# WEEKLY EPIDEMIOLOGY BULLETIN NATIONAL EPIDEMIOLOGY UNIT, MINISTRY OF HEALTH & WELLNESS, JAMAICA

# The Plague

Plague is an infectious disease caused by the bacteria *Yersinia pestis*, usually found in small mammals and their fleas. The disease is transmitted between animals via their fleas and, as it is a zoonotic bacteria, it can also transmit from animals to humans.

Humans can be contaminated by the bite of infected fleas, through direct contact with infected materials, or by inhalation. Plague can be a very severe disease in people, particularly in its septicaemic and pneumonic forms, with a case-fatality ratio of 30% - 100% if left untreated.

Although plague has been responsible for widespread pandemics throughout history, including the so-called Black Death that caused over 50 million deaths in Europe during the fourteenth century, today it can be easily treated with antibiotics and the use of standard preventative measures.

Plague is found on all continents except Oceania but most human cases since the 1990s have occurred in Africa. Democratic Republic of Congo, Madagascar and Peru are the three most endemic countries.

People infected with plague usually develop influenza-like symptoms after an incubation period of 3–7 days. Symptoms include fever, chills, aches, weakness, vomiting and nausea.

There are 3 main forms of plague.

Bubonic plague is the most common and is caused by the bite of an infected flea. The plague bacillus, *Y. pestis*, enters at the bite and travels to the nearest lymph node to replicate. The lymph node becomes inflamed, tense and painful, and is called a bubo. With advanced infections, the inflamed lymph nodes can turn into suppurating open sores. Bubonic plague cannot be transmitted from human to human.

Septicaemic plague occurs when infection spreads through the bloodstream. It may result from flea bites or from direct contact with infective materials through cracks in the skin. Advanced stages of the bubonic form of plague will also lead to direct spread of *Y. pestis* in the blood.

Pneumonic plague – or lung-based plague – is the most virulent and least common form of plague. Typically, it is caused by spread to the lungs from advanced bubonic plague. However, a person with secondary pneumonic plague may form aerosolized infective droplets and transmit plague to other humans. This is usually fatal.

### The plague: a profile

1 3 1			
Region: China	Transmission medium: Infected rodents and fleas		
-	Plague affects rodents, such as rats. People are most commonly infected by being bitten by a flea that is infected with the plague bacteria. The pneumonic form of the plague can be transmitted by		
	cough droplets.		
The first recorded pandemic	Incubation: 2-6 days		
was the Justinian Plague, which began in 541 AD.	Someone infected through the air could become ill within 1 to 3 days.		
Origin: Infected rodents	Transmission rate: Ro of 1.3 for pneumonic plague		
	$R_0\left(\text{basic reproduction number}\right)$ is an approximate measure of how many new infections one person will generate during their infectious period.		
black rats and fleas	Fatality ratio: 8-10% (It was over 60% pre antibiotics)		
Infection agent: Bacteria	Death toll: Over 100 million deaths		
<b>Sec.</b>	Medication status: No vaccine		
Bacteria Yersinia pestis	Plague vaccines are in development but are not expected to be commercially available in the immediate future.		
O theconversation.com	Sources: WHO & CDC		



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SENTINEL SYNDROMIC SURVEILLANCE Sentinel Surveillance in





**Parish health departments** submit reports weekly by 3 p.m. on Tuesdays. **Reports submitted after 3** p.m. are considered late.

A syndromic surveillance system is good for early detection of and response to public health events.

Sentinel surveillance occurs when selected health facilities (sentinel sites) form a network that reports on certain health conditions on a regular basis, for example, weekly. Reporting is mandatory whether or not there are cases to report.

Jamaica's sentinel surveillance system concentrates on visits to sentinel sites for health events and syndromes of national importance which are reported weekly (see pages 2 -4). There are seventy-eight (78) reporting sentinel sites (hospitals and health centres) across Jamaica.



# **REPORTS FOR SYNDROMIC SURVEILLANCE**

#### **FEVER**

Temperature of >38°C /100.4<sup>o</sup>F (or recent history of fever) with or without an obvious diagnosis or focus of infection.



### KEY VARIATIONS OF **BLUE** SHOW CURRENT WEEK







All clinical sites



**INVESTIGATION REPORTS-** Detailed Follow up for all Class One Events



HOSPITAL ACTIVE SURVEILLANCE-30 sites. Actively pursued



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# FEVER AND NEUROLOGICAL

Temperature of >38°C /100.4°F (or recent history of fever) in a previously healthy person with or without headache and vomiting. The person must also have meningeal irritation, convulsions, altered consciousness, altered sensory manifestations or paralysis (except AFP).



### FEVER AND HAEMORRHAGIC

Temperature of  $>38^{\circ}C$ /100.4°F (or recent history of fever) in a previously healthy person presenting with at least one haemorrhagic (bleeding) manifestation with or without jaundice.



### **FEVER AND JAUNDICE**

Temperature of  $>38^{\circ}C/100.4^{\circ}F$ (or recent history of fever) in a previously healthy person presenting with jaundice.

The epidemic threshold is used to confirm the emergence of an epidemic in order to implement control measures. It is calculated using the mean reported cases per week plus 2 standard deviations.





Weekly visits to Sentinel Sites for Fever and Haemorrhagic 2019 and 2020 vs Weekly Threshold; Jamaica





3 NOTIFICATIONS-All clinical sites INVESTIGATION REPORTS- Detailed Follow up for all Class One Events HOSPITAL ACTIVE SURVEILLANCE-30 sites. Actively pursued





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# CLASS ONE NOTIFIABLE EVENTS

## Comments

	CLASS 1 EVENTS		Confirmed YTD		AFP Field Guides
			CURRENT YEAR 2020	PREVIOUS YEAR 2019	from WHO indicate that for an effective surveillance system, detection rates for AFP should be 1/100,000 population under 15 years old (6 to 7) cases annually.
AL	Accidental Poisoning		5	21	
EST	Cholera		0	0	
	Dengue Hemorrhagic Fever*		NA	NA	
	Hansen's Disease (Leprosy)		0	0	
INT	Hepatitis B		0	11	
AL /	Hepatitis C		0	2	Pertussis-like
ATION	HIV/AIDS		NA	NA	syndrome and Tetanus are clinically confirmed classifications.
	Malaria (Imported)		0	0	
Z	Meningitis (Clinically confirmed)		1	10	
EXOTIC/ UNUSUAL	Plague		0	0	* Dengue Hemorrhagic Fever
H IGH MORBIDIT/ MORTALIY	Meningococcal Meningitis		0	0	data include Dengue related deaths;
	Neonatal Tetanus		0	0	
	Typhoid Fever		0	0	** Figures include
	Meningitis H/Flu		0	0	all deaths associated with pregnancy reported for the period. * 2019 YTD figure was updated.
SPECIAL PROGRAMMES	AFP/Polio		0	0	
	Congenital Rubella Syndrome		0	0	
	Congenital Syphilis		0	0	
	Fever and Rash	Measles	0	0	positive cases
		Rubella	0	0	
	Maternal Deaths**		21	29	**** Zika PCR positive cases
	Ophthalmia Neonatorum		23	105	
	Pertussis-like syndrome		0	0	
	Rheumatic Fever		0	0	-
	Tetanus		0	0	-
	Tuberculosis		0	27	
	Yellow Fever		0	0	
	Chikungunya***		0	0	
	Zika Virus <sup>****</sup>		0	0	NA- Not Available



All clinical sites



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HOSPITAL ACTIVE SURVEILLANCE-30 sites. Actively pursued



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# NATIONAL SURVEILLANCE UNIT INFLUENZA <u>REPORT</u>

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NOTIFICATIONS-All clinical sites



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HOSPITAL ACTIVE SURVEILLANCE-30 sites. Actively pursued



# **Dengue Bulletin**

#### June 21, 2020-June 27, 2020 Epidemiological Week 26

Epidemiological Week 26







#### Critical phase hypotension pleural effusion ascites

gastrointestinal bleeding

#### **Recovery phase**

altered level of consciousness

seizures

itchina

# slow heart rate

#### Suspected dengue cases for 2018 and 2019 versus monthly mean, alert, and epidemic thresholds



#### Points to note:

- \*\* figure as at July 3, 2020
- **Only PCR positive dengue cases** are reported as confirmed.
- IgM positive cases are classified as presumed dengue.



7 NOTIFICATIONS-All clinical sites



**INVESTIGATION REPORTS-** Detailed Follow up for all Class One Events

SURVEILLANCE-30 sites. Actively pursued



# **RESEARCH PAPER**

#### ABSTRACT

#### Title: The Use of Breadfruit-based Media to Improve the Turnaround Time and Identification of Fungal Specimen.

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Jamaica

**Objective:** To determine the effectiveness of a breadfruit-based media (BFM) for the enhancement of sporulation, growth and identification of fungal pathogens; a feat that would improve the turnaround time currently observed at the mycology laboratory at the University of the West Indies (UWI).

**Methods:** The BFM was pre-prepared using sterile techniques and inoculated with a total of 25 previously identified fungal clinical isolates (eg. *Trichophyton* spp., *Fusarium* spp, *Chaetominum* spp, *Bipolaris* sp, *Curvalaria* sp, and *Aspergillus flavus*). For the purposes of quality control ATTC strains of *E. coli* and *Candida albicans* were inoculated unto the media following standard microbiological procedures. All 27 species were also inoculated unto other standard media in use in the laboratory to allow for observation and comparison of the key features ie: enhancements to growth rate, sporulation characteristics, texture, colour etc. The isolates from resulting cultures were then identified using routine mycological tests. The observer was blinded as to the type of media in use.

Results: All 27 species of organisms grew within 18-48 hours and showed enhanced characteristic features.

**Conclusion**: Breadfruit, a sustainable Jamaican food staple, when prepared appropriately, can be used to supplement media for enhanced fungal isolation and identification. BFM proved to be a superior media that facilitated improved turnaround time, positioning itself as a possible industrial asset to the health sector. Further studies are needed to assess its capacity for improved isolation and identification of bacterial pathogens.



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NOTIFICATIONS All clinical sites



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