



# ESP CASE DETECTION ALGORITHM

## Active Syphilis

Document Version 2.22

Prepared by the Department of Population Medicine at Harvard Medical School and Harvard Pilgrim Health Care Institute on behalf of the Massachusetts Department of Public Health.

[esphealth@harvardpilgrim.org](mailto:esphealth@harvardpilgrim.org)

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### Modification History

Version	Date	Modification	By
2.22	10/11/2017	Added ESP Logo and MDPH branding	DPM
2.21	7/6/2017	Transferred to updated algorithm template	DPM
2.2	2/22/2017	<ul style="list-style-type: none"><li>Required all positive RPR values to contain a titer value.</li><li>Modified time window in which to report previous reportable lab results.</li></ul>	DPM/MDPH
2.1	11/10/2016	Updated recurrence interval from 1 year to 6 months.	DPM/MDPH
2.0	5/24/2016	Added appropriated ICD-10-CM diagnosis codes.	DPM/MDPH
1.0	4/17/2013	Original circulated version.	DPM/MDPH



## Section 1. Overview

The purpose of this document is to describe the criteria used to identify and report cases of Active Syphilis from electronic medical records (EMR) using ESP. All patients who meet the criteria will be reported to the Massachusetts Department of Public Health (MDPH). In addition, supplementary information (e.g., demographic information) will be included with the initial case report, as described in [Section 3](#).

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## Section 2. Criteria used to identify cases using ESP data

### I. CASE TYPES

This document includes an algorithm to identify active syphilis.

### II. TIME WINDOW

The recurrence window for syphilis is 6 months (180 days).

### III. CASE CRITERIA

Active syphilis defined as meeting criteria for any one of the following:

1. (Diagnosis code (ICD-9 Code 090-097 or ICD10 A50-A53) or TP-IGM) AND an order/prescription for at least ONE of the following antibiotics within a 14 day period:
  - a. "PENICILLIN G" or "PEN G"
  - b. "DOXYCYCLINE" for  $\geq 7$  days<sup>1</sup>
  - c. "CEFTRIAXONE" dose  $\geq 1$  gram<sup>2</sup>
2. Serum RPR or VDRL<sup>3</sup> value greater than or equal to 1:8<sup>4</sup> and any of the following:
  - a. Lab test TPPA with result "reactive" ever in the past and up to 1 month following the positive RPR or VDRL OR
  - b. Lab test FTA-ABS with result "reactive" ever in the past and up to 1 month following the positive RPR or VDRL OR
  - c. Lab test TP-IGG with result "positive" or "reactive" ever in the past and up to 1 month following positive RPR or VDRL
3. Positive CSF test for syphilis. Any of the following:
  - a. VDRL-CSF value "reactive" or  $\geq 1:1$ <sup>4</sup>
  - b. TPPA-CSF with result "reactive" or "positive" or equivalent
  - c. FTA-ABS-CSF with result "reactive" or "positive" or equivalent

### IV. NOTES

1. If doxycycline prescription does not include a specified duration  $\geq 7$  days or a stop date  $\geq 7$  days following the prescription date then a "quantity" field result of  $\geq 14$  is an adequate proxy for  $\geq 7$  days of doxycycline.



2. If dose value for ceftriaxone is not populated it is sometimes possible to discern the minimum dose by looking at the drug name. For example, in Atrius Health there are some preparations of ceftriaxone that include the terms “1G” or “2G” within the drug name in addition to the word “CEFTRIAZONE”.

Examples:

- a. ROCEPHIN VIAL 2G IJ (CEFTRIAZONE SODIUM)
- b. ROCEPHIN VIAL 1G IJ (CEFTRIAZONE SODIUM)
3. VDRL is distinct from VDRL-CSF. We must ensure that only titer ratios are evaluated. Other result values such as “positive” or “reactive” are ignored.
4. RPR and VDRL results are expressed as dilutions (e.g. 1:2, 1:4, 1:8, etc.). The higher the number in the denominator the more ‘positive’ the test result. An RPR  $\geq$  1:8 means a value of (1:8, 1:16, 1:32, etc.). RPR results of “positive” without a titer value should not be considered positive.

### Section 3. Specifications for reporting diseases/conditions to the Massachusetts Department of Public Health

#### I. INITIAL CASE REPORTING CRITERIA

All cases should be immediately reported to MDPH upon initial detection.

#### II. CASE REPORT UPDATE CRITERIA

Any serum RPR or CSF VDRL results that arise following case identification should be sent to MDPH. This should continue indefinitely.

#### III. DATA TO INCLUDE IN REPORTS TO MDPH

##### A. Demographic

Name	Last, first, middle
Date of birth	yyyy/mm/dd
Age	
Social security number	Last 4 digits
Gender	Male / Female
Race	American Indian / Asian / Black / White / Other / Unknown
Ethnicity	Hispanic / Non-Hispanic / Unknown
Address	Line 1, line 2, city, state, zip, country
Phone	xxx-xxx-xxxx
Language spoken	
Medical record number	
PCP	Name, office address, phone number, email

##### B. Ordering Facility Information

Facility name	
Facility address	Line 1, line 2, city, state, zip
Facility contact person	
Contact person email	
Contact person phone	xxx-xxx-xxxx



Facility phone	XXX-XXX-XXXX
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**C. Encounter Data**

Lab test ordering clinician	Name, office address, phone number, email
Treating clinician	Name, office address, phone number, email

**D. Laboratory Results**

Laboratory results occurring within 7 to 0 days prior to the case date should be reported.  
See Section 4 for LOINC mappings.

RPR	Specimen source, value, date
TPPA	Specimen source, value, date
FTA-ABS	Specimen source, value, date
VDRL-CSF	CSF, value, date
TP-IGM	Specimen source, value, date
TP-IGG	Specimen source, value, date
TPPA-CSF	CSF, value, date
FTA-ABS-CSF	CSF, value, date

**E. Pregnancy**

Pregnancy flag active	Yes or No
Expected Date of Delivery (EDD)	

**F. Disease Stage**

If an ICD-9 or ICD-10 diagnosis code indicating disease stage occurs within 28 days of case being established, then disease stage is reported. See Section 4 for specific diagnosis codes.

**G. Clinical Diagnosis**

If an ICD-9 or ICD-10 diagnosis code indicating a clinical diagnosis for syphilis occurs within 28 days of case being established, then the precise ICD-9/ICD-10 code and ICD-9/ICD-10 text is reported. See Section 4 for specific diagnosis codes.

**H. Treatment**

If patient is prescribed medication for syphilis treatment within 28 days before or after case established, then report the date, medication, dose, route, and duration of treatment. See Section 4 for specific medications.

**IV. CRITERIA TO REVOKE A CASE**

None.



**Section 4. Codes, laboratory tests, and medications used to identify criteria listed in Section 2 and supplementary reporting information in Section 3.**

**I. DIAGNOSIS CODES**

**Table 1. Diagnosis Codes Used to Identify Disease Stage**

Code Type	Code	Description
ICD-9-CM	091.0 to 091.2	Primary
ICD-9-CM	091.3 to 091.9	Secondary
ICD-9-CM	092.x	Early latent
ICD-9-CM	096.x	Late latent
ICD-9-CM	094.x	Neurosyphilis
ICD-9-CM	090.0 to 090.4, 090.9	Congenital (infant)
ICD-9-CM	090.5 to 090.7	Late congenital
ICD-10-CM	A50*	Congenital
ICD-10-CM	A51.0 to A51.2	Primary
ICD-10-CM	A51.3 to A51.4	Secondary
ICD-10-CM	A51.5	Early syphilis, latent
ICD-10-CM	A51.9	Early syphilis, unspecified
ICD-10-CM	A52.0	Cardiovascular syphilis
ICD-10-CM	A52.1 to A52.3	Neurosyphilis
ICD-10-CM	A52.7	Other symptomatic late syphilis
ICD-10-CM	A52.8	Late syphilis, latent
ICD-10-CM	A52.9	Late syphilis, unspecified
ICD-10-CM	A53.0	Latent syphilis, unspecified as early or late
ICD-10-CM	A53.9	Syphilis, unspecified

**Table 2. Diagnosis Codes Used to Identify Clinical Diagnosis**

Code Type	Code	Description
ICD-9-CM	090.x to 097.x	Syphilis
ICD-10-CM	A51 to A53	Syphilis

**II. LABORATORY TEST LOINCS**

**Table 3. Laboratory Test LOINC Mapping**

Test Name	LOINC	LOINC Name
TPPA	11597-2	Treponema pallidum Ab : ACnc : Pt : Ser : Qn
RPR	20507-0	Reagin Ab : ACnc : Pt : Ser : Ord : Rapid test
VDRL	20507-0	Reagin Ab : ACnc : Pt : Ser : Ord : Rapid test
FTA-ABS	34147-9	Treponema pallidum Ab.IgG+IgM : ACnc : Pt : Ser : Ord :
TP-IGG	22592-0	Treponema pallidum Ab.IgG : ACnc : Pt : Ser : Qn :
VDRL-CSF	5290-2	Reagin Ab presence, VDRL, CSF

**III. MEDICATIONS**

**Table 4. Syphilis Medications**



Generic Name
Penicillin G or "Pen G"
Ceftriaxone
Doxycycline

#### IV. CODE MAINTENANCE STRATEGY

Continuously screen all incoming Lx\_Component\_Name fields for the text strings: "SYPH" or "RPR" or "VDRL" or "TP" or "FTA" or "TREP" or "PALLIDUM".