

Candida auris Investigation Form		<i>Public Health Use Only</i>	Confirmed Clinical	Not a case Colonization/screening	Out of jurisdiction	Probable
Patient's name: _____ <div style="display: flex; justify-content: space-between;"> Last First MI </div> Address: _____ Homeless City: _____ State: _____ Zip: _____ County: _____ Home #: () _____ Work #: () _____ Date of birth: _____ Age: _____ Sex: Male Female UNK Country of birth: _____ Country of usual residence: _____ Ethnicity: Hispanic/Latino Not Hispanic/Latino UNK Race: Am.Indian/Alaskan Native Asian Black/African Am. Native Hawaiian/ Pacific Isl. White UNK Refused Not asked Other: _____ Current occupation: _____ Current industry: _____			Jurisdiction: _____ Investigation start date: _____ Investigated by: _____ Phone: () _____ Reporting county: _____ Reporting source type: _____ Reporting organization: _____ Reporting provider: _____ Reported by: _____ Phone: () _____ Date reported: _____			
HOSPITAL/FACILITY INFORMATION						
Was the patient admitted to a healthcare facility (HCF)? Yes, name of HCF: _____ HCF County _____ State _____ No UNK Was visit due to an outpatient/ home health/ ER, etc. visit only? Yes, name of HCF: _____ No HCF admission: _____ HCF discharge: _____ Date of outpatient visit: _____ Were control measures (per EAIDG Guidance) implemented at the admitting HCF? Yes No UNK NA Facility type patient came from: Home Acute care hospital LTAC LTCF/NH Rehab Hospice UNK NA Other Name of facility: _____ Was this facility notified of MDRO? Yes No UNK Were control measures (per EAIDG Guidance) implemented at the facility the patient came from? Yes No UNK NA Discharged to: Home Acute care hospital LTAC LTCF/NH Rehab Hospice UNK NA Other Patient still admitted Patient expired Name of facility: _____ Was this facility notified of MDRO? Yes No UNK Were control measures (per EAIDG Guidance) implemented at the facility the patient was discharged to? Yes No UNK NA						
Other Information			Travel and Healthcare			
Diagnosis date: _____ Date of symptom onset: _____ Did patient die? Yes, date of death: _____ No UNK Did C. auris contribute to death? Yes No UNK Where was the disease acquired?: Imported but unable to determine In state, out of jurisdiction Indigenous International Out of state Unknown Imported country: _____ Imported state: _____			Did the patient receive overnight healthcare within the USA, but outside of the patient's resident state, in the year prior to date of specimen collection? Yes No UNK List state(s): _____ Did the patient travel internationally in the year prior to the date of specimen collection? Yes No UNK List country(ies): _____ Did the patient receive overnight healthcare outside of the USA in the year prior to the date of specimen collection? Yes No UNK List country(ies): _____			
LABORATORY DATA Performing laboratory specimen ID#: _____ State laboratory specimen ID#: _____ WGS ID#: _____						
Pathogen: <i>C. auris</i> <i>C. haemulonii</i> Other: _____ Date collected: _____ Specimen source: _____ Swab site(s), if applicable: NA Axilla Groin Nares Ear Oropharyngeal Rectal Wound Other: _____ Test type: PCR Automatic biochemical/phenotypic test DNA sequencing MALDI-TOF Non-PCR culture-independent diagnostic test Other, specify: _____ Test result: Positive Negative Indeterminate Unsatisfactory Patient status at time of specimen collection (inpatient, LTAC, LTCF, outpatient, etc.): _____ County and state of facility where specimen was collected: _____						
PREVIOUS HISTORY <i>Note: Case ID for Carbapenemase-producing (CP) CRE cases and clinical C. auris cases to be provided by regional HAI epidemiologist or DSHS Laboratory.</i>						
Clinical C. auris only: Patient previously counted as a colonization/screening case? Yes No UNK If yes, case ID: _____ Was patient previously counted as a CP-CRE case? Yes No UNK If yes, provide case ID: _____ Does the patient have a history of infection or colonization with another MDRO? Yes No UNK If yes, specify: _____						
EXPOSURE HISTORY <i>Additional information may be added using the Novel or Emerging Antibiotic Resistant Organisms supplemental investigation form.</i>						
Was the patient admitted to an intensive care unit? Yes No UNK At time of specimen collection, did patient have a tracheostomy tube? Yes No UNK At time of specimen collection, was patient on a ventilator? Yes No UNK Did the patient have a stay in a long-term care facility in the 90 days before specimen collection date? Yes No UNK If yes, facility type: _____ Facility Name: _____ Admission Date: _____ Discharge Date: _____						