National Network of Sexually Transmitted Disease Clinical Prevention Training Centers (NNPTC): Evaluation

TODAY'S DATE	Your confidential ID number is the first two letters of your								
	FIRST name, the first two letters of your LAST name, the	FN	FN	LN	LN	М	М	D	D
M M D D Y Y	MONTH of your birth, and the DAY of your birth.		CON	FIDE	NTIAL	_ IDE	ENTI	FIER	

OMB No. 0920-0995

Attachments 21 & 22

Treatment Guidelines Complete Post-Course Evaluation Instrument

Word version and screenshot

STD Treatment Guidelines Complete Post-Course Evaluation

Public reporting burden of this collection of information is estimated to average 6 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/ATSDR Reports Clearance Officer; 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30333; ATTN: PRA (0920-0995).

S1	1 How satisfied were you with your overall learning experience?													
	very unsatisfied	1	2	3	4	(5)	very satisfied							
S2	How satisfied were	you	with	the c	μalit	y of th	he content?							
	very unsatisfied	0	0	0	0	0	very satisfied							
S3	How satisfied were you with the trainer(s)?													
	very unsatisfied	0	0	0	0	0	very satisfied							
S4	How satisfied were	you	with	the t	each	ing m	ethods?							
	very unsatisfied	0	0	0	0	0	very satisfied							
S5	What could improve	e this	s trai	ningʻ	?									
	As a result of inform setting?	atio	n pre	sente	ed, de	o you	intend to make changes in your practice or at your worksite							
	① Yes													
	© No													
	② Not my job ③ I already use	thes	e nra	ctice	S									
	Other reason		•											



A2 If yes, please list at least one intended change
CE1 Do you believe this activity was influenced by commercial interests?
① Yes ® No
◎ NO
© Yes
No
CE3a Were the learning objectives met?
① Yes
© No
CE3b If the learning objectives were not met, please explain.
The fourthing objectives were not med proude explain
CHLAMYDIA
LOC2bef How confident were you in your ability to describe the current CDC screening recommendations for
chlamydia, including extra-genital screening BEFORE the training?
not at all confident O O O O very confident
not at all confident. O O O Very confident
LOC2aft How confident are you AFTER the training?
not at all confident OOOO very confident
LOC4bef How confident were you in your ability to treat patients diagnosed with chlamydia and related
anogenital syndromes based on the most current CDC treatment recommendations BEFORE the training
not at all confident OOOOvery confident ONA
LOC4aft How confident are you AFTER the training?
not at all confident O O O O very confident O NA
Thot at all confident O O O O Very confident O NA
 What is the CDC recommended regimen for treating asymptomatic uncomplicated chlamydia infection of the cervix, urethra, or rectum? O Acyclovir 1 g twice a day for 7 days O Azithromycin 1 g orally in a single dose or doxycycline 100 mg twice a day for 7 days O Azithromycin 1 g orally in a single dose plus ceftriaxone 250 mg intramuscularly in a single dose
O Ciprofloxacin 500 mg orally in a single dose
KC2 What is the recommended follow-up for a non-pregnant patient after diagnosis and treatment of
chlamydia?
① A test of cure at 2 weeks, and repeat test at 3 months
② A test of cure at 2 weeks, and repeat test at 12 months
Repeat test in 3 months
Repeat test in 12 months
PPC1bef Approximately what % of sexually active asymptomatic female patients under age 25 did you screen
annually for chlamydia BEFORE this training?

NA

0% 1-25% 26-50% 51-75% 76-90% >91%



\cup	U	(9	4)	9		©	77						
PPC1aft	What	t % do y	ou inte	nd to so	reen A	FTEF	R the	training?						
	0%	1-25%	26-50%	51-75%	6 76-90)% >!	91%	NA						
	0	0	0	0	0		0	0						
							G	ONORRHEA						
L OC2bef	Llow	oonfido	nt word	vallin i	vour ok	ailita e	+0 d	accribe the curren	t CDC	ooroonin	a recommendations			
LOGZDei								ning BEFORE this			g recommendations			
	not	at all co	onfident	0 0	0	0	0	very confident						
LOG2aft	Hov	w confid	dent are	you AF	TER th	e tra	ining	g?						
	not	at all co	nfident	0 0	0	0	0	very confident						
LOG4bef	LOG4bef How confident were you in your ability to treat patients with gonorrhea according to current CDC recommendations in light of antibiotic resistance in <i>N. gonorrhoeae</i> BEFORE this training?													
	not	at all co	nfident	0 0	0	0	0	very confident	0	NA				
LOG4aft	Hov	w confid	dent are	you AF	TER th	e tra	ining	g?						
	not	at all co	nfident	0 0	0	0	0	very confident	0	NA				
KG2 W	or rectal gonorrhea? ① Ceftriaxone 250 mg intramuscularly only ② Azithromycin 2 g orally in a single dose only ③ Ceftriaxone 250 mg intramuscularly plus azithromycin 1 g orally in a single dose ④ Cefixime 400 mg orally plus doxycycline 100 mg orally BID for 7 days KG2 What is the recommended follow-up for a non-pregnant patient after diagnosis and treatment of gonorrhea? ① A test of cure at 2 weeks, and repeat test at 3 months ② A test of cure at 2 weeks, and repeat test at 12 months ③ Repeat test in 3 months ④ Repeat test in 12 months													
PPG1bef									le patie	ents unde	er 25 did you screen			
		•	r gonorr					ng? NA						
	0% ①	1-25% ②	20-50% ③	51-75% ④	% 76-90 ⑤		91% ©	77						
				_	_									
PPG1aft			you inte 26-50%					e training? NA						
	O	0	0	0	0		0	Ö						
	annu 0% O What	ually for 1-25% O % do y	r uroger 26-50% O	nital and 51-75% O nd to sc	d extrag 6 76-90 O reen A	genita)% >! FTER	al go 91% O the	ale patients who honorrhea and chlar NA O training?			en did you screen this training?			
	Ö	0	0	0	O		0	Ö						



syphilis BEFORE this training?													
	not	at all co	nfident	0 0	0	0	0	very confident					
LOS2aft	How	confide	ent are y	ou AF1	ER the	traini	ing?						
	not	at all co	nfident	0 0	0	0	0	very confident					
LOS3bef			nt were :					er and interpret th ng?	he CDC	crecomn	nended s	serologic	
	not	at all co	nfident	0 2	3	4	⑤	very confident	77	NA			
LOS3aft	How o	confider	nt are yo	ou AFTE	ER the t	rainin	ng?						
	not	at all co	nfident	0 0	0	0	0	very confident	0	NA			
LOS4bef How confident were you in your ability to clinically manage patients diagnosed with syphilis based on CDC treatment, follow-up, and partner management recommendations BEFORE this training?													
	not	at all co	nfident	0 0	0	0	0	very confident	0	NA			
LOS4aft How confident are you AFTER the training?													
			nfident				o	very confident	0	NA			
PPS1bef	① Acy ② Azii ③ Bei ④ Ber	thromyc nzathine nzathine oximatel	g twice in 1 g or e penici	ally in a Ilin G 2. e penici % of yo	single d 4 millio Ilin 2.4 n ur male	ose n uni nillion	units	in a single dose s in a single dose who have sex with	h men	did you s	screen at	t least onc	ce a year
			26-50%					NA					
	0	0	0	0	0	C		0					
PPS1aft			you inte 26-50%					training?					
	0% O	1-25% O	20-50% O	51-75%	0 76-909 O	0 /9.		NA O					
	_								_				_
PPS2bet			ly what ' 26-50%					nts did you scree NA	en for s	yphilis B	SEFORE 1	this trainii	ng?
	O	0	0	0	0	_	D	Ö					
PPS2aft			26-50%					training? NA					
	0	0	0	0	0	· C		Ö					
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PPS3Der			iy what training		ur patie	nts re	eceni	tly diagnosed wit	n sypn	ilis ala y	ou test i	or HIV	
		1-25%	26-50%		6 76-909			NA					
	0	0	0	0	0	C)	0					
PPS3aft								•					
	0% O		26-50% O	51-75% O	6 76-909 O		1%)	NA O					
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EPT As a result of the information presented do you intend to provide Expedited Partner Therapy (EPT) to heterosexual partners of those diagnosed with gonorrhea and/or chlamydia?



- ① Yes
- @ No
- ② Not applicable to my practice or job
- Not allowed in my state/practice
 My practice/worksite is in the planning stages to offer EPT
 My practice/worksite already offers EPT
- © EPT was not discussed