

## **ESTABLISHING ANTIMICROBIAL STEWARDSHIP PROGRAMME AT FOUR SECONDARY CARE** HOSPITALS IN INDIA THROUGH HUB AND SPOKE MODEL WITH CHRISTIAN MEDICAL COLLEGE, **VELLORE AS GUIDING CENTRE**

Prasannakumar Palanikumar, Naveena Gracelin Princy Zacchaeus, Hanna Alexander, Abi Manesh, Lydia John, Jemin Webster. Indu Kulshrestha, Mahima Sadanshiv, Rincy John, Prasanna Samuel.Priscilla Rupali Department of Infectious Diseases, and Biostatistics Christian Medical College, Vellore, Tamilnadu, India, Baptist Christian Hospital,

Assam; Bangalore Baptist Hospital, Bangalore; Padhar Hospital, Madhya Pradesh; Christian Fellowship Hospital, Tamilnadu

RESULTS



0.001

0.023

0.004

India faces serious threat due to antimicrobial resistance at all levels of healthcare including secondary care level. Thus, Antimicrobial Stewardship (ASP) needs to be extended to all health care levels.

BACKGROUND

## **METHODS**

We adopted hub and spoke model to implement ASP based on our early experience with ID Physician driven Prospective Audit with Intervention and Feedback (PAIF) ASP strategy. We translated our experience into training programme for local physicians and pharmacist/nurse to adapt AMSP strategies in four secondary care hospitals with central support from our centre. The study consisted of three phases. Initial phase was to capture baseline antimicrobial days of therapy DOTs data, followed by intervention phase wherein secondary care physicians of the four chosen hospitals were trained in distance education mode for a year along with development of antibiogram based on local hospital microbial resistance pattern entered through WHONET and hospital specific algorithms and augmentation of the existing laboratory facilities by training microbiologists and technicians at CMC for required period. This was followed by post intervention phase with PAIF strategy and assessment of DOTs.

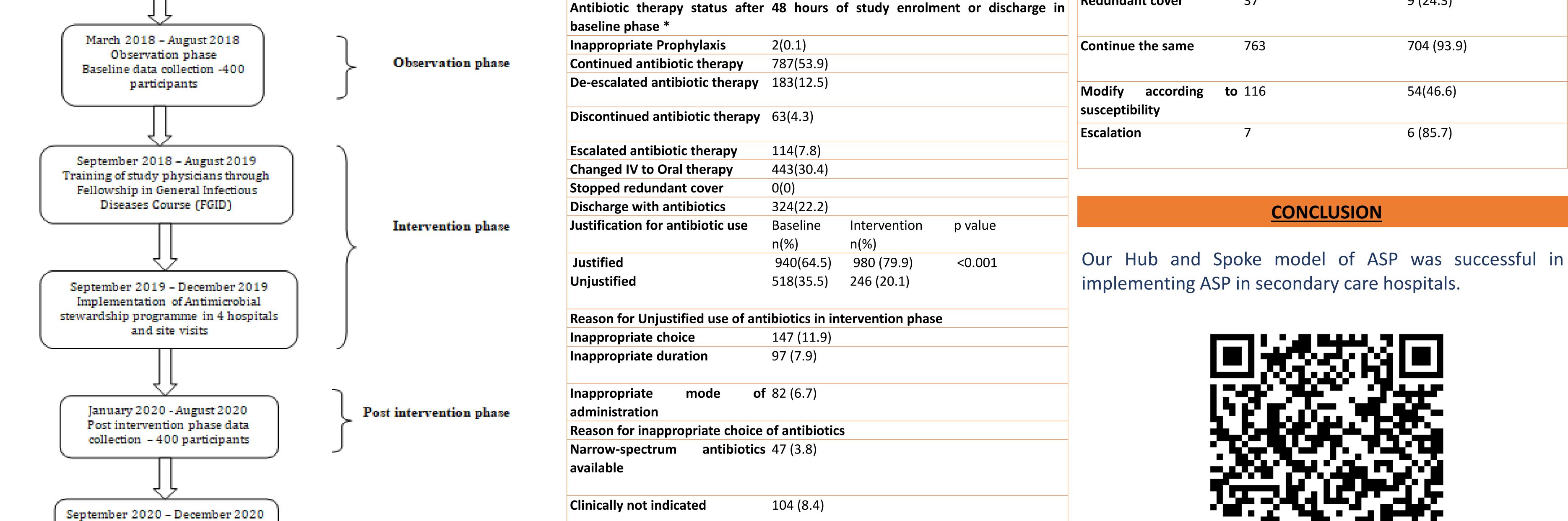
Preparation Phase

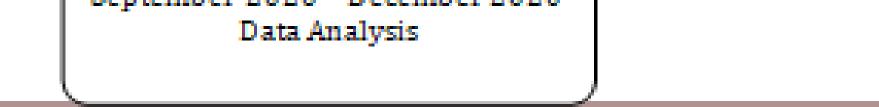
During the baseline phase, 1459 patients from all four sites were enrolled; 1233 patients were enrolled in the intervention phase. Both groups had comparable baseline characters. The key outcome, DOT per 1,000 patient days, was 1952.63 in the baseline phase. The DOT/1000 patient days was significantly lower in the post intervention period, at 1483.06 (P =0.001). Quinolones, macrolides, Cephalosporin's, Clindamycin, and nitroimidazole use were significantly decreased in the post intervention group. Post organism (MDRO) intervention showed 79.9% of antibiotic use was justified. 65 (8.6) Extended-spectrum **β** lactamase 89 (10.6) The reason for unjustified antibiotic use corresponded to 0 (0.0) 1 (0.1) (ESBL) the choice (59.75%), duration (39.4%) and route of **Vancomycin-resistant** Enterococcus 9 (1.1) 15 (2.0) 5 (0.7) 25 (3.0) administration (33.3%), 13.9% of antibiotic use was (VRE) **Carbapenem-resistant** clinically not indicated. The intervention rate was 38%. **Enterobacteriaceae (CRE)** 674 (88.7) 720 (85.4) Overall, the recommendations given by the ID team were **Methicillin-resistant Staphylococcus** fully followed in 946 cases (77.7%), partially followed in 59 aureus (MRSA) None cases (4.8%), and not followed in 137 cases (35.7%). There **Unintended consequences** were no significant adverse events post intervention. No reaction 1455 (99.7) 1221 (99.0) Diarrhea 0 (0.0) 7 (0.6) **Table1. Overview of Antibiotic Therapy use** Table 3. Acceptance of Recommendation Given by the Variables Overall Baseline Intervention p value **Infectious Disease Physician during the Intervention Phase** n(%) n(%) Indication for antimicrobial therapy < 0.001 Type of Recommendation Recommendation Given\* Recommendation Definite 559 (45.3) 1074 (39.9) 515 (35.3) infection Accepted **De-escalation** 50 22 (44.0) 410 (33.3) < 0.001 Probable 1018 (37.8) 608 (41.7) Infection 315 Discontinue 204 (65.8) 600 (22.3) 338 (23.2) 262 (21.2) Prophylaxis 0.234 37 9 (24.3) **Redundant cover** 

Table?	Secondar	y Outcomes
Iancz	Jecondar	y Outcomes

Variable	Baseline Phase (n=1459)	Intervention Phase (n=1233)	P Value
De-escalation rate	186 (12.5)	22 (44.0)	<0.001
Mortality			
Infectious	36 (61.0)	14 (43.8)	0.193
Non-Infectious	8 (13.6)	4 (12.5)	
Both	15 (25.4)	14 (43.8)	
Length of Stay, Median IQR	5.0 (3.0,8.0)	6.0 (4.0,9.0)	<0.001 <sup>\$</sup>
Prevalence of multidrug-resis	stant		

**Fig1.Study Flow** January 2018-February 2018 Preparation and training





www.PosterPresentations.com





