

SYPHILIS CASE REPORT

NORTH DAKOTA DEPARTMENT OF HEALTH DISEASE CONTROL SECTION SFN 61082 (02/2021)

The North Dakota Department of Health (NDDoH) Disease Control Section requires the following information to be reported on all syphilis cases. This form shall be used for all newly diagnosed syphilis cases.

Required Patient Demographic Information:

First Name	Last Name			Date	Date of Birth				
Street Address		City			State		ZIP Code		
Telephone Number:	Assigned sex at birth:			ı: [□ Male □ Female				
Current Gender Identity: ☐ Male ☐ Female ☐ Transgender Male ☐ Transgender Female ☐ Transgender Unspecified ☐ Another Gender ☐ Declined to Answer									
Race: ☐ American Indian/Alaskan Native ☐ Asian ☐ Black/Afri☐ Native Hawaiian/Pacific Islander ☐ White ☐ Refused			can American Ethnicity			•	ity: ☐ Hispanic or Latino ☐ Refused		
Pregnancy Status: ☐ Not Pregnant ☐ P	ry Status: ☐ Not Pregnant ☐ Pregnant ☐ NA If Pr				<u>∍gnant,</u>	gnant, Due Date:			
Was case tested for HIV? ☐ Yes ☐ No	HIV? □ Yes □ No If Y			If Yes: Collection Date:			Result: ☐ Positive ☐ Negative		
Specimen Sources (Check All That Apply):	as case tested for Chlamydia? ☐ Yes ☐ No lecimen Sources (Check All That Apply): Urine ☐ Cervix/Vaginal ☐ Rectum ☐ Pharyngeal			If Yes: Collection Date:			Result: □ Positive □ Negative Positive Source(s):		
Was case tested for Gonorrhea? ☐ Yes ☐ N Specimen Sources (Check All That Apply): ☐ Urine ☐ Cervix/Vaginal ☐ Rectum ☐ N	No I	If Yes: Collection Date:				Result: □ Positive □ Negative Positive Source(s):			
Stage of Diagnosis									
What is Patient's Diagnosed Stage of Sy □ Primary Syphilis (Characterized by a □ Secondary Syphilis (Characterized □ Early Syphilis (No symptoms presen □ Latent Syphilis (No symptoms prese	the presence of o by localized or d nt, initial infection	diffuse mucocu n must have o	utaneous lesions (occurred within the	(e.g. rasi he previo	sh), often ous 12 m	nonths)	! lymphadenopathy)		

Current and Past Symptoms

Did the patient have or ever had any of the following symptoms:		Onset Date	Observed By Healthcare Provider	Duration (# of Days)	Additional Description
Chancre	☐ Yes ☐ No	//	□ Yes □ No		
Sore/Lesion	☐ Yes ☐ No	//	□ Yes □ No		
Skin Rash	☐ Yes ☐ No	//	□ Yes □ No		
Alopecia	☐ Yes ☐ No	//	□ Yes □ No		
Condyloma lata	☐ Yes ☐ No	//	□ Yes □ No		
Mucous Patches	☐ Yes ☐ No	//	□ Yes □ No		
Other Manifestations: ☐ Neurological ☐ Ocula	r □ Otic		□ Yes □ No		
Other Symptoms:			□ Yes □ No		

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Provider Information										
Diagnosing HealthCare Provider:										
Facility:				Teleph	one	Number:				
Testing Information										
Reason Test Conducted: ☐ Infection	□ Scree	en 🗆 Partne								
Did case have history of syphilis testir	ng? □ Y	es □ No	If Yes, coll	s, collection date, test type & results:						
Specimen Collection Date:		Tes	ting Labora	tory:						
Testing Note: Need both a non-trepor					•	is.				
Was a non-treponemal (RPR or VDRL) test			vas performed? RPR Titer 1:						
performed? ☐ Yes ☐ No ☐ RRP ☐ VDRL (VDRL (Sou	'20nce: 🗆 Blood 🗇 C2E)			VDRL Titer 1:			
Was a treponemal (ex TPPA) test	If yes, which test was performed?									
performed? □ Yes □ No	☐ TPPA ☐ FTA-ABS ☐ T			=				□ Reactive □ Non-Reactive □ Reactive □ Non-Reactive		
							тер сім. 🗆 г	reactive \Box	NOII-Reactive	
Treatment Information										
Was treatment given for this infection	? □ Ye	s 🗆 No								
Syphilis Treatment:	\ 2.4 mil	lion unite IM	l in a cinala	☐ Doxycycline, 100 mg PO BID * 14 days ngle dose ☐ Doxycycline, 100 mg PO BID * 28 days						
☐ Benzathine penicillin G (Bicillin L-A☐ Benzathine penicillin G (Bicillin L-A☐						Other	_	7 DID ~ 20	uays	
as 3 doses of 2.4 million units IM e				tered	_	other				
Date of First Dose	Date of	Second Dose	?	Date of Third Dose				Treatment Date (if		
of Bicillin L-A:	of Bicillin L-A:			of Bicillin L-A:				prescribed Doxy):		
If not observed, what pharmacy was prescription sent to?										
Was follow up appointments made? [⊐ Yes □] No	If yes, date	e(s) of fo	llow	у-up:				
Did the patient have or ever ha			wing risk	factor	s?					
Does the patient have a history of STI infections?						Yes		□ No		
Is the patient resident/staff of correctional facility?						Yes	[□ No		
Has patient used intravenous/injection drugs?						Yes	[□ No		
Has patient used non-injection drugs?						Yes	[□ No		
Has the patient had sex while high/intoxicated?						Yes		□ No		
Has the patient had sex with an injection drug user?						Yes	[□ No		
Has the patient traded sex for drugs or money?						Yes	[□ No		
Has the patient had sex with an anonymous sex partner?						Yes	[□ No		
Has the patient ever met sexual partners on the internet?						Yes	[□ No		
Total number of sex partners in last 12 months:										
۸	lumber d	of Female Po	artners							
Number of Male Partners										
Number of Transgender Partners				.,			0 1			
What types of sex has the patient had?				□ Vaginal			□ Oral, uns□ Oral, per	•	☐ Anal, unspecified☐ Anal, top☐	
							□ Oral, per		□ Anal, top	
How frequently does the patient use condoms during sex?				□ Always			□ Half the	□ Never		
				☐ Most of the time			□ Not that	often		

Syphilis Partner History *Duplicate Syphilis Partner History form for additional partners *

Sex partners of persons with syphilis are considered at risk for infection and should be confidentially notified of the exposure and need for evaluation. The NDDoH will notify sex partners. Partners who should be notified include those who have had sexual contact within 1) 3 months plus the duration of symptoms with persons diagnosed with **Primary Syphilis**, 2) 6 months plus duration of symptoms with those diagnosed with **Secondary Syphilis** and 3) 1 year with those diagnosed with **Early or Late Latent Syphilis**.

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Partner Name:			Gender Identity: ☐ Male ☐ Female				
			☐ Transgender Male ☐ Transgender Female				
Address:	City	State:	Another Gender				
Address.	City:	State.	Telephone Number:				
Email Address/Phone Apps/Social Media Identifier (ex. Facebook ID):							
Date of First Exposure:		Frequency of Exposure:					
Date of Last Exposure:		Note for Exposure Dates.	Note for Exposure Dates: Include approximate dates if exact date unknown.				
Did confirmed case recall symptoms (i.e. lesions, rash,							
If yes, describe partner symptoms (include date):							
Partner Specimen Collection Date:	Results:						
Partner Treatment:	Treatment Date:						
Partner Name:	Date of Birth or Appro	oximate Age:	<i>Gender Identity:</i> □ Male □ Female				
			☐ Transgender Male ☐ Transgender Female				
Address:	City:	State:	☐ Another Gender Telephone Number:				
ridaress.	City.	State.	receptione (value).				
Email Address/Phone Apps/Social Media Identifier (ex. Facebook ID):							
Date of First Exposure:	Frequency of Exposure:	ency of Exposure:					
Date of Last Exposure:	Note for Exposure Dates	re Dates: Include approximate dates if exact date unknown.					
Did confirmed case recall symptoms (i.e. lesions, rash, etc) on partner? ☐ Yes ☐ No							
If yes, describe partner symptoms (include date):							
Partner Specimen Collection Date:	Results:						
Partner Treatment:	Treatment Date:						
Partner Name:	Date of Birth or Appro	oximate Age:	Gender Identity: ☐ Male ☐ Female				
			☐ Transgender Male ☐ Transgender Female				
Address:	City:	State:	☐ Another Gender Telephone Number:				
Addiess.	City.	State.	retephone Number.				
Email Address/Phone Apps/Social Media Identifier (ex. Facebook ID):							
Date of First Exposure:	Frequency of Exposure:	re:					
Date of Last Exposure:	: Include approximate dates if exact date unknown.						
Did confirmed case recall symptoms (i.e. lesions, rash, etc) on partner? ☐ Yes ☐ No							
If yes, describe partner symptoms (include date):							
Partner Specimen Collection Date:	Results:						
Partner Treatment:	Treatment Date:						

Please Fax Completed Forms to 701.328.0355. Contact NDDoH at 701.328.2378 for any questions.

