Mother's Name:		Chart No.:	Dhara		
Notrier's Name:	t Delivering		Phone No.: ()	
Infant's Name: Cha	.:: Ph	nysician:	Phone No.: ()	
Pediatrician: — Patient identifier information is not transmitted to CDC –					
DEPARTMENT OF HEALTH & HUMAN SERVICES CENTERS FOR DISEASE CONTROL and PREVENTION INVESTIGATION AND REPORT CONGENITAL SYPHILIS (CS) CASE INVESTIGATION AND REPORT					
PART I. MATERNAL INFORMATION	m Approved OMB No. 09	•	Talan is a second		
1. Report date to health dept. Mo. Day Yr. (8-15)	2. Reporting state FIPS code:	Unl	l	Unk	
4. Reporting city FIPS code:	, ,			6. Country of residence: (leave blank if USA)	
(21-24) Reporting City Name	(25-27)		(28-30) Countr	(28-30) Country of Residence	
7. State FIPS code:	8. Residence county FIPS code:		9. Residence city FIPS code:	Unk	
	<u> </u>				
(31-32) Residence State Name 10. Residence zip code: Unk 11. Mother's date of birth:	(33-35) R Unk 12. Mother's ethni	esidence County Name	(36-39) Re race: (Code 1 for all that apply)	sidence City Name	
///	1 Hispanic or Latino 9 Unk (54) American Indian/Alaska Native (55) Asian (56) Black or African American				
14. Mother's marital status: (60) 1 Separated/ 8 Other	15. Last menstrual period (LMP)	(before delivery) Unk	16. Did mother have prenatal care? (69 1 ☐ Yes 9 ☐ Un	⁹⁾ k <i>(Go to Q19)</i>	
2 Married 4 Widow 9 Unk	/ /		2 No (Go to Q19)	K (40 10 Q 19)	
17. Indicate date of first prenatal visit:	18. Indicate number of prenatal visits:			19. Did mother have a nontreponemal test (e.g., RPR or VDRL) in pregnancy, at delivery, or soon after delivery within 3 days? (80)	
/ / /	(78-79)		1 _ " _ "	9 Unk <i>(Go to Q21)</i>	
20. Indicate dates and results of nontreponemal tests: (list the most recent first) Date Results		21. Did mother have confirmatory trepo (e.g., FTA-ABS or TP-PA)? (133)(Foot			
Mo. Day Yr. a//(81-88) □ Unk 1□ Reactive 2□ Nonreactive 9□ Unk (89) 1: (90-93)			1 Yes, reactive 3 No		
b/			2 Yes, nonreactive 9 Unk 22. Did mother have darkfield or direct flu		
c/			exam of lesions at delivery? (134) (F	ootnote a)	
d/			Yes, positive 3 No te	st of lesions 9 Unk sions present	
23. Before this delivery, when was mother last treated for syphilis? , , Before pregnancy, was mother's treatment adequate? (144) (Footnote b)					
1 Defore pregnancy (Go to Q24) 2 During pregnancy (Go to Q25) 3 No Treatment (Go to Q27) 9 Unk (Go to Q27) 2 No, inadequate (Go to Q27)					
25. During pregnancy, was mother's treatment adequate? (145) (Footnote b) 26. An appropriate serologic response.				priate response: evidence of	
1 ☐ Yes, adequate 3 ☐ No, inadequate: penicillin therapy begun (Go to Q27) 1 ☐ Yes, appropriate response values of the properties of the		ring pregnancy No response was equivocal or could			
2 No, inadequate: non-penicillin therapy (Go to Q27) 4 Unknown (Go to Q27) 4 No, response was equivocal or not be determined from available follow-up serologic titers during pregnancy nontreponemal titer information			mined from available		
PART II. INFANT INFORMATION 27. Date of Delivery:	28. Vital status: (155)	Stillborn (Go to Q31)	29. Indicate date of death	Unk	
1		3 ☐ (Footnote d)	//		
Mo. Day Yr. (147-154) 30. Gender: (164)	31. Birthweight (in grams)		Mo. Day Yr. 32. Estimated gestational age (in weeks	(156-163) Unk	
1 Male 2 Female 9 Unk		(If infant was	stillborn go to Q42)		
33. a)Did infant/child have a reactive b) When was the infant/child's first reactive c) Indicate titer of infant/child's 34. a) Did infant/child have a reactive b) When was the infant/child's first reactive b) When was the infant/child was reactive by when was t					
non-treponemal test for syphilis? first reactive non-treponemal syphilis (e.g., VDRL, RPR)?(171) (172- test for syphilis? (e.				reponemal test for syphilis? (185- (192) (192)	
35. Did the Infant/child have any classic signs of CS? (193) Laboratory Confirmation 36. Did the Infant/child have a		37. Did the infant/child have an IgM-spe	cific treponemal test? (195		
(Footnote e) 1 ☐ Yes 2 ☐ No, asymptomatic 9 ☐ Unk. infant/child	1 Yes, positive 3 No test 2 Yes, negative 9 Unk.		(Footnote f) 1 ☐ Yes, reactive 2 ☐ Yes, nonreactive	3 ☐ No test 9 ☐ Unk.	
Infant/Child Evaluation 38. Did the infant/child have long bone X-rays? 39. Did the infant/ CSF-VDRL?(1s	100, Will				
Yes, changes 1 Yes, changes 1 No xrays 1 Yes, reactive 2 Yes, no signs of CS 9 Unk. 2 Yes, reactive 2 Yes, no signs of CS	re 3 □ No test Yes, one 1 □ or both elevated 3 □ No test 2 □ Yes, with other treatment by Aqueous or Procaine 1 □ Yes, with other treatment 5 □ No treatment				
PART III. Congenital Syphilis Case Classification 42. Classification (200)					
1 Not a case 2 Confirmed case (Laboratory confirmed identification of <i>T.pallidum</i> , e.g., darkfield or direct fluorescent antibody positive lesions) 2 Syphilitic stillbirth (Footnote d) 3 Syphilitic stillbirth (Footnote d)					
Public reporting burden of this collection of information is estimated to average 30 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the					

Public reporting burden of this collection of information is estimated to average 30 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, MS D-24, Atlanta, GA 30333, ATTN: PRA (0920-0128), Do not send the completed form to this address.