Global PPS in South Africa

Surveillance of IPC and AMS metrics

Heather Finlayson
Paeds Infectious Diseases
Specialist TBH





What is a Point Prevalence Study

 Collection of data used to identify the number of people with a disease or condition at a specific point in time." CDC definition



• Easy to conduct, relatively inexpensive and are not time-consuming

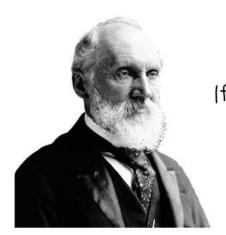
Incidence surveys, sequential are more difficult to perform and more expensive



Why do a PPS?

- Driven by the IPC task working group on the MAC on AMR
- Attempt to get good HAI data
 - Show burden of HAI
 - Buy in from managers
 - Set targets for improvement

- But the Global PPS gives us so much more data!
 - AMS and Quality Assurance
- Use data in order to bring about improvement



To measure is to know.

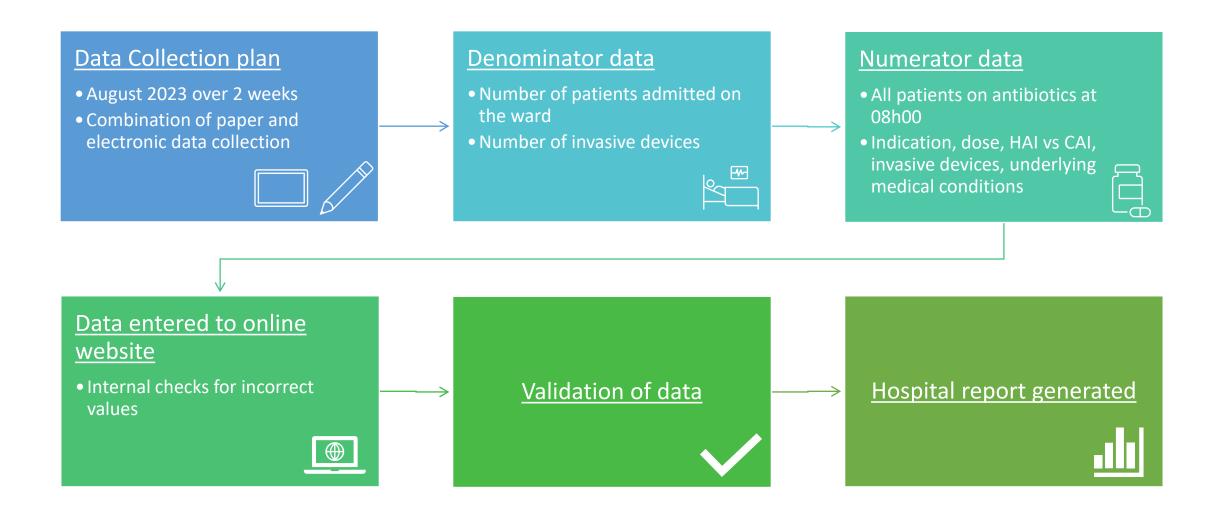
If you can not measure it, you can not improve it.

What is the Global PPS?



- Simple, freely available web-based tool https://www.global-pps.com/
- Measure and monitor antimicrobial prescribing and resistance in hospitals and healthcare centers worldwide.
 - ✓ Evaluate antimicrobial prescribing practices and survey performance indicators in healthcare centers (identify burden),
 - √ Help designing local interventions and identifying targets for quality improvement of antimicrobial prescribing and the prevention of Healthcare-Associated Infections (HAI) (change practice),
 - ✓ Assess the effectiveness of the interventions through repeated PPSs (measure impact).

How is a point prevalence study done?





HOME

PROJECT

DOCUMENTS

NEWS

DISSEMINATION *

EVENTS

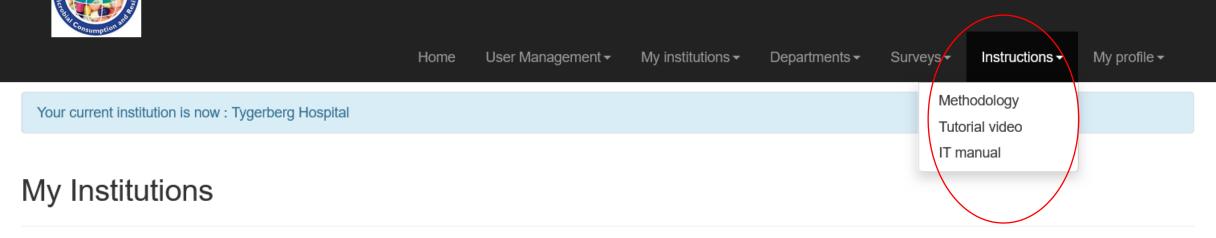
OUR NETWORK ▼

DRIVE-AMS

CONTACT US



Find institution...



IdNameE-mailCountry codeTypeAction374Tygerberg Hospitalfinlayson@sun.ac.zaZATertiary care hospital

Q

×

GLOBAL-PPS PATIENT Form – additional variables for HAI at patient level
(Fill in one form per patient with an active/ongoing antimicrobial at 8am on the day of the PPS – more info on definitions in protocol)

	Activity 1 Patient Age 4							(Current	Neonates (optional)		Sex					
Ward Name/code	(M, S, IC)	Patient Identi	fier ²	Sur	rvey Numb	oer ³		Years	1	Months	Days		Veight*		estatio- al age*	Birth weight*	M, F, U
	, , , ,							≥ 2 years	1-	-23 month	<1 mon	h		11	ur uye	weight	
Date of admission in (dd/mm/yyyy) (optio	•							Surgic hospit	_	ocedure du	ring curre	nt ad	mission	in	☐ Yes	□ No	□ UNK
Previous hospitalizat < 3 months (optional)		Yes, ICU	Yes, o	other	□ No		JNK	Previo	us a	ntibiotic co	urse < 1 n	nonth	(option	al)	☐ Yes	□ No	□ UNK
"Inserted" invasive device present at 8 am on the day of the PPS										Date 1 st inse	ertion/star	t		Cabe	☐ Non-fatal disease		
Indwelling Urinary Cathe	ter (UC)				□ Y	'es	☐ No							Ultimately fatal dis			
Peripheral Vascular / intr	avenous Cath	eter (PVC)			□ Y	'es	□ No	□ UN	K	_/_	_/	☐ Rapidly fatal disease			ease		
Central Vascular Cathete	r (CVC)				□ Y	'es	□ No	UN 🗆	K	_/_	_/	☐ UNK/Not available			ble		
Non-invasive pos. & neg.	mechanical ve	entilation (CPAP,	BiPAP, CN	EP,) ⁱ	□ Y	'es	☐ No	UN 🗆	K	_/_	_/						
Invasive respiratory endo	tracheal intub	oation (IRI) ⁱⁱ			□ Y	'es	□ No	UN 🗆	K	_/_	_/						
Inserted tubes and drains	s (T/D) ⁱⁱⁱ				□ Y	'es	□ No	UN 🗆	K	_/_	_/						
	1-																
Underlying morbidity	☐ Diabete	es mellitus, type 1	or 2			∐ G	enetic c	lisorder				∐ En	End-stage Liver Disease, cirrhosis				
(multiple choice,	☐ AIDS/H	IV (only if last CD	4 count <5	00/mm ³	3)	□ c ₀	ongenit	al heart dis	eases	5		☐ Tr	auma				
maximum 3 choices)		ological or solid ca therapy (<3montl		ent						luding cystic tasis, asthma					gical diseas s, Coeliac d	•	natory
	☐ Stem ce	ell or solid organ t	transplant			□ N	eutrope	enia				☐ Ch	ronic ne	urolo	gical condi	tions ^{iv}	
	Chronic	: Renal Disease (a	ll stages)			Пн	igh dose	e steroids ^v				Ot	her				
	☐ Active t	tuberculosis				□ м	lalnutrit	tion ^{vi}				□ No	one				
						Пьс	ong COV	/ID				Ur	nknown				

Participation 2023



Province	Validated data
Eastern Cape	5
Free State	3
Gauteng	4
KwaZulu-Natal	26
Limpopo	2
Mpumalanga	1
North West	1
Northern Cape	2
Western Cape	8
Total	52

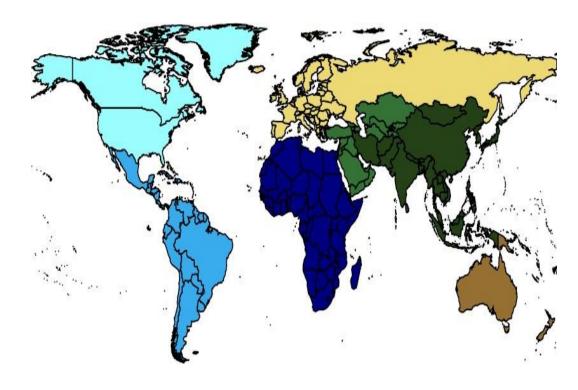


Additional 26 unvalidated 67% validation rate

Participation to Global-PPS by UN macro-geographical regions, year 2023

	Number of countries	Number of hospitals
North America	1	3
South America	2	2
Africa	10	71
Europe	6	17
West & Central Asia	1	2
East & South Asia	6	37
Australia & New Zealand	0	0

73% of Africa participation = South Africa





Latin America

Africa

■ West & Central Asia

■ East & South Asia

Europe

Australia & New Zealand

Prevalence of patients prescribed at least one antimicrobial on day of survey



	Co	untry	Co	ontinent		Eur	rope
		N	%	N	%	N	%
N admitted patients (=denominator)	1	7136		10766		16454	
N patients on antimicrobials	1	2300	32.2	4250	39.5	4888	29.7
N patients with antibacterials for systemic use		2226	31.2	4107	38.1	4687	28.5
N patients with antimycotics or antifungals for systemic use		168	2.4	197	1.8	226	1.4
N patients with drugs for treatment of tuberculosis		33	0.5	90	8.0	45	0.3
N patients with antivirals for systemic use		42	0.6	67	0.6	216	1.3
N patients with antibiotics used as intestinal anti-infectives		27	0.4	35	0.3	135	0.8
N patients with nitroimidazole derivatives	i i	50	0.7	288	2.7	57	0.3
N patients with antimalarials		0	0.0	55	0.5	6	0.0

Reference data: country -2023 (N = 44), continent -2023 (N = 71), EU -2022 (N = 67).



Antimicrobial prevalence (%) by activity

	Country	Continent	Europe
Adults			
Medical	28.8	34.1	26.3
Surgical	24.6	35.5	34.3
ICU	56.8	58.4	51.9
Children			
Medical	51.2	57.4	34.8
Surgical	20.9	35.8	33.9
ICU	78.4	80.4	64.2
Neonates			
GNMW	32.9	37.6	11.5
NICU	44.8	48.9	20.0

Antimicrobial prevalence = 100*(number of treated patients/number of admitted patients)

Antimicrobial prevalence by activity for adults, children and neonates separately for the hospital, country, continent to which the hospital belongs; and the continental results for the hospital type to which the hospital belongs (possible types are primary + secondary level, tertiary level, paediatric and infectious diseases + specialized hospital).

Country: SOUTH AFRICA; Continent: Africa; Hospital type:

Key prescription patterns (adults and children)

		Country		ontinent				Europe
	N	%	N	%			N	%
All patients								
IV therapy	2140	80.1	3791	78.3			3367	72.4
Multiple ATB diagnosis	744	26.5	1655	32.5	_	High IV was	531	11.1
Multiple ATB patient	829	31.0	1796	37.1	•	High IV use	654	14.1
Medical					•	25% more than		
IV therapy	1242	69.0	2127	69.8		one antibiotic for	1905	62.1
Multiple ATB diagnosis	485	28.4	910	31.4		a diagnosis	279	9.1
Multiple ATB patient	543	33.7	999	36.6	•	A third getting	367	12.4
Surgical						more than one		
IV therapy	670	79.0	1370	71.3		antibiotic	1075	81.3
Multiple ATB diagnosis	181	21.5	624	33.5		antiblotic	161	12.3
Multiple ATB patient	195	23.7	659	36.4			186	14.6
ICU								
IV therapy	228	91.2	294	90.2			387	90.2
Multiple ATB diagnosis	78	30.5	121	36.4			91	21.2
Multiple ATB patient	91	38.4	138	45.4			101	24.3

Analyses at patient level. Patients admitted on a NMW and NICU are excluded.

Multiple ATB diagnosis is defined as receiving > 1 antibiotic (J01) for a single identified reason to treat (=diagnose code) at patient level.

Multiple ATB patient is defined as receiving > 1 antibiotic (J01) at patient level.

Country: SOUTH AFRICA; Continent: Africa; Hospital type: Tertiary/Spec/Inf.dis. hosp.

AWaRe is a useful tool to reduce antimicrobial resistance and ensure access.



ACCESS

Which indicates the antibiotic of choice for each of the 25 most common infections. These antibiotics should be available at all times, affordable and quality-assured.



WATCH

Which includes most of the "highestpriority critically important antimicrobials" for human medicine and veterinary use. These antibiotics are recommended only for specific, limited indications

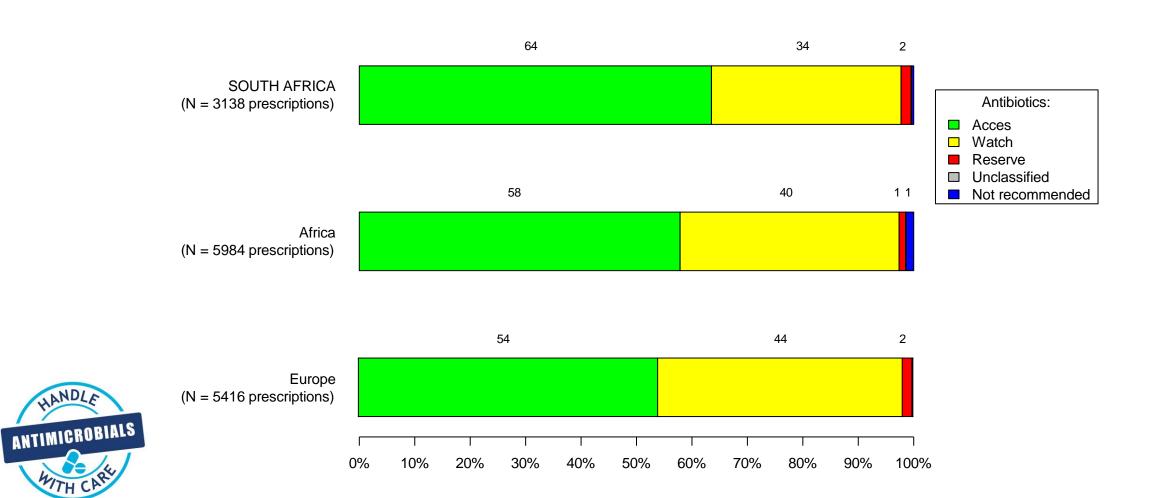




Antibiotics that should only be used as a last resort when all other antibiotics have failed.

Overall antibiotic use (ATC J01) according to the WHO AWaRe classification

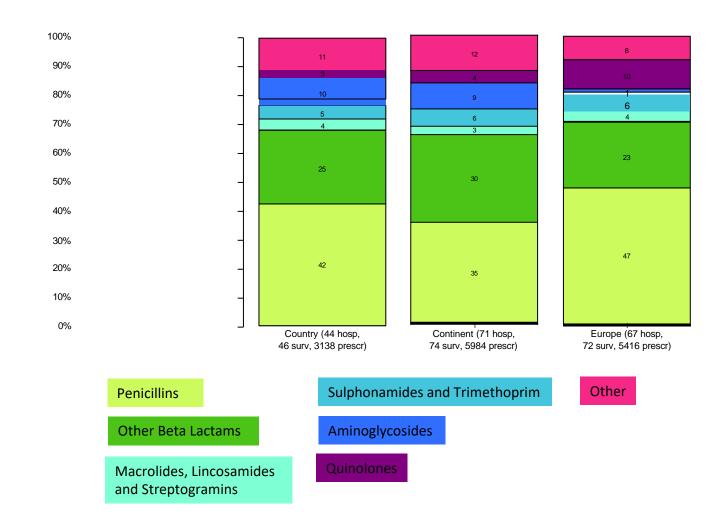
WHO goal: 60% all antibiotics used at a country level are from the ACCESS group by 2030



HANDLE



Overall proportional antibiotic use





Proportional antibiotic use (% of prescriptions)

ATC4	Antibiotics Subgroup	Country	Continent	Europe	
J01CA	Penicillins with extended spectrum	14.8	11.4	5.9	Amoxicillin/Ampicillin
J01CE	Beta-lactamase sensitive penicillins	1.3	2.8	0.9	_
J01CF	Beta-lactamase resistant penicillins	3.2	3.0	2.9	
J01CR	Penicillins incl. beta-lactam. inh.	22.3	17.1	37.3	Amoxicillin Clavulanate
J01DB	First-generation cephalosporins	4.5	2.7	9.6	
J01DD	Third-generation cephalosporins	13.3	20.7	6.2	Ceftriaxone/Cefotaxime
J01DH	Carbapenems	7.0	4.8	3.4	-
J01EE	Comb. Sulfonamides/trimethoprim	3.6	2.8	3.7	
J01FA	Macrolides	4.0	3.7	3.8	
J01FF	Lincosamides	0.7	2.3	2.6	
J01GB	Other aminoglycosides	9.6	9.0	1.1	
J01MA	Fluoroquinolones	2.6	4.3	10.1	
J01XA	Glycopeptide antibacterials	3.1	2.2	2.8	
J01XD J01XX	Imidazole derivatives Other antibacterials	6.1 0.3	8.9 0.2	2.0 1.4	

Our hospital: 494 prescriptions, 379 treated patients; Country: 3138 prescriptions, 44 hospitals, 46 surveys

Continent: 5984 prescriptions, 71 hospitals, 74 surveys

Europe: 5416 prescriptions, 67 hospitals, 72 surveys

Proportional antibiotic use (% of prescriptions) – [Adult] Intensive Care Unit

ATC4	Antibiotics Subgroup	Country	Continent	Europe
J01CA	Penicillins with extended spectrum	6.3	5.2	3.2
J01CF	Beta-lactamase resistant penicillins	0.9	0.7	1.7
J01CR	Penicillins incl. beta-lactam. inh.	22.9	19.9	41.5
J01DB	First-generation cephalosporins	4.9	4.1	8.7
J01DD	Third-generation cephalosporins	11.7	14.4	9.0
J01DE	Fourth-generation cephalosporins	3.6	3.0	0.7
J01DF	Monobactams			0.7
J01DH	Carbapenems	18.4	17.3	7.8
J01EE	Comb. Sulfonamides/trimethoprim	4.5	5.2	2.2
J01FA	Macrolides	2.7	2.6	3.9
J01GB	Other aminoglycosides	4.5	4.4	1.9
J01XA J01XD	Glycopeptide antibacterials Imidazole derivatives	4.9 4.0	4.1 7.4	4.4 0.7



Our hospital: 26 prescriptions, 18 treated patients; Country: 223 prescriptions, 22 hospitals, 24 surveys

Continent: 271 prescriptions, 32 hospitals, 34 surveys

Europe: 412 prescriptions, 55 hospitals, 56 surveys

Ten most common diagnoses treated with therapeutic antimicrobials

		C	ounti	У	Co	ntin	ent	E	ırop	e
Diagnosis	•	N		%	N		%	N		%
Pneumonia	3	396	2	5.0	666		24.3	1034		25.9
Skin and Soft Tissue	2	220	1	3.9	369		13.5	401		10.0
CNS	1	104	(6.6	161		5.9	56		1.4
Intra Abdominal		76	4	4.8	113		4.1	386		9.7
Lower UTI		55	(3.5	99		3.6	329		8.2
Bone and Joint		50	(3.2	82		3.0	209		5.2
TB		40	2	2.5	99		3.6	20		0.5
Gastrointestinal		67	4	1.2	119		4.3	121		3.0
OBS and GYNAE		58	(3.7	111		4.1	17		0.4
Bacteraemia		25	•	1.6	29		1.1	58		1.5



Prevalence (%) of Healthcare Associated Infections: Hospital-wide

		Country	Continent	Europe
		504	700	1000
Numerato	or (N patients)	534	796	1228
Denominator (N admitted patients)		7136	10766	16454
	HAI rate (%)	7.5	7.4	7.5
Post-operative surgical site	e infection (%)	1.5	1.6	1.5
Intervention related	infection (%)	1.6	1.4	1.4
	CDAD (%)	0.2	0.2	0.1
	Other HAI (%)	3.8	3.9	3.6
HAI from another	er hospital (%)	0.6	0.6	0.2
HAI from LTCF or nurs	sing home (%)	0.0	0.1	8.0



Prevalence (%) of Intervention-related versus Other Hospital-Associated Infections Hospital-wide

	Country	Continent	Europe
Numerator (N patients)	534	796	1228
Denominator (N admitted patients)		10766	16454
HAI rate (%)	7.5	7.4	7.5
Intervention-related infections (%)			
Mixed origin	0.9	0.7	0.4
CVC-BSI	0.2	0.1	0.2
PVC-BSI	0.1	0.1	0.0
Ventilator-Associated Pneumonia (VAP)	0.3	0.3	0.2
CAUTI	0.2	0.2	0.5
Other Hospital-Associated Infections (%) HAI of			
mixed or undefined origin	2.1	1.8	1.7
Blood Stream Infection (BSI)	1.2	1.3	0.2
Hospital-Acquired Pneumonia (not VAP)	0.3	0.5	1.1
Urinary Tract Infection (UTI)	0.3	0.4	0.7



CVC-BSI = Central Venous Catheter-related Blood Stream Infection

PVC-BSI = Peripheral Vascular Catheter-related Blood Stream Infection

CAUTI = Catheter-Associated Urinary Tract Infection

Invasive device use hospital-wide

	Cou	Country		inent	Hosp	ital type	Europe	
	N	%	N	%	N	%	N	%
N total admitted patients	9465		11093		6245		5369	
N admitted patients with:								
PVC	3855	40.7	4901	44.2	2859	45.8	2309	43.0
CVC	448	4.7	523	4.7	451	7.2	443	8.3
Indwelling UC	1400	14.8	1594	14.4	971	15.5	753	14.0
Tubes/Drains	488	5.2	535	4.8	415	6.6	410	7.6
IRI	288	3.0	316	2.8	278	4.5	101	1.9
CiPAP-BiPAP	210	2.2	250	2.3	186	3.0	59	1.1

CVC = Central Venous Catheter; PVC = Peripheral Vascular Catheter; UC = Urinary Catheter; IRI = Invasive endotracheal Respiratory Intubation; CiPAP, BiPAP = Non-invasive mechanical ventilation

Therapeutic antimicrobial use for community acquired and healthcare associated infections by type of treatment

	CAI	CAI Empiric		Fargeted	CAI Total		
	N	%	Ν	%	Ν	%	
Country	1596	92.4	132	7.6	1728	67.7	
Continent	2969	92.6	238	7.4	3207	71.6	

High rate of empiric prescription CAI>HAI

	HAI Empiric		HAI Targeted		HAI To	otal		
	Ν	%	Ν	%	Ν	%		
Country	541	65.7	282	34.3	823	32.3	HAI Rate 7.5%	
Continent	942	74.2	328	25.8	1270	28.4		

HAI drives antibiotic Prescribing



Prevalence of patients (%) with previous hospitalisation < 3 months

Hospital (%) Adult wards (%) Paediatric wards (%)

Approx. a quarter patients had a previous hospital admission

Country SOUTH AFRICA - N patients (denominator)	2774	1870	904
Yes, ICU	3.9	3.5	4.5
Yes, Other	18.6	20.5	14.6
No	62.0	56.3	73.7
Unknown	15.6	19.6	7.2
Continent - N patients (denominator)	3568	2460	1108
Yes, ICU	3.4	3.3	3.9
Yes, Other	19.0	20.4	16.0
No	61.8	57.2	72.2
Unknown	15.7	19.2	7.9



Europe - N patients (denominator)	1528	1380	148
Yes, ICU	2.0	2.1	1.4
Yes, Other	21.1	21.9	13.5
No	57.1	57.2	55.4
Unknown	4.0	2.5	18.2

Prevalence of patients (%) with previous antibiotic treatment < 1 month

Hospital (%) Adult wards (%) Paediatric wards (%)

23% patients had received antibiotics in last month

Country SOUTH AFRICA - N patients (denominator)	2759	1863	896
Yes	26.5	24.1	31.4
No	48.7	45.5	55.2
Unknown	24.9	30.4	13.4
Continent - N patients (denominator)	3586	2485	1101
Yes	25.1	23.2	29.5
No	46.5	43.2	54.0
Unknown	28.4	33.6	16.4

Europe - N patients (denominator)	1484	1341	143
Yes	28.2	29.3	17.5
No	59.2	59.0	60.8
Unknown	12.7	11.7	21.7



Proportional antibiotic use (% of prescriptions) – Community Acquired Infections

8.0 1.0
1.0
3.2
46.2
0.5
7.7
2.4
0.1
1.5
2.9
1.3
11.1
2.4 1.4



Our hospital: 294 prescriptions, 231 treated patients; Country: 1577 prescriptions, 42 hospitals, 44 surveys

Continent: 2833 prescriptions, 68 hospitals, 70 surveys Europe: 2987 prescriptions, 64 hospitals, 69 surveys

Proportional antibiotic use (% of prescriptions) – Healthcare Associated Infections

Cost	7
Cost	!

ATC4	Antibiotics Subgroup	Country	Continent	Europe
J01CA	Penicillins with extended spectrum	7.0	7.2	4.7
J01CF	Beta-lactamase resistant penicillins	3.6	3.6	3.9
J01CR	Penicillins incl. beta-lactam. inh.	15.3	12.1	38.1
J01DB	First-generation cephalosporins	0.6	0.6	0.5
J01DC	Second-generation cephalosporins		0.6	2.2
J01DD	Third-generation cephalosporins	6.1	13.1	6.8
J01DF	Monobactams		/	0.2
J01DH	Carbapenems	23.5	18.0	8.0
J01EE	Comb. Sulfonamides/trimethoprim	1.7	1.3	3.1
J01FF	Lincosamides	1.1	1.9	2.0
J01GB	Other aminoglycosides	12.8	13.2	1.4
J01MA	Fluoroquinolones	5.2	5.9	12.2
J01XA	Glycopeptide antibacterials	10.2	7.0	7.5
J01XB	Polymyxins Imidazole derivatives	5.1	3.2 5.5	0.4
J01XD	Imidazole derivatives	2.6	5.5	1.4



Our hospital: 96 prescriptions, 72 treated patients; Country: 727 prescriptions, 38 hospitals, 40 surveys

Continent: 1142 prescriptions, 62 hospitals, 64 surveys

Europe: 1331 prescriptions, 67 hospitals, 72 surveys

Type of antibiotic treatment – Summary

	Cou	ıntry	Con	tinent	Europe	
	N	%	N	%	N	%
All patients						
Empiric	2749	88.8	5489	92.8	3724	70.1
Targeted	347	11.2	429	7.2	1587	29.9
Adults (>= 18 years)						
Empiric	1676	89.7	3403	93.1	3179	68.1
Targeted	192	10.3	254	6.9	1492	31.9
Children (< 18 years)					1	
Empiric	913	89.2	1743	93.3	505	84.7
Targeted	111	10.8	126	6.7	91	15.3
Neonates (NICU)						
Empiric Targeted	160 44	78.4 21.6	343 49	87.5 12.5	40 4	90.9 9.1

Blood Culture Practices?

Blood volumes



Treatment based on microbiology data

	Co	Country		Continent		Hospital type		Europe	
	N	%	N	%	N	%	N	%	
MRSA	22	1.0	27	0.7	20	1.0	16	0.4	
MRCoNS	15	0.7	15	0.4	8	0.4	37	1.0	
VRE	4	0.2	6	0.2	5	0.3	3	0.1	
ESBL	63	2.8	87	2.4	78	4.1	72	1.9	
3GCREB	11	0.5	22	0.6	20	1.0	38	1.0	
CRE	40	1.8	43	1.2	34	1.8	12	0.3	
ESBL-NF	11	0.5	18	0.5	14	0.7	9	0.2	
CR-NF	36	1.6	41	1.1	31	1.6	15	0.4	
Other MDR	0	0.0	0	0.0	0	0.0	0	0.0	
PNSP	0	0.0	0	0.0	0	0.0	0	0.0	
MLS	4	0.2	5	0.1	2	0.1	3	0.1	
Any of the above	186	8.4	236	6.5	189	9.9	190	5.1	

N = the number of patients reported to have received a microbiology-based treatment for the respective pathogen.

% = 100*(the number of patients reported to have received a microbiology-based treatment for the respective pathogen/total number of patients receiving a therapeutic treatment (CAI or HAI) with at least one antibacterial for systemic use (J01)).

Use of laboratory when using antibiotics

Diagnosis	No of prescriptions	%
Pneumonia	1065	63,4
Sepsis	651	74,5
Skin and Soft		
Tissue	558	50,0
Proph OBGY	305	18,7
CNS	290	59,7



WCC



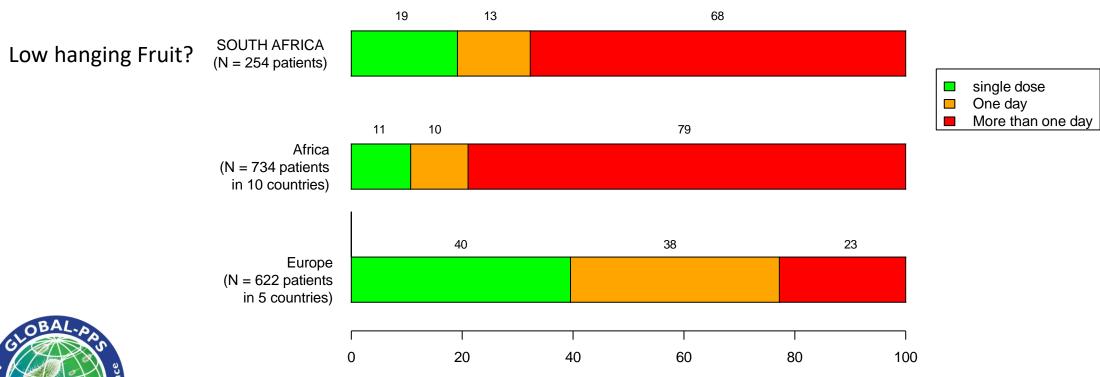
Use of laboratory when using antibiotics

Diagnosis	Number of Prescriptions	Total Cultures to lab (%)	Blood Cultures to lab (%)
Pneumonia	1065	55,1	44,4
Sepsis	651	76,4	70,7
Skin and Soft Tissue	558	37,3	22,4
Proph OBGY	305	4,6	3,3
CNS	290	69,3	41,0



Total Cultures: Blood, urine, stool, CSF, BAL, Sputum, Wound, Other

Duration of surgical prophylaxis in adults and children





Proportional antibiotic use (% of prescriptions) – Surgical Prophylaxis

ATC4	Antibiotics Subgroup	Country	Continent	Europe
J01CA	Penicillins with extended spectrum	8.4	7.8	0.5
J01CF	Beta-lactamase resistant penicillins	0.9	1.7	0.6
J01CR	Penicillins incl. beta-lactam. inh.	33.7	25.2	8.5
J01DB	First-generation cephalosporins	32.0	11.8	76.1
J01DD	Third-generation cephalosporins	5.2	18.0	0.3
J01DE	Fourth-generation cephalosporins	0.6	0.6	
J01GB	Other aminoglycosides	2.6	3.6	0.2
J01MA	Fluoroquinolones	0.6	5.0	3.1
J01XD	Imidazole derivatives	14.2	16.6	2.3



Our hospital: 20 prescriptions, 19 treated patients; Country: 344 prescriptions, 27 hospitals, 29 surveys

Continent: 1022 prescriptions, 51 hospitals, 53 surveys Europe: 645 prescriptions, 59 hospitals, 64 surveys Country: SOUTH AFRICA; Continent: Africa; Hospital type:

Original Article

Optimizing prophylactic antibiotic use among surgery patients in Ethiopian hospitals

Getachew Alemkere ^a, Hailu Tadeg ^a, Workineh Getahun ^a, Wendosen Shewarega ^b, Asrat Agalu ^c, Mohan P. Joshi ^{d,1}, Niranjan Konduri ^{d,*,2}

Table 5Benefit of the SAP intervention between the baseline and intervention phases.

	p value
Patients developing SSI, n (%) 10 (6.1)	0.779 0.190 0.028 *

Journal of Infection and Public Health

^a USAID Medicines, Technologies, and Pharmaceutical Services (MTaPS) Program, Management Sciences for Health, Addis Ababa, Ethiopia

^b Pharmaceuticals and Medical Devices Lead Executive Office, Ministry of Health, Addis Ababa, Ethiopia

^c Department of Pharmacy, Bahir Dar University, Bahir Dar, Ethiopia

^d USAID Medicines, Technologies, and Pharmaceutical Services (MTaPS) Program, Management Sciences for Health, Arlington, VA, USA



Summary of quality indicators for antibiotic use

	Co	Country		nent	Eu	Europe			
	N	%	Ν	%	N	%			
Medical									
Reason in notes	1632	84.3	2609	78.3	3004	88.9			
Guidelines missing	76	3.9	280	8.4	360	10.7			
Guideline	1025	79.0	1618	76.6	2249	83.2			
compliant Stop/review date	642	33.1	1305	39.2	1549	45.8			
documented									
Surgical									
Reason in notes	562	77.4	1411	73.6	1270	85.2			
Guidelines missing	25	3.4	554	28.9	158	10.6			
Guideline	377	69.9	658	67.1	938	81.1			
compliant Stop/review date	284	39.1	955	49.8	912	61.2			
documented									
ICU Reason in notes	409	86.1	637	86.5	500	91.7			
Guidelines missing	24	5.1	77	10.5	54	9.9			
Guideline compliant	253	79.8	345	79.5	332	83.8			
Stop/review date	219	46.1	312	42.4	218	40.0			

⁻ For reason in notes and stop/review date documented: Count at antibacterial level.

⁻ For guidelines missing: Count on NA (= no guideline for an indication) at patient level and diagnosis over total scores for this indicator.

⁻ For guideline compliance: Count at patient level and diagnosis for compliance= yes or no only. For combination therapy with >1 antibiotic: if 1 antibiotic by diagnosis is not compliant, this combination therapy as a whole for this diagnosis will be counted as non-compliant.

Prevalence of missed doses hospital wide

	Country	Continent	Europe	
Hospital (%)				
N antimicrobials	4796	8986	6392	ı
Percentage missed doses	10.15	13.87	3.25	
Mean missed doses	2.31	3.27	1.94	
Median missed doses	1	2	1	
Reason missed doses (%)				_
Stock out	5.1	14.8	7.7	
Could not purchase	0.2	7.7	0.0	•
Declined/refused	0.2	0.3	0.0	
Other reason	19.9	13.1	33.2	
Multiple reasons	2.5	4.0	2.9	
Unknown	72.1	60.1	56.2	

Analyses are performed at antimicrobial level.

% AM with missed doses: 100*(number of reported antimicrobials with at least one missed dose/number of all reported antimicrobials (antimicrobials with unknown number of missed doses are also included in the denominator under the assumption that missing doses equals no missed dose).

Mean and median missed doses are calculated using all antimicrobials with at least one missed dose.

Antimicrobials for which no doses were missed (zero) or reported (missing values) are excluded for these analyses.

Reason missed doses (%): Proportion (%) of reason for missed doses out of all possible reasons for antimicrobials with at least one missed dose. Unknown reason: Counts those antimicrobials with code U + empty/missing values for antimicrobials for which at least one missed dose was reported.

Insert Hospital logo [Hospital] Antibiotic Steward Antibiotic Prescription Chart							p Pro	ogra	amı	_	War	d			
Patient Label							Weigh	nt				Alle	rgie	es	
eGFR															
Infection Episode 1 Diagnosis Pneumonia UTI Meningitis Line infection Other Other															
Source* Community Hospital acquired Indication P = Prophylactic E = Empirical D = Definitive SEND APPROPRIATE CULTURES BEFORE PRESCRIBING ANTIBIOTICS															
	munity acqu	Sent before antibiotics ired: within ≤48h, of d: >48h after admiss		ics LS	lot ent charge	Antibiotic Da	Day 1	2	Review &	Review 2	6	Review 2	8	9	10
Indication	Medicine	Approved Name	e or GE	Dose	Route										
E	Start Date Time	Duratio	on	Frequency	/				<u> </u>					-	
□D	Drs Signa	ture & Name	Contact	Pharmacy				H	+					\dashv	

Conclusions

- Building on previous pilot
 - Increased and repeated participation will give more robust data
 - Specific Hospitals for quality improvement
- Future participation needs a combined approach
 - IPC and AMS
 - All specialities medical, surgical, ICU, general wards
- HAI's drive antibiotic prescription: Increased Cost
- Identify targets for AMS and IPC QI projects for national implementation
 - Oral to IV switch
 - Surgical Prophylaxis
 - Blood Culture Practices
 - Guideline compliance: IPC and AMS









- Ann Versporten and the Global PPS team
- Ruth and Nasreen from MAC secretariat
- Prof Shaheen Methar
- All the survey participants particularly the hospital "admins"