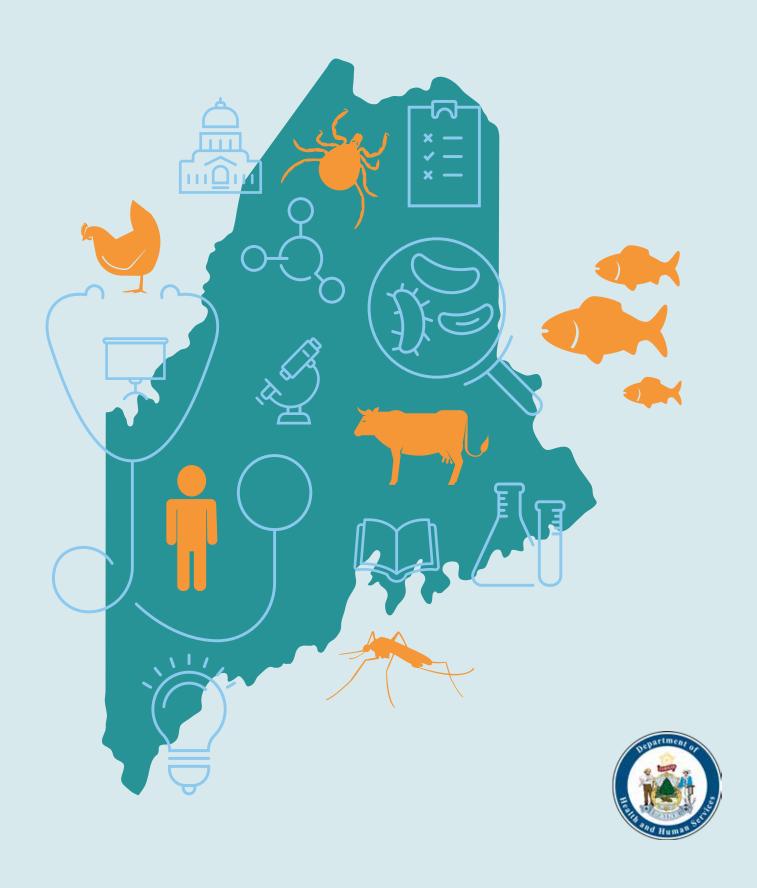
MAINE REPORTABLE INFECTIOUS **DISEASES SUMMARY**





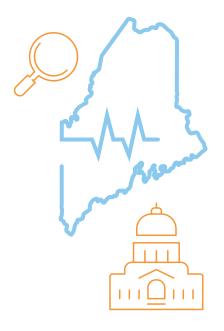
Reportable Infectious Diseases in Maine 2018 Summary

Editors: Sara Robinson, MPH, Catie Peranzi, MPH Contributors: Division of Disease Surveillance

2019 Maine Center for Disease Control and Prevention 286 Water Street State House Station 11 Augusta, ME 04333-0011 www.maine.gov/idepi 800-821-5821



THANK YOU



Maine Center for Disease Control and Prevention (Maine CDC) annually publishes a report on infectious diseases in Maine. This report is prepared by the Division of Disease Surveillance and is intended to provide an overview of notifiable infectious diseases of public health importance in Maine.

We could not produce this report without the continued support of our healthcare and public health partners throughout the state. We greatly appreciate all of the laboratories, healthcare providers, childcare centers, school nurses, veterinarians, and others who provide disease surveillance information. Partners spend considerable time assisting Maine CDC with infectious disease investigations and disease control measures that affect Maine residents. Public health partners' active and critical role in the infectious disease surveillance cycle informs statewide policies and programs that protect our residents from infectious diseases through health promotion, disease prevention, early detection, containment, and treatment.

We appreciate and encourage your vigilance in the effort to protect the people of Maine through timely, complete, and accurate notifiable infectious disease reporting. It is through these collaborative efforts that we are able to respond to emerging infectious disease threats and prevent outbreaks.

We hope you find this report useful as we all work to protect and promote the health of Maine's residents. As always, we welcome your feedback on how we can provide more useful disease information to you, our partners.

For more information on what, when, and how to report infectious diseases please see the Notifiable Diseases and Conditions List (Page 73) of this report, visit our website at www.maine.gov/idepi or call 1-800-821-5821.

Ann Farmer, MS

Associate Director, Division of Disease Surveillance

Maine Center for Disease Control and Prevention

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2018 INFECTIOUS DISEASE

SURVEILLANCE HIGHLIGHTS

4,879

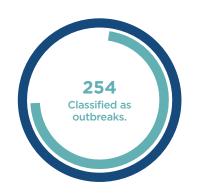
Disease reports investigated by Maine CDC staff including STD cases.

Potential outbreaks investigated by Maine CDC staff.

9,757

Disease reports handled without a full investigation by staff, either through passive surveillance or laboratory reports.*







The main diseases include chlamydia, hepatitis C, Lyme disease, rabies post-exposure prophylaxis, and some varicella and invasive MRSA cases.

People with hepatitis C identified as at risk for hepatitis B.



Maine CDC initiated contact with providers to recommend vaccination.



Maine CDC sponsored a hepatitis A vaccine clinic in response to a confirmed case of hepatitis A in a person experiencing homelessness.

Received a hep A vaccine

Reported cases of infectious syphilis in 2018

Increase from 2017





Bangor is the farthest north Maine detected WNV in a mosquito pool

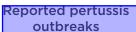
Influenza-related outbreaks during the 2018-2019 flu season

10,300+

Positive influenza reports

Influenza-related hospitalizations

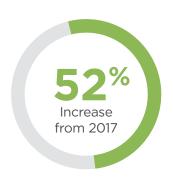
Influenza-related deaths





Animals tested positive for rabies at HETL

Reported cases of Group A Streptococcus



2018 Maine CDC Infectious Disease Program consults



Rabies (32%), influenza, pertussis, tuberculosis, and varicella

of all consults occurred in August

COUNTS OF SELECTED* REPORTABLE DISEASES BY YEAR

MAINE, 2009-2018**

CONDITION	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018
Anaplasma phagocytophilum										
Babesiosis										
Brucellosis										
Campylobacteriosis										
Chikungunya Virus										

NR = not reportable; NA = not available

CONDITION	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018
Listeriosis										
Lyme disease										
Malaria										
Measles (Rubeola)										
Mumps										
										1
Novel Influenza A virus Infections										
Q fever										1
West Nile										

* Maine did not have any cases of the following reportable conditions in the last ten years:

Anthrax

Leptospirosis

Smallpox

Botulism

Plague

Saint Louis Encephalitis

California Serogroup viruses

Polio

Tularemia

Chancroid

 Psittacosis Rabies, human Viral Hemorrhagic Fever

Coronavirus

 Western Equine Encephalitis Yellow Fever

 Diphtheria Hepatitis D, chronic Ricin Rubella

** Counts are updated annually. Data as of 6/5/2018

*** CRE became reportable as of September 8, 2015 so the 2015 numbers do not represent a full year

RATES OF SELECTED* **REPORTABLE DISEASES BY YEAR**

MAINE, 2009-2018** (PER 100,000 PERSONS)

CONDITION	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018

CONDITION	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018
Listeriosis										
Lyme disease										
Malaria										
Measles (Rubeola)										
Mumps										
Neisseria meningitidis, invasive (Mening. disease)										
Novel Influenza A virus Infections										
Pertussis										
Powassan										
Q fever										
Rabies PEP										
Rabies, animal										
S. aureus, methicillin resistant (MRSA), invasive										
S. aureus, vancomycin intermediate resistance (VISA)										
Salmonellosis										
Shellfish poisoning										
Shiga toxin-producing <i>Escherichia coli</i> (STEC)										
Shigellosis										
Spotted Fever Rickettsiosis										
Syphilis										
Tetanus										
Trichinosis (Trichinellosis)										
Tuberculosis										
Varicella (Chickenpox)										
Vibriosis										
West Nile										

* Maine did not have any cases of the following reportable conditions in the last ten years:

Anthrax

Botulism

California Serogroup viruses

Chancroid

Coronavirus

Diphtheria

Hepatitis D, chronic

Leptospirosis

Plague

Polio

Psittacosis

Rabies, human

Ricin

Rubella

Smallpox

Saint Louis Encephalitis

Tularemia

Viral Hemorrhagic Fever

Western Equine Encephalitis

Yellow Fever

** Counts are updated annually. Data as of 6/5/2018

*** CRE became reportable as of September 8, 2015 so the 2015 numbers do not represent a full year

CASES OF REPORTED DISEASES **BY AGE AND GENDER**

MAINE, 2018

CASES OF REPORTED DISEASES BY RACE AND ETHNICITY

MAINE, 2018

		RACE		E	ETHNICIT	Υ
CONDITION						
Anaplasma phagocytophilum						
Babesiosis						
Borrelia miyamotoi						

2018 MAINF OUTBREAKS

Outbreaks are a reportable condition in Maine and are classified into types of outbreak by the potential etiology. All reported outbreaks are assigned out for follow-up with a field epidemiologist. This table only represents those that met an outbreak definition of confirmed, probable, or suspect. Outbreak definitions vary based on the category, setting, and suspected etiology.

OUTBREAK CATEGORIES AND DEFINITIONS

Absenteeism: Absenteeism reports are submitted by schools when they have ≥15% absenteeism due to illness. If there is a single etiology an absenteeism report may also be counted as a disease-specific outbreak.

Airborne and Direct Contact (ADC): Airborne and Direct Contact outbreaks are transmitted through airborne bacteria or viruses or through direct contact. Examples of Airborne and Direct Contact outbreaks include pneumonia, conjunctivitis, hand foot and mouth disease, and MRSA.

Gastrointestinal Illness (GI): GI illness outbreaks are characterized through gastrointestinal symptoms. The most commonly reported GI outbreak is caused by norovirus. Out of state GI outbreaks are when a Maine resident matches a national cluster through Pulsed-field Gel Electrophoresis (PFGE) testing such as Salmonella or Shiga toxin producing E. coli (STEC).

Influenza-like Illness (ILI): Influenza-like illness outbreaks are characterized as a respiratory illness with fever and/or sore throat without another known cause. The majority of ILI outbreaks are confirmed as influenza through laboratory testing.

Other: Outbreaks in this category are not captured in any other group. Examples include C. difficile, multi-drug resistant organisms, or outbreaks caused by contaminated devices.

Vaccine-Preventable Disease (VPD): Vaccine-preventable disease outbreaks are caused by one of the illnesses for which there is a routine vaccine. The most commonly reported VPD outbreak is caused by pertussis. Pertussis was the cause of all VPD outbreaks in 2018.

Varicella: Varicella (chickenpox) outbreaks are caused by chickenpox. An outbreak is defined as three or more confirmed cases in a single setting.

Vector: Vector outbreaks are caused by an organism that spreads infection from one host to another. The most common vectors in Maine are ticks and mosquitoes, but the most common vector outbreak is caused by scabies.

	Androscoggin	Aroostook	Cumberland	Franklin	Hancock	Kennebec	Knox	Lincoln	Out of State	Oxford	Penobscot	Piscataquis	Sagadahoc	Somerset	Waldo	Washington	York	Total
Absenteeism	1	0	1	0	4	1	1	2	0	1	0	0	0	3	5	2	1	22
ADC	1	0	2	0	0	0	0	0	0	0	1	0	0	0	1	0	1	6
Gl	10	2	19	0	0	7	4	2	11	0	4	0	0	1	0	0	9	69
ILI	10	6	34	1	2	11	9	1	0	8	8	0	4	6	0	4	23	127
Other	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	1
VPD	1	0	3	0	4	0	0	0	0	0	1	0	1	0	2	0	9	21
Varicella	0	0	3	0	0	0	0	0	0	0	0	0	0	0	0	0	3	6
Vector	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	1	0	2
Total	23	8	62	1	10	19	14	5	11	9	15	0	5	10	8	7	47	254

FILI outbreaks included here are for the calendar year 2018, so includes outbreaks from the 2017-2018 and 2018-2019 influenza seasons. Any outbreak can be healthcare associated.

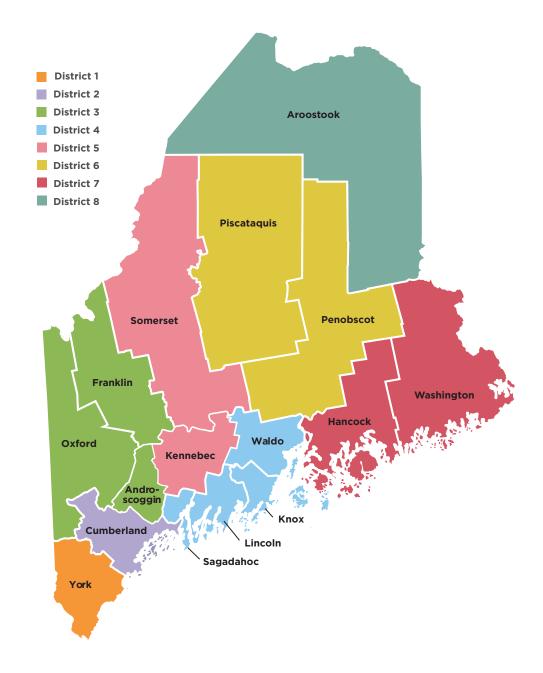
PUBLIC DISTRICT HEALTH MAP

Since 2003, the Infectious Disease Programs of Maine CDC publish an annual summary of infectious disease data. Publishing reports on surveillance activities and data provides the health care community, government agencies, individuals, and groups with important statistical information on Maine's reportable diseases and conditions.

This annual report also includes information on conditions that are investigated that are not explicitly reportable but have public health significance. Examples of these conditions include Borrelia miyamotoi, Jamestown Canyon virus and Zika virus. Maine also follows up on unusual conditions that may not have specific case definitions but potentially have public health significance. These conditions are indicated by "Emerging Infections." In 2018 the emerging infection was a report of alpha-gal allergy that Maine CDC is monitoring. The goal of this annual report is to provide Maine CDC's partners with a helpful resource.

Maine CDC counts cases by their residence, not where they acquired the condition.

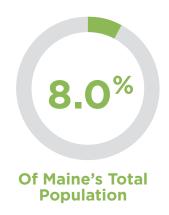
(Population data is from 2018 census estimates.)



ANDROSCOGGIN COUNTY







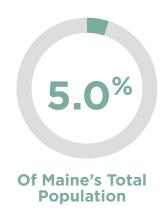
	Cou	ınty	Dist	trict	State			
Condition	Count	Rate	Count	Rate	Count	Rate		
Anaplasma phagocytophilum	31	28.8	48	24.6	476	35.6		
Babesiosis	9	8.4	11	5.6	101	7.5		
Borrelia miyamotoi	0	0.0	0	0.0	8	0.6		
Campylobacteriosis	16	14.9	30	15.4	247	18.5		
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	5	4.6	14	7.2	92	6.9		
Chikungunya Virus	0	0.0	0	0.0	2	0.1		
Chlamydia trachomatis infection	569	528.4	807	413.4	4346	324.7		
Coccidioidomycosis	0	0.0	0	0.0	2	0.1		
Cryptosporidiosis	0	0.0	1	0.5	60	4.5		
Dengue	1	0.9	1	0.5	3	0.2		
Ehrlichiosis, <i>chaffeensis</i>	3	2.8	3	1.5	19	1.4		
Ehrlichiosis/Anaplasmosis, undetermined	1	0.9	2	1.0	9	0.7		
Emerging Infection	0	0.0	0	0.0	1	0.1		
Giardiasis	10	9.3	22	11.3	163	12.2		
Gonorrhea	238	221.0	263	134.7	686	51.3		
Group A <i>Streptococcus</i> , invasive	14	13.0	21	10.8	85	6.4		
Haemophilus influenzae, invasive	2	1.9	5	2.6	24	1.8		
Hepatitis A, acute	2	1.9	2	1.0	9	0.7		
Hepatitis B, acute	1	0.9	1	0.5	52	3.9		
Hepatitis B, chronic	25	23.2	31	15.9	201	15.0		

	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	3	2.8	4	2.0	39	2.9
Hepatitis C, chronic	124	115.2	200	102.5	1872	139.9
HIV	1	0.9	2	1.0		2.2
Invasive Pneumococcal Disease	10	9.3	17	8.7	132	9.9
Jamestown Canyon virus	0	0.0			1	0.1
Legionellosis	3	2.8	7	3.6	34	2.5
Listeriosis	1	0.9	2	1.0	7	
Lyme disease	68	63.2	128	65.6	1405	105.0
Malaria	2	1.9	2	1.0	9	0.7
Mumps	1	0.9	1	0.5	4	0.3
Neisseria meningitidis, invasive (Mening. disease)	1	0.9	1	0.5	1	0.1
Pertussis	25	23.2	34	17.4	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	5	4.6	11	5.6	152	11.4
Rabies, animal	7	NA	15	NA		NA
S. aureus, methicillin resistant (MRSA), invasive	14	13.0	29	14.9	243	18.2
Salmonellosis	8	7.4	14	7.2	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	1	0.9	2	1.0	37	2.8
	4	3.7	4	2.0	7	
Spotted Fever Rickettsiosis	1	0.9	1	0.5	10	0.7
Streptococcal toxic-shock syndrome	0	0.0	1	0.5	12	0.9
Syphilis	7	6.5	13	6.7	104	7.8
Tuberculosis	1	0.9	1	0.5	14	1.0
Varicella (Chickenpox)	9	8.4	15	7.7	252	
Vibriosis	0	0.0	1	0.5	14	1.0
WNV, Non-Human	0	NA		NA		NA
West Nile Fever	0	0.0			2	0.1

AROOSTOOK COUNTY







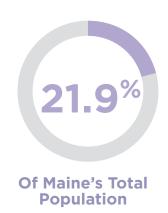
	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	0	0.0	0	0.0	476	35.6
Babesiosis	0	0.0	0	0.0	101	7.5
Borrelia miyamotoi	0	0.0	0	0.0	8	0.6
Campylobacteriosis	23	34.3	23	34.3	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	1	1.5	1	1.5	92	6.9
Chikungunya Virus	0	0.0	0	0.0	2	0.1
Chlamydia trachomatis infection	171	254.8	171	254.8	4346	324.7
Coccidioidomycosis	0	0.0	0	0.0	2	0.1
Cryptosporidiosis	2	3.0	2	3.0	60	4.5
Dengue	0	0.0	0	0.0	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	0	0.0	0	0.0	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	0	0.0	0	0.0	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	4	6.0	4	6.0	163	12.2
Gonorrhea	10	14.9	10	14.9	686	51.3
Group A Streptococcus, invasive	2	3.0	2	3.0	85	6.4
Haemophilus influenzae, invasive	1	1.5	1	1.5	24	1.8
Hepatitis A, acute	0	0.0	0	0.0	9	0.7
Hepatitis B, acute	0	0.0	0	0.0	52	3.9
Hepatitis B, chronic	3	4.5	3	4.5	201	15.0

	Col	unty	District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	3	4.5	3	4.5	39	2.9
Hepatitis C, chronic	55	82.0		82.0	1872	139.9
HIV	2	3.0	2	3.0	30	2.2
Invasive Pneumococcal Disease	10	14.9	10	14.9	132	9.9
Jamestown Canyon virus	0	0.0			1	0.1
Legionellosis	3	4.5	3	4.5	34	2.5
Listeriosis	1	1.5	1	1.5	7	
Lyme disease	4	6.0	4	6.0	1405	105.0
Malaria	0	0.0			9	0.7
Mumps	0	0.0			4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	O.1
Pertussis	1	1.5	1	1.5	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	6	8.9	6	8.9	152	11.4
Rabies, animal	1	NA	1	NA		NA
S. aureus, methicillin resistant (MRSA), invasive	19	28.3	19	28.3	243	18.2
	7	10.4	7	10.4	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	1	1.5	1	1.5	37	2.8
	0	0.0			7	
Spotted Fever Rickettsiosis	0	0.0			10	0.7
Streptococcal toxic-shock syndrome	0	0.0			12	0.9
Syphilis	0	0.0			104	7.8
Tuberculosis	0	0.0			14	1.0
Varicella (Chickenpox)	15	22.4	15	22.4	252	
Vibriosis	0	0.0			14	1.0
WNV, Non-Human	0	NA		NA		NA
West Nile Fever	0	0.0			2	0.1

CUMBERLAND COUNTY







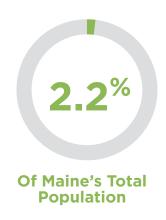
	Co	unty	District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	71	24.2	71	24.2	476	35.6
Babesiosis	20	6.8	20	6.8	101	7.5
Borrelia miyamotoi	1	0.3	1	0.3	8	0.6
Campylobacteriosis	61	20.8	61	20.8	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	33	11.2	33	11.2	92	6.9
Chikungunya Virus	1	0.3	1	0.3	2	0.1
Chlamydia trachomatis infection	1050	357.7	1050	357.7	4346	324.7
Coccidioidomycosis	0	0.0	0	0.0	2	0.1
Cryptosporidiosis	11	3.7	11	3.7	60	4.5
Dengue	1	0.3	1	0.3	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	7	2.4	7	2.4	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	0	0.0	0	0.0	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	46	15.7	46	15.7	163	12.2
Gonorrhea	181	61.7	181	61.7	686	51.3
Group A Streptococcus, invasive	20	6.8	20	6.8	85	6.4
Haemophilus influenzae, invasive	8	2.7	8	2.7	24	1.8
Hepatitis A, acute	4	1.4	4	1.4	9	0.7
Hepatitis B, acute	15	5.1	15	5.1	52	3.9
Hepatitis B, chronic	93	31.7	93	31.7	201	15.0

	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	13	4.4	13	4.4	39	2.9
Hepatitis C, chronic	536	182.6	536	182.6	1872	139.9
HIV	9	3.1	9	3.1	30	2.2
Invasive Pneumococcal Disease	14	4.8	14	4.8	132	9.9
Jamestown Canyon virus	0	0.0			1	0.1
Legionellosis	3	1.0	3	1.0	34	2.5
Listeriosis	0	0.0			7	
Lyme disease	285	97.1	285	97.1	1405	105.0
Malaria	4	1.4	4	1.4	9	0.7
Mumps	0	0.0			4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	0.1
Pertussis	57	19.4	57	19.4	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	22	7.5	22	7.5	152	11.4
Rabies, animal	17	NA	17	NA	76	NA
S. aureus, methicillin resistant (MRSA), invasive	37	12.6	37	12.6	243	18.2
Salmonellosis	26	8.9	26	8.9	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	12	4.1	12	4.1	37	2.8
	0	0.0			7	
Spotted Fever Rickettsiosis	1	0.3	1	0.3	10	0.7
Streptococcal toxic-shock syndrome	3	1.0	3	1.0	12	0.9
Syphilis	34	11.6	34	11.6	104	7.8
Tuberculosis	11	3.7	11	3.7	14	1.0
Varicella (Chickenpox)	73	24.9	73	24.9	252	18.8
Vibriosis	5	1.7	5	1.7	14	1.0
WNV, Non-Human	0	NA		NA		NA
West Nile Fever	2	0.7	2	0.7	2	0.1

FRANKLIN COUNTY







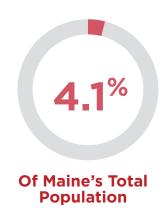
	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	3	10.0	48	24.6	476	35.6
Babesiosis	0	0.0	11	5.6	101	7.5
Borrelia miyamotoi	0	0.0	0	0.0	8	0.6
Campylobacteriosis	4	13.4	30	15.4	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	1	3.3	14	7.2	92	6.9
Chikungunya Virus	0	0.0	0	0.0	2	0.1
Chlamydia trachomatis infection	72	240.8	807	413.4	4346	324.7
Coccidioidomycosis	0	0.0	0	0.0	2	0.1
Cryptosporidiosis	0	0.0	1	0.5	60	4.5
Dengue	0	0.0	1	0.5	3	0.2
Ehrlichiosis, chaffeensis	0	0.0	3	1.5	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	0	0.0	2	1.0	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	4	13.4	22	11.3	163	12.2
Gonorrhea	8	26.8	263	134.7	686	51.3
Group A <i>Streptococcus</i> , invasive	1	3.3	21	10.8	85	6.4
Haemophilus influenzae, invasive	0	0.0	5	2.6	24	1.8
Hepatitis A, acute	0	0.0	2	1.0	9	0.7
Hepatitis B, acute	0	0.0	1	0.5	52	3.9
Hepatitis B, chronic	1	3.3	31	15.9	201	15.0

	Co	unty	District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	0	0.0	4	2.0	39	2.9
Hepatitis C, chronic	16	53.5	200	102.5	1872	139.9
HIV	1	3.3	2	1.0	30	2.2
Invasive Pneumococcal Disease	3	10.0	17	8.7	132	9.9
Jamestown Canyon virus	0	0.0			1	0.1
Legionellosis	3	10.0	7	3.6	34	2.5
Listeriosis	1	3.3	2	1.0	7	
Lyme disease	13	43.5	128	65.6	1405	105.0
Malaria	0	0.0	2	1.0	9	0.7
Mumps	0	0.0	1	0.5	4	0.3
<i>Neisseria meningitidis</i> , invasive (Mening. disease)	0	0.0	1	0.5	1	0.1
Pertussis	7	23.4	34	17.4	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	0	0.0	11	5.6	152	11.4
Rabies, animal	2	NA	15	NA		NA
S. aureus, methicillin resistant (MRSA), invasive	6	20.1	29	14.9	243	18.2
Salmonellosis	3	10.0	14	7.2	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	0	0.0	2	1.0	37	2.8
	0	0.0	4	2.0	7	
Spotted Fever Rickettsiosis	0	0.0	1	0.5	10	0.7
Streptococcal toxic-shock syndrome	0	0.0	1	0.5	12	0.9
Syphilis	5	16.7	13	6.7	104	7.8
Tuberculosis	0	0.0	1	0.5	14	1.0
Varicella (Chickenpox)	4	13.4	15	7.7	252	18.8
Vibriosis	1	3.3	1		14	1.0
WNV, Non-Human	0	NA		NA		NA
West Nile Fever	0	0.0			2	0.1

HANCOCK COUNTY



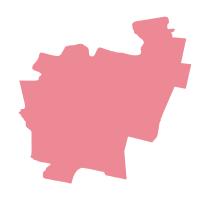




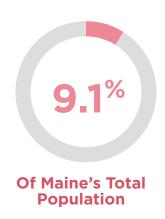
	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	15	27.4	16	18.5	476	35.6
Babesiosis	5	9.1	6	7.0	101	7.5
Borrelia miyamotoi	0	0.0	0	0.0	8	0.6
Campylobacteriosis	12	21.9	16	18.5	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	0	0.0	1	1.2	92	6.9
Chikungunya Virus	0	0.0	0	0.0	2	0.1
Chlamydia trachomatis infection	126	229.9	220	254.9	4346	324.7
Coccidioidomycosis	0	0.0	0	0.0	2	0.1
Cryptosporidiosis	2	3.6	3	3.5	60	4.5
Dengue	0	0.0	0	0.0	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	0	0.0	0	0.0	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	0	0.0	0	0.0	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	7	12.8	8	9.3	163	12.2
Gonorrhea	11	20.1	15	17.4	686	51.3
Group A Streptococcus, invasive	5	9.1	8	9.3	85	6.4
Haemophilus influenzae, invasive	0	0.0	1	1.2	24	1.8
Hepatitis A, acute	0	0.0	0	0.0	9	0.7
Hepatitis B, acute	7	12.8	12	13.9	52	3.9
Hepatitis B, chronic	5	9.1	15	17.4	201	15.0

	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	2	3.6	3	3.5	39	2.9
Hepatitis C, chronic	68	124.1	115	133.3	1872	139.9
HIV	4	7.3	4	4.6	30	2.2
Invasive Pneumococcal Disease	6	10.9	13	15.1	132	9.9
Jamestown Canyon virus		0.0			1	0.1
Legionellosis	2	3.6	2	2.3	34	2.5
Listeriosis	0	0.0			7	
Lyme disease	174	317.5	189	219.0	1405	
Malaria	1	1.8	1	1.2	9	0.7
Mumps	0	0.0			4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	0.1
Pertussis	31	56.6	32	37.1	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	2	3.6	12	13.9	152	11.4
Rabies, animal	3	NA		NA		NA
S. aureus, methicillin resistant (MRSA), invasive	11	20.1	20	23.2	243	18.2
Salmonellosis	7	12.8	10	11.6		
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	0				37	2.8
	0				7	
Spotted Fever Rickettsiosis	1	1.8	2	2.3		0.7
Streptococcal toxic-shock syndrome	1	1.8	2	2.3	12	0.9
Syphilis	2	3.6	4	4.6	104	7.8
Tuberculosis	1	1.8	1	1.2	14	1.0
Varicella (Chickenpox)	9	16.4	13	15.1	252	
Vibriosis	1	1.8	1	1.2	14	1.0
WNV, Non-Human	0	NA		NA		NA
West Nile Fever	0				2	0.1

KENNEBEC COUNTY







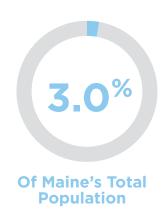
	County		Dis	District		State	
Condition	Count	Rate	Count	Rate	Count	Rate	
Anaplasma phagocytophilum	57	46.7	67	38.8	476	35.6	
Babesiosis	5	4.1	5	2.9	101	7.5	
Borrelia miyamotoi	2	1.6	2	1.2	8	0.6	
Campylobacteriosis	25	20.5	33	19.1	247	18.5	
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	7	5.7	8	4.6	92	6.9	
Chikungunya Virus	0	0.0	0	0.0	2	0.1	
Chlamydia trachomatis infection	364	298.2	518	300.0	4346	324.7	
Coccidioidomycosis	0	0.0	0	0.0	2	0.1	
Cryptosporidiosis	5	4.1	12	6.9	60	4.5	
Dengue	0	0.0	0	0.0	3	0.2	
Ehrlichiosis, <i>chaffeensis</i>	6	4.9	6	3.5	19	1.4	
Ehrlichiosis/Anaplasmosis, undetermined	4	3.3	4	2.3	9	0.7	
Emerging Infection	0	0.0	0	0.0	1	0.1	
Giardiasis	12	9.8	20	11.6	163	12.2	
Gonorrhea	47	38.5	59	34.2	686	51.3	
Group A Streptococcus, invasive	7	5.7	10	5.8	85	6.4	
Haemophilus influenzae, invasive	0	0.0	0	0.0	24	1.8	
Hepatitis A, acute	0	0.0	2	1.2	9	0.7	
Hepatitis B, acute	1	0.8	2	1.2	52	3.9	
Hepatitis B, chronic	4	3.3	6	3.5	201	15.0	

	Co	unty	Dis	trict	State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	0	0.0	3	1.7	39	2.9
Hepatitis C, chronic	147	120.4	212	122.8	1872	139.9
HIV	2	1.6	4	2.3	30	2.2
Invasive Pneumococcal Disease	18	14.7	24	13.9	132	9.9
Jamestown Canyon virus	0	0.0			1	0.1
Legionellosis	3	2.5	5	2.9	34	2.5
Listeriosis	1	0.8	2	1.2	7	
Lyme disease	181	148.3	226	130.9	1405	105.0
Malaria	1	0.8	1	0.6	9	0.7
Mumps	0	0.0	1	0.6	4	0.3
<i>Neisseria meningitidis</i> , invasive (Mening. disease)	0	0.0			1	O.1
Pertussis	15	12.3	26	15.1	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	15	12.3	23	13.3	152	11.4
Rabies, animal	9	NA	10	NA		NA
S. aureus, methicillin resistant (MRSA), invasive	25	20.5	34	19.7	243	18.2
Salmonellosis	7	5.7	10	5.8	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	5	4.1	7	4.1	37	2.8
Shigellosis	0	0.0			7	
Spotted Fever Rickettsiosis	3	2.5	3	1.7	10	0.7
Streptococcal toxic-shock syndrome	1	0.8	1	0.6	12	0.9
Syphilis	9	7.4	14	8.1	104	7.8
Tuberculosis	0	0.0			14	1.0
Varicella (Chickenpox)	12	9.8	22	12.7	252	18.8
Vibriosis	0	0.0			14	1.0
WNV, Non-Human	0	NA		NA		NA
West Nile Fever	0	0.0			2	0.1

KNOX COUNTY







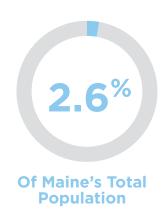
	County		Dist	District		State	
Condition	Count	Rate	Count	Rate	Count	Rate	
Anaplasma phagocytophilum	88	221.3	182	121.8	476	35.6	
Babesiosis	17	42.7	30	20.1	101	7.5	
Borrelia miyamotoi	1	2.5	5	3.3	8	0.6	
Campylobacteriosis	6	15.1	27	18.1	247	18.5	
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	2	5.0	11	7.4	92	6.9	
Chikungunya Virus	0	0.0	1	0.7	2	0.1	
Chlamydia trachomatis infection	114	286.6	385	257.6	4346	324.7	
Coccidioidomycosis	1	2.5	2	1.3	2	0.1	
Cryptosporidiosis	0	0.0	8	5.4	60	4.5	
Dengue	0	0.0	1	0.7	3	0.2	
Ehrlichiosis, <i>chaffeensis</i>	0	0.0	0	0.0	19	1.4	
Ehrlichiosis/Anaplasmosis, undetermined	0	0.0	2	1.3	9	0.7	
Emerging Infection	0	0.0	0	0.0	1	0.1	
Giardiasis	7	17.6	24	16.1	163	12.2	
Gonorrhea	7	17.6	32	21.4	686	51.3	
Group A Streptococcus, invasive	1	2.5	4	2.7	85	6.4	
Haemophilus influenzae, invasive	0	0.0	1	0.7	24	1.8	
Hepatitis A, acute	0	0.0	0	0.0	9	0.7	
Hepatitis B, acute	2	5.0	2	1.3	52	3.9	
Hepatitis B, chronic	4	10.1	17	11.4	201	15.0	

Condition	County		District		State	
	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	1	2.5	1	0.7	39	2.9
Hepatitis C, chronic	96	241.4	214	143.2	1872	139.9
HIV	2	5.0	4	2.7	30	2.2
Invasive Pneumococcal Disease	4	10.1	15	10.0	132	9.9
Jamestown Canyon virus	1	2.5	1	0.7	1	0.1
Legionellosis	1	2.5	2	1.3	34	2.5
Listeriosis	0	0.0			7	
Lyme disease	105	264.0	292	195.4	1405	105.0
Malaria	0	0.0			9	0.7
Mumps	0	0.0			4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	0.1
Pertussis	3	7.5	87	58.2	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	11	27.7	27	18.1	152	11.4
Rabies, animal	2	NA	16	NA		NA
S. aureus, methicillin resistant (MRSA), invasive	7	17.6	26	17.4	243	18.2
Salmonellosis	3	7.5	19	12.7	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	0	0.0	6	4.0	37	2.8
	0	0.0	1	0.7	7	
Spotted Fever Rickettsiosis	3	7.5	3	2.0	10	0.7
Streptococcal toxic-shock syndrome	0	0.0			12	0.9
Syphilis	2	5.0	9	6.0	104	7.8
Tuberculosis	0	0.0			14	1.0
Varicella (Chickenpox)	4	10.1	20	13.4	252	18.8
Vibriosis	2	5.0	2	1.3	14	1.0
WNV, Non-Human	0	NA		NA	5	NA
West Nile Fever	0	0.0			2	0.1

LINCOLN COUNTY







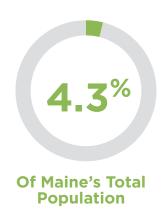
	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	44	128.1	182	121.8	476	35.6
Babesiosis	5	14.6	30	20.1	101	7.5
Borrelia miyamotoi	3	8.7	5	3.3	8	0.6
Campylobacteriosis	9	26.2	27	18.1	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	5	14.6	11	7.4	92	6.9
Chikungunya Virus	1	2.9	1	0.7	2	0.1
Chlamydia trachomatis infection	92	267.9	385	257.6	4346	324.7
Coccidioidomycosis	0	0.0	2	1.3	2	0.1
Cryptosporidiosis	1	2.9	8	5.4	60	4.5
Dengue	0	0.0	1	0.7	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	0	0.0	0	0.0	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	1	2.9	2	1.3	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	8	23.3	24	16.1	163	12.2
Gonorrhea	4	11.6	32	21.4	686	51.3
Group A Streptococcus, invasive	0	0.0	4	2.7	85	6.4
Haemophilus influenzae, invasive	1	2.9	1	0.7	24	1.8
Hepatitis A, acute	0	0.0	0	0.0	9	0.7
Hepatitis B, acute	0	0.0	2	1.3	52	3.9
Hepatitis B, chronic	5	14.6	17	11.4	201	15.0

Condition	County		District		State	
	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	0	0.0	1	0.7	39	2.9
Hepatitis C, chronic	38	110.7	214	143.2	1872	139.9
HIV	1	2.9	4	2.7	30	2.2
Invasive Pneumococcal Disease	2	5.8	15	10.0	132	9.9
Jamestown Canyon virus	0	0.0	1	0.7	1	0.1
Legionellosis	1	2.9	2	1.3	34	2.5
Listeriosis	0	0.0			7	
Lyme disease	62	180.5	292	195.4	1405	105.0
Malaria	0	0.0			9	0.7
Mumps	0	0.0			4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	O.1
Pertussis	5	14.6	87	58.2	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	6	17.5	27	18.1	152	11.4
Rabies, animal	2	NA	16	NA		NA
S. aureus, methicillin resistant (MRSA), invasive	7	20.4	26	17.4	243	18.2
Salmonellosis	6	17.5	19	12.7	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	0	0.0	6	4.0	37	2.8
	0	0.0	1	0.7	7	
Spotted Fever Rickettsiosis	0	0.0	3	2.0	10	0.7
Streptococcal toxic-shock syndrome	0	0.0			12	0.9
Syphilis	1	2.9	9	6.0	104	7.8
Tuberculosis	0	0.0			14	1.0
Varicella (Chickenpox)	5	14.6	20	13.4	252	18.8
Vibriosis	0	0.0	2	1.3	14	1.0
WNV, Non-Human	0	NA		NA	5	NA
West Nile Fever	0	0.0			2	0.1

OXFORD COUNTY







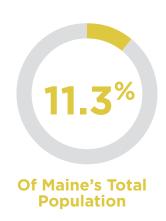
	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	14	24.3	48	24.6	476	35.6
Babesiosis	2	3.5	11	5.6	101	7.5
Borrelia miyamotoi	0	0.0	0	0.0	8	0.6
Campylobacteriosis	10	17.4	30	15.4	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	8	13.9	14	7.2	92	6.9
Chikungunya Virus	0	0.0	0	0.0	2	0.1
Chlamydia trachomatis infection	166	288.1	807	413.4	4346	324.7
Coccidioidomycosis	0	0.0	0	0.0	2	0.1
Cryptosporidiosis	1	1.7	1	0.5	60	4.5
Dengue	0	0.0	1	0.5	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	0	0.0	3	1.5	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	1	1.7	2	1.0	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	8	13.9	22	11.3	163	12.2
Gonorrhea	17	29.5	263	134.7	686	51.3
Group A Streptococcus, invasive	6	10.4	21	10.8	85	6.4
Haemophilus influenzae, invasive	3	5.2	5	2.6	24	1.8
Hepatitis A, acute	0	0.0	2	1.0	9	0.7
Hepatitis B, acute	0	0.0	1	0.5	52	3.9
Hepatitis B, chronic	5	8.7	31	15.9	201	15.0

	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	1	1.7	4	2.0	39	2.9
Hepatitis C, chronic	60	104.1	200	102.5	1872	139.9
HIV	0	0.0	2	1.0	30	2.2
Invasive Pneumococcal Disease	4	6.9	17	8.7	132	9.9
Jamestown Canyon virus	0	0.0			1	0.1
Legionellosis	1	1.7	7	3.6	34	2.5
Listeriosis	0	0.0	2	1.0	7	
Lyme disease	47	81.6	128	65.6	1405	105.0
Malaria	0	0.0	2	1.0	9	0.7
Mumps	0	0.0	1	0.5	4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0	1	0.5	1	0.1
Pertussis	2	3.5	34	17.4	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	6	10.4	11	5.6	152	11.4
Rabies, animal	6	NA	15	NA		NA
S. aureus, methicillin resistant (MRSA), invasive	9	15.6	29	14.9	243	18.2
Salmonellosis	3	5.2	14	7.2	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	1	1.7	2	1.0	37	2.8
	0	0.0	4	2.0	7	
Spotted Fever Rickettsiosis	0	0.0	1	0.5	10	0.7
Streptococcal toxic-shock syndrome	1	1.7	1	0.5	12	0.9
Syphilis	1	1.7	13	6.7	104	7.8
Tuberculosis	0	0.0	1	0.5	14	1.0
Varicella (Chickenpox)	2	3.5	15	7.7	252	18.8
Vibriosis	0	0.0	1	0.5	14	1.0
WNV, Non-Human	0	NA		NA		NA
West Nile Fever	0	0.0			2	0.1

PENOBSCOT COUNTY







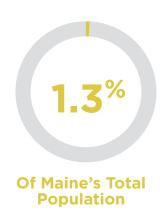
	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	8	5.3	8	4.8	476	35.6
Babesiosis	2	1.3	2	1.2	101	7.5
Borrelia miyamotoi	0	0.0	0	0.0	8	0.6
Campylobacteriosis	24	15.9	28	16.7	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	6	4.0	6	3.6	92	6.9
Chikungunya Virus	0	0.0	0	0.0	2	0.1
Chlamydia trachomatis infection	558	369.3	601	358.0	4346	324.7
Coccidioidomycosis	0	0.0	0	0.0	2	0.1
Cryptosporidiosis	11	7.3	14	8.3	60	4.5
Dengue	0	0.0	0	0.0	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	2	1.3	2	1.2	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	0	0.0	0	0.0	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	19	12.6	22	13.1	163	12.2
Gonorrhea	49	32.4	49	29.2	686	51.3
Group A <i>Streptococcus</i> , invasive	15	9.9	15	8.9	85	6.4
Haemophilus influenzae, invasive	4	2.6	4	2.4	24	1.8
Hepatitis A, acute	0	0.0	0	0.0	9	0.7
Hepatitis B, acute	10	6.6	12	7.1	52	3.9
Hepatitis B, chronic	20	13.2	20	11.9	201	15.0

	Col	unty	District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	7	4.6		4.8	39	2.9
Hepatitis C, chronic	237	156.9	265	157.8	1872	139.9
HIV	2	1.3	2	1.2	30	2.2
Invasive Pneumococcal Disease	18	11.9	21	12.5	132	9.9
Jamestown Canyon virus	0	0.0			1	0.1
Legionellosis	5	3.3	7	4.2	34	2.5
Listeriosis	1	0.7	1	0.6	7	
Lyme disease	77	51.0		47.6	1405	105.0
Malaria	0	0.0			9	0.7
Mumps	1	0.7	1	0.6	4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	O.1
Pertussis	35	23.2	36	21.4	446	33.3
Q fever	0	0.0	1	0.6	1	0.1
Rabies PEP	13	8.6	13	7.7	152	11.4
Rabies, animal	3	NA	5	NA	76	NA
S. aureus, methicillin resistant (MRSA), invasive	44	29.1	46	27.4	243	18.2
Salmonellosis	8	5.3	9	5.4	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	1	0.7	1	0.6	37	2.8
	2	1.3	2	1.2	7	
Spotted Fever Rickettsiosis	0	0.0			10	0.7
Streptococcal toxic-shock syndrome	2	1.3	2	1.2	12	0.9
Syphilis	7	4.6		4.8	104	7.8
Tuberculosis	1	0.7	1	0.6	14	1.0
Varicella (Chickenpox)	39	25.8	40	23.8	252	18.8
Vibriosis	1	0.7	1	0.6	14	1.0
WNV, Non-Human	2	NA	2	NA		NA
West Nile Fever	0	0.0			2	0.1

PISCATAQUIS COUNTY







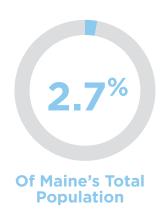
	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	0	0.0	8	4.8	476	35.6
Babesiosis	0	0.0	2	1.2	101	7.5
Borrelia miyamotoi	0	0.0	0	0.0	8	0.6
Campylobacteriosis	4	23.8	28	16.7	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	0	0.0	6	3.6	92	6.9
Chikungunya Virus	0	0.0	0	0.0	2	0.1
Chlamydia trachomatis infection	43	256.0	601	358.0	4346	324.7
Coccidioidomycosis	0	0.0	0	0.0	2	0.1
Cryptosporidiosis	3	17.9	14	8.3	60	4.5
Dengue	0	0.0	0	0.0	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	0	0.0	2	1.2	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	0	0.0	0	0.0	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	3	17.9	22	13.1	163	12.2
Gonorrhea	0	0.0	49	29.2	686	51.3
Group A Streptococcus, invasive	0	0.0	15	8.9	85	6.4
Haemophilus influenzae, invasive	0	0.0	4	2.4	24	1.8
Hepatitis A, acute	0	0.0	0	0.0	9	0.7
Hepatitis B, acute	2	11.9	12	7.1	52	3.9
Hepatitis B, chronic	0	0.0	20	11.9	201	15.0

	Сог	unty	District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	1	6.0		4.8	39	2.9
Hepatitis C, chronic	28	166.7	265	157.8	1872	139.9
HIV	0	0.0	2	1.2	30	2.2
Invasive Pneumococcal Disease	3	17.9	21	12.5	132	9.9
Jamestown Canyon virus	0	0.0			1	0.1
Legionellosis	2	11.9	7	4.2	34	2.5
Listeriosis	0	0.0	1	0.6	7	
Lyme disease	3	17.9		47.6	1405	105.0
Malaria	0	0.0			9	0.7
Mumps	0	0.0	1	0.6	4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	0.1
Pertussis	1	6.0	36	21.4	446	33.3
Q fever	1	6.0	1	0.6	1	0.1
Rabies PEP	0	0.0	13	7.7	152	11.4
Rabies, animal	2	NA	5	NA		NA
S. aureus, methicillin resistant (MRSA), invasive	2	11.9	46	27.4	243	18.2
	1	6.0	9	5.4	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	0	0.0	1	0.6	37	2.8
	0	0.0	2	1.2	7	
Spotted Fever Rickettsiosis	0	0.0			10	0.7
Streptococcal toxic-shock syndrome	0	0.0	2	1.2	12	0.9
Syphilis	1	6.0		4.8	104	7.8
Tuberculosis	0	0.0	1	0.6	14	1.0
Varicella (Chickenpox)	1	6.0	40	23.8	252	18.8
Vibriosis	0	0.0	1	0.6	14	1.0
WNV, Non-Human	0	NA	2	NA		NA
West Nile Fever	0	0.0			2	0.1

SAGADAHOC COUNTY







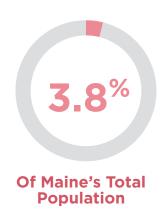
	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	27	75.8	182	121.8	476	35.6
Babesiosis	2	5.6	30	20.1	101	7.5
Borrelia miyamotoi	1	2.8	5	3.3	8	0.6
Campylobacteriosis	6	16.8	27	18.1	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	2	5.6	11	7.4	92	6.9
Chikungunya Virus	0	0.0	1	0.7	2	0.1
Chlamydia trachomatis infection	101	283.4	385	257.6	4346	324.7
Coccidioidomycosis	1	2.8	2	1.3	2	0.1
Cryptosporidiosis	3	8.4	8	5.4	60	4.5
Dengue	0	0.0	1	0.7	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	0	0.0	0	0.0	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	1	2.8	2	1.3	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	3	8.4	24	16.1	163	12.2
Gonorrhea	9	25.3	32	21.4	686	51.3
Group A <i>Streptococcus</i> , invasive	1	2.8	4	2.7	85	6.4
Haemophilus influenzae, invasive	0	0.0	1	0.7	24	1.8
Hepatitis A, acute	0	0.0	0	0.0	9	0.7
Hepatitis B, acute	0	0.0	2	1.3	52	3.9
Hepatitis B, chronic	3	8.4	17	11.4	201	15.0

	Сог	ınty	District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	0	0.0	1	0.7	39	2.9
Hepatitis C, chronic	24	67.4	214	143.2	1872	139.9
HIV	0	0.0	4	2.7	30	2.2
Invasive Pneumococcal Disease	5	14.0	15	10.0	132	9.9
Jamestown Canyon virus	0	0.0	1	0.7	1	0.1
Legionellosis	0	0.0	2	1.3	34	2.5
Listeriosis	0	0.0			7	
Lyme disease	47	131.9	292	195.4	1405	105.0
Malaria	0	0.0			9	0.7
Mumps	0	0.0			4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	0.1
Pertussis	15	42.1	87	58.2	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	7	19.6	27	18.1	152	11.4
Rabies, animal	7	NA	16	NA		NA
S. aureus, methicillin resistant (MRSA), invasive	6	16.8	26	17.4	243	18.2
	7	19.6	19	12.7	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	4	11.2	6	4.0	37	2.8
	1	2.8	1	0.7	7	
Spotted Fever Rickettsiosis	0	0.0	3	2.0	10	0.7
Streptococcal toxic-shock syndrome	0	0.0			12	0.9
Syphilis	3	8.4	9	6.0	104	7.8
Tuberculosis	0	0.0			14	1.0
Varicella (Chickenpox)	1	2.8	20	13.4	252	18.8
Vibriosis	0	0.0	2	1.3	14	1.0
WNV, Non-Human	0	NA		NA		NA
West Nile Fever	0	0.0			2	0.1

SOMERSET COUNTY







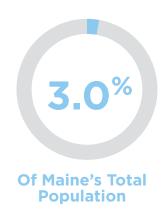
	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	10	19.8	67	38.8	476	35.6
Babesiosis	0	0.0	5	2.9	101	7.5
Borrelia miyamotoi	0	0.0	2	1.2	8	0.6
Campylobacteriosis	8	15.8	33	19.1	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	1	2.0	8	4.6	92	6.9
Chikungunya Virus	0	0.0	0	0.0	2	0.1
Chlamydia trachomatis infection	154	304.4	518	300.0	4346	324.7
Coccidioidomycosis	0	0.0	0	0.0	2	0.1
Cryptosporidiosis	7	13.8	12	6.9	60	4.5
Dengue	0	0.0	0	0.0	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	0	0.0	6	3.5	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	0	0.0	4	2.3	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	8	15.8	20	11.6	163	12.2
Gonorrhea	12	23.7	59	34.2	686	51.3
Group A Streptococcus, invasive	3	5.9	10	5.8	85	6.4
Haemophilus influenzae, invasive	0	0.0	0	0.0	24	1.8
Hepatitis A, acute	2	4.0	2	1.2	9	0.7
Hepatitis B, acute	1	2.0	2	1.2	52	3.9
Hepatitis B, chronic	2	4.0	6	3.5	201	15.0

	Cor	unty	Dis	trict	State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	3	5.9	3	1.7	39	2.9
Hepatitis C, chronic	65	128.5	212	122.8	1872	139.9
HIV	2	4.0	4	2.3	30	2.2
Invasive Pneumococcal Disease	6	11.9	24	13.9	132	9.9
Jamestown Canyon virus	0	0.0			1	0.1
Legionellosis	2	4.0	5	2.9	34	2.5
Listeriosis	1	2.0	2	1.2	7	
Lyme disease	45	88.9	226	130.9	1405	105.0
Malaria	0	0.0	1	0.6	9	0.7
Mumps	1	2.0	1	0.6	4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	0.1
Pertussis	11	21.7	26	15.1	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	8	15.8	23	13.3	152	11.4
Rabies, animal	1	NA	10	NA	76	NA
S. aureus, methicillin resistant (MRSA), invasive	9	17.8	34	19.7	243	18.2
Salmonellosis	3	5.9	10	5.8	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	2	4.0	7	4.1	37	2.8
	0	0.0			7	
Spotted Fever Rickettsiosis	0	0.0	3	1.7	10	0.7
Streptococcal toxic-shock syndrome	0	0.0	1	0.6	12	0.9
Syphilis	5	9.9	14	8.1	104	7.8
Tuberculosis	0	0.0			14	1.0
Varicella (Chickenpox)	10	19.8	22	12.7	252	18.8
Vibriosis	0	0.0			14	1.0
WNV, Non-Human	0	NA		NA		NA
West Nile Fever	0	0.0			2	0.1

WALDO COUNTY







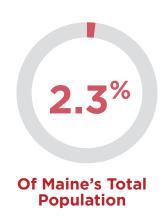
	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	23	57.9	182	121.8	476	35.6
Babesiosis	6	15.1	30	20.1	101	7.5
Borrelia miyamotoi	0	0.0	5	3.3	8	0.6
Campylobacteriosis	6	15.1	27	18.1	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	2	5.0	11	7.4	92	6.9
Chikungunya Virus	0	0.0	1	0.7	2	0.1
Chlamydia trachomatis infection	78	196.5	385	257.6	4346	324.7
Coccidioidomycosis	0	0.0	2	1.3	2	0.1
Cryptosporidiosis	4	10.1	8	5.4	60	4.5
Dengue	1	2.5	1	0.7	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	0	0.0	0	0.0	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	0	0.0	2	1.3	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	6	15.1	24	16.1	163	12.2
Gonorrhea	12	30.2	32	21.4	686	51.3
Group A Streptococcus, invasive	2	5.0	4	2.7	85	6.4
Haemophilus influenzae, invasive	0	0.0	1	0.7	24	1.8
Hepatitis A, acute	0	0.0	0	0.0	9	0.7
Hepatitis B, acute	0	0.0	2	1.3	52	3.9
Hepatitis B, chronic	5	12.6	17	11.4	201	15.0

Condition	County		District		State	
	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	0	0.0	1	0.7	39	2.9
Hepatitis C, chronic	56	141.1	214	143.2	1872	139.9
HIV	1	2.5	4	2.7	30	2.2
Invasive Pneumococcal Disease	4	10.1	15	10.0	132	9.9
Jamestown Canyon virus	0	0.0	1	0.7	1	0.1
Legionellosis	0	0.0	2	1.3	34	2.5
Listeriosis	0	0.0			7	
Lyme disease	78	196.5	292	195.4	1405	105.0
Malaria	0	0.0			9	0.7
Mumps	0	0.0			4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	0.1
Pertussis	64	161.2	87	58.2	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	3	7.6	27	18.1	152	11.4
Rabies, animal	5	NA	16	NA		NA
S. aureus, methicillin resistant (MRSA), invasive	6	15.1	26	17.4	243	18.2
Salmonellosis	3	7.6	19	12.7	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	2	5.0	6	4.0	37	2.8
	0	0.0	1	0.7	7	
Spotted Fever Rickettsiosis	0	0.0	3	2.0	10	0.7
Streptococcal toxic-shock syndrome	0	0.0			12	0.9
Syphilis	3	7.6	9	6.0	104	7.8
Tuberculosis	0	0.0			14	1.0
Varicella (Chickenpox)	10	25.2	20	13.4	252	18.8
Vibriosis	0	0.0	2	1.3	14	1.0
WNV, Non-Human	0	NA		NA		NA
West Nile Fever	0	0.0			2	0.1

WASHINGTON COUNTY







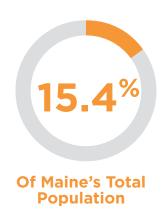
	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	1	3.2	16	18.5	476	35.6
Babesiosis	1	3.2	6	7.0	101	7.5
Borrelia miyamotoi	0	0.0	0	0.0	8	0.6
Campylobacteriosis	4	12.7	16	18.5	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	1	3.2	1	1.2	92	6.9
Chikungunya Virus	0	0.0	0	0.0	2	0.1
Chlamydia trachomatis infection	94	298.5	220	254.9	4346	324.7
Coccidioidomycosis	0	0.0	0	0.0	2	0.1
Cryptosporidiosis	1	3.2	3	3.5	60	4.5
Dengue	0	0.0	0	0.0	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	0	0.0	0	0.0	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	0	0.0	0	0.0	9	0.7
Emerging Infection	0	0.0	0	0.0	1	0.1
Giardiasis	1	3.2	8	9.3	163	12.2
Gonorrhea	4	12.7	15	17.4	686	51.3
Group A Streptococcus, invasive	3	9.5	8	9.3	85	6.4
Haemophilus influenzae, invasive	1	3.2	1	1.2	24	1.8
Hepatitis A, acute	0	0.0	0	0.0	9	0.7
Hepatitis B, acute	5	15.9	12	13.9	52	3.9
Hepatitis B, chronic	10	31.8	15	17.4	201	15.0

	Cor	unty	District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	1	3.2	3	3.5	39	2.9
Hepatitis C, chronic	47	149.3	115	133.3	1872	139.9
HIV		0.0	4	4.6	30	2.2
Invasive Pneumococcal Disease	7	22.2	13	15.1	132	9.9
Jamestown Canyon virus		0.0			1	0.1
Legionellosis		0.0	2	2.3	34	2.5
Listeriosis		0.0			7	
Lyme disease	15	47.6	189	219.0	1405	
Malaria		0.0	1	1.2	9	0.7
Mumps	0	0.0			4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	0.1
Pertussis	1	3.2	32	37.1	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	10	31.8	12	13.9	152	11.4
Rabies, animal	5	NA		NA		NA
S. aureus, methicillin resistant (MRSA), invasive	9	28.6	20	23.2	243	18.2
Salmonellosis	3	9.5	10	11.6		
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	0				37	2.8
	0				7	
Spotted Fever Rickettsiosis	1	3.2	2	2.3		0.7
Streptococcal toxic-shock syndrome	1	3.2	2	2.3	12	0.9
Syphilis	2	6.4	4	4.6	104	7.8
Tuberculosis	0		1	1.2	14	1.0
Varicella (Chickenpox)	4	12.7	13	15.1	252	
Vibriosis	0		1	1.2	14	1.0
WNV, Non-Human	0	NA		NA		NA
West Nile Fever	0				2	0.1

YORK COUNTY







	County		District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Anaplasma phagocytophilum	84	40.7	84	40.7	476	35.6
Babesiosis	27	13.1	27	13.1	101	7.5
Borrelia miyamotoi	0	0.0	0	0.0	8	0.6
Campylobacteriosis	29	14.1	29	14.1	247	18.5
Carbapenem-resistant <i>Enterobacteriaceae</i> (CRE)	18	8.7	18	8.7	92	6.9
Chikungunya Virus	0	0.0	0	0.0	2	0.1
Chlamydia trachomatis infection	594	288.0	594	288.0	4346	324.7
Coccidioidomycosis	0	0.0	0	0.0	2	0.1
Cryptosporidiosis	9	4.4	9	4.4	60	4.5
Dengue	0	0.0	0	0.0	3	0.2
Ehrlichiosis, <i>chaffeensis</i>	1	0.5	1	0.5	19	1.4
Ehrlichiosis/Anaplasmosis, undetermined	1	0.5	1	0.5	9	0.7
Emerging Infection	1	0.5	1	0.5	1	0.1
Giardiasis	17	8.2	17	8.2	163	12.2
Gonorrhea	77	37.3	77	37.3	686	51.3
Group A Streptococcus, invasive	5	2.4	5	2.4	85	6.4
Haemophilus influenzae, invasive	4	1.9	4	1.9	24	1.8
Hepatitis A, acute	1	0.5	1	0.5	9	0.7
Hepatitis B, acute	8	3.9	8	3.9	52	3.9
Hepatitis B, chronic	16	7.8	16	7.8	201	15.0

	Cou	unty	District		State	
Condition	Count	Rate	Count	Rate	Count	Rate
Hepatitis C, acute	4	1.9	4	1.9	39	2.9
Hepatitis C, chronic	275	133.3	275	133.3	1872	139.9
HIV	3	1.5	3	1.5	30	2.2
Invasive Pneumococcal Disease	18	8.7	18	8.7	132	9.9
Jamestown Canyon virus	0	0.0			1	0.1
Legionellosis	5	2.4	5	2.4	34	2.5
Listeriosis	1	0.5	1	0.5	7	
Lyme disease	201	97.5	201	97.5	1405	105.0
Malaria	1	0.5	1	0.5	9	0.7
Mumps	1	0.5	1	0.5	4	0.3
Neisseria meningitidis, invasive (Mening. disease)	0	0.0			1	0.1
Pertussis	173	83.9	173	83.9	446	33.3
Q fever	0	0.0			1	0.1
Rabies PEP	38	18.4	38	18.4	152	11.4
Rabies, animal	4	NA	4	NA		NA
S. aureus, methicillin resistant (MRSA), invasive	32	15.5	32	15.5	243	18.2
Salmonellosis	24	11.6	24	11.6	119	8.9
Shiga toxin-producing <i>Escherichia coli</i> (STEC)	8	3.9		3.9	37	2.8
	0	0.0			7	
Spotted Fever Rickettsiosis	0	0.0			10	0.7
Streptococcal toxic-shock syndrome	3	1.5	3	1.5	12	0.9
Syphilis	22	10.7	22	10.7	104	7.8
Tuberculosis	0	0.0			14	1.0
Varicella (Chickenpox)	54	26.2	54	26.2	252	18.8
Vibriosis	4	1.9	4	1.9	14	1.0
WNV, Non-Human	3	NA	3	NA		NA
West Nile Fever	0	0.0			2	0.1

WORKGROUP **SUMMARIES**



DATA QUALITY WORKGROUP

Maine's Data Quality Workgroup meets every other week to review data quality of laboratory reports that are sent to Maine Center for Disease Control and Prevention (Maine CDC) with attention paid to those facilities that only send electronic lab reports (ELR). The Workgroup is chaired by Maine CDC's Infectious Disease Informatician and other attendees are the Infectious Disease Program Director, Informatics Epidemiologist, and Influenza Surveillance Coordinator. The Data Quality Workgroup ensures that lab volumes are consistent and that all data feeds are running correctly. The group also reviews documents submitted by the labs that are looking to stop faxing paper lab reports and move to ELR only. In 2018, Maine CDC approved an additional facility to report only electronically bringing the total paper off reporters to six.



FOOD SAFETY WORKGROUP

The Maine Interagency Food Safety Workgroup is led by Maine CDC's Foodborne Disease Epidemiologist and is comprised of representatives from state agencies, federal agencies, and other organizations involved in improving food safety in Maine (including, but not limited to, Maine Department of Marine Resources (Maine DMR), Maine Department of Agriculture, Conservation, and Forestry (Maine DACF), Maine Department of Education (Maine DOE), United States Department of Agriculture (USDA), the Food and Drug Administration (FDA), and the University of Maine Cooperative Extension). These organizations and agencies collaborate to reduce the incidence of foodborne and waterborne infectious diseases in the state, respond to foodborne and waterborne outbreaks, and work together to advance food safety initiatives. The Workgroup meets quarterly during the year to discuss the latest developments and cooperate to improve response and prevention. It occasionally holds trainings for its member agencies and invites industry members to meetings to highlight common issues.

Members of the Workgroup and Maine CDC infectious disease epidemiologists investigated a large waterborne outbreak of norovirus during the summer of 2018. Members of the Workgroup collaborated on several other outbreak investigations over the course of the year. Several members of the Workgroup presented at various conferences and meetings throughout the year, including the FDA Northeast Annual Food Protection Seminar in Portland in September 2018.

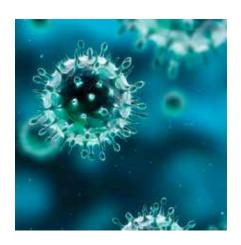
HEALTH EDUCATION WORKGROUP

Maine CDC's Health Education Workgroup started meeting in December of 2018 and meets every other month to address health education topics in public health. The Workgroup aims to strengthen skills such as health literacy, plain language, provider outreach, and the use of social media and infographics. This meeting also provides an opportunity to share resources and allows Maine CDC's programs to be more interconnected. The Workgroup is chaired by Maine CDC's Infectious Disease Health Educator and includes health educators and similar roles from Maine CDC's Environmental and Occupational Health Program, HIV, STD, and Viral Hepatitis Program, Immunization Program, and Tobacco and Substance Use Prevention and Control Program. The Workgroup plans to expand in 2019 to include representation from other programs including Women, Infants, and Children Nutrition Program and the Suicide Prevention Program.



INFLUENZA WORK GROUP

Maine's Influenza Workgroup meets quarterly to address current topics in influenza and other viral respiratory pathogens. The Workgroup is chaired by the Influenza Surveillance Coordinator and includes representatives from Infectious Disease Epidemiology, Public Health Preparedness, the Maine Immunization Program, Public Health Nursing, Maine's Health and Environmental Testing Laboratory (HETL), Maine DACF, and other relevant partners. The Workgroup coordinates surveillance and response to influenza and maintains and updates the Pandemic Influenza Operations Plan. The Influenza Workgroup also sponsors a start of influenza season conference call for healthcare providers and laboratories to update them on new guidance, reporting requirements, and assistance available from the State.



WORKGROUP SUMMARIES



RABIES WORKGROUP

The Maine Rabies Workgroup meets quarterly to address current topics in statewide rabies prevention and management. The Workgroup, co-chaired by the State Epidemiologist and the State Veterinarian, is comprised of animal and human health representatives from local, state, and federal agencies whose mission is to control the spread of rabies, a fatal zoonotic disease that is endemic in Maine. Agencies and organizations that participate in the Workgroup include, but are not limited to: Maine CDC, Maine DACF, HETL, Maine Department of Inland, Fisheries, and Wildlife (Maine IF&W), USDA, Maine Veterinary Medical Association, Maine Federation of Humane Societies, and the Maine Animal Control Association.

Members of the Workgroup provide training to town animal control officers and game wardens regarding rabies biology and prevention and control of the disease in Maine. The USDA's Animal and Plant Health Inspection Service distributes oral rabies vaccines in northern and eastern areas of the state with the goal to reduce the incidence of raccoon rabies. In 2018, the Workgroup introduced and piloted the Mail-A-Bat program at five vet clinics across the state. The Workgroup provided clinics with shipping material to mail bats to HETL for rabies testing. After a successful trial, the Workgroup hopes to expand this program to more vet clinics in future years.

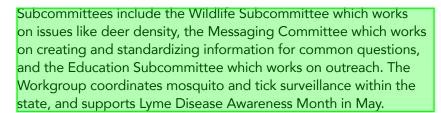


SYNDROMIC SURVEILLANCE USER GROUP

Maine's Syndromic Surveillance User Group meets every other month to address topics related to syndromic surveillance such as opioid overdose and influenza like illness (ILI) related emergency department visits as well as provide training on the use of available tools such as the Electronic Surveillance System for the Early Notification of Community-based Epidemics (ESSENCE). The User Group is chaired by the Informatics Epidemiologist and includes partners from within Maine CDC, Substance Abuse and Mental Health Services Administration (SAMSHA), Emergency Medical Services (EMS) as well as hospitals around Maine. The Group started meeting in April 2018 and provided several trainings for ESSENCE. The Group also worked together to create dashboards which have trend-over-time graphs and maps for topics that are of interest to the Group such as influenza-like illness and drug overdose syndromes.

VECTORBORNE WORKGROUP

Maine's Vectorborne Workgroup meets every other month to address current topics in vectorborne diseases including illnesses spread by ticks, mosquitoes, and vectors of other medical importance like browntail moths. The Workgroup is chaired by the Infectious Disease Epidemiology Program Director at Maine CDC and includes representatives from epidemiology, environmental health, HETL, Maine DACF, Maine DOE, Maine Department of Environmental Protection, Maine Medical Center Research Institute, University of Maine Cooperative Extension, Maine IF&W, the Biodiversity Research Institute, pest control companies, and other relevant individuals.



The committee also launched a new mosquito messaging website with commonly asked questions, their answers, and the references that back the answers up. This website can be viewed at www.maine.gov/dhhs/mosquitofaq.



REMOVING BARRIERS TO HEPATITIS C TREATMENT IN MAINE



MaineCare clients are no longer required to prove sobriety to receive treatment for hepatitis C.

On July 10, 2018, a group of doctors and hepatitis specialists from Maine sent a letter to MaineCare, Maine's Medicaid program, advocating for the removal of the sobriety barrier for hepatitis C treatment for MaineCare patients. This letter prompted MaineCare to request Maine CDC's Viral Hepatitis Prevention Program to present information on studies related to sobriety barriers for hepatitis C treatment to MaineCare's Drug Utilization Review Committee. The Committee voted unanimously to remove the sobriety barrier, and on October 1, 2018, MaineCare clients were no longer required to prove sobriety to receive treatment for hepatitis C.

Previously, MaineCare required clients to be sober from alcohol and illicit drugs for six months before receiving hepatitis C treatment. However, multiple studies show that sobriety restrictions are unnecessary and undermine public health efforts to end the hepatitis C epidemic.^{1,2} In addition, removing sobriety barriers to hepatitis C treatment can result in lower medical costs to the state.3 The cost of treating hepatitis C when diagnosed is far less expensive than treating liver disease or performing liver transplants, for which hepatitis C is the leading cause.4,5

Studies show that hepatitis treatment medication is just as effective in people who use drugs (PWUD) or alcohol as it is in people who do not, and PWUD adhere to treatment at the same rates as non-drug using individuals.^{6,7,8,9,10,11} Also, hepatitis C treatment for PWUD can lead to significant reductions in hepatitis C prevalence and reduced transmissions. Furthermore, the risk of reinfection for PWUD is less than 4%.^{9,12}

Sobriety restrictions come with legal and financial risks. Several states with sobriety barriers for hepatitis C treatment in their Medicaid programs have lost costly lawsuits for not providing this treatment.^{13,14,15,16} Currently, at least half of all state Medicaid programs do not require individuals to meet sobriety criteria before obtaining treatment.¹⁷ Lessons learned from the HIV epidemic of the 1980s and 1990s showed that having health insurance coverage and access to treatment was crucial in ending those epidemics. 18,19 Maine CDC's Viral Hepatitis Prevention Program continues to provide information to healthcare providers, state agencies and the public to help end the hepatitis C epidemic.



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Aspinall EJ, Corson S, Doyle JS, et al., Treatment of hepatitis C virus infection among people who are actively injecting drugs: a systematic review and meta-analysis, 57 CLIN INFECT DIS.S80 (2013).

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LYME DISEASE **SOCIAL MEDIA CAMPAIGN**

The Public Health Corps (PHC), within Maine CDC's Division of Disease Surveillance, designed and launched a Lyme disease social media campaign aimed at adults 65 years and older. The campaign targeted this specific age group because adults ages 65 years and older have the highest rate of Lyme disease in Maine. The PHC conducted a literature review and found that best practices for public health social media campaigns include the use of videos less than 44 seconds, pictures, and mascots. Mascots are beneficial in messaging since they help brand the campaign and allow users to quickly recognize the material. Maine CDC worked with a graphic designer to create four video slideshows, a static photo advertisement, and a mascot for the campaign (Image 1).



Image 1

In August 2018, Maine CDC posted each advertisement for one week as Facebook boosted posts, Facebook sponsored advertisements, and YouTube paid instream video advertisements. Maine CDC spent \$9,298.42 in total on the campaign. To determine the number of views and reach of each advertisement, Maine CDC looked at Google and Facebook analytics.

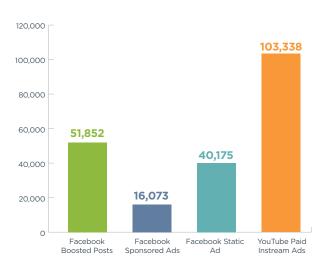


Chart 1: Average Number of Advertisement Views by Dissemination Method

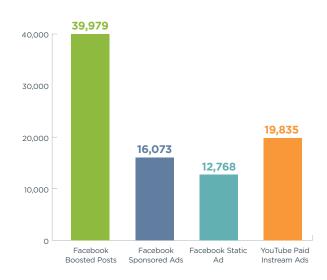


Chart 2: Average Number of Viewers Within the Target Population

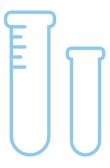


Facebook boosted posts generated an average of 51,852 views and reached 46,370 people. Facebook sponsored advertisements generated an average of 16,073 views and reached 25,304 people. The Facebook sponsored static advertisement reached 40,175 people. There are no data available on the average number of views for Facebook static advertisements since they are viewed differently than videos. YouTube paid instream advertisements generated an average of 46,377 views and reached 104,021 people. Maine CDC found that YouTube paid instream advertisements and Facebook boosted posts reached more people and generated the most views (Chart 1). YouTube paid instream advertisements generated the most views when Maine CDC selected to advertise to both the

target age group and users with an unknown age. Facebook boosted posts and Facebook sponsored advertisements reached the most people within the target population (Chart 2).

Maine CDC found that each advertisement method had strengths and weaknesses. YouTube paid instream advertisements work best to reach the general public, while Facebook boosted posts work best to reach an age-specific target population. The Facebook static advertisement had more views than Facebook sponsored advertisements, although the static advertisement reached less of the target population. Maine CDC will use this information to inform future social media campaigns to maximize the reach of the information to target populations.

EXTRAGENITAL TESTING: POTENTIAL SOLUTIONS TO CURB THE **INCIDENCE OF STI INFECTIONS IN MAINE**



Rapid detection of STIs is crucial in reducing the incidence of STIs in Maine.

There is an increase in cases of sexually transmitted infections (STIs) in the United States, as well as in the state of Maine. From 2016-2017, Maine saw an increase of 30% for gonorrhea and 77% for infectious syphilis. There was another increase from 2017-2018, with a 19% increase in gonorrhea and a 25% increase in infectious syphilis. There are many hypotheses of why STIs are increasing nationally, including: better surveillance, increased testing, reduced condom use, and an upsurge in popularity of dating apps which may increase one's sexual network. Transmission is further amplified by the high rates of undiagnosed asymptomatic STIs, or STIs that are left untreated/ untested because they are in exposure sites that are not typically tested (pharyngeal or rectal). Rapid detection of STIs is crucial in reducing the incidence of STIs in Maine. Some health consequences of untreated STIs include: stillbirths, ectopic pregnancy, pelvic inflammatory disease, infertility and an increased risk for HIV transmission.



It is possible that providers are missing asymptomatic

STIs that present in extragenital sites of exposure. The federal CDC's current recommendations state that men who have sex with men (MSM) who have had receptive anal or oral sex be screened using the Nucleic Acid Amplification Test (NAAT) annually¹. Medical providers should take thorough sexual health histories to assess if an individual should be tested more frequently. Studies indicate that genitourinary screening alone for MSM has the potential to miss 80% of chlamydia and 77% of gonorrhea infections². Research shows that medical providers are screening regularly for syphilis among HIV positive MSM but are screening less than 10% of HIV positive MSM for gonorrhea and chlamydia (rectal and pharyngeal).1 Anal chlamydia and gonorrhea infections are associated with a greater risk of acquiring HIV. One-in-ten new HIV cases among MSM are due to the presence of chlamydia and gonorrhea infection³. Maine 2018 data indicate



that rectal and pharyngeal STI testing only represent about 1% of total STI testing (8% of STI testing had no specimen description listed). Potential barriers to lack of extragenital testing could be physicians' comfort level in taking sexual history and the fact that extragenital testing is not yet FDA approved.

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EMERGING THREAT:

CARBAPENEMASE-PRODUCING ORGANISMS

BACKGROUND:

While most bacterial resistance is acquired from prior exposure to antibiotics, there are some bacteria that can share their resistance genes directly with other bacteria, including bacteria of other genus and species. This type of bacterial resistance is known as plasmid-mediated resistance.

THE THREAT:

- Carbapenemase-producing organisms (CPO) are a subset of carbapenemase-resistant organisms (CRO) that are capable of plasmid-mediated resistance.
- CPOs are highly resistant to multiple classes of antibiotics, leaving clinicians with few treatment options.
- Infections with CPOs are associated with high mortality rates.
- A person can be colonized with a CPO and not be aware of it, potentially spreading it.
- Hospitalization outside of the USA, including surgery and dialysis, in an CPO endemic country, is a significant risk factor for acquiring a CPO.
- CPOs are now found in every state in the USA, including Maine.

CARBAPENEM-RESISTANCE DEFINED:

CRO

Carbapenem-resistant

Carbapenem-resistant Organisms (CRO) are bacteria in the family Enterobacteriaceae (e.g. Citrobacter spp., Enterobacter spp., Escherichia coli, Klebsiella spp., Morganella morganii, Proteus spp., Providencia spp., Serratia spp., Salmonella spp., Shigella spp., Yersinia spp.), Pseudomonas aeruginosa and Acinetobacter baumannii that test resistant to one or more carbapenem antibiotics, regardless of the method of resistance.

Non-CPO Acquired resistance

Non carbapenemase-producing organisms (non-CPO), are those CROs that have undergone further laboratory testing and were found not to contain a carbapenemase-producing enzyme that breaks down the carbapenem antibiotics or the gene (e.g. IMP, KPC, NDM, OXA-48, VIM) that produces the carbapenemase enzyme. Therefore, resistance to the carbapenem antibiotic(s) is acquired resistance.

CPO Plasmid-mediated resistance

Carbapenemase-producing organisms (CPO), are CROs that test positive for the carbapenemaseproducing enzyme or gene, signifying that resistance to the carbapenem antibiotic(s) is plasmid-mediated. CPOs can share their resistance genes directly with other bacteria and are considered a public health threat.

PREVENTION:

- Hand hygiene is still the best prevention.
- Clean and disinfect patient rooms and equipment.
- Follow contact precautions for known patients colonized or infected with a CPO.
- Practice good antimicrobial stewardship.

HEALTHCARE FACILITIES SHOULD NOTIFY MAINE CDC AT 1-800-821-5821 WHEN:

- A CRO is identified (and send the isolate to the state public health laboratory)
- Healthcare associated transmission of a CRO is suspected

Carbapenemaseproducing organisms are emerging multidrug resistant bacteria that present a serious health threat for the people of Maine.



MAINE'S RYAN WHITE PART B AND AIDS DRUG ASSISTANCE PROGRAM

The Ryan White Part B Program helps low-income people living with HIV (PLWH) in Maine fill gaps in care and treatment by providing a variety of services, depending on individual need, with the ultimate goal being the achievement and maintenance of viral suppression. The Ryan White Part B Program helps low-income PLWH access food, dental care, and housing. The Program also supports medical case management for those who do not qualify for other types of case management, and administers the AIDS Drug Assistance Program (ADAP), which helps low-income PLWH obtain and maintain access to prescription drugs to treat HIV and its related conditions. ADAP services include assistance with health insurance premiums, copays, deductibles, coinsurance, HIV-related lab tests, and the full cost of HIVrelated drugs for those without insurance.

People Living with HIV Utilizing Ryan White Part B Services, 2015-2018

Service	2015	2016	2017	2018
Dental assistance	183	180	279	293
Food assistance	497	522	579	584
Full-cost drugs	110	120	106	118
Housing assistance	168	199	257	304
Insurance premiums	208	190	240	299
Lab tests	14	20	24	25
Medical case management	87	90	97	101
Prescription wrap-around	626	602	544	560
Total utilizing members	882	923	939	987

PLWH who are virally suppressed (defined as a viral load test result less than 200 copies/mL) are less likely to develop HIV-related complications, so they lead longer, healthier lives and require less costly care and treatment. PLWH who are virally suppressed are much less likely to transmit the virus to others. The National HIV/AIDS Strategy calls for viral suppression among 80 percent of all PLWH in the U.S. by 2020. In 2018, 90 percent of Part B Program enrollees were virally suppressed as of the last result reported in 2018, compared to 85 percent of enrollees in 2017.

Viral Suppression Among Ryan White Part B Enrollees by Public Health District, 2018

District	Number Virally Suppressed	Number Enrolled	% Virally Suppressed
Aroostook	27	29	93%
Central	123	142	87%
Cumberland	354	386	92%
Downeast	70	78	90%
Mid Coast	64	69	93%
Penquis	81	86	94%
Western	126	151	83%
York	133	146	91%
Overall	978	1,087	90%



90%

of people living with HIV enrolled in Maine's Ryan White Part B Program were virally suppressed in 2018.

POND OUTBREAK **CUMBERLAND COUNTY JULY 2018**

On Friday July 6, 2018 a member of the public contacted Maine CDC about family and friends who became ill with gastrointestinal symptoms after visiting a public beach on Woods Pond, operated by the Town of Bridgton, Maine. Given the gastrointestinal (GI) symptoms and the epidemiology described in the initial report, an outbreak team quickly formed to investigate. Maine CDC notified and worked closely with local hospitals and officials to investigate this outbreak. Based on information provided to town officials, the Town of Bridgton closed the public beach on July 6.

Using the initial information provided, Maine CDC's Cumberland Field Epidemiologist conducted open ended interviews with seven heads of households to generate a list of questions for a wider attendee survey. The Foodborne and Waterborne Epidemiologist added exposures of note to a modified version of the Division of Disease Surveillance's Foodborne/Enteric Disease survey template. The Maine CDC interview team consisted of five epidemiologists and one public health educator. The interview team used this survey to collect information about individuals' exposures, symptoms, and duration of illness.

Using the Maine CDC Facebook page, Maine CDC issued a call for cases and identified additional individuals who visited the pond between July 1 – July 6, 2018. In many circumstances, the heads of households provided answers for multiple family members. The interview team entered interview data into a spreadsheet, and the Foodborne and Waterborne Epidemiologist analyzed these using Epi Info 7 statistical software. Maine CDC asked ill respondents to submit stool specimens for viral and bacterial testing at Maine's Health and Environmental Testing Laboratory (HETL). New York State Wadsworth Center Virology Laboratory assisted in confirming test results as needed.

Maine CDC consulted with the federal Centers for Disease Control and Prevention (CDC) regarding testing the pond's water if the outbreak team deemed it appropriate. The outbreak team also discussed additional testing and epidemiology strategies. Maine CDC decided testing of the pond water would not be useful given the circumstances of the outbreak.

After the call for additional cases, Maine CDC interviewed a total of thirty-four (34) heads of household by phone and collected data on 148 people who visited the pond or were exposed to ill individuals at home. 139 people reported attending the pond between July 1 – July 6. Of those who visited the pond, 97 people reported being sick (attack rate: 69.8%). Pond visitors who went under the water or swallowed the water while swimming were over three times more likely to be sick than those who did not (relative risk = 3.19, p < 0.001). There were no significant relative risks for other exposures such as using the playground and eating or drinking at the beach.

69.8% Attack Rate

Swimmers who went under water were 3 times more likely to become ill.



Health care providers collected four (4) human stool specimens for bacterial and viral testing that was performed at HETL. Two out of four human stool specimens tested positive for Norovirus GI. Based on collected data and two positive human specimens, Maine CDC confirmed this as a continuous common source waterborne outbreak of norovirus with secondary person-to-person transmission.

Although Maine CDC associated the outbreak with swimming underwater at the beach, Maine CDC was unable to determine the source of the norovirus. There may have been numerous sources for contamination that resulted in a high number of ill individuals. The town of Bridgton reopened the public beach on July 10, 2018 and no additional individuals reported illnesses associated with this outbreak.

Maine CDC confirmed this as a continuous common source waterborne outbreak of norovirus with secondary person-toperson transmission.

CAKE OUTBREAK **ANDROSCOGGIN COUNTY JUNE 2018**

On Monday June 25, 2018 a Hospital Infection Preventionist (IP) notified Maine CDC of a gastrointestinal outbreak seemingly stemming from a work party event held on Thursday, June 21 at a private medical practice in Androscoggin County. Maine CDC's Western District Field Epidemiologist began collecting information about the outbreak and notified the Foodborne Epidemiologist who convened an outbreak investigation team.

The work party consisted of approximately 40 attendees; 21 had been or were currently ill with nausea, vomiting, and diarrhea. Many of the ill individuals had symptom onset on Friday, June 22 and Saturday, June 23. Average duration of illness was approximately two days. Employees attending the party brought in food for the potluck including: taco salad, homemade cookie cake, ice cream, assorted dips and chips, seven-layer dip, meatballs, strawberries stuffed with cheesecake filling, rotisserie chicken, potato salad, pasta salad, and soda.



Using this list, epidemiologists designed and administered a customized survey to better collect exposure and illness information. The Western District Field Epidemiologist interviewed 23 employees in person at the office, and other epidemiologists interviewed another 15 attendees by telephone. The Foodborne Epidemiologist analyzed the data collected from the surveys to look for statistically significant exposures that may have led to these illnesses. Those who ate or came in to contact with the cookie cake were more than eight (8) times more likely to be ill (relative risk = 8.34 [1.27 - 54.69], p < 0.0002) than those who did not. Contact included eating, serving, cooking, preparing, and disposing of the cake. Epidemiologists found no

Number of attendees who had been or were currently ill.





Attendees who came in contact with the cake were 8 times more likely to become ill.

other food exposures significantly associated with illness. The employee who brought the homemade cookie cake to the potluck said that multiple family members were ill during the previous week.

One attendee submitted a stool specimen to Maine's Health and Environmental Testing Laboratory (HETL) for bacterial and viral testing. Epidemiologists collected leftover food from the office and brought it to HETL, and laboratorians sampled parts of the cookie cake for norovirus testing. All of the cake components tested negative for norovirus, but the one human specimen submitted tested positive for norovirus GII. Testing for norovirus in food is challenging, so the absence of norovirus in the cake sample did not mean that other parts of the cake were not contaminated.

In conclusion, epidemiologists classified this outbreak as a probable norovirus outbreak associated with consuming or coming into contact with the cookie cake at the party. Epidemiologists were able to provide a strong epidemiological link between illness and consumption of the cake despite the negative results on the cake samples tested. The Western District Field Epidemiologist shared these findings with the office and provided guidance on safe food handling practices, staying out of work while ill, not preparing food for others while ill, and general sanitation practices.

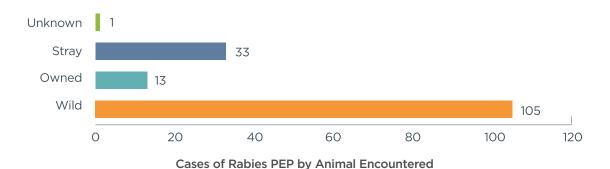
RABIES POST-EXPOSURE PROPHYLAXIS (PEP) **IN MAINE, 2018**

Animal rabies is endemic in Maine and every year humans are exposed to rabid or potentially rabid animals. Rabies is 100% fatal if left untreated, but can be prevented when treated promptly after an exposure to the rabies virus. Recommendations for individuals to receive rabies treatment after an exposure to the rabies virus depend on the circumstances surrounding the exposure. The treatment is called rabies postexposure prophylaxis (PEP), and it involves a series of rabies vaccines and rabies immune globulin (RIG) shots. Healthcare providers are required by Maine law to report rabies PEP to Maine CDC within 48 hours of administration, but many providers fall short of this requirement. For this reason, the reported cases of rabies PEP in Maine does not accurately represent the true number of rabies PEP given statewide.

Maine CDC considers a case of rabies PEP when a person either receives a treatment recommendation, begins treatment, or both. In 2018, Maine reported 152 cases of rabies PEP. 96% (146 reported cases) went to a healthcare provider or emergency department to begin their treatment. Six individuals received recommendations, but did not begin rabies PEP. One individual refused treatment. One individual no longer needed treatment because the attacking animal was ruled rabies free. Four individuals were lost to follow-up.

Rabies PEP is considered a medical urgency, but not a medical emergency. In many cases, rabies PEP can wait until the response measure (either quarantine or testing) for the attacking animal is completed. Of the 146 individuals who began rabies PEP, 37% (57 individuals) received rabies PEP because the attacking animal could not be located after the exposure and 23% (34 individuals) received rabies PEP because the attacking animal tested positive or unsatisfactory for rabies. Other reasons for the administration for rabies PEP include, but are not limited to, possible exposures to bats, contact with mucous membranes, or a severe attack to the head or neck.

Rabies PEP is readily available, but often in short supply. For this reason, healthcare providers must individually evaluate each suspected human exposure to a potentially rabid animal. Rabies exposures are rarely clear-cut issues, so treatment decisions must consider a variety of factors. These factors include type of exposure, extent or severity of exposure, type of animal species involved, animal vaccination status, circumstances leading to exposure, availability of animal for confinement and observation or testing, and the epidemiology of rabies in the region.



Generally, Maine CDC will recommend rabies PEP in cases where an animal tests positive for rabies, a high-risk animal is unavailable for quarantine or testing, or when serious attacks occur to the head, face, or neck. This is not a hard-set rule, but a guide as most people who are exposed to a potentially rabid animal still require a thorough evaluation. 37% of the 152 exposures in 2018 occurred to a hand or finger while 11% occurred to a leg, 8% to an arm, and 7% to the head or neck. In 7% of cases, exposures occurred to two or more regions of the body. About 30% of exposure sites were unknown.

Of the 152 animal exposures, bats were by far the most frequent animal encountered accounting for almost 40% of all cases of rabies PEP. Wild animal encounters made up 69% of all exposures followed by stray animal exposures at 22% and owned animal exposures just under 9%.

Rabies PEP is typically not required when pets bite humans because pets are typically available for confinement and observation. In 2018, 13 individuals received rabies treatment after a bite from an owned pet. In 31% of these cases, the pet's owner fled with the pet after the attack and in another 31% because the pet died within the days of exposure. Other situations why people received treatment after an exposure to a pet include, but are not limited to, owners provoking their pets, people exposed by someone else's pet while on a trip or camping, or pet could not be identified. Maine CDC normally does not recommend rabies PEP for encounters with small rodents like mice and squirrels because it is generally understood that small mammals are highly unlikely to survive an attack from a commonly rabid animal. Thus, it is unlikely for this group of mammals to transmit the rabies virus. Only in one rabies PEP case was the animal type and status unknown.

Maine CDC and Maine Department of Agriculture, Conservation, and Forestry, together with the Maine Rabies Workgroup, produce the Maine Rabies Management Guidelines. To review these guidelines and other resources, please visit www.maine.gov/dhhs/rabies.

Cases of Rabies PEP by Animal Encountered Reported to Maine CDC, 2018

Status	Animal	Rabies PEP	
	Bat	60	
	Bear	1	
	Ermine	1	
NACT -	Fox	15	
Wild	Mouse	1	
	Raccoon	20	
	Skunk	4	
	Squirrel	3	
	Cat	4	
Owned	Dog	9	
	Cat	23	
Stray	Dog	10	
Unknown	Unknown	1	
Total		152	

EVALUATION OF THE MAINE VARICELLA SURVEILLANCE SYSTEM

BACKGROUND

In 2002, federal CDC encouraged states to develop case-based varicella surveillance systems by 2005 to help monitor changing trends in varicella epidemiology following the implementation of a routine varicella immunization schedule. Varicella surveillance aims to document the impact of varicella vaccine on varicella related morbidity and mortality, evaluate the effectiveness of prevention strategies, and evaluate vaccine effectiveness.

The Maine varicella surveillance system relied on passive case reports from healthcare providers, school nurses, and laboratories. Maine investigated all cases of varicella reported by a provider and all hospitalized or invasive cases of varicella zoster in patients over the age of 18 (determined by lab specimen source and/or patient status in lab report). Maine does not investigate other laboratory reports of varicella zoster in patients over the age of 18 but counts them as cases in the surveillance system. Maine classifies those with PCR or gM positive labs as cases. Maine considers these cases of varicella to be "non-investigated varicella cases".

METHODS

In 2018, the Maine vaccine preventable disease (VPD) epidemiologist reviewed varicella surveillance as part of the VPD program evaluation. The VPD epidemiologist validated every reporting source and created a process to test the accuracy of case classification for non-investigated varicella cases. The VPD epidemiologist pulled all non-investigated cases of varicella from Maine's electronic case database. An epidemiologist followed up on selected cases by calling the associated physician to get clinical information about the case and determine if the patient met the surveillance case classification for varicella.

RESULTS

Since 2012, Maine averaged 117 non-investigated varicella cases per year. The proportion of non-investigated varicella cases averaged 57% of all cases (Table 1).

Table 1. Investigated and non-investigated varicella cases.	by yoar

	2012	2013	2014	2015	2016	2017	2018	
Investigated varicella cases	170	47	58	92	113	85	132	
	(65.9 %)	(33.6 %)	(28.0 %)	(39.5 %)	(49.6 %)	(42.9 %)	(52.4 %)	
Non-investigated varicella cases	88	93	149	141	115	113	120	
	(34.1 %)	(66.4 %)	(72.0 %)	(60.5 %)	(50.4 %)	(57.1 %)	(47.6 %)	
Total varicella cases	258	140	207	233	228	198	252	
	(100%)	(100%)	(100%)	(100%)	(100%)	(100%)	(100%)	



In 2018, the VPD epidemiologist identified 120 non-investigated varicella cases and randomly selected 63 cases for follow-up. None of the 63 cases investigated met the varicella case definition. Forty-six of the 63 cases (73%) had shingles, six (10%) had titers drawn to check for immunity, and ten (16%) had other non-varicella diagnoses without a diffuse rash.

CONCLUSIONS

The results of the investigation showed that classifying cases as varicella based only on laboratory results was inappropriate. Therefore, as of January 1, 2019, epidemiologists will review and document all varicella PCR positive or IgM positive laboratory results reported in adults >18 years but will not investigate or count the reports as cases unless providers specifically report the patients as cases or varicella-related deaths. This will greatly impact Maine varicella data, especially when comparing to previous years, but will improve accuracy and the Maine varicella surveillance system will reflect a better understanding of the true burden of varicella in the population.

INFLUENZA EDUCATION AMONG YOUTH IN AGRICULTURE

During the 2017-2018 influenza season Maine participated in a federally funded project aimed at educating youth in agriculture on influenza and other zoonotic diseases. This was done as a collaboration between Maine Center for Disease Control and Prevention (Maine CDC) and Maine Department of Agriculture, Conservation, and Forestry (Maine DACF). The goal of this project was to educate youth about zoonotic infections, deliver prevention messages to keep people and animals safe and healthy, and strengthen One Health networks among state human and animal health departments and agricultural communities.





TRAIN-THE-TRAINER EVENT:

In January 2018, Maine CDC and Maine DACF held a train-the-trainer event for 39 youth agricultural leaders. Attendees included representatives from 14 out of 16 counties in Maine and representatives from 22 out of the 25 Maine annual agricultural fairs.

This event provided direct instruction to youth agriculture leaders about diseases that can be transmitted between animals and humans, signs and symptoms of the diseases, and prevention strategies to reduce the risk of transmission. Maine CDC and Maine DACF delivered this information through presentations, interactive activities and group discussions. Maine encouraged attendees to teach and share the information with their local youth agriculture groups after the train-the-trainer event.

As of July 1, 2018, 37 of the 39 (95%) participants completed or planned a teaching event for youth. Of the completed events, the average number of participants was 22 with a range of 6-60 with over 700 estimated total participants.

Events completed or planned

Total Participants

Maine Center for Disease Control and Prevention

NOTIFIABLE DISEASES AND CONDITIONS LIST

24 Hours A Day, 7 Days A Week Disease Reporting:

Telephone: 1-800-821-5821 Fax: 1-800-293-7534

Conditions are reportable immediately by telephone on recognition or strong suspicion of disease

All others are reportable by telephone, fax, electronic lab report, or mail within 48 hours of recognition or strong suspicion of disease

→ ☑ Directors of laboratories are to submit isolates or clinical specimens, as well as any isolates or clinical specimens as requested by Maine CDC, to the Maine Health and Environmental Testing Laboratory for confirmation, typing, and/or antibiotic sensitivity

	Acid-Fast Bacillus → ⊠		Malaria
	Acquired Immunodeficiency Syndrome (AIDS)	~	Measles → ⋈ (Rubeola virus)
	Anaplasmosis		Meningococcal Disease, invasive → ⋈ (Neisseria meningitidis)
A	Anthrax → ⊠ (Bacillus anthracis)		Mumps → ⊠
	Babesiosis	~	Pertussis
	Botulism → ☑ (Clostridium botulinum)		Plague → ⊠ (Yersinia pestis)
	Brucellosis → 区 (Brucella species)		Poliomyelitis → ⋈ (Polio virus)
	California Serogroup Viruses		Powassan Virus
	Campylobacteriosis		Psittacosis
	Carbapenem-resistant Enterobacteriaceae (CRE)	~	Q Fever
	Carbon Monoxide Poisoning ²	200	Rabies (human and animal) → ☑ (Rabies virus)
	Chancroid		Rabies Post-Exposure Prophylaxis
	Chlamydia	~	Ricin Poisoning → ⊠
	Chickenpox (Varicella)	~	Rubella (including congenital) → ☑ (Rubella virus)
	Chikungunya		Salmonellosis → ☑ (Salmonella species)
A	Coronavirus, Novel and SARS → ⊠	~	Shellfish Poisoning
	Creutzfeldt-Jakob disease, <55 years of age		Shigellosis → ⊠ (Shigella species)
	Cryptosporidiosis		Smallpox → ☑ (Variola virus)
	Cyclosporiasis		Spotted Fever Rickettsiosis
	Dengue		St. Louis Encephalitis
æ	Diphtheria → ☑ (Corynebacterium diphtheriae)		Staphylococcus aureus, Methicillin-Resistant (MRSA), invasive
	E. coli, Shiga toxin-producing (STEC) → ⊠	~	Staphylococcus aureus with resistance to Vancomycin (VRSA) → ⊠
	Eastern Equine Encephalitis		Streptococcus Group A, invasive
	Ehrlichiosis		Streptococcus pneumoniae, invasive
	Giardiasis		Syphilis
	Gonorrhea	~	Tetanus → ⊠ (Clostridium tetani)
	Haemophilus influenzae, invasive → ⊠		Trichinosis
	Hantavirus, pulmonary and non-pulmonary syndromes		Tuberculosis (active and presumptive) → ⊠ (Mycobacterium tuberculosis)
	Hemolytic-uremic syndrome (post-diarrheal)	~	Tularemia → ☑ (Francisella tularensis)
	Hepatitis A, B, C, D, E (acute)		Vibrio species, including Cholera → ⊠ (Vibrio species)
	Hepatitis B, C, D (chronic)	**	Viral Hemorrhagic Fever
	Human Immunodeficiency Virus (HIV) 3		West Nile Virus
	Influenza-associated pediatric death		Western Equine Encephalitis
~	Influenza A, Novel → ⊠		Yellow Fever
	Influenza-associated hospitalizations, laboratory-confirmed		
	Legionellosis	*** **	Any Case of Unusual Illness of Infectious Cause
	Leptospirosis	*** **	Any Cluster/Outbreak of Illness with Potential Public Health Significance
	Listeriosis → ⊠ (Listeria monocytogenes)		
	Lyme Disease		
Mho	must report: Health Care Providers Medical Laboratories		Footnotes:

Health Care Facilities, Administrators, Health Officers, Veterinarians

What to report: Disease reports must include as much of the following as is known:

- Disease or condition diagnosed or suspected
- Patient's name, date of birth, address, phone number, occupation, race, and ethnicity
- Diagnostic laboratory findings and dates of test relevant to the notifiable condition
- Health care provider name, address and phone number
- Name and phone number of person making the report

- 1. Carbapenem-resistant Enterobacteriaceae (CRE): See current definition as adopted by the United States Centers for Disease Control and Prevention
- 2. Carbon Monoxide, including clinical signs, symptoms or known exposure consistent with diagnosis of carbon monoxide poisoning and/or: a carboxyhemoglobin (COHb) level >5%
- 3. Human Immunodeficiency Virus (HIV), including:
 - Confirmed, positive antibody tests
 - Viral load tests, all results
 - CD4 lymphocyte counts, all results



Maine Center for Disease Control and Prevention

Department of Health and Human Services

Complete Rules for the Control of Notifiable Diseases and Conditions:

http://www.maine.gov/dhhs/mecdc/infectious-disease/epi/disease-reporting/index.shtml



Department of Health and Human Services Maine Center for Disease Control and Prevention

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