

Microbiological and epidemiological aspects of endocarditis caused by *Rothia* spp.: a systematic review

ABSTRACT

Background and aim: *Rothia*, a genus comprising pleomorphic Gram-positive bacteria found in the human oral, intestinal, and skin microbiota, is recognized as an opportunistic pathogen. Case reports of *Rothia* spp. endocarditis in the scientific literature are scarce, with limited knowledge of relevant data on *Rothia* endocarditis and clinical aspects of its treatment. The objective of this literature review is to compile clinical aspects of endocarditis caused by *Rothia* species, analyzing results and clinical practices to elucidate the most important risk factors, comorbidities, prognostic factors, and appropriate antibiotic treatment options.

Methods: Employing the PRISMA model, a systematic review was conducted utilizing PubMed, SciELO, and Google Scholar databases, encompassing articles published from 1978 to November 2023. Pertinent aspects were systematically recorded and summarized for subsequent analysis.

Results: *Rothia dentocariosa* (48.69%), *Rothia mucilaginosa* (22.37%), *Rothia aerea* (14.47%), and *Rothia kristinae* (14.47%) were recognized as the agents of endocarditis cases. Patients exhibited an average age of 48.5 years, with a notable male preponderance (71.6%). Clinical manifestations of *Rothia* spp. endocarditis presented similar features compared to other Gram-positive bacterial endocarditis cases. The mortality rate was notably lower than observed in other infectious endocarditis instances (11.84%). Predominant risk factors included preexisting cardiovascular diseases (50%), followed by odontological procedures, caries, and precarious oral hygiene (17.1%), immunocompromised status (14.47%), injectable illicit drug use (11.84%), and diabetes (9.21%). Embolic events were documented in 35.53% of patients, predominantly in Central Nervous System (28.95%). Mycotic aneurysms were identified in 6.58% of cases. Resistance to antibiotics was identified in only 13.16% of strains causing endocarditis, although certain strains displayed characteristics indicative of multidrug resistance.

Conclusion: Despite its rarity, *Rothia* spp. endocarditis exhibits clinical parallels with endocarditis caused by other Gram-positive bacteria, but with a comparatively lower mortality rate. Challenges in identifying *Rothia* spp. species based on cultural and microscopic characteristics, associated to early resolution in antibiotic therapy, seems to contribute to the underreporting of endocarditis caused by these bacteria.

Keywords: *Rothia* spp., Endocarditis, Opportunistic infection, Rare endocarditis agents

1. INTRODUCTION

Rothia is a genus of Gram-positive bacteria. Species within this genus exhibit pleomorphism, manifesting in coccoid, spherical, rod-shaped, or filamentous forms. They typically have a diameter of approximately 1.0 µm, with occasional irregular dilations and rounded ends reaching up to 5.0 µm in diameter. The coexistence of various forms within a single colony is commonly observed. Non-motile and lacking endospore formation, *Rothia* ferment carbohydrates, primarily producing lactic acid, and are generally catalase-positive [1,2].

The taxonomy of the *Rothia* genus was initially proposed based on biological and metabolic characteristics attributed primarily to *Rothiadentocariosa*, previously classified as a member of the *Nocardia* genus within the Actinomycetaceae family. Unlike *Nocardia*, *Rothia* bacteria are not acid-alcohol-resistant [3]. *Rothiamucilaginosus*, initially classified as *Stomatococcum mucilaginosus*, was included as a second species after genetic and molecular analyses [1]. Recent genetic studies led to the reclassification of three *Kocuria* species (*K. kristinae*, *K. halotolerans*, and *K. koreensis*) into the *Rothia* genus [2]. Currently, *Rothia* is classified within the Micrococcaceae family, comprising fourteen recognized species [4], with ongoing research indicating potential undiscovered species [5].

Rothia bacteria are part of the normal human microbiota in the skin [6,7,8], intestine [9,10,11], and oral cavity [12,13,14,15,16]. Considered opportunistic pathogens with low virulence [17], *Rothia* spp. infections are associated with invasive medical procedures [17,18] and conditions leading to immunocompromise [18,19]. Comparative studies on microbiome composition revealed alterations due to COVID-19, with a notable increase in *Rothia* spp. relative abundance [20,21,22].

Infections by *Rothia* spp. are considered rare. The relative abundance of *Rothia* spp. in the oral microbiota is often associated with better periodontal health [23,24,25]. However, an increased relative abundance of *Rothia* spp. in the oral microbiota is also observed in underlying diseases, especially in individuals with cancer and acute neutropenia [26,27,28]. Dysbiosis, with an imbalance favoring the abundance of a particular opportunistic pathogen, poses a risk to the host. Regarding the oral microbiota, this imbalance may increase the risk of dental caries, periodontal disease, and even traverse local immune barriers, leading to infections in other parts of the body or systemic infections [29]. *Rothia* spp. have been implicated in opportunistic infections affecting the respiratory tract, meningitis, cerebral empyema, infectious arthritis, endophthalmitis, keratitis, skin infections, pyelonephritis, infectious ascites, peritoneal dialysis-related peritonitis, catheter-associated infections, bacteremia, sepsis, and endocarditis [30-33].

Infective endocarditis is a systemic and potentially life-threatening condition characterized by bacterial infection affecting native or prosthetic heart valves, the endocardial surface, or cardiac devices. This clinical entity is associated with prolonged hospitalizations, significant morbidity, and elevated mortality rates [34,35]. Currently, approximately 130 pathogens, predominantly bacteria and fungi, are recognized as causative agents of infective endocarditis [36]. The majority of cases, ranging from 80% to 90%, are attributed to Gram-positive cocci, mainly from the *Staphylococcus*, *Streptococcus*, and *Enterococcus* genera [35,37-42].

The pathogenesis of infective endocarditis involves the adherence of microorganisms, particularly on cardiac valves or prosthetic valves, leading to the formation of bacterial vegetations. These vegetations can induce lesions on heart valves and facilitate the dissemination of infection to other organs. The intricate infectious dynamics of endocarditis pose substantial health risks, often culminating in fatal outcomes. Endocardial infection typically arises from direct colonization or, more commonly, during episodes of bacteremia. Asymptomatic bacteremia, frequently associated with oral flora following dental procedures, is prevalent in humans. Nosocomial infections commonly arise following surgical prosthetic valve implantation, vascular catheterization, hemodialysis, or other invasive medical interventions. Community-acquired infections are often linked to immunosuppression, intravenous drug usage, poor oral hygiene, degenerative valve disease, or rheumatic fever resulting from infections with Lancefield Group A *Streptococcus pyogenes* [41]. On a global scale, the incidence and mortality rates of infective endocarditis have experienced a substantial increase over the past three decades. Chen et al. [37] estimated a case count of 478,000 and 28,750 deaths in 1990, escalating to 1,090,530 cases and 66,320 deaths in 2019. During the period from the 1970s to 2000, the annual incidence of infective endocarditis worldwide was estimated at 5 to 7 cases per 100,000 person-years [43]. In the last two decades, incidence rates have averaged 10 to 15 cases per 100,000 people per year, indicating a notable surge in cases globally [37,44].

According to Graevenitz [45], reports of infective endocarditis caused by *Rothia* spp. in the scientific literature are very scarce, and relevant data on *Rothia* endocarditis and clinical aspects of its treatment are not well-known. The objective of this literature review is to gather clinical aspects of endocarditis caused by *Rothia* species, analyzing results and clinical practices that help elucidate the most important risk factors and comorbidities, prognostics, and suitable antibiotic treatment options.

2. METHODS

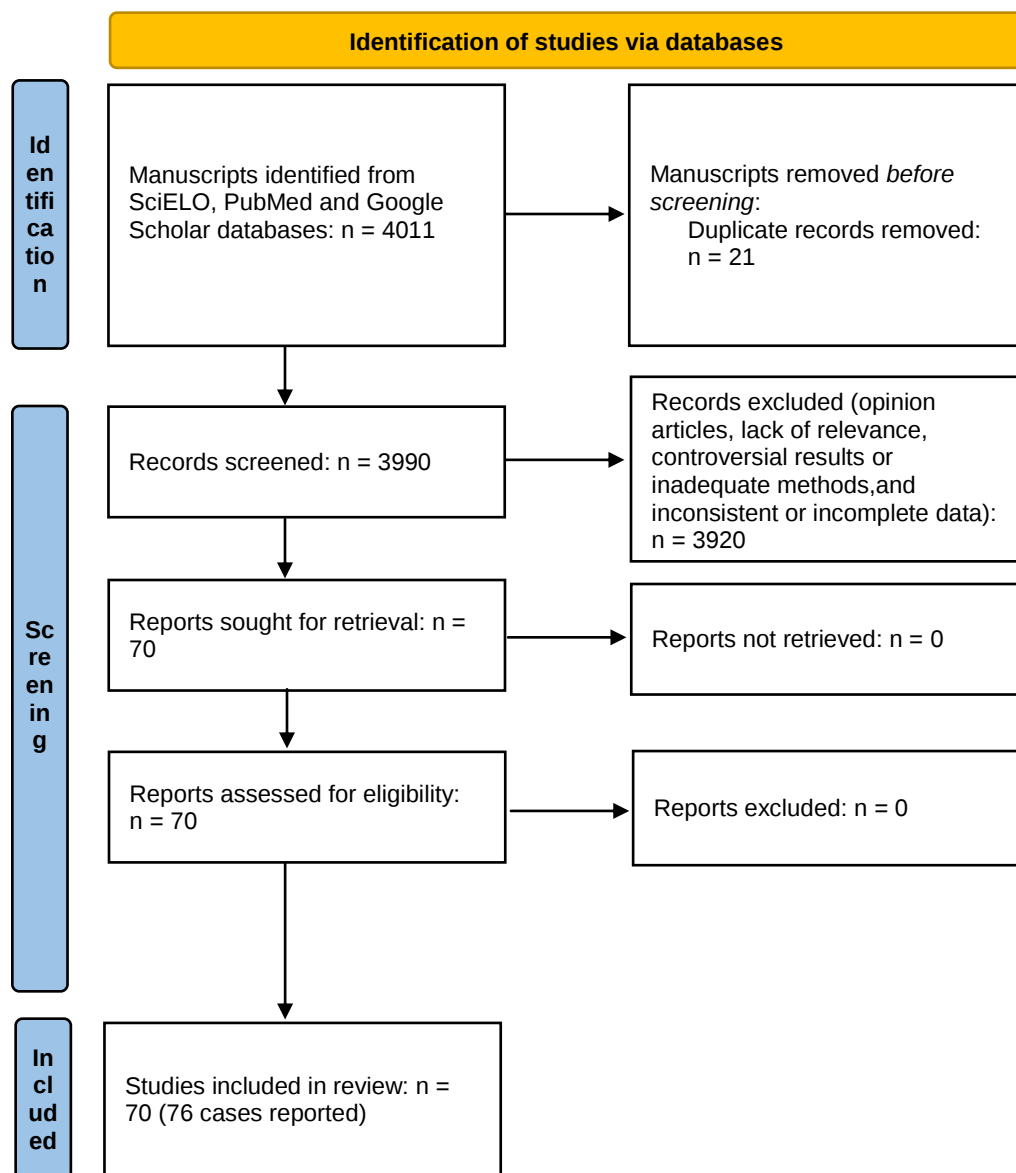
A systematic review was conducted following the methodological guidelines proposed by Moher et al. [46], as updated by Page et al. [47], utilizing the PRISMA model. The objective of the review was to investigate the etiology of Infective Endocarditis caused by *Rothia* species, including associated comorbidities or baseline conditions, prognosis, and antibiotic treatment. The review utilized sources from databases including PubMed, SciELO, and Google Scholar. The search employed descriptors such as "*Rothia*", "*Stomatococcum mucilaginosus*", "*Micrococcus mucilaginosus*", "*Actinomyces dentocariosus*", "*Nocardia salivae*", "*Nocardia dentocariosa*", "*Kocuria halotolerans*", "*Kocuria koreensis*", "*Kocuria kristinae*"; "Endocarditis"; "valve", "Prosthetic valve", and "Cardiac". The reviewed manuscripts spanned publications in English, Portuguese, and Spanish, covering the period from 1978 to 2023.

Studies and case reports of Infective Endocarditis caused by *Rothia* species were evaluated, and cases were included in the review if they met the Updated Modified Duke Criteria [48]. In this context, the incorporated investigations needed to satisfy a minimum of two criteria: either a culture derived from hemoculture or culture from cardiac biopsy samples, identified through biochemical methods, mass spectrometry, genetic sequencing analysis, or an alternative method

tailored for the precise determination of *Rothia* genus species. Confirmation through diagnostic methodologies affirming the existence of vegetation in cardiac tissue, explanted prosthetic valve or sewing ring, ascending aortic graft, endovascular intracardiac implantable electronic device, or originating from an arterial embolus was also required. Although the inclusion filter adheres to the Updated Modified Duke Criteria, some clinical aspects and the diagnostic trajectory of endocarditis will not be examined, as well as the clinical assessment of the cardiological evaluation within the article's scope. This constrained focus allows for a more detailed exploration of the epidemiology, identification, and secondary complications associated with *Rothia* species implicated in endocarditis cases, along with an evaluation of the efficacy of antibiotic therapy employed in this pathogenic context.

After reading the full text of each article and case report, relevant information was selected and evaluated. The most crucial aspects were recorded and summarized for analysis. Review articles, opinion articles, and articles with incomplete data or controversial diagnostic interpretations were excluded. A flowchart (Figure 1) was provided to illustrate the rationale for selecting the reference material for this review. Summarized results are categorized by species (Tables 1, 2, 3, and 4) and antibiotic resistance profile of the strains (Table 5), subsequently analyzed in the discussion section

Fig. 1. Flowchart of the screening process of publications according to the PRISMA model.



*Source: the authors

3. RESULTS

The systematic review of the currently available scientific literature revealed a total of 70 articles describing 76 cases of endocarditis caused by *Rothia* spp. The most frequently attributed species was *Rothia dentocariosa*, accounting for 37 cases, followed by *Rothia mucilaginosa* with 17 cases, *Rothia kristinae* with 11 cases, and *Rothia aerea*, also with 11 cases. Antibiotic resistance profiles were assessed in only 10 cases, where strains were identified as resistant to some antibacterial agents. Clinical manifestations were recorded as described by the manuscript authors.

Table 1. Key clinical aspects of endocarditis cases attributed to *Rothiaaeria*.

Reference	Gender	Age	Comorbidities or underlying conditions	Suspected infection sources	Other infection sites	Antibiotic treatment	Outcome
Tarumoto et al. [49]	Male	40	Allergic conjunctivitis. smoking	Not reported	Stroke	Ceftriaxone,Gentamicin and Imipenem	Deceased (stroke)
Holleran and Kasiah [50]	Not reported	58	-	Not reported	Multiple infective intracranialaneurysms. Subarachnoid haemorrhage	Not reported	Death (subarachnoid haemorrhage)
Thiyagarajan et al. [51]	Male	61	Hypertension and renal calculi	Not reported	Ceebralseptic emboli	Benzympenicillin, Rifampicin and Gentamicin	Discharged after 4 weeks
Crowe et al. [52]	Male	48	Hypertension, ex-smoker, IgM positive for <i>Mycoplasma pneumoniae</i>	Not reported	Multiple infective intracranial aneurysms; Cerebral septic embolization.	Benzympenicillin, Gentamicin, Ceftriaxone, Rifampicin, and Ciprofloxacin	Discharged
Hiraiwaand Izumi [53]	Male	63	Renal transplantation. Immunosuppressive therapy. Dental caries with gingival pain	Not reported	Cerebral septic emboli	Levofloxacin, Vancomycin, Ceftriaxone,Trimethoprim/ sulfamethoxazole, and Penicillin G	Discharged after 8 weeks
Nicodemo et al. [54]	Male	25	-	Not reported	-	Levofloxacin, Ampicillinand Vancomycin	Discharged after 5 weeks
Kim et al. ([55]	Male	53	History of ankylosing spondylitis, aortic valvuloplasty, tricuspid valvuloplasty, and a Maze operation owing to severe aortic valve regurgitation with atrial fibrillation. Four dental implant placements. Immunosuppressive therapy	Not reported	-	Ceftriaxone and Doxycycline	Discharged after 4 weeks with sequels (complete atrioventricular block occurred after surgery; permanent pacemaker implanted 10 days after the surgery)
Collarino et al. [56]	Male	57	History of of right sub-thalamic ischaemic stroke and severe mitral insufficiency without ventricular dysfunction for dystrophic mitral valve	Not reported	Subarachnoid haemorrhage. Femoral mycotic aneurysms	Amoxicillin and Gentamicin	Discharged after 4 weeks
Aoiyagi et al. [57]	Male	53	Smoking	Cut on his left thumb and wound licking	Multiple systemic embolisms.Subarachnoid hemorrhage.Mycotic	Cefmetazole, PenicillinG	Discharged

					aneurysms of the peripheral and visceral arteries		
Greve et al. [58]	Female	18	Phenylketonuria. Interstitial viral pneumonia.	Viral pneumonia	Skin rashes.	Gentamycin (resistant), Ampicillin and Ceftriaxone	Discharged
Zeng et al. [59]	Not reported	-	Not reported	Not reported	-	Imipenem, Cilastatin sodium and Vancomycin	Discharged after 36 days

*Source: the authors

Table 2. Key clinical aspects of endocarditis cases attributed to *Rothiadentocariosa*.

Reference	Gender	Age	Comorbidities or underlying conditions	Suspected infection sources	Other infection sites / clinical signs	Antibiotic treatment	Outcome
Pape et al. [60]	Male	61	-	Not reported	-	Gentamicin and Penicilin	Discharged
Schafer et al. [61]	Male	57	Refitting of dental bridges one month before	Oral infection	-	Fenoximetilpenicilina, Streptomycin and Penicilin G	Discharged after 4 months
Broeren and Peel [62]	Male	53	Impaired function of the mitral valve as a consequence of previous attacks of rheumatic fever. Dental carie.	Nor reported	-	Rifampicin, Penicillin and Gentamycin	Discharged after 8 weeks
Shands [63]	Male	41	Previous mitral valve prolapse, myxomatous degeneration, and regurgitant flow	Not reported	-	Penicilin and Vancomycin	Discharged
Isaacson and Grenko [64]	Male	27	Migraine headaches	Not reported	Brain abscess	Penicilin G	Discharged
Ruben [65]	Male	71	Previous valve prosthesis implant	Previous invasive cardiac interventions (14 years late)	-	Vancomycin, Gentamicin, Penicilin and Ceftriaxone	Discharged
Sudduth and Farrar [66]	Male	35	"Heart murmur" diagnosed fourteen years before. Alcohol abuse	Not reported	Perivalvular Abscess	Nafcillin, Gentamicin and Vancomycin	Discharged
Weersink et al. [67]	Male	17	Rheumatic with consequent cardiac valve disease	Not reported	Abdominal aneurysm	Ceftriaxone, Amoxicillin/Clavulanate, Gentamicin and Penicilin	Discharged
Binder et al. [68]	Female	70	Dental carie and Alcohol abuse	Oral infection	-	Penicilin, Netilmycin and Vancomycin	Deceased (mitral valve reconstruction surgery complications)
Binder et al. [68]	Male	67	Periodontal disease and residual root of a lost molar. Aortic valve	Oral infection	Bacteremia with maculopapular rash	Rifampicin and Ciprofloxacin	Discharged after nine weeks

replacement with a bioprosthesis 30 years before									
Binder [68]	et al.	Male	50	Extensive periodontitis	Oral infection	Abscess between aortic wall and composite graft. Skin rashes	Rifampicin and Ceftriaxone	Discharged after six weeks	
Kong ([69]	et al.	Male	37	Alcohol abuse. Several dental caries. Hepatic carcinoma diagnosed two months after endocarditis	Oral infection	-	Netilmicin, Metronidazole and Amoxicillin	Discharged	
Ferraz [70]	et al.	Male	54	valvular heart disease diagnosed 20 years late.	Not reported	Aortic root abscess	Penicilin, Gentamycin and Cloxacillin	Deceased (cardiac failure)	
Braden [71]	et al.	Female	7	Perimembranous ventricular septal defect with congestive heart failure	Not reported	-	Cefaclor, Amoxicilin, Imipenem-Cilastatin and Ceftriaxone	Discharged after 9 days	
Llopis and Carratalà[72]		Male	62	Alcohol abuse. Several dental caries.	Oral infection	Vertebral osteomyelitis	Penicilin, Gencamycin and Ceftriaxone	Discharged	
Nguyen al.[73]	et	Male	15	Poor dentalhygiene	Oral infection	-	Penicilin and Gentamicin	Discharged	
Ricaurte al.[74]	et	Male	49	“Heart murmur” since his childhood. Odontologic intervention (root canal) six months before. Cocaine and marijuana abuse.	Oral infectiion	Ischaemic stroke	Cefotaxime, Gentamycin and Penicilin G	Discharged	
Larkin [75]	et al.	Female	61	liliac embolectomy.Chronic atrial fibrillation and long- standing mitral stenosis	Not reported	-	Penicilin G	Discharged	
Salamon Prag [76]	and	Female	72	Inoperable rectal cancer and extensive abdominal metastasis.	Not reported	-	Antibiotic therapy not performed	Deceased (Deteriorated health status. No septic shock signals)	
Salamon Prag [76]	and	Male	51	Arterial hypertension. Dental abscesses.	Not reported	-	Penicillin G and Dicloxacillin	Discharged	
Salamon Prag [76]	and	Male	78	Acute myocardial Infarction. Admitted with angina pectoris. multiple diverticulosis and bleeding in the sigmoid colon.	Not reported	-	Penicillin G	Discharged	
Boudewijns al. [77]	et	Female	17	Impaired function of the aortic valve as a consequence of a congenital	Previous invasive cardiac interventions or	Subarachnoid hemorrhage. infective intracranial aneurysms.	Penicillin and Amikacin	Discharged after 20 days	

				bicuspid valve. Valvotomy at the age of 4 and percutaneous dilatation at the age of 13. Orthodontic treatment.	orthodontic treatment.	Skin rashes.		
Almuzara et al. [78]	Female	22	-		Not reported	Subarachnoid hemorrhage. Multiple septic embolisms in the SNC	Ceftriaxone, Vancomycin, Gentamicin, Penicilin	Discharged
Morris et al. [79]	Female	41	Bilateral nephrectomy and failed renal transplant. Peritoneal dialysis peritonitis eleven years before presentation.		Catheter contamination	-	Vancomycin, Gentamicin and Trimethoprim-sulfamethoxazole	Discharged
Sadhu et al. [80]	Male	55	Type 2 diabetes mellitus, essential hypertension, and liver cirrhosis due to hepatitis C. Several dental extractions without antibiotic prophylaxis.		Oral infection	Multiple cerebellar hemorrhages	Penicillin G and Gentamicin	Discharged
Shakoor et al. [81]	Male	37	Cholecystectomy after an episode of acute cholecystitis two months prior to presentation. Smoking.		Not reported	Osteomyelitis	Vancomycin, Penicillin-G and Ceftriaxone	Discharged
Chowdhary et al. [82]	Male	34	Methamphetamine abuse		Not reported	Ischaemic stroke	Vancomycin, Ceftriaxone and Doxycycline	Discharged after 2 weeks
Fridman et al. [83]	Male	58	-		Nor reported	Endophthalmitis. Cerebral infarct, with intracranial hemorrhaging	Amoxicillin, Clavulanate and Penicilin G	Discharged after 2 weeks
Willner et al. [84]	Female	62	Not reported		Not reported	Brain embolic infarction	Vancomycin, Ceftriaxone and Penicilin G	Discharged after 6 weeks
Doddapaneni et al. [85]	Male	65	Aortic valve stenosis, hypertension, and migraines		Not reported	Ischaemic stroke	Penicillin G and Ceftriaxone	Discharged
Myadam et al. [86]	Male	37	Three previous episodes of pneumonia with antibiotic treatment. Congenital bicuspid aortic valve stenosis with bioprosthetics. Dental abscesses		Oral infection	-	Rifampim	Discharged
Kisilevsky et al. [87]	Male	62	hypertension, type 2 diabetes mellitus and dyslipidaemia. Poor dental hygiene and inconsistent		Oral infection	Multiple infective intracranial aneurysms and ischaemic stroke	Ceftriaxone	Discharged after three months

			dental care. Dental caries.				
Elkattawy et al. [88]	Male	69	Diabetes mellitus, hypertension, hepatitis-C, and end-stage renal disease on hemodialysis	Not reported	Multiple embolic hemorrhagic strokes	Ceftriaxone and Penicillin	Discharged
Greve et al. [58]	Male	46	Previous valve prosthesis implant	Valve prosthesis implant	-	Vancomycin, Rifampicin, Gentamicin and Fenoximetilpenicilina	Discharged (no follow-up)
Franconieri et al. [32]	Female	21	Coarctation of the aorta repaired 18 days after birth and a persistent but asymptomatic ventricular septal defect	Not reported	-	Amoxicillin and Gentamicin	Discharged
Obi et al. [89]	Male	33	Implanted cardioverter defibrillator five years before	Cardioverter defibrillator implant surgery	-	Vancomycin, Gentamycin, Rifampin,	Discharged after 6 weeks
Jianjian et al. [90]	Male	40	Not reported	Not reported	Subarachnoid hemorrhage. Brain abscess. Thrombocytopenic purpura	Piperacillin Tazobactam, Tigecycline, Penicillin, Amikacin, and Meropenem	Discharged

*Source: the authors

Table 3. Key clinical aspects of endocarditis cases attributed to *Rothia kristinae*.

Reference	Gender	Age	Comorbidities or underlying conditions	Suspected infection sources	Other infection sites	Antibiotic treatment	Outcome
Lai et al. [91]	Female	89	Ischaemic bowel disease, , short bowel syndrome, Total parenteral nutrition, Port-A Catheter	Not reported	-	Vancomycin, Teicoplanin and Oxacillin	Discharged
Bastidas et al. [92]	Male	42	-	Accidental left hand incision	-	Gentamicin and Oxacillin	Discharged
Seyman et al. [93]	Male	65	Diabetes mellitus	Not reported	-	Ampicillin and Sulbactam	Discharged after 6 weeks
Citro et al. [94]	Male	74	Systemic hypertension. Diabetes mellitus.Diabetic foot.	Not reported	Pneumonia	Ceftriaxone, Metromidazole, Sulbactam/Ampicillin, Gentamicin and Rifampicin	Deceased (multiple organ failure due to sepsis)
Hollanda et al. [95]	Female	27	Systemic Lupus Erythematosus	Not reported	-	Not described	Discharged
Aleksic et al. [96]	Female	35	Chronic hepatitis C. Intravenous Illicit drug user.	Not reported	Stroke	Vancomycin, Gentamicin and Ceftriaxone	Discharged
Robles-	Male	56	Aortic-	Not reported	-	Cloxacillin and Amoxicillin	Discharged

Marhuenda et al. [97]			bifemoralbypassurgerysecondary toLeriche'ssyndrome					
Horino et al. [98]	Male	61	Hemodialysis patient. Diabetes mellitus	Not reported	Septic arthritis	Piperacillin, Ampicillin/Sulbactam and Gentamicin	Discharged	
Rojas-Molina et al. [99]	Female	44	-	Not reported	Endophtalmitis	Vancomycin, Ampicillin	Discharged	
Ali et al. [100]	Female	-	Ventricular septal defect. Poor dental hygiene and carious teeth.	Oral infection	-	Ampicillin, Tetracycline, Azithromycin, Ciprofloxacin, Linezolid, Vancomycin, Amikacin, Moxifloxacin, Gentamicin and Daptomycin	Discharged after 2 weeks	
Dewi et al. [101]	Female	25	-	Not reported	-	Cefotaxime, Gentamicin	Deceased (Cardiac arrest. Endocarditis by coinfection of <i>Rothia</i> <i>kristinae</i> and <i>Streptococcus alactolyticus</i>)	

*Source: the authors

Table 4. Key clinical aspects of endocarditis cases attributed to *Rothiamucilaginos*.

Reference	Gender	Age	Comorbidities or underlying conditions	Suspected infection sources	Other infection sites / clinical signs	Antibiotic treatment	Outcome
Rubin et al. [102]	Male	63	-	Not reported	-	Cephalosporin, Penicillin	Discharged after 5 weeks
Prag et al. [103]	Male	46	“heart murmur” diagnosed two years before	Not reported	-	Ampicillin, Penicilin G, Methicilin	Discharged
Coudron et al. [104]	Male	29	Intravenous drug abuse. Aortic valveendocarditiscaused by <i>Aspergillus fumigatus</i> . Aortic valve replacement with a porcine bioprosthesis four months before.	Not reported	-	Rifampin	Deceased (cardio-respiratory arrest)
Relman et al. [105]	Female	34	Illicit intravenous drug abuse.	Not reported	-	Nafcillin,Piperacillin,Gentamicin and Penicillin G	Discharged after 6 weeks
Pinsky et al. [106]	Male	35	Intravenous drug abuse. <i>Streptococcus mitis</i> Endocarditis 5 months before;	Not reported	-	Vancomycin and Gentamicin	Not reported

Aortic and mitral insufficiency							
Castaño et al. [107]	Male	79	Chronic heart failure;Chronic obstructive pulmonary disease.Cor pulmonale.Arterial hypertension.Right adrenal myolipomadiagnosed 4 years before. Immunossuppressive treatment.	Nor reported	-	Rifampicin, Vancomycin andTetracilin	Deceased (Severecardiac insufficiency)
Pérez-Vega et al. [108]	Male	44	Mitral valve prolapse diagnosed 10 years before	Not reported	-	Ampicillin, Gentamicin, Penicillin G	Discharge
Rolland and Wallet [109]	Male	73	Hypercholesterolemia. Arterial hypertension. Pulmonary embolism 4 years before.	Not reported	nodular liver lesions compatible with secondary abscesses.	Ceftriaxone, Gentamicin and Rifampin	Not reported
Faiad et al. [110]	Female	52	Intravenous drug user. Mitral and aortic infective endocarditis resulting in bioprosthetic replacement of both valves seven years before.	Not reported	Brain multiple small subcortical and pontine embolic infarct. Periaortic abscess	Vancomycin ,Gentamicin, Rifampin, Ampicillin and Penicillin G	Deceased
Faiad et al. [110]	Male	21	Pre-B acute lymphocytic leukemia diagnosed two years before. Immunossuppressive therapy.	Not reported	-	Vancomycin, Ceftriaxone, Rifampin,	Discharged
Bruminhent et al. [111]	Male	36	Previous <i>Streptococcus mitis</i> mitral valve endocarditis and prosthetic valve replacement five years before.	Not reported	Left popliteal artery thrombosis	Vancomycin, Piperacillin-tazobactam	Discharged
Ramanan et al. [17]	Female	28	Intravenous drug user. Prosthetic aortic valve. native tricuspid and aortic valve. Previous endocarditis by <i>Staphylococcus aureus</i>	Not reported	-	Ceftriaxone, Vncomicycn and Rifampin	Discharged
Sugunesegran et al. [112]	Female	-	Two previous sternotomies.	Not reported	-	Antibiotic therapy not described	Discharged
Song et al. [113]	Male	65	Aortic valvoplasty two decades before. Three episodes of prosthetic valve endocarditis in the last 11 years by <i>Enterococcus avium</i> , <i>Enterococcus faecalis</i> , and	Not reported	Multiple brain hemorrhages suggesting septic emboli on both hemisphere, corticomedullary junction area and cerebellum	Vancomycin, Rifampin, Ampicillin and Ceftriaxone	Discharged (Right side motor weakness remained as a sequela to the

			<i>Streptococcus gallolyticus</i> . Diabetes mellitus, hypertension, dyslipidemia, and alcoholic liver.				patient)
Haddad et al. [114]	Male	80	Coronary artery disease, mild intermittent asthma and moderate aortic stenosis	Not reported	-	Ceftriaxone, Vancomycin	Discharged
Abdelmaseih et al. [115]	Male	46	Chronic hepatitis C. Intravenous Illicit drug user.	Not reported	-	Vancomycin	Discharged after 6 weeks
López et al. [116]	Female	68	Extraction of a third molar	Oral infection	Embolic splenic infarction due to septic embolism	Antibiotic therapy not described	Discharged

*Source: the authors

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Table 5. Antibiotic sensitivity profile of *Rothia* spp. strains isolated from patients with endocarditis showing some degree of antimicrobial resistance.

Reference	Species	Antibiogram results		
		Resistant	Intermediary resistance	Sensible
Greve et al. [58]	<i>R. aeria</i>	Gentamycin		Penicilin, Ampicillin, Amoxicillin-clavulanic acid, Piperacillin+Tazobactam, Meropenem, Moxifloxacin, Clindamycin, Vancomycin, Rifampicin, Linezolid, Tigecycline
Ruben [65]	<i>R. dentocariosa</i>	Amikacin, Ciprofloxacin, Kanamycin, Sulfamethoxazole-trimethoprim	Erythromycin, Gentamycin, Rifampin, Tetracycline.	Ceftizoxime, Ceftriaxone, Cephalothin, Chloramphenicol, Imipenem, Penicillin, Vancomycin,
Kong et al. [69]	<i>R. dentocariosa</i>	Tobramycin, Gentamicin, Doxycycline, Pristinamycin, Sulfonamide, Pefloxacin		Penicillin, Amoxicillin, Imipenem, Erythromycin, Spiramycin, Rifampin.
Greve et al. [58]	<i>R. dentocariosa</i>	Gentamycin		Penicillin, Cefotaxime, Vancomycin, Rifampicin
Ali et al. [100]	<i>R. kristinae</i>	Erythromycin, Clindamycin, Tobramycin, Vancomycin, Teicoplanin, Tetracycline, Rifampicin, Cefoxitin, Penicillins and Cephalosporins	Levofloxacin, Gentamycin	Moxifloxacin, Sulphamethoxazole -Trimethoprim
Prag et al. [103]	<i>R. mucilaginosa</i>	Nalidixic acid	Mecillinam	Cephalothin, Cefoxitin, Cefotaxime, Carbenicillin, Sulphosomidine, Trimethoprim, Tobramycin, Erythromycin, Tetracycline, Chloramphenicol, Clindamycin, Nitrofurantoin, Piperacillin
Relman et al. [105]	<i>R. mucilaginosa</i>		Tetracycline	Penicillin, Methicillin, Cephalothin, Vancomycin, Chloramphenicol, Clindamycin, Erythromycin

Castañó et al. [107]	<i>R. mucilaginosa</i>	Ciprofloxacin, Cotrimoxazol, Gentamycin, Penicilin		Rifampicin, Vancomycin, Nitrofurantoin, Tetraciclín, Teicoplanin, Cloranfenicol
Song et al. [113]	<i>R. mucilaginosa</i>	Clindamycin, Sulfamethoxazole- Trimethoprim, Tobramycin		Penicilin, Ampicillin-sulbactam, Erythromycin, Rifampin, Tetracycline, Levofloxacin, Vancomycin
Haddad et a. [114]	<i>R. mucilaginosa</i>	Sulfamethoxazole-Trimethoprim	Clindamycin, erythromycin	Penicilin, Vancomycin, Levofloxacin

4. DISCUSSION

The scientific literature on endocarditis caused by *Rothia* spp. is scarce, and currently, there are no systematic studies addressing this clinical condition. Systematic review studies may not necessarily provide a reliable overview, particularly for pathogenies considered rarer, as case studies are often outlined based on peculiarities, circumstances, or specificities of medical practice [117,118,119]. However, the investigative approach of systematic literature review has the potential to identify particular situations, formulate hypotheses, compare situations to consolidated knowledge, and highlight trends, associations, or clinical reasoning that can guide future systematic research [117].

The incidence of endocarditis caused by *Rothia* spp. is likely underreported. The species of this genus are challenging to identify and are often confused with *Staphylococcus* spp. [108,120,121,122], *Streptococcus* spp. [108,120,121,122], *Micrococcus* spp. [55,120], or *Nocardia* spp. [31,55] due to the pleomorphism of the genus. The high sensitivity of most *Rothia* spp. strains causing endocarditis to antibiotics may contribute to the lack of specific diagnosis and registration as an infectious agent. The resolution of the infectious clinical condition with empirical antibiotic therapy does not stimulate greater interest on the part of the medical team in the effort to precisely identify the pathogenic agent.

The taxonomic resemblance among species within the *Rothia* genus adds complexity to the analysis of pathogenesis caused by this group of microorganisms. *Rothiadentocariosa* and *Rothiaaeria* exhibit similar biochemical profiles, both manifesting positivity for nitrate reduction, α -glucosidase, alanine-phenylalanine-proline arylamidase, and esculin hydrolysis, while demonstrating negativity for urease [55,123]. Some strains exhibit even more closely aligned biochemical profiles, testing positive for alanine-phenylalanine-proline arylamidase, α -glucosidase, and esculin hydrolysis, and negative for urease. Notably, certain strains of *Rothiadentocariosa* display biochemical profiles akin to those of *Rothiamucilaginoso*[55]. The differentiation among various *Rothia* species through biochemical reactions poses challenges due to their similar profiles [52,124]. Sequencing comparison of 16S rRNA between *Rothiadentocariosa* and *Rothiamucilaginoso* reveals homology at 98.0% and 96.4%, respectively, to the genetic material of *Rothiaaeria*[123]. These attributes substantially contribute to the inherent lack of precision in laboratory identification tests. Michon et al. [125] reveal a unique microorganism was identified as *Rothiadentocariosa* or *Rothiakristinae* by the Api CORYNE and ID 32 STAPH strips method from BioMérieux with 99.9% similarity; as *Rothiakristinae* by the VITEK 2 system from BioMérieux with 99.9% similarity; until genetic analysis of 16S rRNA sequencing identified it as *Rothiaaeria*. The 16S rRNA sequencing of this strain also showed little divergence from *Rothiamucilaginoso*[125]. Fatahi-Bagfi[31] reports that a strain of *Rothiadentocariosa* was distributed by the Swiss External Quality Assessment Scheme in Bacteriology and Mycology for analysis in 50 laboratories, and only 36 (72%) were able to provide an accurate result for species identification. In this context, the reliability of species identification in various previous studies cannot be guaranteed, especially in those where 16S rRNA sequencing was not performed. It is still likely that many cases of endocarditis identified as caused by *Rothiadentocariosa* may have actually been caused by *Rothiaaeria*, described more recently. Therefore, the retrospective analysis of infectious endocarditis cases by *Rothia* species should be conducted without considering species distinction as a baseline for comparison.

Among the *Rothia* species attributed as agents of endocarditis, *Rothiadentocariosa* was the most prevalent. Considering the year 2012, when *Rothiaaeria* was first identified as a causative agent of endocarditis, the number of reported cases of endocarditis caused by *Rothiadentocariosa* and *Rothiaaeria* are equivalent, with 11 cases reported for each species. This parity suggests that up to 50% of cases previously categorized as *Rothiadentocariosa* may have been misclassified. Disregarding alternative possibilities of misidentification, the combined *Rothiadentocariosa* (48.69%) and *Rothiaaeria* (14.47%) accounted for 63.16% of cases, followed by *Rothiamucilaginoso* with 22.37% and *Rothiakristinae* with 14.47%. The comprehensive examination of endocarditis cases involving *Rothia* species reveals an average patient age of 48.5 years, with male predominance at 71.6%.

Within the set of reported cases of *Rothia* spp. endocarditis, 11.84% (9/76) progressed to mortality. This observed mortality rate is notably lower when compared to documented rates in studies investigating infectious endocarditis caused by other pathogens, where mortality rates typically fall within the range of 18% to 32% for individuals affected by this clinical condition [126,127]. The diminished mortality rate may be attributed to the heightened susceptibility of a substantial proportion of *Rothia* spp. strains to a broad spectrum of antibiotics, a lower inherent pathogenicity intrinsic to the genus, or a potential bias arising from suboptimal sampling. Notably, all patients in fatal cases exhibited multiple comorbidities, immunosuppression, or succumbed to mortality due to complications arising from surgical procedures.

Only 27.6% of the scrutinized case reports provided insights into the putative primary focus of infection or contamination. Within this specific subset, an overwhelming 61.9% implicated dental procedures or substandard oral hygiene practices, conceivably linked to periodontal disease, as the primary focus. It is noteworthy that two cases within this subset explicitly mentioned a history of dental caries.

Despite assertions by numerous researchers that infections caused by *Rothia* spp. are strongly linked to individuals with compromised immune systems [17,56,85,89,108,111,113,114,115,122,125], only seven patients diagnosed with endocarditis were explicitly characterized as immunocompromised. Additionally, four individuals could be presumed immunocompromised due to a combination of comorbidities, constituting 14.47% of the total cases. Endocarditis patients

attributed to *Rothia* spp. presented a diverse range of comorbidities and underlying conditions, with a higher association with pre-existing cardiac issues, dental caries, inadequate oral hygiene, and dental procedures.

A study conducted by Tsuzukibashi et al. [128] reported *Rothiamucilaginos*a as the most prevalent species in human saliva (74%), followed by *Rothiadentocariosa* (16%) and *Rothiaaeria* (10%). Conversely, AlEraky et al. [129] investigated bacterial species associated with the progression of dental caries, revealing that *Rothiamucilaginos*a and *Rothiaaeria* predominate in individuals with a low incidence of caries, while *Rothiadentocariosa* is more prevalent in individuals with a high incidence of caries. Moreover, the absence of *Rothiamucilaginos*a and *Streptococcus salivarius* was noted in individuals with a high incidence of caries, in contrast to their coexistence in individuals with a low incidence of caries. Concerning *Rothia* spp. endocarditis, 29.7% (11/37) of cases were ascribed to *Rothiadentocariosa* in patients with a clinical history of dental caries, periodontal disease, and poor oral hygiene. In contrast, only one case (1/17) was attributed to *Rothiamucilaginos*a following the extraction of a molar tooth. These findings suggest that inadequate oral hygiene, periodontitis, and dental caries may play a significant role in the prevalence of *Rothiadentocariosa* as an etiological agent of endocarditis through direct invasive processes. In the context of cariogenic processes and subsequent dysbiosis of the oral microbiota, bacteria with invasive potential, such as *Streptococcus* spp., may access blood vessels in periodontal tissues and dental pulp, ultimately colonizing the endocardium after bacteremia [130,131]. Considering the parallels between the pathogenic processes leading to endocarditis caused by *Streptococcus* spp. and *Rothia* spp., it is noteworthy that species from both genera possess the ability to produce adhesins. These adhesins play a pivotal role in biofilm formation in both the oral cavity and cardiac valves [10,40,41]. In this context, it is plausible that *Rothiadentocariosa*, prevalent in the oral cavity of patients with a high incidence of caries and periodontal diseases, may exhibit a more prominent invasive potential compared to other species within the genus, following an infectious dynamic akin to that observed in *Streptococcus* spp. for the entry of bacteria into the bloodstream and subsequent colonization of the endocardium. This hypothesis highlights the predominance of endocarditis linked to deteriorating oral health and subsequent invasive infection by *Rothiadentocariosa*, as observed in our literature analysis. The subset of endocarditis cases in patients with a history of dental procedures, dental caries, and poor oral hygiene constituted 17.1%.

Recent research outlines new trends in the factors of contamination by infectious agents triggering endocarditis, highlighting a significant increase in the incidence of this cardiac condition in individuals who use intravenous illicit drugs [132-138], a considerable proportion of whom are carriers of the hepatitis C virus [132-136]. The investigation of case reports pertaining to endocarditis caused by *Rothia* spp. substantiates this inclination, wherein nine patients (11.84%) acknowledged intravenous illicit drug use. This incidence aligns with findings by Marques et al. [127], who reported that 13.4% of patients exhibiting infectious endocarditis caused by other pathogens were users of intravenously administered illicit drugs. In the examination of cases involving endocarditis due to *Rothia* spp., four patients (5.26%) were diagnosed with chronic hepatitis C, indicative of potential syringe and needle sharing. The likelihood of underestimating the incidence of endocarditis resulting from needle contamination during illicit drug use exists, given that drug users may not consistently disclose such substance use during the anamnesis process [139].

The prevalence of pre-existing cardiovascular diseases is prominent among patients who develop endocarditis caused by *Rothia* spp., occurring in 50% (38/76) of the reported cases. Within this subset of patients with cardiovascular diseases, 28.9% (11/38) were identified as hypertensive, 31.5% (12/38) had undergone prosthetic valve implantation, and 18.4% (7/38) exhibited valvular insufficiency. Six cases (15.79%) involved individuals with a history of previous episodes of endocarditis, encompassing one instance of fungal endocarditis (*Aspergillus fumigatus*) and five instances of bacterial endocarditis caused by Gram-positive cocci (*Streptococcus mitis*, *Streptococcus gallolyticus*, *Staphylococcus aureus*, *Enterococcus avium*, and *Enterococcus faecalis*). Bussani et al. [140] emphasize that the persistence of risk factors contributes to the recurrence of infectious endocarditis. In the study conducted by Marques et al. [127], the majority of cases were associated with native valves (71.6%), while the remainder were linked to prosthetic heart valves (25.4%), a distribution analogous to that observed in *Rothia* spp. endocarditis cases (71.1% in patients with natural heart valves and 28.9% in patients with prosthetic heart valves). The incidence of diabetes was reported in 9.21% (7/76) of patients. Overall, the prevalence of cardiovascular diseases, arterial hypertension, prosthetic valve implants, valvular insufficiency, prior episodes of endocarditis, and diabetes mirrors findings in other studies on infectious endocarditis [127,140,141], suggesting that risk factors for *Rothia* spp. endocarditis align closely with those associated with endocarditis caused by other pathogens.

Embolisms are commonly recognized as complications in the context of infectious endocarditis. The incidence of embolic events, occurring in roughly 40% of cases, is a notable feature. Events related to embolism in the Central Nervous System are identified in approximately 25% of patients [126,142]. Among individuals with *Rothia* spp. endocarditis, embolic events were documented in 35.53% (27/76) of cases, with 28.95% (22/76) specifically involving Central Nervous System embolisms—incidences consistent with those observed across endocarditis cases in general. Mycotic aneurysms were discerned in 6.58% (5/76) of patients. Although constituting a small proportion of all intracranial aneurysms, these aneurysms carry a heightened mortality risk upon rupture [143,144,145]. One fatal case of *Rothiaaeria* endocarditis was linked to subarachnoid hemorrhage resulting from such an aneurysm. Despite the noted association between strokes and intravenous illicit drug use in infectious endocarditis cases reported by Ridha et al. [146] for the period between 2005 and 2015, this specific type of embolism was documented in only two case reports of *Rothia* spp. endocarditis.

The presence of abscesses in *Rothia* spp. endocarditis indicates that bacteria of this genus have the potential to disseminate hematogenously, leading to the establishment of focal infections in diverse anatomical sites. In the compilation of case reports, three aortic abscesses, one perivalvular abscess, one cerebral abscess, and one hepatic abscess were described. The presence of aortic abscesses in patients with endocarditis is considered a high-risk factor [147,148,149], and it was diagnosed in two cases of *Rothia* spp. endocarditis (one aortic root abscess and one peri-aortic abscess). Remarkably, these cases accounted for 22.22% of the total fatalities documented in the study. Additionally, two cases of osteomyelitis induced by *Rothia* spp. endocarditis were observed, presumably as a secondary consequence of endocardial infection. Additionally, two cases of osteomyelitis induced by *Rothia* spp. endocarditis were observed, presumably as a secondary consequence of endocardial infection.

In a comprehensive literature review, Franconieri et al. [32] emphasized a robust association between *Rothia mucilaginosa* and endocarditis, attributing 70% of *Rothia* genus-induced endocarditis cases to this particular species. In contrast, our systematic analysis of the current scientific literature reveals a higher proportion of endocarditis cases associated with *Rothia dentocariosa*. With an examination of an extended case report sample (76 cases) compared to the study conducted by Franconieri et al. [32] (50 cases), each species presenting distinct incidence rates. It emphasizes the need for systematic studies, including control groups, comprehensive patient histories, and standardized genetic sequencing methodology to determine the genuine epidemiological landscape of *Rothia* spp. endocarditis.

Despite the absence of systematic investigations concerning endocarditis caused by *Rothia* spp., studies on bacteremias caused by species within this genus reveal notable disparities in epidemiological patterns. In a retrospective 12-year analysis of *Rothia* spp. bacteremias at a Hungarian teaching hospital, Gajdács et al. [122] documented 37 cases, being *Rothia dentocariosa* responsible for 28 instances (75.67%) and *Rothia mucilaginosa* for the remaining 9 cases (24.33%). A slight female predominance (56.76%) was discerned, and the average age of affected patients stood at 57 years. Conversely, Odeberg et al. [150], in a retrospective study spanning from 2012 to 2021 at a Swedish clinic, identified 108 cases of *Rothia* spp. bacteremia, classifying 24 as infections and the rest as contamination. *Rothia mucilaginosa* accounted for 53 cases, with 16 deemed genuine infections. *Rothia dentocariosa* was detected in 26 cases, of which two were considered authentic infections. *Rothia kristinae* featured in two cases classified as contamination, while *Rothia aerea* was identified in a lone case, also categorized as contamination. Unidentified *Rothia* species were associated with 27 cases, of which six were classified as true infections. Distinctively, half of the cases considered genuine infections occurred in immunocompromised individuals. The substantial epidemiological variances between the findings of Gajdács et al. [122] and Odeberg et al. [150] highlight the imperative for timely and systematic research to elucidate such disparities.

The majority of *Rothia* spp. strains responsible for endocarditis displayed susceptibility to most or nearly all antibiotics (86.84%). However, a subset of strains (13.16%) manifested resistance to antibiotics across various classes, encompassing Aminoglycosides (Gentamicin, Amikacin, Kanamycin, and Tobramycin), Quinolones (Ciprofloxacin, Pefloxacin, and Nalidixic acid), Sulfonamides (Sulfamethoxazole-trimethoprim), Tetracyclines (Doxycycline and Tetracycline), Macrolides (Erythromycin), Lincosamides (Clindamycin), Glycopeptides (Vancomycin and Teicoplanin), Streptogramins (Pristinamycin), Rifamycins (Rifampicin), Cephamycins (Cefoxitin), Penicillins, and First and Second-generation Cephalosporins. The *Rothia kristinae* strain, as isolated by Ali et al. (2020), is an illustrative example of a multidrug-resistant strain, lacking sensitivity to antibiotics from diverse classes (Macrolides, Lincosamides, Aminoglycosides, Glycopeptides, Tetracyclines, Rifamycins, Penicillins, and First and Second-generation Cephalosporins).

The acquisition of antimicrobial resistance by pathogens is widely acknowledged as one of the foremost contemporary threats to human health. Antibiotic utilization exerts selective pressure not only on the target pathogens but also on the entire microbiome, wherein resilient microorganisms persist as commensals post-treatment. Resistance in commensal bacteria entails risks, as these microorganisms function as reservoirs of resistance genes that may be horizontally transferred to pathogens or precipitate opportunistic infections in subsequent instances. This predicament assumes even greater gravity when antibiotic therapy is administered incompletely or inadequately [151,152,153]. In instances of opportunistic infections by *Rothia* spp., apart from the selection of resistant strains, antibiotic therapy can induce dysbiosis, diminishing organisms that modulate the abundance of normal microbiota species. This, in turn, fosters the proliferation of *Rothia* spp., heightening the probability of this bacteria's involvement in inflammatory, lytic, and cariogenic processes that precede invasive infection [154,155,156,157,158,159].

While the primary focus and empirical antibiotic therapy planning for infectious endocarditis target bacteria more prevalent in these infections, such as *Staphylococcus* spp., *Streptococcus* spp., and *Enterococcus* spp., the effectiveness of currently recommended regimens can be extrapolated to the analysis of *Rothia* spp. endocarditis. Recent research suggests that the use of aminoglycosides, once considered part of standard treatment, should be restricted due to the high incidence of strains resistant to these antibiotic compounds [160,161,162,163]. This recommendation aligns with the resistance patterns exhibited by *Rothia* spp., wherein 7 out of the 10 strains resistant to antibiotics were not sensitive to at least one aminoglycoside, notably gentamicin, with four resistant strains and two moderately sensitive. It is worth mentioning that gentamicin was part of the antibiotic therapy regimens in 38.16% (29/76) of *Rothia* spp. endocarditis cases.

In a consensus article authored by 51 researchers from ten different countries and published in the Journal of the American Medical Association, the recommended antibiotic therapy regimen for managing cases of infectious endocarditis involves a combination of Vancomycin or Daptomycin and beta-lactams, specifically Ceftriaxone or Cefazolin. The authors advocate reserving aminoglycosides and Rifampin for deployment in definitive antibiotic therapy, contingent upon the identification of the etiological agent [164]. By aligning the proposed antibiotic therapy regimen by McDonald et al. [164] with the antibiotic susceptibility profiles of *Rothia* spp. strains responsible for endocarditis, it becomes evident that this regimen holds efficacy in curbing the disease's progression caused by this particular pathogen. The latest recommendations from the European Society of Cardiology propose an empirical treatment approach for infectious endocarditis, encompassing Ampicillin in conjunction with Ceftriaxone or Flucloxacillin, and Gentamicin, all administered prior to ascertaining the causative agent [163]. When considering the antibiotic susceptibility patterns in *Rothia* spp. strains triggering endocarditis, this antibiotic therapy regimen proves effective, encompassing substances to which these strains exhibited susceptibility, with the sole exception being Gentamicin.

4. CONCLUSION

Species belonging to the *Rothia* genus pose challenges in identification due to pleomorphism and genetic proximity. Regarding species implicated as causative agents in endocarditis cases, prevalent strains include *Rothiadentocariosa* (48.69%), *Rothiamucilaginoso* (22.37%), *Rothiaaeria* (14.47%), and *Rothiakristinae* (14.47%). The average age of affected patients was 48.5 years, with a male predominance of 71.6%.

Endocarditis induced by *Rothia* spp. lacks distinctive clinical features specifying its uniqueness in comparison to Gram-positive bacterial endocarditis cases. The observed lower mortality rate (11.84%) compared to endocarditis caused by other pathogens may be attributed to heightened susceptibility of most strains to a diverse range of antibiotics, facilitating resolution during empirical antibiotic therapy. This characteristic, combined with the challenge of *Rothia* spp. species identification based on cultural and microscopic characteristics, contributes to significant underreporting of endocarditis involving these bacteria.

The predominant risk factor in *Rothia* spp. endocarditis is the preexistence of cardiovascular diseases (50%). Other relevant factors included dental procedures, caries, and poor oral hygiene (17.1%), immunocompromise (14.47%), intravenous drug use (11.84%), and diabetes (9.21%). Among extra-cardiac consequences, embolic events were documented in 35.53% of patients, mainly manifesting as Central Nervous System embolisms, observed in 28.95% of total cases. Mycotic aneurysms were identified in 6.58%.

Merely 13.16% of strains causing endocarditis exhibited resistance to antibiotics, although some strains presented multidrug resistance. We caution that projections about the efficacy of antibiotic therapy regimens in *Rothia* spp. endocarditis cases based on the percentage of resistant strains in this review study should be considered in conjunction with current recommendations from systematic studies and patient histories. Given the limited number of reported cases, antibiotic resistance testing, regional and individual patient microbiota variations, the results provide a static overview spanning almost four decades, and antibiotic susceptibility patterns have decreased for almost all pathogenic species, especially those constituting the normal human microbiota. We suggest that empirical antibiotic therapy be based on the recommendations of the European Society of Cardiology, offering a broader antibiotic spectrum, also considering the patients' history of antibiotics used in previous infections to establish a personalized regimen with a better likelihood of success.

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