

Supplementary table 3. Overview of clinical and environmental samples, sample sources, number of isolates, geographic distribution, ESKAPE pathogens, isolation methods and identification techniques in the included studies.

Article identification	Types of samples	Number of isolates	Country	Targeted ESKAPE pathogen(s)	Isolation and presumptive identification	Typing methods and Identification
28	Clinical - Urine	Clinical: 57 Environmental: 59	Italy	<i>Enterococcus faecium</i>	Clinical The isolates were obtained from the clinical microbiology department.	Clinical Methodology: Biochemical identification - Automated: SCEPTOR system
	Environmental - Domestic refuse composting - Public swimming pool water - Marine brackish - Effluent - Sediments from harbour dredging activity				Environmental Methodology: Most probable number (MPN) method - Azide Dextrose Broth [S] - Esculin vancomycin azide broth [S] - Bile Esculin Agar [S, D] - Catalase - Gram-staining	Environmental Methodology: Presumptive phenotypic and Biochemical identification Manual and automated identification (SCEPTOR system)
29	Clinical - Blood, urine, sputum, pus, cow, rabbit, wild boar	Clinical: 68 Environmental: 23	Japan	<i>Pseudomonas aeruginosa</i>	Clinical The isolates were previously isolated	Clinical and environmental Methodology: Biochemical and Genotypic identification - Automated: BD Phoenix Automated microbiology system - Serotyping
	Environmental - River water - Lake, ponds and marine water - Surface seawater				Environmental Methodology: Membrane filtration and culture - Nalidixic acid cetrime (NAC) agar [S] - Nutrient broth agar (NA) [E] Cetrime kanamycin nalidixic acid agar [S]	PFGE
30	Clinical - Blood cultures of patients hospitalized	Clinical: 10 Environmental: 53	Greece	<i>Enterococcus faecium</i>	Clinical Methodology: Biochemical and culture - Automated system Bactec 9240 - Blood agar [E]	Clinical Methodology: Biochemical and Molecular fingerprint identification - Automated: API Strep test - Automated: VITEK2 automated system
	Environmental - Wastewater (raw) - Effluent (treated) - Secondary treated and disinfected wastewater from effluents of				Environmental Methodology: Direct plating - Kanamycin aesculin azide agar [S] supplemented with	Environmental Methodology: Biochemical and Molecular fingerprint identification

	- hospitals - Pig sample (rectum and faeces)				- vancomycin - Catalase - Gram-staining	- Manual: Biochemical - PFGE
	Clinical - Urine and wound				Clinical It was not described	Clinical and environmental Methodology: Genotypic and phenotypic identification - PCR - PhP-RF system - Ribotyping - PFGE
31	Environmental - Wastewater (raw)	Clinical: 48 Environmental: 58	Iran	<i>Enterococcus faecium</i>	Environmental Methodology: Membrane filtration and culture - Brain heart infusion (BHI) agar [E] - m-<i>Enterococcus</i> agar [S] supplemented with gentamicin.	
	Clinical - Pus, urine, sputum, bronchial wash solution	Clinical: 6 Environmental: 12	South Korea	<i>Pseudomonas aeruginosa</i>	Clinical and Environmental Methodology: Direct plating - <i>Pseudomonas</i> isolation agar [S]	Clinical and Environmental Methodology: Presumptive phenotypic and Molecular fingerprint identification RAPD
	Environmental - Spring water					
	Clinical - Clinical samples (it was not specified)				Clinical and Environmental Methodology: Biochemical and Culture - Trypticase soy broth (TSB) [E] - Glutamate starch <i>Pseudomonas</i> agar (GSP) [S, D] - Gram staining - Positive oxidase reaction - O/F +/-	Clinical and Environmental Methodology: Phenotypic and genotypic identification - SDS-PAGE - PFGE RAPD
33	Environmental - Water (tap water) - Garden	Clinical: 56 Environmental: 20	Spain	<i>Pseudomonas aeruginosa</i>		
	Clinical - Obtained from Pathology Queensland and QUT culture collection [did not specify the sample site origin]	Clinical: 27 Environmental: 47	Australia	<i>Enterococcus faecium</i>	Clinical The isolated were obtained from culture collection Environmental Methodology: Membrane filtration and culture - <i>Enterococcus</i> agar [S]	Clinical The bacterial species were identified according to the classification provided by the reference culture collection, since the isolates were sourced directly from it. Environmental Methodology: Presumptive phenotypic and Genotypic identification Real-time PCR using <i>ddlE. faecalis</i> and <i>ddlE. faecium</i>
34	Environmental - River water					

35	Clinical - Blood, cerebrospinal fluid, eyes, livers, peripheral venous catheters, pleural fluid, sputum, urine, wounds, respiratory system Environmental - Groundwater for human use (wells) - Wetland (lake) - Wastewater (treated)	Clinical: 14 Environmental: 35	Mexico	<i>Enterococcus faecium</i> , <i>Enterococcus faecalis</i>	Clinical Method: Culture - Sheep blood agar [E] - Grain-staining - Catalase - Esculin hydrolysis - BHI 6.5% NaCl Environmental Method: Membrane filtration and culture - K-F agar [S, D] - Grain-staining - Catalase - Esculin hydrolysis - BHI 6.5% NaCl	Clinical Methodology: Genotypic and phenotypic identification - Multiplex PCR PFGE
37	Clinical - Hospitalized patients: urinary and respiratory tract infections Environmental - Wastewater (raw) - River water	Clinical: 20 Environmental: 214	Australia	<i>Pseudomonas aeruginosa</i>	Clinical Methodology: Culture - Previously isolated by the pathology unit of the hospital Environmental Methodology: Membrane filtration and culture Cetrimide agar [S]	Clinical and Environmental Genotypic fingerprinting and molecular detection - PCR RAPD
38	Clinical - Blood Environmental - Tap water	Clinical: 49 Environmental: 6	Ethiopia	<i>Klebsiella pneumoniae</i>	Clinical Methodology: Culture - Tryptic soy broth (TSB) [E] - Sheep blood [E, D] - Chocolate [E] - MacConkey [D] Environmental Method: Culture - MacConkey agar [D] - Xylose lysine deoxycholate (XLD) agar [S, D] - Cystine lactose electrolyte deficient agar (CLED) [D] - Sheep blood agar [E, D]	Clinical and Environmental Methodology: Presumptive phenotypic and biochemical identification - Fermentation method - Urea 40% broth [D] - Simmons citrate agar [S, D] - Sulfide indole motility medium [D] - Triple sugar iron agar [D] - Klinger iron agar [D] - Oxidase test reagents

Chocolate agar [E]						
39	Clinical - Tracheal aspirates Environmental - Wastewater (hospital)	Clinical: 10 Environmental: 10	Croatia	<i>Acinetobacter baumannii</i>	Clinical - Previously isolated Environmental Methodology: Membrane filtration and culture - CHROMagar Acinetobacter supplemented [S] with CR102 and cefsulodin sodium salt hydrate Nutrient agar (NA) [E]	Clinical Methodology: Genotypic fingerprint - Previously identified - PFGE Environmental Methodology: Phenotypic and genotypic fingerprint - MALDI TOF MS PFGE
40	Clinical - Catheters, tracheobronchial, gastric, rectal, urine Environmental - Tap water	Clinical: 139 Environmental: 19	Besançon, France	<i>Pseudomonas aeruginosa</i>	Clinical and environmental Methodology: It was not specified	Clinical and Environmental Methodology: Genotypic identification PFGE
41	Clinical - Nasal, ear, pus, wound, skin Environmental - Water samples: tap, borehole, tank, reservoir, well water	Clinical: 120 Environmental: 45	Nigeria	<i>Staphylococcus aureus</i>	Clinical Methodology: Culture - Blood agar [E] - Mannitol salt agar (MSA) [S, D] Environmental Methodology: Culture Mannitol salt agar (MSA) [S, D]	Clinical and Environmental Only the culture selective (presumptive phenotypic identification on selective and differential culture)
42	Clinical - They only say that samples were from Austrian hospitals Environmental - River water	Clinical: 4 Environmental: 5	Australia	<i>Klebsiella pneumoniae</i>	Clinical It was not specified Environmental Methodology: membrane filtration and culture - BBL fluid thioglycolate medium [E] - Chromogenic media [S, D] ChromID ESBL [S, D]	Clinical - It was not specified Environmental Methodology: Phenotypic and genotypic fingerprint - MALDI TOF MS - WGS
43	Clinical - Diarrheal stool	Clinical: 15 Environmental: 49	India	<i>Klebsiella pneumoniae</i>	Clinical Methodology: Direct plating - Nutrient agar (NA) [E]	Clinical and environmental: Methodology: Presumptive phenotypic identification

44	Environmental - Well water (residential, market, hospital) - Effluent (Fish market)	Clinical: 108 Environmental: 11	China	<i>Enterococcus</i> (including <i>Enterococcus faecium</i>)	- MacConkey (MA) [D]	Selective and differential medium.
	Clinical - Feces from healthy individuals and farmed animals - Fish samples - Pork and poultry meat from supermarkets				Environmental Methodology: Direct plating - Tryptic soy agar (TSA) [E] - MacConkey (MA) [D] Nutrient agar (NA) [E]	
	Environmental - River and laker waters				Clinical Methodology: Culture - Luria-Bertani (LB) broth [E] - Columbia agar base supplemented with sheep blood and florfenicol [S] - Gram-staining	Clinical and environmental Methodology: Phenotypic and Genotypic identification - MALDI-TOF MS - WGS
45	Clinical - Patients hospitalized	Clinical: 110 Environmental: 231	Czech Republic	<i>Klebsiella</i> spp.	Clinical Methodology: Culture - MacConkey agar [S] supplemented with cefotaxime or meropenem.	Clinical and Environmental Methodology: Phenotypic and Genotypic identification - MALDI-TOF WGS
	Environmental - Raw hospital wastewater, inflow and outflow from hospital - River water				Environmental Methodology: Membrane filtration and culture - Klebsiella ChromoSelect [S] MacConkey [S] supplemented with cefotaxime or meropenem	