Form Approved OMB No. 0920-1385 Exp. Date: 3/31/26

## Antifungal-resistant dermatophytosis case report form

Unique patient ID (DCIPHER):	
ARLN specimen ID:	ARLN isolate ID:   ARLN patient ID:
Form completion data	
Name of person completing this form: Institution: Email: Telephone: Date form completed:	
A. Patient demographics	
<ol> <li>Age at DISC:</li> <li>(use months or days if patient was aged</li> <li>years)</li> </ol>	□ Years □ Months □ Days □ Unknown
2. Sex	□ Male □ Female
2 M/bet is your rose and /or atherists 2	American Indian ar Alacka Nativa
3. What is your race and/or ethnicity? (select all that apply and enter additional details in the spaces provided)	□ American Indian or Alaska Native Enter, for example, Navajo Nation, Blackfeet Tribe of the Blackfeet Indian Reservation of Montana, Native Village of Barrow Inupiat Traditional Government, Nome Eskimo Community, Aztec, Maya, etc.
	□ Asian – provide details below □ Chinese □ Asian Indian □ Filipino □ Vietnamese □ Korean □ Japanese Enter, for example, Pakistani, Hmong, Afghan, etc.
	□ Black or African American – provide details below □ African American □ Jamaican □ Haitian □ Nigerian □ Ethiopian □ Somali Enter, for example, Trinidadian and Tobagonian, Ghanaian, Congolese, etc. ————
	□ Hispanic or Latino – provide details below □ Mexican □ Puerto Rican □ Salvadoran □ Cuban □ Dominican □ Guatemalan Enter, for example, Colombian, Honduran, Spaniard, etc. ————————————————————————————————————
	□ Middle Eastern or North African – provide details below □ Lebanese □ Iranian □ Egyptian □ Syrian □ Iraqi □ Israeli Enter, for example, Moroccan, Yemeni, Kurdish, etc.
	□ Native Hawaiian or Pacific Islander – provide details below □ Native Hawaiian □ Samoan □ Chamorro □ Tongan □ Fijian □ Marshallese Enter, for example, Chuukese, Palauan, Tahitian, etc.
	□ White – provide details below □ English □ German □ Irish □ Italian □ Polish □ Scottish

CDC estimates the average public reporting burden for this collection of information as 30 minutes per response, including the time for reviewing instructions, searching existing data/information sources, gathering and maintaining the data/information 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/ATSDR CDC/ATSDR Information Collection Review Office, 1600 Clifton Road NE, MS H21-8, Atlanta, Georgia 30333; ATTN: PRA (0920-1385).

	Enter, for example, French, Swedi	ish, Norwegian, etc.
4. Patient's country of primary residence		
(e.g., USA)	□ Unknown	
5. Patient's state, jurisdiction, or		
territory of primary residence		Inknown
6. Patient's county of primary residence (Please do not write the word "County"; for example, write "Cook" instead of "Cook County"):	= U	Inknown
7. Patient's city of primary residence		Inknown
8. Patient's ZIP code of primary		
residence		Inknown
9. Patient's type of health insurance at	□ Private □ Medicare □ Medicaid	/state assistance program □ Military □ Indian Health Service
DISC	☐ Incarcerated ☐ Uninsured ☐ Un	
	1	
D. Dationt and onlying viels footone C modic	al aanditiana muaaant dumina tha 2 .	vesus before DISC (vinless of the or time frame and offert)
1. Cancer   Yes   No  Unknown	al conditions present during the 2	years before DISC (unless other timeframe specified)  3. Other immunocompromising conditions   Yes   No   Unknown
☐ Hematologic malignancy		☐ Transplant in the last 2 years
specify type:		☐ Hematologic
☐ Solid organ malignancy		□ Solid organ
specify type:		□ Chemotherapy
		☐ Chronic use of steroids
2. HIV infection ☐ Yes ☐ No ☐ Unknown		☐ Medications/therapies that weaken the immune system
If yes, choose one of the below		☐ TNF-alpha inhibitors (e.g., infliximab, adalimumab,
Ever had CD4 < 200 cells/mm³ within  ☐ Yes ☐ No ☐ Unknown	past 6 months	etanercept) □ Other (specify):
☐ Yes ☐ No ☐ OTIKITOWIT		Li other (specify).
		□ Cirrhosis
4. Other conditions		5. Other potentially relevant underlying conditions?
□ Liver disease		□ Yes (specify below) □ No □ Unknown
□ Cirrhosis		
□ Diabetes	vois	
☐ History of stroke, plegia, paral <sup>,</sup> ☐ Chronic kidney disease	ysis	
□ Chronic respiratory failure		
□ Cardiac disease		
□ Other, specify:	_	
C. Incident specimen data		
1. Date of incident specimen		
collection (DISC)*: (mm-dd-	· <sup>-</sup>	
yyyy)		
*This is the earliest date that a		
patient had a positive test for		
antifungal-resistant		
dermatophytosis		

2. Test type	□ Culture □ PCR
3. Body site	□ Tinea capitis (scalp, hair)
	□ Tinea barbae (beard) or faciei (face)
	□ Tinea manuum (hands)
	□ Tinea unguium (toenails)
	□ Tinea unguium (fingernails)
	□ Tinea genitalis (genitals)
	□ Tinea corporis (other parts of body such as arms or legs), specify:
	□ Tinea cruris (groin, inner thighs, or buttocks)
	□ Tinea pedis (feet)
	□ Other body site specify:
4 Canus and anasias	
4. Genus and species	□ Trichophyton mentagrophytes
	□ Genotype VIII ( <i>T indotineae</i> )
	□ Other genotype, specify:
	□ Unknown genotype
	□ Trichophyton rubrum
	□ Other Trichophyton species
	Species: species unknown
	· — ·
	□ Microsporum
	Species:   species unknown
	a species I species uninown
	□ Epidermophyton
	Species: species unknown
	□ Other genus (specify)
	Species: species unknown
5. Antifungal susceptibility testing	Drug, minimum inhibitor concentration (MIC), mg/L (μg/mL)
testing	Terbinafine (Lamisil)
	Itraconazole (Sporanox)
	Amphotericin B
	Anidulafungin (Eraxis)
	Caspofungin (Cancidas)
	Fluconazole (Diflucan)
	Flucytosine (5FC)
	Ibrexafungerp (Brexafemme)
	Isavuconazole (Cresemba)
	Micafungin (Mycamine)
	Posaconazole (Noxafil)
	Voriconazole (Vfend)
Molecular determinant of	
resistance (e.g., SQLE):	□ Unknown
	1

D. Patient diagnosis and outcomes		
1. Patient location at time of incident	specimen	
collection:		
□ Hospital inpatient	□ Outpatient	□ Long-term care facility (LTCF)
□ Intensive care unit	□ Emergency room	□ Long-term acute care hospital (LTACH)
□ Surgery/OR	□ Clinic/Provider's office (specify)	□ Autopsy
□ Radiology	□ Dermatologist	□ Unknown
□ Other inpatient	□ Infectious Diseases	□ Other

_ F	Podiatrist
_ F	Primary care (adult)
	Primary care (pediatrics)
	Other provider type, specify
	Jnknown provider type
	alysis center
□ Sur	
	gent care
	servational/clinical decision unit
□ Oth	ner outpatient
2. Rash onset date (mm/dd/yyyy)://	
3. Indicate body site(s) affected.	
☐ Tinea capitis (scalp, hair)	
□ Tinea capitis (scaip, hair) □ Tinea barbae (beard)	
☐ Tinea manuum (hands)	
□ Tinea unguium (toenails)	
□ Tinea unguium (fingernails)	
□ Tinea genitalis (genitals)	
☐ Tinea corporis (other parts of body such as arms or legs)	), specify:
☐ Tinea cruris (groin, inner thighs, or buttocks)	
□ Tinea pedis (feet)	
□ Other body site, specify:	
□ Unknown	
Unknown	
1.5 1.6 1.4 11.5 1.4 11.1 2.5 1	
4. Date of most recent follow-up for rash (within 90 days a	after DISC) (mm/dd/yyyy):/
Compared with the patient's rash on DISC, what was the s	tatus of the patient's rash at most recent follow-up?
□ Worse	
□ Neither better nor worse	
☐ Improving, but not fully resolved	
□ Fully resolved	
□ Unknown	

E. Antifungal treatment: Did the patient receive antifungal drugs during the 90 days before to 60 days after the DISC?			
☐ Yes ☐ No ☐ Unknown (If yes, please complete the ta	able below for each drug receiv	ed)	
Systemic antifungals			
Amphotericin B lipid complex (ABLC)	Fluconazole (FLC)	Micafungin (MFG)	Unknown drug (UNK-S)
Liposomal Amphotericin B (L-AmB)	Flucytosine (5FC)	Terbinafine (TRB-S)	
Amphotericin B colloidal dispersion (ABCD)	Griseofulvin (GSF)	Posaconazole (PSC)	
Anidulafungin (ANF)	Ibrexafungerp (IBR)	Voriconazole (VRC)	
Caspofungin (CAS)	Isavuconazole (ISA)	Other systemic drug	
	Itraconazole (ITC)	(specify) (OTH-S):	
Topical antifungals			
Butenafine (BTF)	Econazole (ECZ)	Naftifine (NFT)	Tavaborole (TVB)
Ciclopirox (CPX)	Efinaconazole (EFZ)	Nystatin-	Terbinafine (TRB-T)
Clotrimazole (CTZ)	Ketoconazole (KTC)	triamcinolone (NTC)	Terconazole (TCZ)
Clotrimazole-betamethasone dipropionate (CBM)	Luliconazole (LCZ)	Oxiconazole (OCZ)	Other topical antifungal

	Miconazole (MCZ)	Sertaconazole (STC)	(specify) (OTH-T): Unknown drug (UNK-T)
Drug Abbrev	b. First date given (mm-dd-yyyy)	c. Last date given (mm-dd-yyyy)	e. Therapeutic drug monitoring (TDM)
	□ Start date unknown □ Start date was >60 days before DISC	☐ Still on treatment at time CRF completed Stop date unknown	□ Yes Date of earliest TDM: TDM level:  Date of second TDM: TDM level: □ No
	□ Start date unknown □ Start date was >60 days before DISC	☐ Still on treatment at time CRF completed ☐ Stop date unknown	□ Yes Date of earliest TDM: TDM level:  Date of second TDM: TDM level: □ No

E. Supplemental patient interview form:	
Note that "you" in these questions refers to the p	
Have you traveled internationally during the two years before rash onset?	☐ Yes  If yes, specify country/city/cities/dates:
	□ No □ Unknown
2. Have you had any known exposures to possible ringworm during the month before rash onset?	☐ Yes  If yes, specify country/city/cities/dates: If yes, select all that apply  ☐ Other person with possible ringworm ☐ Animal with possible ringworm If yes, what type of animal? ☐ Cat ☐ Dog ☐ Other, specify: ☐ Environment (e.g., public showers, gyms, shared equipment), specify: ☐ Other, specify: ☐ Other, specify: ☐ Other, specify: ☐ Provide any details of exposure that you might be relevant and are not captured above:
3. How many people are in your household (including yourself) and how many developed signs symptoms of ringworm?	Number of people in the household   Unknown  Number of people in the household who developed possible ringworm   Unknown
4. Did you use topical steroids before this diagnosis?	☐ Yes If yes, name of drug(s), dose(s), duration(s):

5. Did you use topical and/or systemic antibacterial medications before this diagnosis (including those purchased over-the-counter)?*	☐ Yes  If yes, name of drug(s), method(s) of administration (e.g., oral, topical), dose, duration:  ☐ No
6. Over the last week, how itchy, sore, painful, or stinging has your skin been?*	□ Very much □ A lot □ A little □ Not at all
7. Over the last week, how embarrassed or self-conscious have you been because of your skin?*	□ Very much □ A lot □ A little □ Not at all
8. Over the last week, how much has your skin interfered with you going shopping or looking after your home or garden?*	□ Very much □ A lot □ A little □ Not at all □ Not relevant
9. Over the last week, how much has your skin influenced the clothes you wear?*	□ Very much □ A lot □ A little □ Not at all □ Not relevant
10. Over the last week, how much has your skin affected any social or leisure activities?*	□ Very much □ A lot □ A little □ Not at all □ Not relevant
11. Over the last week, how much has your skin made it difficult for your to do any sport?*	□ Very much □ A lot □ A little □ Not at all □ Not relevant
12. Over the last week, has your skin prevented you from working or studying?*	□ Yes □ No  If no, over the last week, how much has your skin been a problem at work or studying? □ A lot □ A little □ Not at all □ Not relevant □ Not at all □ Not relevant
13. Over the last week, how much has your skin created problems with your partner or any of your close friends or relatives?*	□ Very much □ A lot □ A little □ Not at all □ Not relevant
14. Over the last week, how much has your skin caused any sexual difficulties?*	□ Very much □ A lot □ A little □ Not at all □ Not relevant
15. Over the last week, how much of a problem has the treatment for your skin been, for	□ Very much □ A lot

example by making your home messy, or by	□ A little
taking up time?*	□ Not at all
	□ Not relevant
*Questions were adapted from the Dern	natology Life Quality Index (DLQI); approval obtained from DLQI Administrator.
Additional comments:	