

Long Term Care Facility Component—Annual Facility Survey

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Instructions for this form can be accessed: https://www.cdc.gov/nhsn/forms/instr/57.137-toi-annual-facility-survey.pdf	
*Required for saving	Tracking #:
Facility ID:	*Survey Year:
*National Provider ID:	State Provider #:
Facility Characteristics	
*Ownership (check one):	
<input type="checkbox"/> For profit <input type="checkbox"/> Not for profit, including church <input type="checkbox"/> Government (not VA) <input type="checkbox"/> Veterans Affairs	
*Certification (check one):	
<input type="checkbox"/> Dual Medicare/Medicaid <input type="checkbox"/> Medicare only <input type="checkbox"/> Medicaid only <input type="checkbox"/> State only	
*Affiliation (check one):	
<input type="checkbox"/> Independent, free-standing <input type="checkbox"/> Independent, continuing care retirement community <input type="checkbox"/> Multi-facility organization (chain) <input type="checkbox"/> Hospital system, attached <input type="checkbox"/> Hospital system, free-standing	
<i>In the previous calendar year:</i>	
*Average daily census: _____	
*Total number of short-stay residents: _____ Average length of stay for short-stay residents: _____	
*Total number of long-stay residents: _____ Average length of stay for long-stay residents: _____	
*Total number of new admissions: _____	
*Number of Beds: _____ *Number of Pediatric Beds (age <21): _____	
*Indicate which of the following primary service types are provided by your facility. On the day of this survey, indicate the number of residents receiving those services (list only one service type per resident, i.e. total should sum to resident census on day of survey completion):	
<u>Primary Service Type</u>	<u>Service provided?</u>
a. Long-term general nursing:	<input type="checkbox"/> _____
b. Long-term dementia:	<input type="checkbox"/> _____
c. Skilled nursing/Short-term (subacute) rehabilitation:	<input type="checkbox"/> _____
d. Long-term psychiatric (non-dementia):	<input type="checkbox"/> _____
e. Ventilator:	<input type="checkbox"/> _____
f. Bariatric:	<input type="checkbox"/> _____
g. Hospice/Palliative:	<input type="checkbox"/> _____
h. Other:	<input type="checkbox"/> _____
<p>Assurance of Confidentiality: The voluntarily provided information obtained in this surveillance system that would permit identification of any individual or institution is collected with a guarantee that it will be held in strict confidence, will be used only for the purposes stated, and will not otherwise be disclosed or released without the consent of the individual, or the institution in accordance with Sections 304, 306 and 308(d) of the Public Health Service Act (42 USC 242b, 242k, and 242m(d)).</p> <p>Public reporting burden of this collection of information is estimated to average 2 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering, and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC, Reports Clearance Officer, 1600 Clifton Rd., MS D-74, Atlanta, GA 30333, ATTN: PRA (0920-0666).</p>	
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Facility Microbiology Laboratory Practices

*1. Does your facility have its own laboratory that performs microbiology/antimicrobial susceptibility testing?

- Yes No

If No, where is your facility's antimicrobial susceptibility testing performed? (check one)

- Affiliated medical center, within same health system Medical center, contracted locally
 Commercial referral laboratory

*2. Indicate whether your facility screens new admissions for any of the following multidrug-resistant organisms (MDROs): (check all that apply)

- We do not screen new admissions for MDROs
- Methicillin-resistant *Staphylococcus aureus* (MRSA)
If checked, indicate the specimen types sent for screening: (check all that apply)
 Nasal swabs Wound swabs Sputum Other skin site
- Vancomycin-resistant *Enterococcus* (VRE)
If checked, indicate the specimen types sent for screening: (check all that apply)
 Rectal swabs Wound swabs Urine
- Multidrug-resistant gram-negative rods (includes carbapenemase resistant Enterobacteriaceae; multidrug-resistant *Acinetobacter*, etc.)
If checked, indicate the specimen types sent for screening: (check all that apply)
 Rectal swabs Wound swabs Sputum Urine
- Candida Auris* (*C. Auris*)
If checked, indicate the specimen types sent for screening: (check all that apply)
 Skin Nares Other site
(axilla/groin)

*3. What is the primary testing method for *C. difficile* used most often by your facility's laboratory or the outside laboratory where your facility's testing is performed? (check one)

- | | |
|--|---|
| <input type="checkbox"/> Enzyme immunoassay (EIA) for toxin | <input type="checkbox"/> GDH plus NAAT (2-step algorithm) |
| <input type="checkbox"/> Cell cytotoxicity neutralization assay | <input type="checkbox"/> GDH plus EIA for toxin, followed by NAAT for discrepant results |
| <input type="checkbox"/> Nucleic acid amplification test (NAAT) (e.g., PCR, LAMP) | <input type="checkbox"/> Culture (<i>C. difficile</i> culture followed by detection of toxins) |
| <input type="checkbox"/> NAAT plus EIA, if NAAT positive (2-step algorithm) | <input type="checkbox"/> Other (specify): _____ |
| <input type="checkbox"/> Glutamate dehydrogenase (GDH) antigen plus EIA for toxin (2-step algorithm) | |

("Other" should not be used to name specific laboratories, reference laboratories, or the brand names of *C. difficile* tests; most methods can be categorized accurately by selecting from the options provided. Please ask your laboratory, refer to the Tables of Instructions for this form, or conduct a search for further guidance on selecting the correct option to report.)

*4. Does your laboratory provide a report summarizing the percent of antibiotic resistance seen in common organisms identified in cultures sent from your facility (often called an antibiogram)?

- Yes No

If Yes, how often is this summary report or antibiogram provided to your facility? (check one)

- Once a year Every 2 years Other (specify): _____

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Infection Prevention and Control Practices			
*5. Total staff hours per week dedicated to infection prevention and control activity in facility: _____			
a. Total hours per week performing surveillance: _____			
b. Total hours per week for infection prevention and control activities other than surveillance: _____			
*6. Is it a policy in your facility to routinely use gown/gloves for care of residents infected or colonized with a multidrug-resistant organism (MDRO)? <input type="checkbox"/> Yes <input type="checkbox"/> No (<i>If "No", continue to question #7</i>)			
If yes, please select the option that is applicable to your facility for each MDRO. (<i>"No" should only be selected if your facility does not have a policy for the MDRO listed.</i>)			
<u>Multidrug-resistant organism (MDRO)</u>	<u>All infected or colonized with?</u>	<u>Certain characteristics that make them high risk for transmission (e.g., wounds, presence of an indwelling device)</u>	<u>No</u>
a. MRSA:	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
b. VRE:	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
c. CRE:	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
d. ESBL or extended spectrum cephalosporin resistant Enterobacteriaceae	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<u>Novel and/or CDC-targeted MDROs</u>			
e. Pan-resistant organisms	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
f. Carbapenemase-producing organisms (e.g., Carbapenemase-producing Enterobacterales)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
g. Candida auris	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
*7. Is it a policy in your facility to use gowns/gloves for care of residents with certain characteristics that make them high-risk for transmission or acquisition of an MDRO (e.g., wounds, presence of an indwelling device) regardless of MDRO status? <input type="checkbox"/> Yes <input type="checkbox"/> No			
*8. When a resident colonized or infected with an MDRO is transferred to another facility, does your facility communicate the resident's MDRO status to the receiving facility at the time of transfer? <input type="checkbox"/> Yes <input type="checkbox"/> No			
*9. Among residents with an MDRO admitted to your facility from other healthcare facilities, what percentage of the time does your facility receive information from the transferring facility about the resident's MDRO status? _____%			

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Antibiotic Stewardship Practices

*10. Are there one or more individuals responsible for the impact of activities to improve use of antimicrobials at your facility? Yes No

If Yes, what is the position of the individual(s)? (select all that apply)

- Medical director Director of Nursing Infection Preventionist
 Consultant Pharmacist Other (please specify): _____

*11. Does your facility have a policy that requires prescribers to document an indication for all antimicrobials in the medical record or during order entry? Yes No

If Yes, has adherence to the policy to document an indication been monitored? Yes No

*12. Does your facility provide treatment recommendations for common infections based on national guidelines to assist with antimicrobial decision making? Yes No

If Yes, has adherence to facility-specific treatment recommendations been monitored? Yes No

*13. Is there a formal procedure for performing a follow-up assessment 2-3 days after a new antimicrobial start to determine whether the antimicrobial is still indicated and appropriate (e.g. antibiotic time out)? Yes No

*14. Is there a formal procedure for reviewing courses of antimicrobial therapy and communicating with prescribers on antimicrobial selection, dosing, or duration of therapy (i.e., audit and feedback) at your facility? Yes No

*15. Does your facility have a system for tracking antimicrobial use?
If yes, what is the source of the antimicrobial use report provided? Yes No

- Pharmacy services Electronic Health Records
 Manual reporting (i.e., facility infection control log) Other (please specify): _____

*16. Has your facility provided education to clinicians and other facility staff on improving antimicrobial use in the past 12 months? Yes No

*17. Does your facility have a written statement of support from leadership that supports efforts to improve antimicrobial use? Yes No

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|--|------------------------------|-----------------------------|------------------------------|-----------------------------|
| Temperature | <input type="checkbox"/> Yes | <input type="checkbox"/> No | | |
| If Yes, do you have a plan for corrective actions when temperatures are not within acceptable limits as determined by your water management program? | | | <input type="checkbox"/> Yes | <input type="checkbox"/> No |
| Heterotrophic plate counts | <input type="checkbox"/> Yes | <input type="checkbox"/> No | | |
| If Yes, do you have a plan for corrective actions when heterotrophic plate counts are not within acceptable limits as determined by your water management program? | | | <input type="checkbox"/> Yes | <input type="checkbox"/> No |
| Specific tests for <i>Legionella</i> | <input type="checkbox"/> Yes | <input type="checkbox"/> No | | |
| If Yes, do you have a plan for corrective actions when specific tests for <i>Legionella</i> are not within acceptable limits as determined by your water management program? | | | <input type="checkbox"/> Yes | <input type="checkbox"/> No |