

Office of Healthcare Inspections

Report No. 16-00112-267

Combined Assessment Program Review of the James H. Quillen VA Medical Center Mountain Home, Tennessee

April 21, 2016

To Report Suspected Wrongdoing in VA Programs and Operations
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Glossary

AD advance directive

CAP Combined Assessment Program

CSP compounded sterile product

CT computed tomography
EHR electronic health record

EOC environment of care

facility James H. Quillen VA Medical Center

FY fiscal year
MH mental health
NA not applicable

NM not met

OIG Office of Inspector General

OR operating room

QSV quality, safety, and value

RRTP residential rehabilitation treatment program

VHA Veterans Health Administration

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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 March 14, 2016.

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

- Quality, Safety, and Value
- Environment of Care
- Medication Management
- Coordination of Care
- Computed Tomography Radiation Monitoring
- Advance Directives

The facility's reported accomplishments were its Wound Care Program and culture of patient safety.

Recommendations: We made recommendations in the following two activities:

Suicide Prevention Program: Ensure the Suicide Prevention Coordinator consistently provides at least five community outreach activities every month.

Mental Health Residential Rehabilitation Treatment Program: Ensure that Domiciliary Care for Homeless Veterans Program employees consistently perform and document weekly inspections of a minimum of 10 percent of resident rooms for contraband and that Mental Health Residential Rehabilitation Treatment Program employees consistently perform and document daily resident room inspections for unsecured medications.

Comments

The Veterans Integrated Service Network Director and Acting Facility Director agreed with the Combined Assessment Program review findings and recommendations and provided acceptable improvement plans. (See Appendixes C and D, pages 27–29, for the full text of the Directors' comments.) We will follow up on the planned actions until they are completed.

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

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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:

- QSV
- EOC
- Medication Management
- Coordination of Care
- CT Radiation Monitoring
- ADs
- Suicide Prevention Program
- MH RRTP

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 2015 and FY 2016 through March 14, 2016, 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 James H. Quillen VA Medical Center, Mountain Home, Tennessee*, Report No. 13-02643-20, November 22, 2013).

During this review, we presented crime awareness briefings for 73 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 576 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

Excellence in Wound Care Program

The facility has a nationally recognized Wound Care Program that provides lymphedema¹ management and wound and ostomy care for patients in acute care, primary care, the community living center, and the outpatient wound clinic and through the Tele-Wound Program.² The facility is one of only five nationally recognized sites approved to perform Store-and-Forward Telehealth³ for wound care. Additionally, the facility participates in cutting-edge pressure ulcer research and is a two-time recipient of the VHA Innovation Award for its aggressive approach to decrease the incidence of hospital-acquired pressure ulcers. The facility's hospital-acquired pressure ulcer rate decreased from 3.2 percent in 2011 to 0.25 percent in FY 2015.

Performance Excellence and Patient Safety Culture

The facility continues its journey to engrain performance excellence and a culture of patient safety as the foundation of all care provided. This has been accomplished through employee training and programs such as "The Good Catch" program, which

¹ Lymphedema is an abnormal collection of high-protein fluid just beneath the skin. This swelling, or edema, occurs most commonly in the arm or leg.

² In the Tele-Wound Program, specialists in wound care use EHRs and images sent electronically for patients in nursing homes, outpatient clinics, and home care.

³ Store-and-Forward Telehealth is the acquisition and storage of clinical information (data, images, sound, video) that is then forwarded to (or retrieved by) another site for clinical evaluation. http://www.telehealth.va.gov/sft/ Accessed March 23, 2016.

encourages and positively acknowledges reporting close calls and near miss incidents that could have had a negative patient outcome. This journey of excellence received recognition through the Robert W. Carey Performance Excellence Trophy Award and State of Tennessee Baldrige award. In addition, the facility's Patient Safety Program has received a gold award from the VA National Center for Patient Safety every year since 2010 for a strong root cause analysis process.

Results and Recommendations

QSV

The purpose of this review was to determine whether the facility complied with selected QSV program requirements.^a

We conversed with senior managers and key QSV employees, and we evaluated meeting minutes, 20 licensed independent practitioners' profiles, 10 protected peer reviews, 5 root cause analyses, and other relevant documents. 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
	There was a senior-level committee responsible for key QSV functions that met at least quarterly and was chaired or co-chaired by the Facility Director. The committee routinely reviewed aggregated data.		
	Credentialing and privileging processes met selected requirements: • Facility policy/by-laws addressed a frequency for clinical managers to review practitioners' Ongoing Professional Practice Evaluation data. • Facility clinical managers reviewed Ongoing Professional Practice Evaluation data at the frequency specified in the policy/by-laws. • The facility set triggers for when a Focused Professional Practice Evaluation for cause would be indicated. • The facility followed its policy when employees' licenses expired.		

Areas Reviewed (continued)	Findings	Recommendations
Protected peer reviews met selected	_	
requirements:		
 Peer reviewers documented their use of 		
important aspects of care in their review		
such as appropriate and timely ordering of		
diagnostic tests, timely treatment, and		
appropriate documentation.		
 When the Peer Review Committee 		
recommended individual improvement		
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	Protected peer reviews met selected requirements: • Peer reviewers documented their use of important aspects of care in their review such as appropriate and timely ordering of diagnostic tests, timely treatment, and appropriate documentation. • When the Peer Review Committee	Protected peer reviews met selected requirements: Peer reviewers documented their use of important aspects of care in their review such as appropriate and timely ordering of diagnostic tests, timely treatment, and appropriate documentation. When the Peer Review Committee recommended individual improvement actions, clinical managers implemented the actions. Utilization management met selected requirements: The facility completed at least 75 percent of all required inpatient reviews. Physician Utilization Management Advisors documented their decisions in the National Utilization Management Integration database. The facility had designated an interdisciplinary group to review utilization management data. Patient safety met selected requirements: The Patient Safety Manager entered all reported patient incidents into the WEBSPOT database. The facility completed the required minimum of eight root cause analyses. The facility provided feedback about the root cause analysis findings to the individual or department who reported the incident. At the completion of FY 2015, the Patient Safety Manager submitted an annual

NM	Areas Reviewed (continued)	Findings	Recommendations
	Overall, if QSV reviews identified significant issues, the facility took actions and evaluated them for effectiveness.		
	Overall, senior managers actively participated in QSV activities.		
	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 the dental clinic and the OR.^b

We inspected the medical/surgical/telemetry, intensive care, progressive care, post-anesthesia care, and locked MH units; two community living center units; the Emergency Department; the Physical Medicine and Rehabilitation Service; the OR; and the dental and cardiology clinics. Additionally, we reviewed relevant documents and 21 employee-training records, and we conversed with key employees and managers. 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 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 between patients.		
	The facility conducted required fire drills in		
	buildings designated for health care		
	occupancy and documented drill critiques.		
	The facility had a policy/procedure/guideline		
	for identification of individuals entering the		
	facility, and units/areas complied with		
	requirements.		
	The facility met fire safety requirements.		

NM	Areas Reviewed for General EOC	Findings	Recommendations
	(continued)		
	The facility met environmental safety		
	requirements.		
	The facility met infection prevention		
	requirements.		
	The facility met medication safety and		
	security requirements.		
	The facility met privacy requirements.		
	The facility complied with any additional		
	elements required by VHA, local policy, or		
	other regulatory standards.		
	Areas Reviewed for Dental Clinic		
	Dental clinic employees completed		
	bloodborne pathogens training within the		
	past 12 months.		
	Dental clinic employees received hazard		
	communication training on chemical		
	classification, labeling, and safety data		
	sheets.		
	Designated dental clinic employees received		
	laser safety training in accordance with local		
	policy.		
	The facility tested dental water lines in		
	accordance with local policy.		
	The facility met environmental safety and		
	infection prevention requirements in the		
	dental clinic.		
	The facility met laser safety requirements in		
	the dental clinic.		
	The facility complied with any additional		
	elements required by VHA, local policy, or		
	other regulatory standards.		

NM	Areas Reviewed for the OR	Findings	Recommendations
	The facility had emergency fire		
	policy/procedures for the OR that included		
	alarm activation, evacuation, and equipment		
	shutdown with responsibility for turning off		
	room or zone oxygen.		
	The facility had cleaning policy/procedures		
	for the OR and adjunctive areas that		
	included a written cleaning schedule and		
	methods of decontamination.		
	OR housekeepers received training on OR		
	cleaning/disinfection in accordance with local		
	policy.		
	The facility monitored OR temperature,		
	humidity, and positive pressure.		
	The facility met fire safety requirements in		
	the OR.		
	The facility met environmental safety		
	requirements in the OR.		
	The facility met infection prevention		
	requirements in the OR.		
	The facility met medication safety and		
	security requirements in the OR.		
	The facility met laser safety requirements in		
	the OR.		
	The facility complied with any additional		
	elements required by VHA, local policy, or		
	other regulatory standards.		

Medication Management

The purpose of this review was to determine whether the facility complied with selected requirements for the safe preparation of CSPs.c

We reviewed relevant documents and the competency assessment/testing records of 10 pharmacy employees (3 pharmacists and 7 technicians). Additionally, we inspected the area where sterile products are compounded. 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 policy on preparation of		
	CSPs that included required components:		
	Pharmacist CSP preparation or		
	supervision of preparation except in urgent		
	situations		
	Hazardous CSP preparation in an area		
	separate from routine CSP preparation or		
	in a compounding aseptic containment		
	isolator		
	Environmental quality and control of ante		
	and buffer areas		
	Hood certification initially and every		
	6 months thereafter		
	Cleaning procedures for all surfaces in the		
	ante and buffer areas		
	The facility established competency		
	assessment requirements for employees		
	who prepare CSPs that included required		
	elements, and facility managers assessed		
	employee competency at the required		
	frequency based on the facility's risk level.		

NM	Areas Reviewed (continued)	Findings	Recommendations
	If the facility used an outsourcing facility for		
	CSPs, it had a policy/guidelines/a plan that		
	included required components for the		
	outsourcing facility:		
	 Food and Drug Administration registration 		
	Current Drug Enforcement Agency		
	registration if compounding controlled		
	substances		
	The facility had a safety/competency		
	assessment checklist for preparation of		
	CSPs that included required steps in the		
	proper order to maintain sterility.		
	All International Organization for		
	Standardization classified areas had		
	documented evidence of periodic surface		
	sampling, and the facility completed required		
	actions when it identified positive cultures.		
	The facility had a process to track and report		
	CSP medication errors, including near		
	misses.		
	The facility met design and environmental		
	safety controls in compounding areas.		
	The facility used a laminar airflow hood or		
	compounding aseptic isolator for preparing		
	non-hazardous intravenous admixtures and		
	any sterile products.		
	The facility used a biological safety cabinet		
	in a physically separated negative pressure		
	area or a compounding aseptic containment		
	isolator for hazardous medication		
	compounding and had sterile chemotherapy		
	type gloves available for compounding these		
	medications.		

NM	Areas Reviewed (continued)	Findings	Recommendations
	If the facility prepared hazardous CSPs, a		
	drug spill kit was available in the		
	compounding area and during transport of		
	the medication to patient care areas.		
	Hazardous CSPs were physically separated		
	or placed in specially identified segregated		
	containers from other inventory to prevent		
	contamination or personnel exposure.		
	An eyewash station was readily accessible		
	near hazardous medication compounding		
	areas, and there was documented evidence		
	of weekly testing.		
	The facility documented cleaning of		
	compounding areas, and employees		
	completed cleaning at required frequencies.		
	During the past 12 months, the facility initially certified new hoods and recertified all		
	hoods minimally every 6 months.		
	Prepared CSPs had labels with required		
	information prior to delivery to the patient		
	care areas:		
	Patient identifier		
	Date prepared		
	Admixture components		
	Preparer and checker identifiers		
	Beyond use date		
-	The facility complied with any additional		
	elements required by VHA, local policy, or		
	other regulatory standards.		
	other regulatory standards.		

Coordination of Care

The purpose of this review was to evaluate selected aspects of the facility's patient flow process over the inpatient continuum (admission through discharge).^d

We reviewed relevant documents and conversed with key employees. Additionally, we reviewed the EHRs of 34 randomly selected patients who had an acute care inpatient stay of at least 3 days from July 1, 2014, through June 30, 2015. 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 policy that addressed		
	patient discharge and scheduling discharges		
	early in the day.		
	The facility had a policy that addressed		
	temporary bed locations, and it included:		
	Priority placement for inpatient beds given		
	to patients in temporary bed locations		
	Upholding the standard of care while		
	patients are in temporary bed locations		
	Medication administration		
	Meal provision		
	The Facility Director had appointed a Bed		
	Flow Coordinator with a clinical background.		
	Physicians or acceptable designees		
	completed a history and physical exam		
	within 1 day of the patient's admission or		
	referenced a history and physical exam		
	completed within 30 days prior to admission.		
	 When resident physicians completed the 		
	history and physical exams, the attending		
	physicians provided a separate admission		
	note or addendum within 1 day of the		
	admission.		

NM	Areas Reviewed (continued)	Findings	Recommendations
	 When the facility policy and/or scopes of practice allowed for physician assistants or nurse practitioners to complete history and physical exams, they were properly documented. 		
	Nurses completed admission assessments within 1 day of the patient's admission.		
	 When patients were transferred during the inpatient stay, physicians or acceptable designees documented transfer notes within 1 day of the transfer. When resident physicians wrote the transfer notes, attending physicians documented adequate supervision. Receiving physicians documented transfers. 		
	When patients were transferred during the inpatient stay, sending and receiving nurses completed transfer notes.		
	Physicians or acceptable designees documented discharge progress notes or instructions that included patient diagnoses, discharge medications, and follow-up activity levels. • When resident physicians completed the discharge notes/instructions, attending physicians documented adequate supervision. • When facility policy and/or scopes of practice allowed for physician assistants or nurse practitioners to complete discharge notes/instructions, they were properly documented.		

NM	Areas Reviewed (continued)	Findings	Recommendations
	Clinicians provided discharge instructions to		
	patients and/or caregivers and documented		
	patients and/or caregiver understanding.		
	The facility complied with 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 eight CT technologists and CT scanner inspection reports, and we 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		
	prior to ocario		l

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.		

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 34 randomly selected patients who had an acute care admission July 1, 2014, through June 30, 2015. 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 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.		
	Employees asked inpatients if they would		
	like to discuss creating, changing, and/or		
	revoking ADs.		
	When inpatients requested a discussion,		
	employees documented the discussion		
	and used the required AD note titles.		
	The facility met any additional elements		
	required by VHA or local policy.		

Suicide Prevention Program

The purpose of this review was to evaluate the extent the facility's MH providers consistently complied with selected suicide prevention program requirements.⁹

We reviewed relevant documents and conversed with key employees. Additionally, we reviewed the EHRs of 40 patients assessed to be at risk for suicide during the period October 1, 2014–September 30, 2015, plus those who died from suicide during this same timeframe. We also reviewed the training records of 15 new employees. 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	Recommendation
	The facility had a full-time Suicide Prevention		
	Coordinator.		
	The facility had a process for responding to		
	referrals from the Veterans Crisis Line and		
	for tracking patients who are at high risk for		
	suicide.		
	The facility had a process to follow up on		
	high-risk patients who missed MH		
	appointments.		
	The facility provided training within required		
	timeframes:		
	 Suicide prevention training to new 		
	employees		
	Suicide risk management training to new		
	clinical employees		
X	The facility provided at least five suicide	 In the 3 months prior to the site visit, the 	1. We recommended that the Suicide
	prevention outreach activities to community	Suicide Prevention Coordinator provided	Prevention Coordinator consistently provide
	organizations each month.	evidence of only two outreach activities	at least five community outreach activities
		for 1 month and four outreach activities	every month and that facility managers
		for another month.	monitor compliance.
	The facility completed required reports and		
	reviews regarding patients who attempted or		
	completed suicide.		

NM	Areas Reviewed (continued)	Findings	Recommendation
	Clinicians assessed patients for suicide risk		
	at the time of admission.		
	Clinicians appropriately placed Patient		
	Record Flags:		
	High-risk patients received Patient Record		
	Flags.		
	Moderate- and low-risk patients did not		
	receive Patient Record Flags.		
	Clinicians documented Suicide Prevention		
	Safety Plans that contained the following		
	required elements:		
	Identification of warning signs		
	 Identification of internal coping strategies 		
	Identification of contact numbers of family		
	or friends for support		
	Identification of professional agencies		
	Assessment of available lethal means and		
	how to keep the environment safe		
	Clinicians documented that they gave		
	patients and/or caregivers a copy of the		
	safety plan.		
	The treatment team evaluated patients as		
	follows:		
	At least four times during the first 30 days		
	after discharge		
	Every 90 days to review Patient Record		
	Flags		
	The facility complied with any additional		
	elements required by VHA or local policy.		

MH RRTP

The purpose of this review was to determine whether the facility's Domiciliary Care for Homeless Veterans Program and the general domiciliary RRTP complied with selected EOC requirements.^h

We reviewed relevant documents, inspected the Domiciliary Care for Homeless Veterans Program female and male units and the general domiciliary RRTP units P and PG, and conversed with key employees. 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.

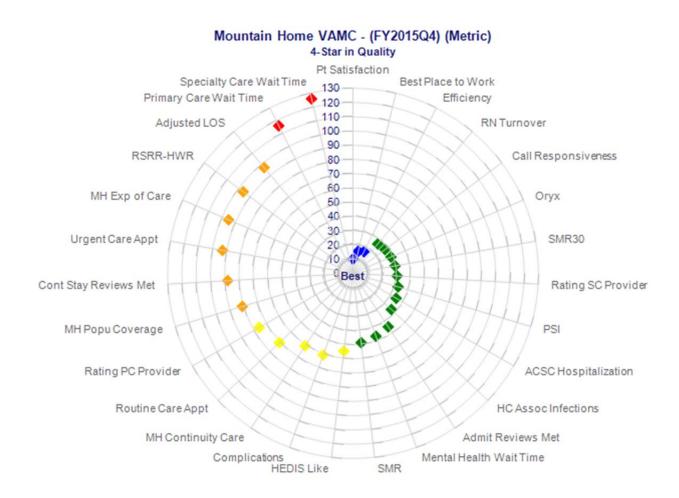
NM	Areas Reviewed	Findings	Recommendation
	The residential environment was clean and in good repair.		
	Appropriate fire extinguishers were available near grease producing cooking devices.		
	There were policies/procedures that addressed safe medication management and contraband detection.		
	MH RRTP employees conducted and documented monthly MH RRTP self-inspections that included all required elements, submitted work orders for items needing repair, and ensured correction of any identified deficiencies.		
X	MH RRTP employees conducted and documented contraband inspections, rounds of all public spaces, daily bed checks, and resident room inspections for unsecured medications.	 For the month of February 2016, Domiciliary Care for Homeless Veterans Program employees did not document weekly inspections of a minimum of 10 percent of resident rooms for contraband. For the 13-day period 	2. We recommended that Domiciliary Care for Homeless Veterans Program employees consistently perform and document weekly inspections of a minimum of 10 percent of resident rooms for contraband and that program managers monitor compliance.
		For the 13-day period February 8–20, 2016, MH RRTP employees did not consistently document daily resident room inspections for unsecured medications.	3. We recommended that Mental Health Residential Rehabilitation Treatment Program employees consistently perform and document daily resident room inspections for unsecured medications and that program managers monitor compliance.

NM	Areas Reviewed (continued)	Findings	Recommendation
	The MH RRTP had written agreements in		
	place acknowledging resident responsibility		
	for medication security.		
	MH RRTP main point(s) of entry had keyless		
	entry and closed circuit television monitoring,		
	and all other doors were locked to the		
	outside and alarmed.		
	The MH RRTP had closed circuit television		
	monitors with recording capability in public		
	areas but not in treatment areas or private		
	spaces and signage alerting veterans and		
	visitors of recording.		
	There was a process for responding to		
	behavioral health and medical emergencies,		
	and MH RRTP employees could articulate		
	the process.		
	In mixed gender MH RRTP units, women		
	veterans' rooms had keyless entry or door		
	locks, and bathrooms had door locks.		
	Residents secured medications in their		
	rooms.		
	The facility complied with any additional		
	elements required by VHA or local policy.		

Facility Profile (Mountain Home/621) FY 2016 through February 2016		
Type of Organization	Secondary	
Complexity Level	1c-High complexity	
Affiliated/Non-Affiliated	Affiliated	
Total Medical Care Budget in Millions	\$396.7	
Number of:		
Unique Patients	45,301	
Outpatient Visits	281,263	
Unique Employees ⁴	2,056	
Type and Number of Operating Beds:		
Hospital	102	
Community Living Center	120	
Domiciliary	150	
Average Daily Census:		
Hospital	79	
Community Living Center	68	
Domiciliary	105	
Number of Community Based Outpatient Clinics ⁵ 7		
Location(s)/Station Number(s)	Knoxville/621BY	
	Rogersville/621GA	
	Norton/621GC	
	Morristown/621GG	
	Sevierville/621GI	
	Bristol/621GJ	
	LaFollette/621GK	
Veterans Integrated Service Network Number 9		

 ⁴ Unique employees involved in direct medical care (cost center 8200).
 ⁵ We have omitted 621QA (Norton), 621QB (Marion), 621QC (Vansant), 621QD (Knoxville), and 621BU (Mountain Home) as no workload/encounters or services were reported.

Strategic Analytics for Improvement and Learning (SAIL)⁶



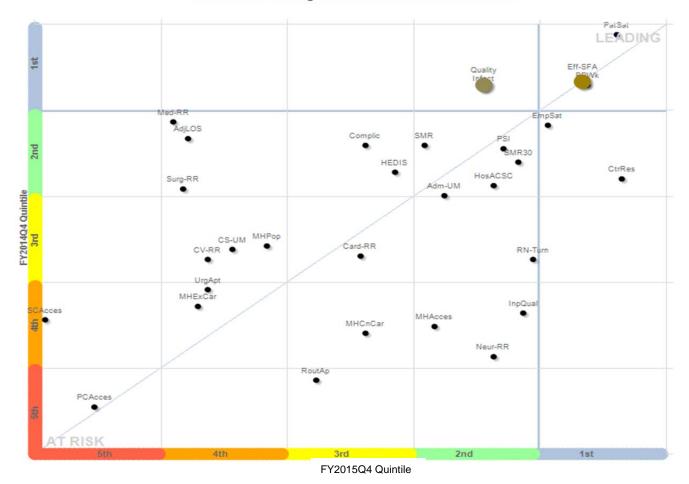
Marker color: Blue - 1st quintile; Green - 2nd; Yellow - 3rd; Orange - 4th; Red - 5th quintile.

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⁶ Metric definitions follow the graphs.

Scatter Chart

FY2015Q4 Change in Quintiles from FY2014Q4



DESIRED DIRECTION =>

NOTE

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

Veterans Integrated Service Network Director Comments

Department of Veterans Affairs

Memorandum

Date: April 6, 2016

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

Subject: CAP Review of the James H. Quillen VA Medical Center,

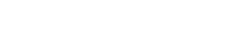
Mountain Home, TN

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

Director, Management Review Service (VHA 10E1D MRS OIG CAP

CBOC)

- 1. I concur with the findings and recommendations of this Office of Inspector General Combined Assessment Program Review of the James H. Quillen VA Medical Center, Mountain Home, Tennessee, as well as the action plan developed by the facility.
- 2. If you have any questions or need additional information from the Network, please do not hesitate to contact me at 615-695-2206.



Acting Facility Director Comments

Department of Veterans Affairs

Memorandum

Date: April 4, 2016

From: Acting Director, James H. Quillen VA Medical Center (621/00)

Subject: CAP Review of the James H. Quillen VA Medical Center,

Mountain Home, TN

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

 On behalf of the James H. Quillen VA Medical Center, Mountain Home, Tennessee, I concur with the findings and recommendations of this Office of Inspector General report.

2. Included herein is an outline of improvement actions taken, in progress, or planned in response to these findings. We believe these changes will further enhance key systems and processes throughout our healthcare system.

Daniel Snyder, P.E., FACHE

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 the Suicide Prevention Coordinator consistently provide at least five community outreach activities every month and that facility managers monitor compliance.

Concur

Target date for completion: September 30, 2016

Facility response: The Suicide Prevention Coordinators are assigned to ensure at least five community outreach activities are completed every month with a summation to be reported to Mental Health Executive Council as a standing agenda item to ensure 100 percent compliance. These outreach activities are to be recorded and reported to the Mental Health Social Work Supervisor on a weekly basis.

Recommendation 2. We recommended that Domiciliary Care for Homeless Veterans Program employees consistently perform and document weekly inspections of a minimum of 10 percent of resident rooms for contraband and that program managers monitor compliance.

Concur

Target date for completion: September 30, 2016

Facility response: The Mental Health Residential Rehabilitation Treatment Program weekly inspection sheets have been revised to include verbiage of no contraband found. The documentation of weekly contraband inspections will be monitored by Chief, Domiciliary and reported to Mental Health Executive Council as standing agenda item to ensure 100 percent compliance.

Recommendation 3. We recommended that Mental Health Residential Rehabilitation Treatment Program employees consistently perform and document daily resident room inspections for unsecured medications and that program managers monitor compliance.

Concur

Target date for completion: September 30, 2016

Facility response: The Mental Health Residential Rehabilitation Treatment Program daily room inspection sheets have been revised to require Chief/Assistant Chief Domiciliary or designated Administrator on Duty signature and daily approval to ensure documentation of unsecured medication checks. These daily monitors are required to be 100% and will be reported to Mental Health Executive Council as a standing agenda item.

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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Jim Cooper; Scott DesJarlais; John J. Duncan, Jr.; Stephen Fincher;

Chuck Fleischmann; Phil Roe

This report is available at www.va.gov/oig.

Endnotes

- ^a The references used for this topic were:
- VHA Directive 1026, VHA Enterprise Framework for Quality, Safety, and Value, August 2, 2013.
- VHA Directive 1117, Utilization Management Program, July 9, 2014.
- VHA Directive 2010-025, Peer Review for Quality Management, June 3, 2010.
- VHA Handbook 1050.01, VHA National Patient Safety Improvement Handbook, March 4, 2011.
- VHA Handbook 1100.19, Credentialing and Privileging, October 15, 2012.
- ^b The references used for this topic included:
- VHA Directive 2005-037, *Planning for Fire Response*, September 2, 2005.
- VHA Directive 2009-026; Location, Selection, Installation, Maintenance, and Testing of Emergency Eyewash and Shower Equipment; May 13, 2009.
- 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, National Fire Protection Association, Association of periOperative Registered Nurses, U.S. Pharmacopeial Convention, American National Standards Institute.
- ^c The references used for this topic included:
- VHA Handbook 1108.06, Inpatient Pharmacy Services, June 27, 2006.
- VHA Handbook 1108.07, Pharmacy General Requirements, April 17, 2008.
- Various requirements of VA Pharmacy Benefits Management Services, The Joint Commission, the United States Pharmacopeial Convention, the American Society of Health-System Pharmacists, the Institute for Safe Medication Practices, the Food and Drug Administration, and the American National Standards Institute.
- ^d The references used for this topic included:
- VHA Directive 1009, Standards for Addressing the Needs of Patients Held in Temporary Bed Locations, August 28, 2013.
- VHA Directive 1063, Utilization of Physician Assistants (PA), December 24, 2013.
- VHA Handbook 1400.01, Resident Supervision, December 19, 2012.
- VHA Handbook 1907.01, Health Information Management and Health Records, March 19, 2015.
- ^e The 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 The references used for this topic included:
- VHA Directive 2010-025, Peer Review for Quality Management, June 3, 2010.
- VHA Directive 2010-053, Patient Record Flags, December 3, 2010 (corrected 2/3/11).
- VHA Handbook 1050.01, VHA National Patient Safety Improvement Handbook, March 4, 2011.
- VHA Handbook 1160.01, *Uniform Mental Health Services in VA Medical Centers and Clinics*, September 11, 2008.
- VHA Handbook 1160.06, Inpatient Health Services, September 16, 2013.
- Various Deputy Under Secretary for Health for Operations and Management memorandums and guides.
- VA Suicide Prevention Coordinator Manual, August 2014.
- Various requirements of The Joint Commission.

VA OIG Office of Healthcare Inspections

^h The references used for this topic were:

[•] VHA Handbook 1162.02, *Mental Health Residential Rehabilitation Treatment Program (MH RRTP)*, December 22, 2010.

[•] VHA Handbook 1330.01, Health Care Services for Women Veterans, May 21, 2010.

[•] Requirements of the VHA Center for Engineering and Occupational Safety and Health and the National Fire Protection Association.