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

Diphtheria

- Diphtheria is an infectious disease caused by the bacterium Corynebacterium diphtheria, which primarily infects the throat and upper airways, and produces a toxin affecting other organs. The illness has an acute onset and the main characteristics are sore throat, low fever and swollen glands in the neck, and the toxin may, in severe cases, cause myocarditis or peripheral neuropathy. The diphtheria toxin causes a membrane of dead tissue to build up over the throat and tonsils, making breathing and swallowing difficult. The disease is spread through direct physical contact or from breathing in the aerosolized secretions from coughs or sneezes of infected individuals.
- Vaccination against diphtheria has reduced the mortality and morbidity of diphtheria dramatically, however diphtheria is still a significant child health problem in countries with poor EPI coverage. In countries endemic for diphtheria, the disease occurs mostly as sporadic cases or in small outbreaks. Diphtheria is fatal in 5 - 10% of cases, with a higher mortality rate in young children. Treatment involves administering diphtheria antitoxin to neutralize the effects of the toxin, as well as antibiotics to kill the bacteria.
- Diphtheria vaccine is a bacterial toxoid, ie. a toxin whose toxicity has been inactivated. The vaccine is normally given in combination with other vaccines as DTwP/DTaP vaccine or pentavalent vaccine. For adolescents and adults the diphtheria toxoid is frequently combined with tetanus toxoid in lower concentration (Td vaccine).
- WHO recommends a 3-dose primary vaccination series with diphtheria containing vaccine • followed by 3 booster doses. The primary series should begin as early as 6-week of age with subsequent doses given with a minimum interval of 4 weeks between doses. The 3 booster doses should preferably be given during the second year of life (12-23 months), at 4-7 years and at 9-15 years of age. Ideally, there should be at least 4 years between booster doses.
- To further promote immunity against diphtheria, combined diphtheria and tetanus toxoid vaccine (Td or TD) should be used rather than tetanus toxoid alone. This can be used in pregnancy as well as following injuries.

WEEK 33 SYNDROMES PAGE 2 CLASS 1 DISEASES PAGE 4 INFLUENZA PAGE 5 **DENGUE FEVER** PAGE 6 GASTROENTERITIS PAGE 7 **RESEARCH PAPER**

PAGE 8

DIPHTHERIA

- A bacterial infection usually affecting the mucous membranes of nose and throat In advanced stages, diphtheria can damage heart, kidneys, nervous system SYMPTOMS A thick, gray membrane covering throat, tonsils A sore throat and hoarseness Swollen glands (enlarged)
- lymph nodes) in neck Difficulty in breathing or rapid breathing
- Nasal discharge Fever and chills



- Contaminated personal items Contaminated
- household items We have been shocked by

the Koottayi case. We will monitor the neighbourhood and distribute medicines and vaccines." **District Deputy Medical Officer**

Released August 28, 2020

SENTINEL SYNDROMIC SURVEILLANCE Sentinel Surveillance in





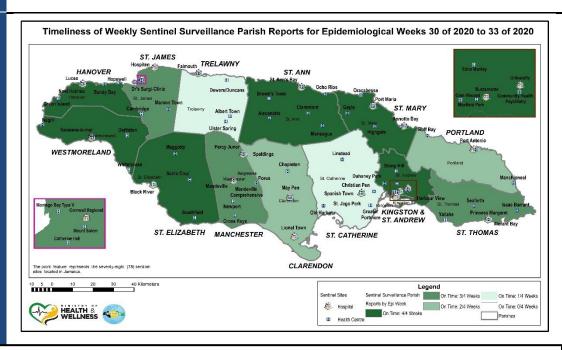
Timeliness of Weekly Sentinel Surveillance Parish Reports for the Four Most Recent Epidemiological Weeks -30 to 33 of 2020

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.



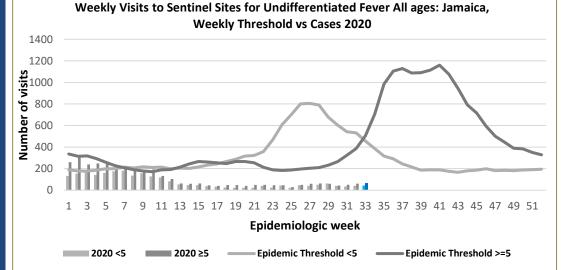
REPORTS FOR SYNDROMIC SURVEILLANCE

FEVER

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



KEY VARIATIONS OF **BLUE** SHOW CURRENT WEEK



2 NOTIFICATIONS-All clinical sites



INVESTIGATION REPORTS- Detailed Follow up for all Class One Events



HOSPITAL ACTIVE SURVEILLANCE-30 sites. Actively pursued



Released August 28, 2020

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

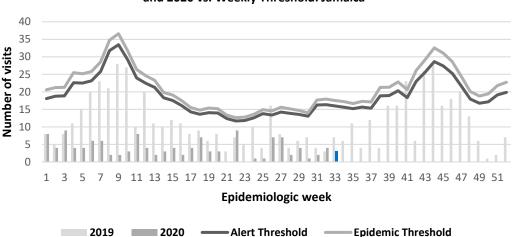
Temperature of $>38^{\circ}C/100.4^{\circ}F$ (or recent history of fever) in a previously healthy person presenting with jaundice.

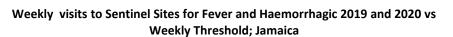
The epidemic threshold is used to confirm the emergence of an epidemic in order to implement control measures. It is calculated using the mean reported cases per week plus 2 standard deviations.

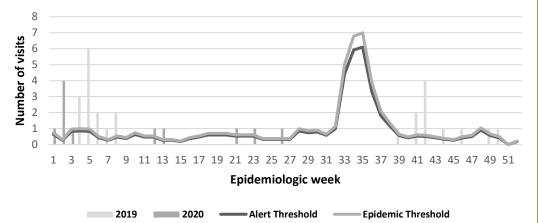


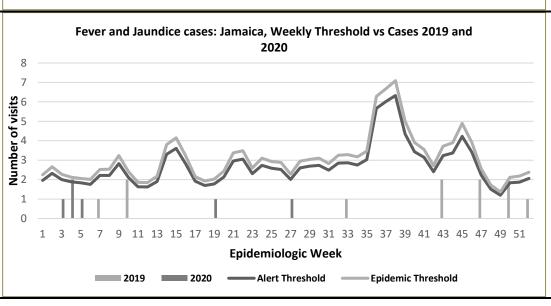
Weekly Visits to Sentinel Sites for Fever and Neurological Symptoms 2019 and 2020 vs. Weekly Threshold: Jamaica

ISSN 0799-3927











3 NOTIFICATIONS-All clinical sites

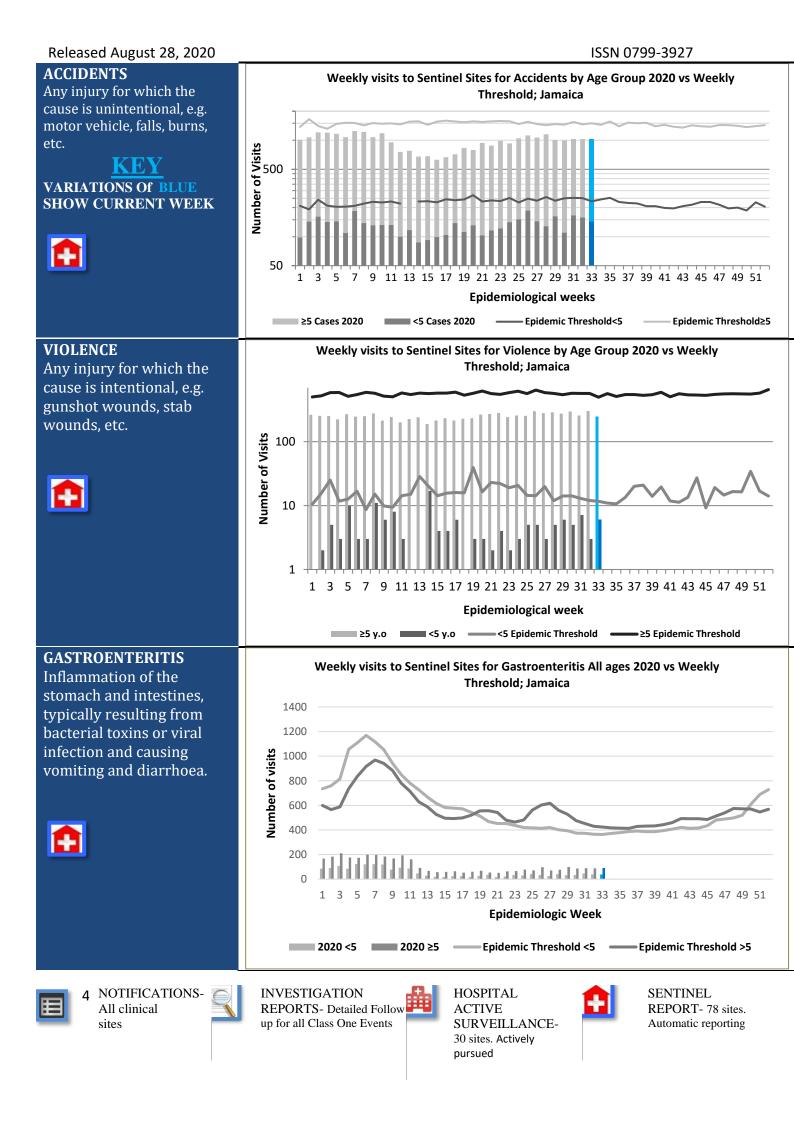


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HOSPITAL ACTIVE SURVEILLANCE-30 sites. Actively pursued





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CLASS ONE NOTIFIABLE EVENTS

Comments

			Confirmed YTD		AFP Field Guides
	CLASS 1 EVENTS		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	40	
NATIONAL /INTERNATIONAL INTEREST	Cholera		0	0	
	Dengue Hemorrhagic Fever*		NA	NA	
	Hansen's Disease (Leprosy)		0	0	
	Hepatitis B		0	11	
	Hepatitis C		0	2	Pertussis-like syndrome and Tetanus are clinically confirmed classifications.
	HIV/AIDS		NA	NA	
	Malaria (Imported)		0	0	
	Meningitis (Clinically confirmed)		1	15	
EXOTIC/ UNUSUAL	Plague		0	0	 * Dengue Hemorrhagic Fever data include Dengue related deaths; ** Figures include all deaths associated with pregnancy reported for the period. * 2019 YTD figure was updated. *** CHIKV IgM
H IGH MORBIDIT/ MORTALIY	Meningococcal Meningitis		0	0	
	Neonatal Tetanus		0	0	
	Typhoid Fever		0	0	
	Meningitis H/Flu		0	0	
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**		28	42	PCR positive cases
	Ophthalmia Neonatorum		23	161	
	Pertussis-like syndrome		0	0	
	Rheumatic Fever		0	0	
	Tetanus		0	0	
	Tuberculosis		6	33	
	Yellow Fever		0	0	
	Chikungunya ^{***}		0	0	
	Zika Virus ^{****}		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>

EW 33

ISSN 0799-3927

August 09, 2020-August 15, 2020 Epidemiological Week 33 Weekly visits to Sentinel Sites for Influenza-like Illness (ILI) All EW 33 **YTD** ages 2020 vs Weekly Threshold; Jamaica SARI cases 18 393 2500 Number of visits **Total** 2000 Influenza 0 69 1500 positive Samples 1000 45 Influenza A 0 0 4 H3N2 500 H1N1pdm09 0 38 0 0 3 Not subtyped 7 9 11 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 51 3 5 1 **Epidemiologic week** Influenza B 0 24 2020 2020 2020 Parainfluenza 0 0 Epidemic Threshold <5 Epidemic Threshold 5-59 ■ Epidemic Threshold ≥60 **Epi Week Summary** Jamaica: Percentage of Hospital Admissions for Severe Acute Respiratory Illness (SARI 2020) (compared with 2011-2019) During EW 33, 18 (eighteen) Percentage of hospitalizations for SARI 6.0% SARI admissions were reported. 5.0% 4.0% 3.0% 2.0% 1.0% 0.0% 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 51 1 3 13 **Epidemiological Week** SARI 2020 Alert Threshold Average epidemic curve (2011-2019) Seasonal Threshold Epidemic Threshold Caribbean Update EW 33 DISTRIBUTION OF INFLUENZA AND OTHER **RESPIRATORY VIRUSES IN SURVEILLANCE BY EW** Caribbean: Influenza and other 15 110% **NUMBER OF POSITIVE SAMPLES** 100% respiratory virus activity 90% 80% PERCENT POSITIVITY remained low in the subregion. 10 70% In Haiti and Suriname, detections 60% 50% 40% of SARS-CoV-2 continue elevated 5 30% and increasing. 20% 10% 0 0% 1 3 5 9 11131517192123252729313335373941434547495153 EPIDEMIOLOGIC WEEK A not subtyped A(H1N1)pdm09 A no subty A(H1) A(H3) Metha Parainfluenza RSV Adenovirus Bocavirus Rhinovirus Cord



6

NOTIFICATIONS-All clinical sites



INVESTIGATION REPORTS- Detailed Follow up for all Class One Events



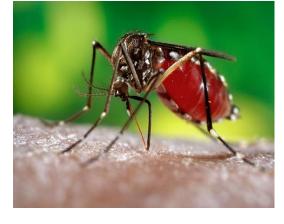
HOSPITAL ACTIVE SURVEILLANCE-30 sites. Actively pursued

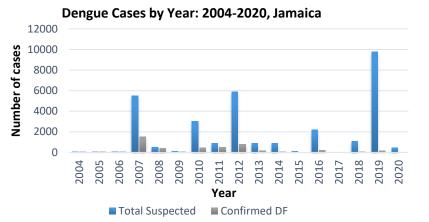


Dengue Bulletin

August 09, 2020-August 15, 2020 Epidemiological Week 33

Epidemiological Week 33





Reported suspected and confirmed dengue with symptom onset in week 33 of 2020 2020 EW YTD 33 **Total Suspected Dengue** 736** 0** Cases Lab Confirmed Dengue 1** 0** cases

0**

** figure as at August 17, 2020

Only PCR positive dengue cases

IgM positive cases are classified

are reported as confirmed.

as presumed dengue.

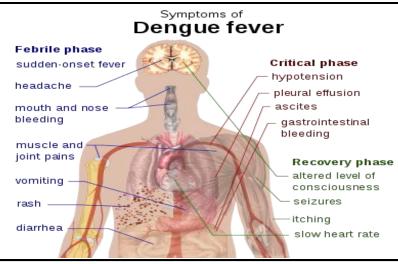
1**

CONFIRMED

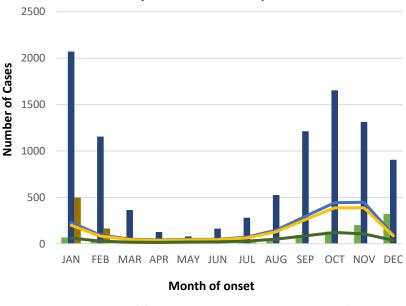
Dengue Related Deaths

Points to note:

0



Suspected dengue cases for 2018, 2019 and 2020 versus monthly mean, alert, and epidemic thresholds



2018 suspected dengue

2019 Suspected Dengue **Epidemic threshold**

sites



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2020

HOSPITAL ACTIVE SURVEILLANCE-30 sites. Actively pursued



RESEARCH PAPER

The Occurrence of Chronic Sorrow and Coping Strategies Employed by Adult **Oncology Patients in Western Jamaica**

VH Waugh-Brown, GA Wright

The University of the West Indies, Mona, Kingston, Jamaica Email Address: veronica.waughbrown02@uwimona.edu.jm or veraugh@gmail.com

Objective: To explore the occurrence of chronic sorrow and describe the coping strategies used by patients diagnosed with cancer.

Method: A phenomenological study was conducted among adult patients attending oncology clinic in western Jamaica. Purposive sampling was used to select eight participants who met the criteria for a Focus Group Discussion. Informed consent and demographic data were obtained. A Focus Group Discussion Guide aided the exploration of participants' feelings and coping mechanisms. The discussion was audiotaped. Data were transcribed verbatim and checked for accuracy. Common themes were connected, inter-relationships identified and narrative constructed.

Results: Eight persons diagnosed with cancer and receiving treatment at the Oncology Clinic participated in the focus group discussion. The chronicity of the illness, negative shift in the equilibrium of life and financial challenges caused major stress which contributed to chronic sorrow. Strong spiritual belief was the major common element expressed that helped persons to cope. Keeping physically active and volunteerism were other coping mechanisms that emerged. Participants with greater family and financial supports expressed greater ability to cope with the illness than those with poor family or financial support. Psychological / emotional therapy from a professional source was lacking.

Conclusion: Persons diagnosed with cancer experience chronic sorrow resulting from emotional strain and stress. Spiritual and psychological support forms the bed-rock of their mental well-being and coping ability. The magnitude of the impact of chronic sorrow experienced by cancer patients can be reduced by integrating these critical components in the patient's medical management plan.



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All clinical sites



INVESTIGATION REPORTS- Detailed Follow up for all Class One Events



HOSPITAL ACTIVE SURVEILLANCE-30 sites. Actively pursued

