

# **WEEKLY EPIDEMIOLOGICAL REPORT**

# A publication of the Epidemiology Unit Ministry of Health

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Antimicrobial Resistance-(Part I)

## Vol. 42 No. 05

## 24<sup>th</sup> – 30<sup>th</sup> January 2015

This is the first in a series of two article on antimicrobial resistance.

### Key facts

Antimicrobial resistance (AMR) threatens the effective prevention and treatment of an ever-increasing range of infections caused by bacteria, parasites, viruses and fungi.

It is an increasingly serious threat to global public health that requires action across all government sectors and society.

AMR is present in all parts of the world. New resistance mechanisms emerge and spread globally.

In 2012, there were about 450 000 new cases of multidrug-resistant tuberculosis (MDR-TB). Extensively drug-resistant tuberculosis (XDR-TB) has been identified in 92 countries. MDR-TB requires treatment courses that are much longer and less effective than those for non-resistant TB.

- Resistance to earlier generation antimalarial drugs is widespread in most malariaendemic countries. Further spread, or emergence in other regions, of artemisininresistant strains of malaria could jeopardize important recent gains in control of the disease.
- There are high proportions of antibiotic resistance (ABR) in bacteria that cause common infections (e.g. urinary tract infections,

pneumonia, bloodstream infections) in all regions of the world. A high percentage of hospital-acquired infections are caused by highly resistant bacteria such as methicillinresistant *Staphylococcus aureus* (MRSA) or multidrug-resistant Gram-negative bacteria.

- Treatment failures due to resistance to treatments of last resort for gonorrhoea (third-generation cephalosporins) have now been reported from 10 countries. Gonorrhoea may soon become untreatable as no vaccines or new drugs are in development.
- Patients with infections caused by drugresistant bacteria are generally at increased risk of worse clinical outcomes and death, and consume more healthcare resources than patients infected with the same bacteria that are not resistant.

### What is antimicrobial resistance?

Antimicrobial resistance (AMR) is resistance of a microorganism to an antimicrobial drug that was originally effective for treatment of infections caused by it.

Resistant microorganisms (including bacteria, fungi, viruses and parasites) are able to withstand attack by antimicrobial drugs, such as antibacterial drugs (e.g., antibiotics), antifungals, antivirals, and antimalarials, so that standard treatments become ineffective and infections persist, increasing the risk of spread to others.

The evolution of resistant strains is a natural phenomenon that occurs when microorganisms

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replicate themselves erroneously or when resistant traits are exchanged between them. The use and misuse of antimicrobial drugs accelerates the emergence of drug-resistant strains. Poor infection control practices, inadequate sanitary conditions and inappropriate food-handling encourages the further spread of AMR.

# What is the difference between antibiotic and antimicrobial resistance?

Antibiotic resistance refers specifically to the resistance to antibiotics that occurs in common bacteria that cause infections. Antimicrobial resistance is a broader term, encompassing resistance to drugs to treat infections caused by other microbes as well, such as parasites (e.g. malaria), viruses (e.g. HIV) and fungi (e.g. *Candida*).

### Why is antimicrobial resistance a global concern?

New resistance mechanisms emerge and spread globally threatening our ability to treat common infectious diseases, resulting in death and disability of individuals who until recently could continue a normal course of life.

Without effective anti-infective treatment, many standard medical treatments will fail or turn into very high risk procedures.

### AMR kills

Infections caused by resistant microorganisms often fail to respond to the standard treatment, resulting in prolonged illness, higher health care expenditures, and a greater risk of death. As an example, the death rate for patients with serious infections caused by common bacteria treated in hospitals can be about twice that of patients with infections caused by the same non-resistant bacteria. For example, people with MRSA (methicillin-resistant *Staphylococcus aureus*, another common source of severe infections in the community and in hospitals) are estimated to be 64% more likely to die than people with a non-resistant form of the infection.

#### AMR hampers the control of infectious diseases

AMR reduces the effectiveness of treatment; thus patients remain infectious for a longer time, increasing the risk of spreading resistant microorganisms to others. For example, the emergence of *Plasmodium falciparum* resistance to artemisinin in the Greater Mekong subregion is an urgent public health concern that is threatening global efforts to reduce the burden of malaria. Although MDR-TB is a growing concern, it is still largely underreported, compromising control efforts.

### AMR increases the costs of health care

When infections become resistant to first-line drugs, more expensive therapies must be used. A longer duration of illness and treatment, often in hospitals, increases health care costs as well as the economic burden on families and societies.

#### AMR jeopardizes health care gains to society

The achievements of modern medicine are put at risk by AMR. Without effective antimicrobials for prevention and treatment of infections, the success of organ transplantation, cancer chemotherapy and major surgery would be compromised.

# AMR has the potential to threaten health security, and damage trade and economies

The growth of global trade and travel allows resistant microorganisms to spread rapidly to distant countries and continents through humans and food. Estimates show that AMR may give rise to losses in Gross Domestic Product of more than 1% and that the indirect costs affecting society may be more than 3 times the direct health care expenditures. It affects developing economies proportionally more than developed ones.

Name of Bacte- rium/resistance	Examples of typical diseases	No. out of 194 Member states providing data	No. WHO regions with national reports of 50% resistance or more			
Escherichia coli	Urinary tract infections, blood					
cephalosporins	Inections	86	5/6			
nolones		92	5/6			
Klebsiella pneu- moniae	Pneumonia, blood stream infections,					
-vs 3 <sup>rd</sup> generation	tions	87	6/6			
-vs fluoroqui- nolones		71	2/6			
Staphylococcus aureus	Wound infections, blood stream infections					
-vs methicil- lin"MRSA"		85	5/6			

Bacteria mainly causing infections in the community

#### Sources

WHO-antimicrobial resistance global report on surveillance, available at <u>http://www.who.int/mediacentre/factsheets/fs194/en/</u>

Compiled by Dr. C U D Gunasekara of the epidemiology unit.

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 Table 1: Selected notifiable diseases reported by Medical Officers of Health
 17th - 23rd
 Jan 2015 (04th Week)

# 24th-January 30th 2015

Source: Weekly Returns of Communicable Diseases (WRCD). •T=Timeliness refers to returns received on or before 23<sup>d</sup> January , 2015 Total number of reporting units 337 Number of reporting units data provided for the current week: 278 C\*\*\*-Completeness

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RDHS Division		Colombo	Gampaha	Kalutara	Kandy	Matale	NuwaraEliya	Galle	Hambantota	Matara	Jaffna	Kilinochchi	Mannar	Vavuniya	Mullaitivu	Batticaloa	Ampara	Trincomalee	Kurunegala	Puttalam	Anuradhapura	Polonnaruwa	Badulla	Monaragala	Ratnapura	Kegalle	Kalmune	SRILANKA	
Page	e 3																												

## 24th-January 30th 2015

## Table 2: Vaccine-Preventable Diseases & AFP

## 17th - 23rd Jan 2015 (04th Week)

Disease			N	lo. of Cas	es by P	rovince				Number of cases         Number of cases         Total number of during         Total num- ber of during         Differ betwee           during         during         cases to cases to         cases to cases to         number cases to           current         same         date in         date in         cases for								
	w	С	S	N	E	NW	NC	U	Sab	week in 2015	week in 2014	2015	2014	in 2014& 2015				
AFP*	00	00	00	00	00	00	00	00	00	00	0 01 06 04		04	-50%				
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0%				
Mumps	04	01	00	01	00	01	01	01	02	11	12	27	79	-65.9%				
Measles	04	03	08	01	00	05	05	02	04	32	74	74 108 362		-70.2%				
Rubella	00	00	00	00	00	00	00	00	00	00	00	02	00	0%				
CRS**	00	00	00	00	00	00	00	00	00	00	00	00	00	0%				
Tetanus	01	00	00	00	00	00	00	00	00	01	01	01	01	0%				
Neonatal Teta- nus	00	00	00	00	00	00	00	00	00	00	) 00 00		00	0%				
Japanese En- cephalitis	00	00	00	00	00	00	00	00	00	00	03	02	08	-75%				
Whooping Cough	01	00	00	01	00	00	00	00	00	02	02	08	03	+166.7%				
Tuberculosis	254	22	28	12	17	15	01	08	72	429	261	795	946	-16.1%				

### Key to Table 1 & 2

Provinces: W: Western, C: Central, S: Southern, N: North, E: East, NC: North Central, NW: North Western, U: Uva, Sab: Sabaragamuwa.

RDHS Divisions: CB: Colombo, GM: Gampaha, KL: Kalutara, KD: Kandy, ML: Matale, NE: Nuwara Eliya, GL: Galle, HB: Hambantota, MT: Matara, JF: Jaffna,

KN: Killinochchi, MN: Mannar, VA: Vavuniya, MU: Mullaitivu, BT: Batticaloa, AM: Ampara, TR: Trincomalee, KM: Kalmunai, KR: Kurunegala, PU: Puttalam, AP: Anuradhapura, PO: Polonnaruwa, BD: Badulla, MO: Moneragala, RP: Ratnapura, KG: Kegalle.

Data Sources:

Weekly Return of Communicable Diseases: Diphtheria, Measles, Tetanus, Neonatal Tetanus, Whooping Cough, Chickenpox, Meningitis, Mumps., Rubella, CRS, Special Surveillance: AFP\* (Acute Flaccid Paralysis), Japanese Encephalitis

CRS\*\* =Congenital Rubella Syndrome

AFP and all clinically confirmed Vaccine Preventable Diseases except Tuberculosis and Mumps should be investigated by the MOH

## **Dengue Prevention and Control Health Messages**

Look for plants such as bamboo, bohemia, rampe and banana in your surroundings and maintain them

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Comments and contributions for publication in the WER Sri Lanka are welcome. However, the editor reserves the right to accept or reject items for publication. All correspondence should be mailed to The Editor, WER Sri Lanka, Epidemiological Unit, P.O. Box 1567, Colombo or sent by E-mail to chepid@sltnet.lk. Prior approval should be obtained from the Epidemiology Unit before publishing data in this publication

# **ON STATE SERVICE**

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