

Department of Veterans Affairs Office of Inspector General

Office of Healthcare Inspections

Report No. 15-00619-515

Combined Assessment Program Review of the Robley Rex VA Medical Center Louisville, Kentucky

September 17, 2015

Washington, DC 20420

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Glossary

AD	advance directive
CAP	Combined Assessment Program
СТ	computed tomography
EAM	emergency airway management
EHR	electronic health record
EOC	environment of care
facility	Robley Rex VA Medical Center
FY	fiscal year
LNET	lung nodule evaluation team
MH	mental health
NA	not applicable
NM	not met
OIG	Office of Inspector General
QM	quality management
SCI	spinal cord injury
VHA	Veterans Health Administration
VISN	Veterans Integrated Service Network

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Executive Summary

Review Purpose: The purpose of the review was to evaluate selected health care facility operations, focusing on patient care quality and the environment of care, and to provide crime awareness briefings. We conducted the review the week of August 3, 2015.

Review Results: The review covered eight activities. We made no recommendations in the following four activities:

- Medication Management
- Coordination of Care
- Computed Tomography Radiation Monitoring
- Surgical Complexity

The facility's reported accomplishments were the lung nodule evaluation team and the Women's Health Program.

Recommendations: We made recommendations in the following four activities:

Quality Management: Require that licensed practitioners who perform emergency airway management have the appropriate training. Ensure the Surgical Work Group meets monthly.

Environment of Care: Conduct fire drills in all health care occupancy buildings with the frequency required. Ensure negative pressure systems in the medicine primary care clinic are functional. Require locked mental health unit stationary panic alarm testing to include documentation of VA Police response time. Secure equipment on the locked mental health unit.

Advance Directives: Offer inpatients the opportunity to discuss advance directives, hold the requested discussions, and document the discussions.

Emergency Airway Management: Ensure that initial clinician emergency airway management competency assessment includes all required elements. Revise the local out of operating room emergency airway management policy to include requirements for successful demonstration of procedural skills. Document the review of provider-specific emergency airway management data in Cardiopulmonary Review Committee meeting minutes.

Comments

The Veterans Integrated Service Network and Facility Directors agreed with the Combined Assessment Program review findings and recommendations and provided acceptable improvement plans. (See Appendixes C and D, pages 26–31, for the full text of the Directors' comments.) We consider recommendation 6 closed. We will follow up on the planned actions for the open recommendations until they are completed.

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JOHN D. DAIGH, JR., M.D. Assistant Inspector General for Healthcare Inspections

Objectives and Scope

Objectives

CAP reviews are one element of the OIG's efforts to ensure that our Nation's veterans receive high quality VA health care services. The objectives of the CAP review are to:

- Conduct recurring evaluations of selected health care facility operations, focusing on patient care quality and the EOC.
- Provide crime awareness briefings to increase employee understanding of the potential for program fraud and the requirement to refer suspected criminal activity to the OIG.

Scope

The scope of the CAP review is limited. Serious issues that come to our attention that are outside the scope will be considered for further review separate from the CAP process and may be referred accordingly.

For this review, we examined selected clinical and administrative activities to determine whether facility performance met requirements related to patient care quality and the EOC. In performing the review, we inspected selected areas, conversed with managers and employees, and reviewed clinical and administrative records. The review covered the following eight activities:

- QM
- EOC
- Medication Management
- Coordination of Care
- CT Radiation Monitoring
- ADs
- Surgical Complexity
- EAM

We have listed the general information reviewed for each of these activities. Some of the items listed may not have been applicable to this facility because of a difference in size, function, or frequency of occurrence.

The review covered facility operations for FY 2014 and FY 2015 through August 3, 2015, and inspectors conducted the review in accordance with OIG standard operating procedures for CAP reviews. We also asked the facility to provide the status on the recommendations we made in our previous CAP report (*Combined Assessment Program Review of the Robley Rex VA Medical Center, Louisville, Kentucky,* Report No. 13-00433-199, May 20, 2013).

During this review, we presented crime awareness briefings for 255 employees. These briefings covered procedures for reporting suspected criminal activity to the OIG and included case-specific examples illustrating procurement fraud, conflicts of interest, and bribery.

Additionally, we surveyed employees regarding patient safety and quality of care at the facility. We distributed an electronic survey to all facility employees and received 311 responses. We shared summarized results with facility managers.

In this report, we make recommendations for improvement. Recommendations pertain to issues that are significant enough for the OIG to monitor until the facility implements corrective actions.

Reported Accomplishments

The LNET

The LNET is a multidisciplinary group of health care providers who evaluate and manage patients with lung nodules (a small round or oval-shaped growth in the lung). Once a nodule is identified, an electronic consult notifies the LNET. The LNET then reviews the consults and determines the appropriate management disposition—surveillance, diagnostics, or therapy. On average, 189 LNET consults are entered each month. This has decreased the time from initial imaging to a definitive management plan by a specialist from 49.1 days to only 3.7 days.

Women's Health Program

The facility's Women's Health Program has expanded and flourished in the last year. The facility offered more than 3,000 female veterans reassignment to designated women's health providers, and to accommodate the increased demand, it added two designated women's health providers and a community provider for mammograms. In addition, the facility offers comprehensive evening women's primary care and gynecology services. A Women's Health Nurse Navigator coordinates maternity patients, mammography, Pap tracking, and non-VA referrals. In February, the facility held a Go Red Event to provide education on women's heart health and established a women-only yoga class. On April 11, 2015, in collaboration with the State Department of Veterans Affairs, the facility held the first ever state conference for women veterans in addition to the annual women's fair held for the last 4 years.

Results and Recommendations

QM

The purpose of this review was to determine whether facility senior managers actively supported and appropriately responded to QM efforts and whether the facility met selected requirements within its QM program.^a

We conversed with senior managers and key QM employees, and we evaluated meeting minutes, 22 credentialing and privileging folders, and other relevant documents. The table below shows the areas reviewed for this topic. The areas marked as NM did not meet applicable requirements and needed improvement. Any items that did not apply to this facility are marked NA.

NM	Areas Reviewed	Findings	Recommendations
	 There was a senior-level committee responsible for key quality, safety, and value functions that met at least quarterly and was chaired or co-chaired by the Facility Director. The committee routinely reviewed aggregated data. QM, patient safety, and systems redesign appeared to be integrated. 		
	 Peer reviewed deaths met selected requirements: Peers completed reviews within specified timeframes. The Peer Review Committee reviewed cases receiving initial Level 2 or 3 ratings. Involved providers were invited to provide input prior to the final Peer Review Committee determination. 		

NM	Areas Reviewed (continued)	Findings	Recommendations
X	 Credentialing and privileging processes met selected requirements: Facility managers reviewed privilege forms annually and ensured proper approval of revised forms. Facility managers ensured appropriate privileges for licensed independent practitioners. Facility managers removed licensed independent practitioners' access to patients' EHRs upon separation. Facility managers properly maintained licensed independent practitioners' folders. 	 None of the 22 licensed practitioners whose folders we reviewed had EAM privileges that were appropriate for their training. 	1. We recommended that facility managers ensure that licensed practitioners who perform emergency airway management have the appropriate training.
	 Observation bed use met selected requirements: The facility gathered data regarding appropriateness of observation bed usage. The facility reassessed observation criteria and/or utilization if conversions to acute admissions were consistently 25–30 percent or more. 		
	 The process to review resuscitation events met selected requirements: An interdisciplinary committee reviewed episodes of care where resuscitation was attempted. Resuscitation event reviews included screening for clinical issues prior to events that may have contributed to the occurrence of the code. The facility collected data that measured performance in responding to events. 		

NM	Areas Reviewed (continued)	Findings	Recommendations
Х	The surgical review process met selected	The Surgical Work Group only met	2. We recommended that the Surgical Work
	requirements:	10 times over the past 12 months.	Group meet monthly.
	An interdisciplinary committee with		
	appropriate leadership and clinical membership met monthly to review		
	surgical processes and outcomes.		
	• •		
	 The Surgical Work Group reviewed surgical deaths with identified problems or 		
	opportunities for improvement.		
	 The Surgical Work Group reviewed 		
	 The Surgical Work Group reviewed additional data elements. 		
	Clinicians appropriately reported critical		
	incidents.		
	The safe patient handling program met		
	selected requirements:		
	 A committee provided program oversight. 		
	 The committee gathered, tracked, and 		
	shared patient handling injury data.		
	The process to review the quality of entries		
	in the EHR met selected requirements:		
	A committee reviewed EHR quality.		
	A committee analyzed data at least		
	quarterly.		
	Reviews included data from most services		
	and program areas.		
	The policy for scanning internal forms into		
	EHRs included the following required items:		
	Quality of the source document and an		
	alternative means of capturing data when		
	the quality of the document is inadequate.		
	 A correction process if scanned items 		
	have errors.		

NM	Areas Reviewed (continued)	Findings	Recommendations
	 A complete review of scanned documents 		
	to ensure readability and retrievability of		
	the record and quality assurance reviews		
	on a sample of the scanned documents.		
	Overall, if QM reviews identified significant		
	issues, the facility took actions and		
	evaluated them for effectiveness.		
	Overall, senior managers actively		
	participated in performance improvement		
	over the past 12 months.		
	Overall, the facility had a comprehensive,		
	effective QM program over the past		
	12 months.		
	The facility met any additional elements		
	required by VHA or local policy.		

EOC

The purpose of this review was to determine whether the facility maintained a clean and safe health care environment in accordance with applicable requirements. We also determined whether the facility met selected requirements in emergency management.^b

We inspected the 6N medical intensive care, 5N hospice/chemotherapy, 6S medical surgical, and locked MH units; the Emergency Department; and the medicine primary care, 3S hematology/oncology, and wound care clinics. Additionally, we reviewed relevant documents, including 10 employee training records, and conversed with key employees and managers. The table below shows the areas reviewed for this topic. The areas marked as NM did not meet applicable requirements and needed improvement. Any items that did not apply to this facility are marked NA.

NM	Areas Reviewed for General EOC	Findings	Recommendations
	EOC Committee minutes reflected sufficient		
	detail regarding identified deficiencies,		
	corrective actions taken, and tracking of		
	corrective actions to closure for the facility		
	and the community based outpatient clinics.		
	The facility conducted an infection		
	prevention risk assessment.		
	Infection Prevention/Control Committee		
	minutes documented discussion of identified		
	high-risk areas, actions implemented to		
	address those areas, and follow-up on		
	implemented actions and included analysis		
	of surveillance activities and data.		
	The facility had established a process for		
	cleaning equipment.		
Х	The facility conducted required fire drills in	Past 2 quarters of fire drill documentation for	3. We recommended that facility managers
	buildings designated for health care	health care occupancy buildings reviewed:	ensure all health care occupancy buildings
	occupancy and documented drill critiques.	All applicable buildings did not have at	have at least one fire drill per shift per
		least one fire drill per shift per quarter.	quarter and monitor compliance.
	The facility had a policy/procedure/guideline		
1	for identification of individuals entering the		
1	facility, and units/areas complied with		
	requirements.		

NM	Areas Reviewed for General EOC (continued)	Findings	Recommendations
	The facility met fire safety requirements.		
	The facility met environmental safety		
X	requirements. The facility met infection prevention requirements.	 Neither of the two negative air pressure systems in the medicine primary care clinic airborne infection isolation rooms 	4. We recommended that facility managers ensure negative air pressure systems in the medicine primary care clinic are functional
	The facility met medication safety and security requirements.	was functional.	and monitor compliance.
	The facility met privacy requirements.		
X	The facility complied with any additional elements required by VHA, local policy, or other regulatory standards.	 VA National Center for Patient Safety MH EOC Checklist reviewed, which requires testing of panic alarms, including VA Police response time, on a periodic basis at a frequency determined by the facility. It also requires that furniture and equipment be secured or heavy enough to prevent it from being picked up, thrown, moved, or overturned. Although employees conducted stationary panic alarm testing on the locked MH unit, they did not document VA Police response time January–March 2015. A television bolted to a rolling cart on the locked MH unit was not secured or heavy enough to prevent it from being picked up, thrown, moved, or overturned. 	 5. We recommended that facility managers ensure locked mental health unit stationary panic alarm testing includes documentation of VA Police response time. 6. We recommended that equipment on the locked mental health unit is secured and heavy enough to prevent it from being picked up, thrown, moved, or overturned.
	Areas Reviewed for SCI Center		
NA	The facility completed and documented required inspection checklists of all ceiling mounted patient lifts.		

NM	Areas Reviewed for SCI Center (continued)	Findings	Recommendations
NA	The facility met fire safety requirements in the SCI Center.		
NA	The facility met environmental safety requirements in the SCI Center.		
NA	The facility met infection prevention requirements in the SCI Center.		
NA	The facility met medication safety and security requirements in the SCI Center.		
NA	The facility met patient privacy requirements in the SCI Center.		
NA	The facility complied with any additional elements required by VHA, local policy, or other regulatory standards.		
	Areas Reviewed for Emergency Management		
	The facility had a documented Hazard Vulnerability Assessment and reviewed the assessment annually.		
	The facility maintained a list of resources and assets it may need during an emergency.		
	The facility had a written Emergency Operations Plan that addressed key components.		
	The facility had a written description of how it will respond to an influx of potentially infectious patients and a plan for managing them over an extended period of time.		
	Employees received training and competency assessment on use of emergency evacuation devices.		
	Evacuation devices were immediately accessible and in good repair.		

NM	Areas Reviewed for Emergency Management (continued)	Findings	Recommendations
	The facility complied with any additional		
	elements required by VHA, local policy, or		
	other regulatory standards.		
	Areas Reviewed for Construction Safety		
NA	The facility met selected dust control,		
	temporary barrier, storage, and security		
	requirements for the construction site		
	perimeter.		
NA	The facility complied with any additional		
	elements required by VHA or local policy, or		
	other regulatory standards.		

Medication Management

The purpose of this review was to determine whether the facility had established safe medication storage practices in accordance with VHA policy and Joint Commission standards.^c

We reviewed relevant documents, the training records of 20 nursing employees, and pharmacy monthly medication storage area inspection documentation for the past 6 months. We inspected the Emergency Department and the surgical intensive care, post-anesthesia care, and 4N medical/surgical/telemetry units. Additionally, for these areas, we reviewed documentation of narcotic wastage from automated dispensing machines and inspected crash carts containing emergency medications. The table below shows the areas reviewed for this topic. Any items that did not apply to this facility are marked NA. The facility generally met requirements. We made no recommendations.

NM	Areas Reviewed	Findings	Recommendations
	Facility policy addressed medication receipt		
	in patient care areas, storage procedures		
	until administration, and staff authorized to		
	have access to medications and areas used		
	to store them.		
	The facility required two signatures on		
	controlled substances partial dose wasting.		
	The facility defined those medications and		
	supplies needed for emergencies and		
	procedures for crash cart checks, checks		
	included all required elements, and the		
	facility conducted checks with the frequency		
	required by local policy.		
	The facility prohibited storage of potassium		
	chloride vials in patient care areas.		
NA	If the facility stocked heparin in		
	concentrations of more than 5,000 units per		
	milliliter in patient care areas, the Chief of		
	Pharmacy approved it.		

NM	Areas Reviewed (continued)	Findings	Recommendations
	The facility maintained a list of the look-alike		
	and sound-alike medications it stores,		
	dispenses, and administers; reviewed this		
	list annually and ensured it was available for		
	staff reference; and had labeling/storage		
	processes to prevent errors.		
	The facility identified in writing its high-alert		
	and hazardous medications, ensured the		
	high-alert list was available for staff		
	reference, and had processes to manage		
	these medications.		
	The facility conducted and documented		
	inspections of all medication storage areas		
	at least monthly, fully implemented corrective		
	actions, and monitored the changes.		
	The facility/Pharmacy Service had a written		
	policy for safe use of automated dispensing		
	machines that included oversight of		
	overrides and employee training and		
	minimum competency requirements for		
	users, and employees received training or		
	competency assessment in accordance with		
	local policy.		
	The facility employed practices to prevent		
	wrong-route drug errors.		
	Medications prepared but not immediately		
	administered contained labels with all		
	required elements.		
	The facility removed medications awaiting		
	destruction or stored them separately from		
	medications available for administration.		

NM	Areas Reviewed (continued)	Findings	Recommendations
	The facility met multi-dose insulin pen		
	requirements.		
	The facility complied with any additional		
	elements required by VHA or local policy.		

Coordination of Care

The purpose of this review was to evaluate the consult management process and the completion of inpatient clinical consults.^d

We reviewed relevant documents, and we conversed with key employees. Additionally, we reviewed the EHRs of 39 randomly selected patients who had a consult requested during an acute care admission from January 1 through June 30, 2014. The table below shows the areas reviewed for this topic. Any items that did not apply to this facility are marked NA. The facility generally met requirements. We made no recommendations.

NM	Areas Reviewed	Findings	Recommendations
	A committee oversaw the facility's consult		
	management processes.		
	Major bed services had designated		
	employees to:		
	 Provide training in the use of the 		
	computerized consult package		
	 Review and manage consults 		
	Consult requests met selected requirements:		
	 Requestors included the reason for the consult. 		
	 Requestors selected the proper consult title. 		
	 Consultants appropriately changed consult statuses, linked responses to the requests, and completed consults within the specified timeframe. 		
	The facility met any additional elements required by VHA or local policy.		

CT Radiation Monitoring

The purpose of this review was to determine whether the facility complied with selected VHA radiation safety requirements and to follow up on recommendations regarding monitoring and documenting radiation dose from a 2011 report, *Healthcare Inspection – Radiation Safety in Veterans Health Administration Facilities*, Report No. 10-02178-120, March 10, 2011.^e

We reviewed relevant documents, including qualifications and dosimetry monitoring for 14 CT technologists and CT scanner inspection reports, and conversed with key managers and employees. We also reviewed the EHRs of 50 randomly selected patients who had a CT scan January 1–December 31, 2014. The table below shows the areas reviewed for this topic. Any items that did not apply to this facility are marked NA. The facility generally met requirements. We made no recommendations.

NM	Areas Reviewed	Findings	Recommendations
	The facility had a designated Radiation		
	Safety Officer responsible for oversight of		
	the radiation safety program.		
	The facility had a CT/imaging/radiation		
	safety policy or procedure that included:		
	A CT quality control program with program		
	monitoring by a medical physicist at least		
	annually, image quality monitoring, and CT		
	scanner maintenance		
	 CT protocol monitoring to ensure doses 		
	were as low as reasonably achievable and		
	a method for identifying and reporting		
	excessive CT patient doses to the		
	Radiation Safety Officer		
	A process for managing/reviewing CT		
	protocols and procedures to follow when		
	revising protocols		
	 Radiologist review of appropriateness of 		
	CT orders and specification of protocol		
	prior to scans		

NM	Areas Reviewed (continued)	Findings	Recommendations
	A radiologist and technologist expert in CT		
	reviewed all CT protocols revised during the		
	past 12 months.		
	A medical physicist tested a sample of CT		
	protocols at least annually.		
	A medical physicist performed and		
	documented CT scanner annual inspections,		
	an initial inspection after acquisition, and		
	follow-up inspections after repairs or		
	modifications affecting dose or image quality		
	prior to the scanner's return to clinical		
	service.		
	If required by local policy, radiologists		
	included patient radiation dose in the CT		
	report available for clinician review and		
	documented the dose in the required		
	application(s), and any summary reports		
	provided by teleradiology included dose		
	information.		
	CT technologists had required certifications		
	or written affirmation of competency if		
	"grandfathered in" prior to January 1987, and		
	technologists hired after July 1, 2014, had		
	CT certification. There was documented evidence that CT		
	technologists had annual radiation safety training and dosimetry monitoring.		
	If required by local policy, CT technologists		
	had documented training on dose		
	reduction/optimization techniques and safe		
	procedures for operating the types of CT		
	equipment they used.		
	The facility complied with any additional		
	elements required by VHA or local policy.		
	elements required by VIA or local policy.		

ADs

The purpose of this review was to determine whether the facility complied with selected requirements for ADs for patients.^f

We reviewed relevant documents and conversed with key employees. Additionally, we reviewed the EHRs of 48 randomly selected patients who had an acute care admission January 1–December 31, 2014. The table below shows the areas reviewed for this topic. The area marked as NM did not meet applicable requirements and needed improvement. Any items that did not apply to this facility are marked NA.

NM	Areas Reviewed	Findings	Recommendations
	 The facility had an AD policy that addressed: AD notification, screening, and discussions Proper use of AD note titles 		
	Employees screened inpatients to determine whether they had ADs and used appropriate note titles to document screening.		
	 When patients provided copies of their current ADs, employees had scanned them into the EHR. Employees correctly posted patients' AD status. 		
X	 Employees asked inpatients if they would like to discuss creating, changing, and/or revoking ADs. When inpatients requested a discussion, employees documented the discussion 	• Five of the 48 EHRs (10 percent), did not contain documentation that employees asked patients whether they wished to discuss creating, changing, and/or revoking ADs.	7. We recommended that employees ask inpatients whether they would like to discuss creating, changing, and/or revoking advance directives and that facility managers monitor compliance.
	and used the required AD note titles.	 Four of the six applicable EHRs did not contain documentation that employees held the discussions requested. . 	8. We recommended that employees hold advance directive discussions requested by inpatients and document the discussions and that facility managers monitor compliance.
	The facility met any additional elements required by VHA or local policy.		

Surgical Complexity

The purpose of this review was to determine whether the facility provided selected support services appropriate to the assigned surgical complexity designation.⁹

We reviewed relevant documents and the training records of 20 employees, and we conversed with key managers and employees. The table below shows the areas reviewed for this topic. Any items that did not apply to this facility are marked NA. The facility generally met requirements. We made no recommendations.

NM	Areas Reviewed	Findings	Recommendations
	Facility policy defined appropriate availability		
	for all support services required by VHA for		
	the facility's surgical designation.		
	Employees providing selected tests and		
	patient care after operational hours had		
	appropriate competency assessments and		
	validation.		
NA	The facility properly reported surgical		
	procedures performed that were beyond the		
	facility's surgical complexity designation.		
	 The facility reviewed and implemented 		
	recommendations made by the VISN Chief		
	Surgical Consultant.		
	The facility complied with any additional		
	elements required by VHA or local policy.		

EAM

The purpose of this review was to determine whether the facility complied with selected VHA out of operating room airway management requirements.^h

We reviewed relevant documents, including competency assessment documentation of 22 clinicians applicable for the review period January 1–June 30, 2014, and we conversed with key managers and employees. The table below shows the areas reviewed for this topic. The areas marked as NM did not meet applicable requirements and needed improvement. Any items that did not apply to this facility are marked NA.

NM	Areas Reviewed	Findings	Recommendations
	The facility had a local EAM policy or had a		
	documented exemption.		
NA	If the facility had an exemption, it did not		
	have employees privileged to perform		
	procedures using moderate or deep sedation		
	that might lead to airway compromise.		
	Facility policy designated a clinical subject		
	matter expert, such as the Chief of Staff or		
	Chief of Anesthesia, to oversee EAM.		
	Facility policy addressed key VHA		
	requirements, including:		
	 Competency assessment and 		
	reassessment processes		
	Use of equipment to confirm proper		
	placement of breathing tubes		
	A plan for managing a difficult airway		
Х	Initial competency assessment for EAM	 None of the 22 clinicians had 	9. We recommended that the facility ensure
	included:	documentation of all required subject	initial clinician emergency airway
	 Subject matter content elements and 	matter content elements.	management competency assessment
	completion of a written test		includes all required subject matter content
	Successful demonstration of procedural		elements and that facility managers monitor
	skills on airway simulators or mannequins		compliance.
	Successful demonstration of procedural		
	skills on patients		

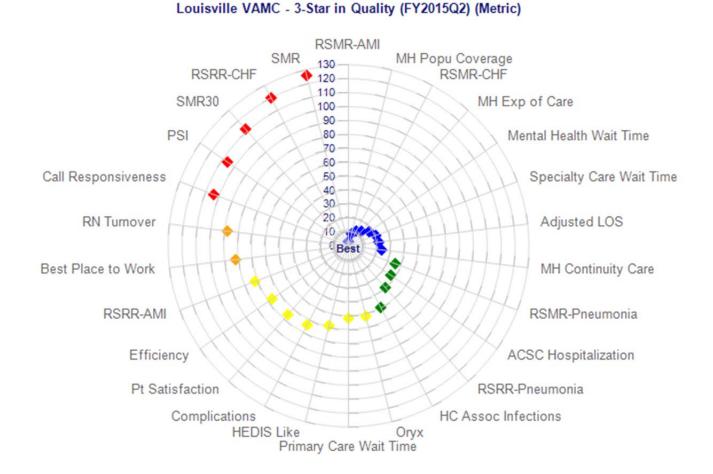
NM	Areas Reviewed (continued)	Findings	Recommendations
NA	 Reassessments for continued EAM competency were completed at the time of renewal of privileges or scope of practice and included: Review of clinician-specific EAM data Subject matter content elements and completion of a written test Successful demonstration of procedural skills on airway simulators or mannequins At least one occurrence of successful airway management and intubation in the preceding 2 years, written certification of competency by the supervisor, or successful demonstration of skills to the subject matter expert A statement related to EAM if the clinician was not a licensed independent practitioner 		
	The facility had a clinician with EAM privileges or scope of practice or an anesthesiology staff member available during all hours the facility provided patient care. Video equipment to confirm proper placement of breathing tubes was available for immediate clinician use.		

NM	Areas Reviewed (continued)	Findings	Recommendations
X	The facility complied with any additional elements required by VHA or local policy.	 Facility policy on out of operating room EAM reviewed: The facility did not require simulator training for reassessments. Facility oversight committee meeting minutes did not reflect a review of provider-specific EAM data as required by local policy. 	 10. We recommended that the facility revise the local policy for out of operating room emergency airway management to include successful demonstration of all required procedural skills on airway simulators for providers seeking renewal of privileges. 11. We recommended that the facility document the review of provider-specific emergency airway management data in Cardiopulmonary Review Committee meeting minutes.

Facility Profile (Louisville/603) FY 2015 through	ugh July 2015 ¹
Type of Organization	Secondary
Complexity Level	1c-High complexity
Affiliated/Non-Affiliated	Affiliated
Total Medical Care Budget in Millions	\$309
Number of:	
Unique Patients	41,891
Outpatient Visits	524,823
Unique Employees ²	1,694
Type and Number of Operating Beds:	
Hospital	107
Community Living Center	NA
• MH	16
Average Daily Census:	
Hospital	65
Community Living Center	NA
• MH	13
Number of Community Based Outpatient Clinics	8
Location(s)/Station Number(s)	Ft. Knox/603GA
	New Albany/603GB
	Louisville/603GC
	Louisville/603GD
	Louisville/603GE
	Clarkson/603GF
	Scottsburg/603GG
	Carrollton/603GH
VISN Number	9

 ¹ All data is for FY 2015 through July 2015 except where noted.
 ² Unique employees involved in direct medical care (cost center 8200).

Appendix B



Strategic Analytics for Improvement and Learning (SAIL)³

Marker color: Blue - 1st quintile; Green - 2nd; Yellow - 3rd; Orange - 4th; Red - 5th quintile.

³ Metric definitions follow the graphs.

Scatter Chart

CHE FADING 1st SCAcces AdjLOS . PatSat 2nd Complic RN-Turn Quality . NEU-MR MHAcces PCAcces FY2014Q2 Quintile 3rd AMI-RR InpQual Eff-SFA . Infect PNEU-RR HosACSC SMR30 HEDIS -. • CallRes BPWk • SMR • CHF-RR PSI . RISK 4th 3rd 2nd 1st FY2015Q2 Quintile

FY2015Q2 Change in Quintiles from FY2014Q2

NOTE

DESIRED DIRECTION =>

Quintiles are derived from facility ranking on z-score of a metric among 128 facilities. Lower quintile is more favorable.

DESIRED DIRECTION =>

Metric Definitions

Measure	Definition	Desired direction
ACSC Hospitalization	Ambulatory care sensitive condition hospitalizations (observed to expected ratio)	A lower value is better than a higher value
Adjusted LOS	Acute care risk adjusted length of stay	A lower value is better than a higher value
Best Place to Work	Overall satisfaction with job	A higher value is better than a lower value
Call Center Responsiveness	Average speed of call center responded to calls in seconds	A lower value is better than a higher value
Call Responsiveness	Call center speed in picking up calls and telephone abandonment rate	A lower value is better than a higher value
Complications	Acute care risk adjusted complication ratio	A lower value is better than a higher value
Efficiency	Overall efficiency measured as 1 divided by SFA (Stochastic Frontier Analysis)	A higher value is better than a lower value
Employee Satisfaction	Overall satisfaction with job	A higher value is better than a lower value
HC Assoc Infections	Health care associated infections	A lower value is better than a higher value
HEDIS	Outpatient performance measure (HEDIS)	A higher value is better than a lower value
MH Wait Time	MH wait time for new and established patients (top 50 clinics; FY13 and later)	A higher value is better than a lower value
MH Continuity Care	MH continuity of care (FY14Q3 and later)	MH Continuity Care
MH Exp of Care	MH experience of care (FY14Q3 and later)	A higher value is better than a lower value
MH Popu Coverage	MH population coverage (FY14Q3 and later)	A higher value is better than a lower value
Oryx	Inpatient performance measure (ORYX)	A higher value is better than a lower value
Primary Care Wait Time	Primary care wait time for new and established patients (top 50 clinics; FY13 and later)	A higher value is better than a lower value
PSI	Patient safety indicator (observed to expected ratio)	A lower value is better than a higher value
Pt Satisfaction	Overall rating of hospital stay (inpatient only)	A higher value is better than a lower value
RN Turnover	Registered nurse turnover rate	A lower value is better than a higher value
RSMR-AMI	30-day risk standardized mortality rate for acute myocardial infarction	A lower value is better than a higher value
RSMR-CHF	30-day risk standardized mortality rate for congestive heart failure	A lower value is better than a higher value
RSMR-Pneumonia	30-day risk standardized mortality rate for pneumonia	A lower value is better than a higher value
RSRR-AMI	30-day risk standardized readmission rate for acute myocardial infarction	A lower value is better than a higher value
RSRR-CHF	30-day risk standardized readmission rate for congestive heart failure	A lower value is better than a higher value
RSRR-Pneumonia	30-day risk standardized readmission rate for pneumonia	A lower value is better than a higher value
SMR	Acute care in-hospital standardized mortality ratio	A lower value is better than a higher value
SMR30	Acute care 30-day standardized mortality ratio	A lower value is better than a higher value
Specialty Care Wait Time	Specialty care wait time for new and established patients (top 50 clinics; FY13 and later)	A higher value is better than a lower value

VISN Director Comments

Department of Veterans Affairs

Memorandum

Date: August 28, 2015

From: Director, VA Mid South Healthcare Network (10N9)

Subject: CAP Review of the Robley Rex VA Medical Center, Louisville, KY

To: Director, Bay Pines Office of Healthcare Inspections (54SP)

Director, Management Review Service (VHA 10AR MRS OIG CAP CBOC)

- 1. Attached, please find the comments and corrective action plan for the Combined Assessment Review (CAP) of the Robley Rex VA Medical Center, Louisville, Kentucky.
- 2. I have reviewed and concur with the responses and action plan submitted by the medical center.
- 3. If you have any questions or require additional information, please contact Ms. Cynthia L. Johnson, VISN 9 Quality Management Officer at 615-695-2143.

(original signed by:) John E. Patrick Network Director

Facility Director Comments

Department of Veterans Affairs

Memorandum

Date: August 28, 2015

From: Director, Robley Rex VA Medical Center (603/00)

Subject: CAP Review of the Robley Rex VA Medical Center, Louisville, KY

- To: Director, VA Mid South Healthcare Network (10N9)
- 1. I want to express my appreciation to the Office of the Inspector General (OIG) survey team for their comprehensive review of the Robley Rex VAMC. We appreciated the professional and consultative nature of the review.
- 2. I have reviewed the report for the Robley Rex VAMC and I concur with the findings and recommendations.
- 3. Should you have any questions, please do not hesitate to contact Randy Johnson, Chief, Quality Management at 502-287-5331.

(original signed by:) Martin Traxler Medical Center Director

Comments to OIG's Report

The following Director's comments are submitted in response to the recommendations in the OIG report:

OIG Recommendations

Recommendation 1. We recommended that facility managers ensure that licensed practitioners who perform emergency airway management have the appropriate training.

Concur-Yes

Target date for completion: November 30, 2015

Facility response: The updated Medical Center Memorandum has been sent out for concurrence. A checklist is being used to monitor compliance with training. An Associate Chief of Education for residents and medical students was hired this year and will enhance communication between the program and the academic affiliate.

Recommendation 2. We recommended that the Surgical Work Group meet monthly.

Concur-Yes

Target date for completion: December 31, 2015

Facility response: The work group has met monthly since December, 2014 and they will continue to do so.

Recommendation 3. We recommended that facility managers ensure all health care occupancy buildings have at least one fire drill per shift per quarter and monitor compliance.

Concur-Yes

Target date for completion: December 31, 2015

Facility response: Facility managers will promote and ensure better coordination with Services to increase fire drill participation. The Occupancy Matrix and Fire Drill Frequency will be updated and added to the Annual Review to ensure compliance. The data will be presented to the Environment of Care committee quarterly.

Recommendation 4. We recommended that facility managers ensure negative air pressure systems in the medicine primary care clinic are functional and monitor compliance.

Concur-Yes

Target date for completion: October 31, 2015

Facility response: The new monitor has been installed on room D-124, alarm points have been programmed and is functioning as designed. A new lock and key has been ordered for D-122.

Recommendation 5. We recommended that facility managers ensure locked mental health unit stationary panic alarm testing includes documentation of VA Police response time.

Concur-Yes

Target date for completion: October 31, 2015

Facility response: The VA Police's response time to the panic alarm testing will now be recorded as required.

Recommendation 6. We recommended that equipment on the locked mental health unit is secured and heavy enough to prevent it from being picked up, thrown, moved, or overturned.

Concur-Yes

Target date for completion: August 11, 2015

Facility response: The previously mobile TV cart has been bolted to the dining room wall.

Recommendation 7. We recommended that employees ask inpatients whether they would like to discuss creating, changing, and/or revoking advance directives and that facility managers monitor compliance.

Concur-Yes

Target date for completion: August 26, 2015

Facility response: A new clinical reminder will be implemented, to be completed by the nurse upon admission. The clinical reminder will include the following three yes or no questions: 1. Do you have an advanced directive?; 2. Do you have a copy of your advanced directive?; 3. Do you wish to discuss creating, changing and/or revoking your Advanced Directive? A yes response to questions 2 or 3 will generate an Advanced Directive consult to Social Work Service. Compliance will be monitored by the Chief of

Social Work Service or designee and the Chief of Health Information Management (HIMS). Results will be reported monthly to the HIMS Committee.

Recommendation 8. We recommended that employees hold advance directive discussions requested by inpatients and document the discussions and that facility managers monitor compliance.

Concur-Yes

Target date for completion: August 26, 2015

Facility response: A new clinical reminder is being implemented which will allow an Advanced Directive consult to be generated to Social Work Service when the Veteran indicates they have a copy of their Advanced Directive or if they request an Advanced Directive discussion. Compliance will be monitored by the Chief of Social Work Service or designee and the Chief of HIMS. Results will be reported monthly to the HIMS Committee.

Recommendation 9. We recommended that the facility ensure initial clinician emergency airway management competency assessment includes all required subject matter content elements and that facility managers monitor compliance.

Concur-Yes

Target date for completion: September 1, 2015

Facility response: The memorandum was updated and sent out for concurrence on August 18, 2015. The Medical Staff Office will monitor all providers with Out-of-OR Airway Management (OOORAM) privileges for compliance before approving the privilege/certification initially and at renewal which occurs every 2 years. The residents will be reviewed initially and at recertification by the Medical Staff Coordinator and followed by the Resident Education Office.

Recommendation 10. We recommended that the facility revise the local policy for out of operating room emergency airway management to include successful demonstration of all required procedural skills on airway simulators for providers seeking renewal of privileges.

Concur-Yes

Target date for completion: September 30, 2015

Facility response: The Medical Center Memorandum has been updated and is out for concurrence. The process is already taking place.

Recommendation 11. We recommended that the facility document the review of provider-specific emergency airway management data in Cardiopulmonary Review Committee meeting minutes.

Concur-Yes

Target date for completion: December 31, 2015

Facility response: Provider-specific complications have been analyzed and will be provided to each Service for ongoing professional practice evaluations beginning with the August 20, 2015 meeting.

Office of Inspector General Contact and Staff Acknowledgments

Contact	For more information about this report, please contact the OIG at (202) 461-4720.
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This report is available at <u>www.va.gov/oig</u>.

Endnotes

^a References used for this topic included:

- VHA Directive 1026, VHA Enterprise Framework for Quality, Safety, and Value, August 2, 2013.
- VHA Handbook 1050.01, VHA National Patient Safety Improvement Handbook, March 4, 2011.
- VHA Directive 2010-025, Peer Review for Quality Management, June 3, 2010.
- VHA Directive 2010-032, Safe Patient Handling Program and Facility Design, June 28, 2010.
- VHA Directive 1036, Standards for Observation in VA Medical Facilities, February 6, 2014.
- VHA Handbook 1100.19, Credentialing and Privileging, October 15, 2012.
- VHA Handbook 1102.01, National Surgery Office, January 30, 2013.
- VHA Directive 2008-063, Oversight and Monitoring of Cardiopulmonary Resuscitative Events and Facility Cardiopulmonary Resuscitation Committees, October 17, 2008.
- VHA Handbook 1907.01, *Health Information Management and Health Records*, July 22, 2014. ^b References used for this topic included:
- VHA Directive 2008-052, Smoke-Free Policy for VA Health Care Facilities, August 26, 2008.
- VHA Directive 2010-032, Safe Patient Handling Program and Facility Design, June 28, 2010.
- VHA Directive 2011-007, Required Hand Hygiene Practices, February 16, 2011.
- VA National Center for Patient Safety, "Issues continue to occur due to improper ceiling mounted patient lift installation, maintenance and inspection," Addendum to Patient Safety Alert 14-07, September 3, 2014.
- Various requirements of The Joint Commission, the Occupational Safety and Health Administration, the International Association of Healthcare Central Service Materiel Management, the Health Insurance Portability and Accountability Act, Underwriters Laboratories, VA Master Specifications.

^c References used for this topic included:

- VHA Directive 2008-027, The Availability of Potassium Chloride for Injection Concentrate USP, May 13, 2008.
- VHA Directive 2010-020, Anticoagulation Therapy Management, May 14, 2010.
- VHA Handbook 1108.01, Controlled Substances (Pharmacy Stock), November 16, 2010.
- VHA Handbook 1108.05, Outpatient Pharmacy Services, May 30, 2006.
- VHA Handbook 1108.06, Inpatient Pharmacy Services, June 27, 2006.
- VHA Handbook 1108.07, Pharmacy General Requirements, April 17, 2008.
- Various requirements of The Joint Commission.
- ^d The reference used for this topic was:
- Under Secretary for Health, "Consult Business Rule Implementation," memorandum, May 23, 2013.
- ^e References used for this topic included:
- VHA Directive 1129, Radiation Protection for Machine Sources of Ionizing Radiation, February 5, 2015.
- VHA Handbook 1105.02, Nuclear Medicine and Radiation Safety Service, December 10, 2010.
- VHA Handbook 5005/77, *Staffing*, Part II, Appendix G25, Diagnostic Radiologic Technologist Qualifications Standard GS-647, June 26, 2014.
- The Joint Commission, "Radiation risks of diagnostic imaging," Sentinel Event Alert, Issue 47, August 24, 2011.
- VA Radiology, "Online Guide," updated October 4, 2011.
- The American College of Radiology, "ACR–AAPM TECHNICAL STANDARD FOR DIAGNOSTIC MEDICAL PHYSICS PERFORMANCE MONITORING OF COMPUTED TOMOGRAPHY (CT) EQUIPMENT, Revised 2012.

^f The references used for this topic included:

- VHA Handbook 1004.02, Advance Care Planning and Management of Advance Directives, December 24, 2013.
- VHA Handbook 1907.01, Health Information Management and Health Records, July 22, 2014.
- ^g References used for this topic included:
- VHA Directive 2009-001, Restructuring of VHA Clinical Programs, January 5, 2009.
- VHA Directive 2010-018, Facility Infrastructure Requirements to Perform Standard, Intermediate, or Complex Surgical Procedures, May 6, 2010.
- ^h References used for this topic included:
- VHA Directive 2012-032, Out of Operating Room Airway Management, October 26, 2012.
- VHA Handbook 1101.04, Medical Officer of the Day, August 30, 2010.