with CLIP by nurses than by doctors.

It has been reported that the application of maximal sterile barriers (MSBs) used before CVC insertion could lower medical costs and decrease the incidence of catheter colonization, catheter-related bloodstream infections, and death,⁶ which reflects the importance of MSBs in the prevention of CLABSI. However, compliance with MSB protocols for CVC insertions was only 50.4% in this investigation, especially for sterile gowns (55.0%) and large sterile drapes (76.7%). This finding suggests that enough and timely access to adequate supplies and personal protective equipment (PPE) for CLABSI prevention would greatly affect compliance with CLIP, and these costs should be fully supported in departmental budgets. In stratified analyses by department, the worst compliance with MSB use in CVC insertion occurred for anesthesia and operating rooms (1.6%), radiotherapy units (40%), and neurosurgery units (47.6%), which strongly suggests the need to improve supervision, to strengthen training, and to increase feedback in the use of MSBs.

In conclusion, our data indicate that compared to PICC insertions, there was significantly less adherence with hand hygiene, complete drying of the skin disinfectant, and poor adherence with MSBs with CVC insertions. CLIP adherence should also be monitored daily to optimize patient safety.

ACKNOWLEDGMENTS

Financial support: This study was supported by grants from the 2017 Hospital Management Innovation Research Project (grant no. JSYGY-2-2017-205) and the Priority Academic Program Development of Jiangsu Higher Education Institutions (PAPD grant no. JX10231801).

Potential conflicts of interest: All authors declare that they have no competing interests.

Wensen Chen, MBBS, MPH;¹ Yiqun Yang, MPH;² Huifen Li, BSN;¹

Xiaoqiang Huang, MD;³ Weihong Zhang, MD¹

Affiliations: 1. Infection Management Office, the First Affiliated Hospital of Nanjing Medical University, Nanjing, Jiangsu, China; 2. Department of Nursing, the First Affiliated Hospital of Soochow University, Suzhou, Jiangsu, China; 3. Department of Infection Management, the Affiliated Xiaolan Hospital, Southern Medical University, Xiaolan People's Hospital, Zhongshan City, Guangdong, China.

Address correspondence to Zhang Wei-Hong, First Affiliated Hospital, Nanjing Medical University, 300 Guangzhou Rd, Nanjing, Jiangsu, China (wensenchen@njmu.edu.cn).

Infect Control Hosp Epidemiol 2018;39:122-123

© 2017 by The Society for Healthcare Epidemiology of America. All rights reserved. 0899-823X/2018/3901-0024. DOI: 10.1017/ice.2017.259

REFERENCES

- 1. Johansson E, Hammarskjöld F, Lundberg D, Arnlind MH. Advantages and disadvantages of peripherally inserted central venous catheters (PICC) compared to other central venous lines: a systematic review of the literature. *Acta Oncol* 2013;52:886–892.
- Guembe M, Pérez-Granda MJ, Capdevila JA, et al. Nationwide study on peripheral-venous-catheter-associated-bloodstream infections in internal medicine departments. J Hosp Infect 2017;97:260–266; pii:S0195-6701(17)30390-0.
- Furuya EY, Dick AW, Herzig CT, Pogorzelska-Maziarz M, Larson EL, Stone PW. Central line-associated bloodstream infection reduction and bundle compliance in intensive care units: a national study. *Infect Control Hosp Epidemiol* 2016;37:805–810.
- 4. Surveillancefor Central Line Insertion Practices (CLIP) Adherence. Centers for Disease Control and Prevention (CDC) website. https:// www.cdc.gov/nhsn/acute-care-hospital/clip/index.html. Updated 2017. Accessed March 10, 2017.
- O'Grady NP, Alexander M, Burns LA, et al. Guidelines for the prevention of intravascular catheter-related infections. *Clin Infect Dis* 2011;52:e162–e193.
- 6. Hu KK, Veenstra DL, Lipsky BA, Saint S. Use of maximal sterile barriers during central venous catheter insertion: clinical and economic outcomes. *Clin Infect Dis* 2004;39:1441–1445.

The Need for Rotavirus Vaccine Introduction in the National Immunization Program of More Than 100 Countries around the World

To the Editor—Despite the existence of the improved health care and health promotion, diarrhea continues to cause 1.7 million deaths each year in children younger than 5 years old worldwide. Roughly, rotavirus is responsible for more than a third of these deaths¹; it is a leading cause of severe diarrhea in children younger than 5 years of age around the world. According to the published report of sentinel rotavirus surveillance of the 35 member states of the World Health Organization (WHO), an average of 40% (range, 34%–45%) of diarrhea cases attributable hospitalization in children under 5 years of age were associated with rotavirus infection.² Similarly, in countries of the eastern Mediterranean region, ~40% of gastroenteritis cases in children ensue from rotavirus infection.³ In Iran, a developing country in the Middle East, the proportion of rotavirus infection in children suffering from gastroenteritis varies from 11.6% to 64.67% across the country by 2009.4

The WHO has recommended the integration of rotavirus vaccine into all national immunization schedule. However, more than 100 countries had not introduced rotavirus vaccine as of April 2016, including Afghanistan, Algeria, Andorra, Antigua and Barbuda, Azerbaijan, Bahamas, Bangladesh, Barbados, Belarus, Belize, Benin, Bhutan, Bosnia and Herzegovina, Brunei Darussalam, Bulgaria, Cabo Verde, Cambodia, Central African Republic, Chad, Chile, China, Comoros, Cook Islands, Costa Rica, Croatia, Cuba, Cyprus, Czech Republic, Democratic People's Republic of Korea, Democratic Republic of the Congo, Denmark, Dominica, Egypt, Equatorial Guinea, France, Gabon, Grenada, Guinea, Hungary, Iceland, Indonesia, Iran (Islamic Republic of), Ireland, Italy, Jamaica, Japan, Kazakhstan, Kuwait, Kyrgyzstan, Lao People's Democratic Republic, Lebanon, Lesotho, Lithuania, Malaysia, Maldives, Malta, Monaco, Mongolia, Montenegro, Myanmar, Nauru, Nepal, Netherlands, Nigeria, Niue, Oman, Pakistan, Papua New Guinea, Poland, Portugal, Republic of Korea, Romania, Russian Federation, Saint Kitts and Nevis, Saint Lucia, Saint Vincent and the Grenadines, Samoa, San Marino, Serbia, Singapore, Slovakia, Slovenia, Solomon Islands, Somalia, South Sudan, Spain, Sri Lanka, Suriname, Switzerland, Syrian Arab Republic, The former Yugoslav Republic of Macedonia, Timor-Leste, Tonga, Trinidad and Tobago, Tunisia, Turkey, Turkmenistan, Tuvalu, Ukraine, United Arab Emirates, Uruguay, Vanuatu and Viet Nam.⁵ Moreover, the results of a meta-analysis study in Iran have provided sufficient evidence to introduce rotavirus vaccine in the routine national immunization program. According to the findings of this meta-analysis, the pooled estimates of rotavirus infection in gastroenteritis cases and gastroenteritis-related hospitalizations were 35% (95% CI, 28%-41%) and 39% (95% CI, 30%-48%), respectively⁶.

Rotarix and RotaTq are 2 types of new oral vaccines available against rotavirus infection. The monovalent rotavirus vaccine (RV1) is implemented in 2 scheduled doses (at 2 and 4 months of age), and the pentavalent rotavirus vaccine (RV5) is implemented in 3 oral doses (at 2, 4, and 6 months of age). Both vaccines listed can be integrated in a national immunization program.⁵ The pooled estimates of rotavirus vaccine efficacy derived from the result of clinical trials suggested that these 2 vaccines can prevent 70% and 83% of all disease cases, respectively. Moreover, these 2 vaccines were 80% and 90% protective against rotavirus gastroenteritis, respectively.⁷ A review study in collaboration with the WHO using 89 observational studies and clinical trials approved the safety and efficacy of rotavirus vaccine. Published studies indicate higher efficacy of vaccine in developed countries than in developing countries. Vaccine efficacy in the prevention of severe cases in America, Europe, and Latin America was estimated to be 89.1%.¹ In Asian and African countries, the pooled estimate of Rotarix vaccine efficacy against severe cases of disease during the first year was 58% (95% CI, 40%-72.3%).8 The rate in developing countries may be improved if relevant preventive measures are considered along with immunization against rotavirus.

In conclusion, rotavirus vaccine should be introduced to the immunization program as a part of a prevention program for diarrheal diseases to maximize its impact. Policy makers should consider additional control measures such as exclusive breastfeeding up to 6 months, healthy water supply, personal hygiene, and sanitation along with the treatment of rotavirus-related illnesses and gastroenteritis cases to achieve an efficient immunization program against rotavirus.⁵

ACKNOWLEDGMENTS

Financial support: No financial support was provided relevant to this article. *Potential conflicts of interest:* All authors report no conflicts of interest relevant to this article.

Manoochehr Karami, PhD;^{1,2} Zeinab Berangi, MSc²

Affiliations: 1. Social Determinants of Health Research Center, Hamadan University of Medical Sciences, Hamadan, Iran; 2. Department of Epidemiology, School of Public Health, Hamadan University of Medical Sciences, Hamadan, Iran.

Address correspondence to Dr Manoochehr Karami, Fahmide St, Department of Epidemiology, School of Public Health, Hamadan University of Medical Sciences, Hamadan, Iran (ma.karami@umsha.ac.ir).

Infect Control Hosp Epidemiol 2018;39:124–125

© 2017 by The Society for Healthcare Epidemiology of America. All rights reserved. 0899-823X/2018/3901-0025. DOI: 10.1017/ice.2017.237

REFERENCES

1. Munos MK, Walker CLF, Black RE. The effect of rotavirus vaccine on diarrhoea mortality. *Int J Epidemiol* 2010;39: i56–i62.

- 2. World Health Organization. Rotavirus vaccines, WHO position paper. *Weekly Epidemiol Rec* 2013;88:49–64.
- 3. Malek MA, Teleb N, Abu-Elyazeed R, et al. The epidemiology of rotavirus diarrhea in countries in the eastern Mediterranean region. *J Infect Dis* 2010;202:S12–S22.
- Zaraei-Mahmoodabadi B, Kargar M, Tabatabaei H, Saedegipour S, Ghaemi A, Nategh R. Determination of annual incidence, age specific incidence rate and risk of rotavirus gastroenteritis among children in Iran. *Iran J Virol* 2009;3:33–36.
- World Health Organization. Rotavirus vaccines WHO position paper: January 2013–recommendations. Vaccine 2013;31:6170–6171.
- 6. Moradi-Lakeh M, Shakerian S, Yaghoubi M, et al. Rotavirus infection in children with acute gastroenteritis in Iran, a systematic review and meta-analysis. *Int J Prevent Med* 2014:5.
- 7. Rostami S, Lakeh M, Esteghamati A, et al. The efficacy and safety of rotavirus vaccine in children under the five years of age; systematic review and meta-analysis. *J Isfahan Med Sch* 2014;32: 1605–1622.
- 8. Breiman RF, Zaman K, Armah G, et al. Analyses of health outcomes from the 5 sites participating in the Africa and Asia clinical efficacy trials of the oral pentavalent rotavirus vaccine. *Vaccine* 2012;30:A24–A29.