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Recent advances in sensor arrays aided by machine learning for pathogen identification

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The development of rapid and accurate pathogen detection methods is of paramount importance for slowing the evolution of antibiotic resistance in bacteria. However, the high similarity between different pathogens, especially between antibiotic-sensitive and antibiotic-resistant strains of the same species, presents great challenges for the precise discrimination of pathogens. In recent years, chemical nose strategies, i.e. sensor arrays, have achieved certain success in pathogen discrimination. Currently, chemical nose strategies for identifying pathogens are mainly designed from two perspectives: the disparity in extrinsic properties (biomolecules, charge, and hydrophobicity of the bacterial surface) and intrinsic properties (processes and products mediated by bacterial enzymes) among different pathogens. Biosensing probes capable of responding to these properties are introduced for pathogen detection. The output signals are then processed and analyzed by machine learning algorithms to visualize the multidimensional detection results and achieve pathogen discrimination. This paper introduces the latest developments in sensor arrays for pathogen identification based on the extrinsic and intrinsic nature of bacteria, highlights the recognition mechanism of probes for bacteria, and outlines the current challenges and prospects of sensor arrays for pathogen discrimination.

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1. Introduction

Pathogens, which can be transported through air, water, and food, are intimately linked to daily life and pose significant threats to human health.^{1,2} Antibiotics inhibit bacteria growth or kill bacteria through various mechanisms such as

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inhibiting bacterial cell wall synthesis, disrupting protein synthesis, etc. While antibiotics have saved countless lives, their overuse and misuse have led to a rise in antimicrobial resistance.³⁻⁵ Therefore, the differentiation and identification of pathogens, especially antibiotic-resistant strains, are crucial for the selection of appropriate antibiotics for the treatment of bacterial infections and slowing down the evolution of antimicrobial resistance. Conventional methods, including bacterial culture, microscopic observation, and gene sequencing techniques, are accurate but time-



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detection of antibiotic-resistant bacteria.

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consuming, which may delay the optimal time for treatment, lead to deterioration of the disease, and endanger patients' lives.⁶⁻⁹ Therefore, the development of rapid and accurate bacterial identification and differentiation methods is of great importance in clinical diagnosis. Emerging bacterial detection techniques, such as polymerase chain reaction (PCR),¹⁰ surface-enhanced Raman scattering (SERS),¹¹ and mass spectrometry (MS),^{12,13} although faster than bacterial culture, require expensive and sophisticated instruments and specialized technicians, making them difficult to implement widely.^{7,14} Therefore, a variety of simple and convenient biosensors for detecting bacteria have been developed based on ligand-receptor-specific recognition. These methods exhibit high specificity, as each probe can only detect a particular bacterial strain, but they are limited in detecting unknown bacterial samples in real-life situations such as food samples and clinical samples.

To address these issues, chemical nose strategies designed based on simulated animal olfaction have been widely proposed.¹⁵ The chemical nose strategy, also known as the sensor array approach, usually adopts multiple recognition probes capable of recognizing targets, and sensitively recognizing the subtle differences among them.^{15–17} These differences are then converted into optical or electrical signals through the probes, processed with statistical methods, and then aided by appropriate machine learning algorithms to realize the classification of different targets (Scheme 1).^{18,19} To be brief, array-based biosensors can spontaneously identify multiple targets, with differences in their physical and chemical properties that can be recognized by the sensor array.

Based on the discrimination principles, the sensor arrays are mainly divided into two types (Scheme 2).^{16,19} (1) Pathogen discrimination through differences in the surface



Scheme 1 Schematic illustrating the workflow of sensor arrays for differentiating bacteria. Scheme 1 was created with https://BioRender.com.



Scheme 2 Schematic overview of the recognition mechanism of sensor arrays for bacteria identification, constructed based on the intrinsic and extrinsic properties of bacteria. Note: VOCs: volatile organic compounds. N-VOCs: non-volatile organic compounds. D-AA-X: D-amino acid-tags. Van: vancomycin. AMPs: antimicrobial peptides. AIE: aggregation-induced emission luminogens. VIE: vibration-induced emission luminogens. Scheme 2 was created with https://BioRender.com.

properties. The bacterial surface contains a variety of biomolecules, including functional groups, proteins, polysaccharides, etc., which can be recognized by probes for identifying bacteria; additionally, these biomolecules codetermine the bacterial surface charge and hydrophobicity. With probes binding to bacteria through electrostatic and hydrophobic interactions, the disparity in the surface charge and hydrophobicity of pathogens can be detected and pathogen discrimination can be achieved. (2) Pathogen discrimination through differences in the intrinsic characteristics of bacteria. The intrinsic characteristics mainly refer to the biochemical processes mediated by enzymes produced by bacteria. As bacteria possess various enzyme systems, their biochemical processes are diverse. Various metabolites are produced by bacterial metabolism, primarily including H⁺, volatile organic compounds, and non-volatile organic compounds; bacteria incorporate exogenous D-amino acids (D-AAs) into their peptidoglycan through the mediation of transpeptidases to label the peptidoglycan; bacteria reduce metal ions into nanoparticles with various shapes, sizes, and crystalline structures through some reductases. Based on these differences among bacteria, each bacterium possesses a unique fingerprint response spectrum. To discriminate bacteria more quickly and accurately, appropriate algorithms are selected to preprocess and analyze the signal output by the sensors. In this review, recent developments of sensor arrays for bacterial recognition

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and differentiation are discussed in terms of the intrinsic and extrinsic properties of bacteria. The working principles, analysis performance, advantages, and disadvantages of the reported chemical nose strategies for pathogen identification in recent years are compared and discussed. Finally, the challenges and perspectives for array-based biosensors in bacterial identification are summarized.

2. Identification mediated by bacterial surface properties

Bacteria are generally categorized into Gram-positive and Gram-negative based on the differences in their cell walls. The surface of Gram-positive bacteria is mainly composed of peptidoglycan (which accounts for 50% to 80% of the total dry weight) and teichoic-acid, and Gram-negative bacterial walls cell mainly consist of lipopolysaccharide, peptidoglycan, phospholipid, and lipoprotein. These composites are mainly responsible for the overall surface charge of bacteria.^{20–22} Due to the genomic difference, the composition of the surface varies among different genera/ species/strains, resulting in differences in the overall surface charge or hydrophobicity. For the same strains, culture environments such as evolutionary pressures also lead to differences in surface compositions.²²⁻²⁴

For sensor arrays based on the difference in bacterial surface properties to achieve bacteria identification, they can be roughly divided into two categories: those based on the overall difference in bacterial surface charge or hydrophobicity through noncovalent recognition between bacteria and probes, and those employing probes that specifically recognize a certain type of composite on the bacteria surface.

2.1. Noncovalent recognition – electrostatic and hydrophobic interaction

The principle to distinguish bacteria via the disparity of the surface charge and hydrophobicity is that, either the difference in bacterial gene or culture environment would lead to the compositional difference in the bacteria surface, and thereby affect the overall surface charge and hydrophobicity of a certain type of bacterium at a specular situation.²⁵⁻²⁸ By sensing the differences in bacterial surface potential and hydrophobicity with a series of probes, rapid distinguishment of bacteria can be achieved. These proposed approaches do not require complex instrumentation or specialized technicians, and the detection coverage is broad. In this section, we will discuss sensor arrays constructed by using various sensing materials (nanomaterials, small molecules, and conjugated polymers) that are capable of identifying bacteria through electrostatic and hydrophobic interactions.

2.1.1. Noncovalent recognition of bacteria by nanomaterials. Nanomaterials, which encompass a range of materials including precious metal nanomaterials, rare earth

nanomaterials, and carbon-based nanomaterials, among others, due to their small size, facile surface modification, and low biotoxicity, have been widely employed in various fields such as biosensing in recent years.²⁹⁻³²

Gold nanomaterials exhibit low biological toxicity, large surface area, ease of functionalization, and tunable optical property. These advantages facilitate fruitful applications of gold nanomaterials for bacteria detection and identification. For instance, a novel three-dimensional optical sensing array based on gold nanoparticle/gold nanocluster (AuNP/AuNC) composites was developed by our group to discriminate various Gram-negative bacteria (Fig. 1A).³³ The nanocomposites consist of cetyltrimethylammonium bromide-coated gold nanoparticles (CTAB-AuNPs, positively charged) and vancomycin templated gold nanoclusters (Van-AuNCs, negatively charged). CTAB imparts positive charges and hydrophobicity to AuNPs. Bacteria with negative charges compete with Van-AuNPs to bind with CTAB-AuNPs, leading to the dissociation of the nanocomposite, and resulting in changes in three types of optical signals. With the assistance of linear discriminant analysis (LDA), the sensor array can selectively identify 10 strains of Gram-negative bacteria (including 3 antibiotic-resistant strains), and is able to analyze bacterial samples with low concentration (OD_{600} = 0.015).

Furthermore, Xianyu's group developed plasmonic nanosensors to determine the antibiotic-sensitive and antibiotic-resistant phenotypes among ESKAPE pathogens based on peptide-functionalized AuNPs (P-AuNPs) differing in surface plasmon resonance (SPR) wavelength, diameter, hydrophobicity, and surface charge (Fig. 1B).³⁴ The six types of P-AuNPs interact with ESKAPE through hydrophobic and electrostatic interactions, which, with different binding forces, result in the attachment of different amounts of P-AuNPs on the bacterial surface. The SPR spectra of the P-AuNPs display a blue shift after incubating with bacteria, resulting in distinct SPR fingerprint profiles among different bacteria. Processing SPR spectra with machine learning algorithms can differentiate antibiotic-sensitive and resistant phenotypes of ESKAPE pathogens within 20 min, with an overall accuracy of 89.74%.

Lanthanide nanoparticles possess excellent photostability, multicolor emission, tunable weak autofluorescence background, and strong resistance to photobleaching, greatly improving the sensitivity and reliability of detection.36 Wang et al. reported a multi-emission sensor array based on lanthanide encoding, where lanthanide nanoparticles with different surface charges and hydrophobicity were prepared as recognition probes to identify biofilms (Fig. 1C).³⁵ Three lanthanide nanoparticles, which emit green fluorescence, respectively, are red, yellow, and through hydrophobic and bound to the biofilm interactions. Different biofilms differ in electrostatic physicochemical properties, leading to variable binding abilities to lanthanide nanoparticles and generating characteristic luminescence signal patterns. Then, when the random forest algorithm was used to train the output



Fig. 1 Bacterial recognition through noncovalent interactions between bacteria and nanoparticles. A) A 3D sensing array constructed by using AuNP/AuNC nanocomposites for discriminating Gram-negative bacteria. Reproduced with permission.³³ Copyright 2019, Royal Society of Chemistry. B) Identification of antibiotic resistance in ESKAPE pathogens by plasmonic nanosensors through electrostatic and hydrophobic interactions. Reproduced with permission.³⁴ Copyright 2023, American Chemical Society. C) Identification of bacterial biofilm based on lanthanide nanoparticles. Reproduced with permission.³⁵ Copyright 2022, American Chemical Society.

signals, biofilm identification could be achieved within a few minutes, with an accuracy exceeding 80%.

2.1.2. Noncovalent recognition of bacteria by small luminescent molecules. Aggregation-induced emission luminogens (AIEgens) and vibration-induced emission luminogens (VIEgens) have been widely applied in recent years due to their promising and unique advantages. AIEgens exhibit significantly enhanced fluorescence in aggregated states, endowing them with advantages of weak background fluorescence and high signal to noise ratio. Chen et al. constructed a sensor array based on fluorescence properties (F-array), which uses a series of tetraphenylethylene derivative AIE probes for bacterial discrimination (Fig. 2A).³⁷ The Farray, built from five AIE probes with distinct electrostatic properties, can identify eight bacteria with different surface potentials. Cross-validation demonstrated a high predictive accuracy of 93.75%.

Tang's group proposed a competitive binding strategy based on AIEgen/graphene oxide (GO) complexes for the identification of different microbial lysates (Fig. 2B).³⁸ The sensor array consists of one negatively charged, five positively charged, and one neutral AIEgen combined with GO. The aggregation induced emission properties of the AIEgens together with the fluorescence quenching properties of GO reduced background fluorescence, thereby greatly enhancing the sensitivity of the sensor.

Another promising luminescent probe, VIEgens, was first reported by Tian and co-workers in 2015, who discovered this interesting phenomenon while studying the dynamic luminescence properties of dihydrodibenzo[a,c]phenazine derivatives.^{40,41} VIEgens emit blue fluorescence in the "saddle"-shaped conformation and emit red fluorescence in the planar conformation. Their conformation is sensitive to the surrounding microenvironment, making VIEgens promising probes for biosensing. Hu et al. proposed a sensor array constructed from eight VIE probes for phenotypic identification of methicillin-resistant Staphylococcus aureus (MRSA) (Fig. 2C).³⁹ Around the VIEgen core, these probes carry two quaternary ammonium salts across different substitution sites. The substituents determine the structure of VIEgens, leading to differences in interactions with negatively charged bacterial cell walls, which in turn determine the molecular conformation of the VIEgens. When VIEgens interact with bacteria, they change into a "saddle" conformation with an enhanced blue fluorescence. After the blue-to-red fluorescence intensity ratio of VIEgens generated by different bacteria was processed by the principal component analysis (PCA) algorithm, this sensor array was able to identify six phenotypes of Staphylococcus aureus, with analysis results that were consistent with those obtained from polymerase chain reaction analysis.

2.1.3. Noncovalent recognition of bacteria by conjugated polymers. Conjugated polymers with tunable structures that endow them with specific chemical modifications (such as fluorescent modification) can be used as sensing elements for sensor construction, facilitating their extensive application in biosensing.^{42,43}

Wang *et al.* developed a single-component multichannel array based on composites of graphene oxide (GO) and polyethyleneimine derivatives (PEI, Fig. 3A).²³ These PEI derivatives altered the surface physicochemical properties of GO, including charge, polarity, and hydrophobicity, resulting



Fig. 2 Bacterial recognition through noncovalent interaction between bacteria and small luminescent molecules. A) Fluorescent array based on AlEgens for bacterial identification. Reproduced with permission.³⁷ Copyright 2014, Wiley-VCH. B) Identification microbial lysate through competitive interactions by AlEgens and GO.³⁸ Copyright 2018, American Chemical Society. C) Ratiometric sensor array for phenotype identification of methicillin-resistant *Staphylococcus aureus* based on VIEgens. Reproduced with permission.³⁹ Copyright 2022, American Chemical Society.

in diverse binding strengths with bacteria. Through electrostatic and hydrophobic interactions with bacteria, the sensor array successfully identified 10 types of strains within minutes. Moreover, the sensor array could effectively recognize bacteria in complex biological samples (such as urine) (OD₆₀₀ = 0.25), achieving an accuracy of 94%.

Roy *et al.* have developed a sensor array capable of differentiating antibiotic-resistant bacteria and grouping bacteria with the same resistance phenotype, based on three cationic benzyl-functionalized polymers and three different fluorescent transducers to build four ratio channels (Fig. 3B).⁴⁴ By interacting with bacteria through electrostatic interactions, the sensor array sensitively recognized subtle differences on the bacterial surface induced by antibiotics within 30 min, generating unique responses and providing useful information for bacteria resistance.

Gut microbiota is closely associated with the development of diseases. Tomita *et al.* developed a sensor array capable of identifying gut microbiota (Fig. 3C). This sensor array was composed of twelve polyethylene glycol-*block*-poly-L-lysine (PEG-*b*-PLL) derivatives bearing AIE fluorophores.⁴⁵ These probes bind through various interactions with the bacterial surface, generating distinctive fluorescent fingerprint responses. By further processing the generated fluorescent responses with machine learning algorithms, this array sensor could categorize gut bacteria into strains, species, and phylums and distinguish the gut microbiota of obese and healthy individuals by detecting artificially simulated fecal samples.

order to improve the sensitivity of bacterial In Han's group developed a dual-channel recognition, fluorescence "turn-on" sensor array consisting of six electrostatic complexes formed by six negatively charged (PPE) and poly(*para*-aryleneethynylene) polymers six positively charged AIE fluorophores (Fig. 3D).⁴⁶ AIE quenches the fluorescence of PPE upon complexing with PPE, whereas bacteria could compete with PPE for binding with AIE and recover the fluorescence of PPE. Meanwhile, as AIE fluorophores aggregated on the bacterial surface, their emission also enhanced. This unique dual "turn-on" response, based on a six-element sensor array, assisted by various machine learning algorithms, enabled the identification of 20 bacteria at low concentrations (OD₆₀₀ = 0.001), achieving 96.7% accuracy in distinguishing urinary tract infection patients from healthy individuals.





Fig. 3 Bacterial recognition through noncovalent interaction between bacteria and polymers. A) One-component multichannel sensor array for rapid identification of bacteria based on polymers.²³ Copyright 2022, American Chemical Society. B) A polymer-based sensor for differentiating the resistant phenotypes of antibiotic-resistant bacteria. Reproduced with permission.⁴⁴ Copyright 2022, American Chemical Society. C) Fluorescent polymer-based chemical-nose systems for recognizing gut microbiota. Reproduced with permission.⁴⁵ Copyright 2022, Royal Society of Chemistry. D) A dual fluorescence turn-on sensor array formed by polymer and AlEgens for recognizing bacteria. Reproduced with permission.⁴⁶ Copyright 2024, Wiley-VCH.

2.1.4. Noncovalent recognition of bacteria by antimicrobial **peptides**. Antimicrobial peptides (AMPs) are short peptides consisting of 10–50 amino acid residues and are part of the biological immune system.^{47,48} Their natural cationic charge and hydrophobicity make them interact easily with negatively-charged bacteria. AMPs may also target specific biomolecules on the surface of bacteria, leading to antimicrobial activity.^{49–52} In 2017, Han *et al.* created a four-unit sensor array based on four electrostatic complexes formed by four positively charged AMPs and negatively charged *para*-phenyleneethynylene (PPE).⁵³ Both excited-state energy

transfer and charge transfer occur in the AMP/PPE complex, leading to fluorescence quenching of PPE. When bacteria were added, AMPs bind to the bacterial cell wall and permeate into the cell, resulting in PPE release and fluorescence recovery. The recognition efficiency of the AMP/PPE array was validated by detecting 14 bacteria (including 6 types of Gramnegative bacteria and 8 types of Gram-positive bacteria) with low-concentration (OD₆₀₀ = 0.01), and the successful identification of bacteria in urine was also achieved.

Similarly, Zhang's group recognized and identified bacteria based on the differences in recognition and

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disintegration by four short antimicrobial peptides (SAMPs) against different bacterial strains.⁵⁴ Processing of the collected unique fluorescence spectra of bacteria by LDA enabled the identification and differentiation of six types of bacteria within 30 min, with a classification accuracy of 100%.

2.2. Specific recognition – recognize biomarkers on bacterial surfaces

Bacterial surfaces contain a variety of biomarkers, such as carbohydrates, proteins, short peptides, *etc.* These biomarkers can serve as target molecules interacting specifically with recognition probes that are functionalized with receptor proteins, carbohydrates, or antibiotics, providing insights for the development of bacterial detection approaches. In this section, we summarize and discuss the identification approaches for bacteria based on various biomarker recognition mechanisms.

2.2.1. Recognition of the cis-diol group on bacterial surfaces. Boronic acid derivatives are widely used for the detection and differentiation of bacteria based on their covalent and reversible binding with cis-diols in carbohydrates or glycoproteins on bacterial surfaces.55-57 Under alkaline conditions, the tetragonal boronic acid anion (sp³) can react with *cis*-diol to form five or six-membered cyclic esters, while under acidic conditions, boronic acid in a trigonal configuration (sp²) limits its binding with *cis*-diol, resulting in the decomposition of the complex.^{56,58,59} As the content or boronic acid-binding ability of cis-diol on the bacteria surface varies among different bacteria, by transferring these differences into response differences of boronic acid-based probes, bacteria discrimination can be achieved.

Yan *et al.* proposed a colorimetric sensor array based on Wulff-type 4-mercaptophenylboronic acid (MPBA)-

mercaptoethylamine (MA) co-functionalized AgNPs (MPBA-MA@AgNPs) for bacterial detection and identification (Fig. 4A).⁶⁰ Wulff-type boronic acid is a unique adjacent amine boronic acid that can form cyclic ester bonds with cis-diol through intramolecular B-N bonds, allowing boronic acid and *cis*-diol to form cyclic esters under neutral conditions. Under neutral and alkaline conditions, MPBA binds to cis-diol on the bacterial surface, leading to the dispersion of AgNPs. Due to the presence of different carbohydrates on the bacterial surface, MPBA exhibits different affinities for the cis-diol in different carbohydrates, resulting in the color change of AgNPs. Under acidic conditions, the B-N bond breaks, and the positively charged MA binds to the negatively charged bacteria via electrostatic interactions, causing AgNPs to aggregate on the bacterial surface, and the color of AgNPs varies due to differences in bacterial surface potentials. Therefore, the MPBA-MA@AgNPs show variable binding ability to bacteria at different pH buffer solutions, and generate distinct color responses. By using LDA for data processing, 17 strains were successfully differentiated. This colorimetric method, based on specific affinity and electrostatic interactions, holds promise for the identification of bacteria in clinical urine and serum specimens; however, the color changes of various bacteria in this method are slight, which suggests that relying on colorimetry alone to differentiate bacteria is not reliable. Additionally, our group proposed a sensor array fabricated by using boronic acid modified g-C₃N₄ (BA-g-CN) nanosheets.61 Under acidic conditions, the positively charged BA-g-CN nanosheets interact with the negatively charged bacteria through electrostatic interaction. Under neutral or alkaline conditions, BA-g-CN nanosheets bond to the bacteria by forming covalent bonds through boronic acid and carbohydrates on the bacterial cell wall. These interactions lead to bacterial sedimentation and



Fig. 4 Bacterial recognition through the interaction between boronic acid and the *cis*-diol group. A) A colorimetric sensor array based on Wulff-type boronate functionalized AgNPs for bacterial identification. Reproduced with permission.⁶⁰ Copyright 2019, American Chemical Society. B) A dual-mode sensor array based on colorimetric and photothermal detection for bacterial identification. Reproduced with permission.⁶² Copyright 2023, American Chemical Society.

fluorescence reduction in the supernatant. By using acidic, neutral, and alkaline conditions as three different sensing channels, the identification of nine strains can be achieved. This sensing strategy focuses on distinguishing Gramnegative bacteria, particularly four strains of Escherichia coli, ensuring timely and targeted treatment of Gram-negative bacteria. Recently, Wang et al. constructed a dual-mode bacterial sensor array based on colorimetric and photothermal detection.⁶² This sensor uses boronic acidfunctionalized Au-Fe₃O₄ nanoparticles (BA-GMNPs) as probes, which not only exhibited localized surface plasmon resonance properties but also demonstrated superparamagnetism. BA-GMNPs formed B-N bonds with cis-diol on the bacterial surface, dispersing on the bacterial surface and weakening the aggregation of BA-GMNPs induced by NaCl. The content of cis-diol on the surface of different bacteria varied, resulting in different quantities of BA-GMNPs bound to bacteria and different amounts of BA-GMNPs aggregated in the supernatant. After magnetic separation, the RGB values and UV-vis spectra of the aggregated BA-GMNPs in the supernatant were measured; the precipitate was irradiated with an 808 nm laser, and temperature changes were monitored using a thermal imager. With the assistance of LDA, six bacteria were successfully distinguished, and even bacteria in real samples, such as milk, were successfully identified, with a recognition accuracy of 93.3%.

2.2.2. Recognition of lectins on bacterial surfaces. Lectins are a type of protein on the bacterial surface that bind specifically to carbohydrates (monosaccharides and

oligosaccharides), playing a crucial role in adhesion during the initial stage of bacterial infection.⁶³⁻⁶⁵ For instance, FimH, a protein at the end of the fimbria of Escherichia coli, can specifically bind to mannose.^{66,67} Otten et al. investigated five bacteria, including E. coli K12, E. coli Top 10, M. marinum, P. putida, and M. smegmatis, demonstrating a platform for profiling bacteria based on the relative binding affinity of bacteria to nine polysaccharide surfaces (Fig. 5A).⁶⁸ Unique barcodes for each bacterium were created based on their relative binding affinities, resulting in five independent clusters after analysis by LDA. Zhang's group proposed a bacterial detection and identification strategy constructed from Cu:CdS quantum dots modified with glucose, stachyose, and raffinose (Fig. 5B).69 The carbohydrate binding strength between glycosylated quantum dots and the bacterial cell wall determines the amount of quantum dots bound to the bacteria, and the carbohydrate varies in the bacterial cell wall, generating a unique fluorescent fingerprint response, allowing six types of bacteria to be identified within 30 min.

2.2.3. Recognition of p-**ala**-p-**ala on bacterial surfaces.** Vancomycin inhibits bacterial cell wall synthesis by binding to p-alanyl-p-alanine (p-Ala-p-Ala) at the terminus of peptidoglycan, exerting bactericidal effects.^{70,71} The cell wall of Gram-positive bacteria is mainly composed of peptidoglycan, while Gram-negative bacteria have an additional outer membrane composed of lipopolysaccharides outside of peptidoglycan.⁷² The dense lipopolysaccharide layer in Gram-negative bacterial cell walls prevents molecules larger than 500 kDa from passing through, making



Fig. 5 Bacterial recognition through the interaction between lectins and glycan. A) Discrimination of bacteria by ratiometric analysis of their carbohydrate binding profile. Reproduced with permission.⁶⁸ Copyright 2016, Royal Society of Chemistry. B) A sensor array consisting of multivalent glycosylated Cu:CdS quantum dots for rapid bacterial discrimination. Reproduced with permission.⁶⁹ Copyright 2017, Elsevier.



Fig. 6 Bacterial recognition through the interaction between vancomycin and D-ala-D-ala. A) Identification of bacteria by a fluorescent sensor array based on boronic acid, vancomycin, and polymyxin functionalized carbon dots. Reproduced with permission.⁷⁵ Copyright 2019, Elsevier. B) A colorimetric sensor array for rapid diagnosis of urinary tract infection by a multi-recognition mechanism, which includes the recognition between vancomycin and D-ala-D-ala. Reproduced with permission.⁷⁶ Copyright 2024, American Chemical Society.

vancomycin specific to Gram-positive bacteria.72-74 Zhang's group constructed a fluorescent sensor array for differentiating various bacteria using carbon dots functionalized with vancomycin (Van), polymyxin (PM), and boronic acid, respectively (BA, Fig. 6A).75 Van specifically recognizes D-Ala-D-Ala on the Gram-positive bacterial surface, PM recognizes phospholipid on the Gram-negative bacteria, and BA forms B-N bonds with the cis-diol on the bacterial surface. Probes modified with Van, PM, and BA can bind to all bacteria with different affinities, and fingerprints specific to each bacterium were obtained from the fluorescent responses of the three probes. The obtained fluorescent responses were further analyzed by LDA, achieving the differentiation of six types of bacteria with an accuracy of 91.6%. Recently, Yang et al. proposed a colorimetric sensor array for bacterial differentiation based on vancomycin, *D*-alanine (*D*-Ala), boronic acid, and cetyltrimethylammonium bromide ligand-(CTAB) functionalized Fe single-atom nanozymes (SANs, Fig. 6B).⁷⁶ In their design, vancomycin is used for the recognition of Gram-positive bacteria; D-Ala is incorporated into

peptidoglycan mediated by enzymes to discriminate bacteria from fungi; boronic acid provides information regarding the *cis*-diols on the surface of microorganisms; and the positively charged CTAB binds to microorganisms by electrostatic interaction. With the combination of noncovalent interactions, specific molecular recognition, and enzyme-mediated metabolism, this sensing method could complete detection within 1 h with an accuracy of up to 97% for identifying more than 10 types of microorganisms in clinical samples with the assistance of the support vector machine (SVM) algorithm.

2.2.4. Recognition of the receptor of protein on bacterial surfaces. Some bacteria-unique components can serve as recognition targets.⁷⁷ As their level varies among different types of bacteria, by constructing a series of probes that separately respond to these recognition targets, sensor arrays can be fabricated for bacteria identification. For example, Ji *et al.* proposed a fluorescent sensor array constructed with gold nanoclusters (AuNCs) modified with human serum proteins (HSA), lactoferrin (Lf), lysozyme (Lys), and vancomycin (Fig. 7A).⁷⁸ The peptide motifs on human serum



fluorescence sensor array for discrimination of bacteria. Reproduced with permission.⁷⁸ Copyright 2018, Wiley-VCH. B) Multichannel sensor array based on muti-functionalized AuNCs for gut microbiota detection. Reproduced with permission.⁸³ Copyright 2023, American Chemical Society.

proteins (HSA) can interact with bacterial cell walls;^{78,79} lactoferrin (Lf) can recognize lactoferrin receptors expressed by bacteria;^{80,81} peptidoglycans of bacterial cell walls are a natural substrate for lysozyme (Lys);^{79,82} vancomycin (Van) can bind with the p-ala-p-ala moiety through hydrogenbonding. By taking advantage of these recognition pairs, six types of bacteria, including kanamycin-resistant *E. coli* and methicillin-resistant *Staphylococcus aureus*, were readily differentiated.⁷⁸

The composition and activity of the gut microbiota are crucial for health management and disease treatment. Liu et al. developed a rapid sensor array for distinguishing gut bacteria (Fig. 7B).⁸³ Gold nanoclusters (AuNCs) were modified with Van, bacitracin, and lysozyme as signal-reporting probes that specifically identified D-ala-D-ala moieties, pyrophosphate groups, and peptidoglycans, respectively. Ti₃C₂ transitionmetal carbides (MXenes) were used as fluorescence quenchers. When three recognition probes were mixed with Ti₃C₂ MXenes, they were tightly bound through hydrogen bonding, leading to energy transfer and quenching the fluorescence of AuNCs. Both bacteria and three functionalized AuNCs could bind with Ti3C2 MXenes but the binding strengths vary. After adding bacteria to Ti₃C₂ MXenes-AuNCs composites, they either restore the fluorescence of AuNCs by competing with AuNCs for binding with Ti₃C₂ MXenes or lead to further fluorescence quenching. Due to the association between gut bacteria and colorectal disease, this strategy coupled with pattern recognition algorithms accurately differentiated colorectal cancer patients from healthy individuals within 30 min, providing a robust and simple platform for the clinical diagnosis of colorectal cancer.

3. Identification mediated by bacterial intrinsic properties

Bacterial metabolism includes catabolism and anabolism, which are the most fundamental processes in bacterial life activity. Enzymatic catalysis is required for bacterial metabolic processes, with different bacteria possessing distinct enzyme systems, leading to diverse metabolic behaviors among different bacteria.^{84–88}

3.1. Bacteria identification by metabolites

Variations exist among bacteria in the enzymes involved in the decomposition of nutrients such as sugars and proteins, resulting in diverse decomposition capabilities, pathways, and products. Methods for the identification of bacteria have been widely proposed based on metabolites among different bacteria.^{87,88} In this section, we will discuss the sensor arrays constructed based on the recognition of various metabolites of bacteria.

3.1.1. Detection of H^+ generated by bacteria metabolic processes. Bacteria have the ability to break down polysaccharides into monosaccharides, which are then converted into pyruvic acid. The latter can be further broken down into carbon dioxide and hydrogen if the bacteria



Fig. 8 Bacterial recognition through metabolites produced by bacterial metabolism. A) The glucose-metabolism-triggered colorimetric sensor array based on MER, BTB, and BCG for differentiation and antibiotic susceptibility testing of bacteria. Reproduced with permission.⁸⁹ Copyright 2023, Elsevier. B) Agar gel-based colorimetric sensor array for identification of bacteria in household refrigerators. Reproduced with permission.⁹⁰ Copyright 2023, Royal Society of Chemistry. MER: methyl red. BTB: bromothymol blue. BCG: bromocresol green.

contain pyruvate decarboxylase, resulting in different sugar decomposition products among bacteria.^{84,87}

Liu's group developed a colorimetric sensor array for microbial discrimination and antimicrobial susceptibility testing, using three pH indicators, including methyl red (MER), bromothymol blue (BTB), and bromocresol green (BCG), as H⁺-sensing probes (Fig. 8A).⁸⁹ The bacterial metabolism of glucose generates acids, including acetates, lactates, and other acidic byproducts, causing changes in the acidity of the bacterial culture microenvironment. Due to variations in the ability of different bacteria to metabolize glucose, the acids produced by different bacteria also vary in terms of both types and concentrations, resulting in significant differences in the color change of the three pH indicators. Combined with the LDA algorithm, this sensor array facilitated the identification of 11 types of bacteria.

3.1.2. Detection of volatile organic compounds generated by bacteria metabolic processes. In addition to acids, bacterial metabolism also generates various volatile organic compounds (VOCs), primarily including carboxylic acids, alcohols, aldehydes, esters, hydrocarbons, and organic sulfur derivatives.⁹¹⁻⁹³ Some bacteria may produce the same VOCs but at various concentrations. For instance, the concentration of formaldehyde and acetaldehyde produced by Gramnegative bacteria is generally higher than that by Grampositive bacteria. Additionally, some VOCs could only be produced during specific metabolic pathways.^{94–96} The rapid differentiation of bacteria can therefore be achieved by detecting bacteria-produced VOCs, as demonstrated by some studies. Xu et al. qualitatively analyzed VOCs released by bacteria that are associated with ventilator-associated pneumonia (VAP) by proton transfer reaction-mass spectrometry (PTR-MS).97 Principal component analysis (PCA) and analysis of similarities (ANOSIM) were employed to analyze the VOCs of bacteria collected at different time points. After 3 hours of in vitro cultivation, significant

differences in VOCs among different bacteria were observed. Six types of common bacteria associated with VAP were effectively distinguished, offering potential assistance to physicians in promptly formulating treatment plans. Mass spectrometers have many benefits in detecting VOCs, but they are sophisticated and expensive. For rapid bacterial detection in daily life, economically portable bacterial identification strategies are necessary. Chen's group developed a colorimetric sensor array based on a multilayer neural network (CSA-NN) for identifying common pathogenic bacteria in household refrigerators. The colorimetric sensor array consists of 16 dye-containing agar gel spots, exhibiting good resistance to moisture and frost (Fig. 8B).⁹⁰ The dyecontaining agar gel array generated colorimetric responses through reactions with VOCs produced by bacteria. After

training with CSA-NN, this sensor array successfully differentiated *Escherichia coli*, *Staphylococcus aureus*, *Salmonella*, and *Listeria monocytogenes* in refrigerated environments. It also accurately identified *Staphylococcus aureus* and *Listeria monocytogenes* on eggshells with an accuracy of 83%.

Real-time monitoring and identification of bacteria are highly elusive goals in the field of bacterial detection. Shaulof *et al.* introduced a capacitive artificial nose for the VOCidentification-based bacteria discrimination method using interdigitated electrodes (IDEs) modified with carbon dots (Cdots) exhibiting different polarities (C-dot-IDE).⁹⁸ VOCs metabolized by bacteria induce rapid changes in the capacitance of the C-dot-IDE, which is intimately related to the polarity of gas molecules. With the application of machine



Fig. 9 Bacterial recognition through nonvolatile compounds produced by bacterial metabolism. A) Gold nanoparticle chemiresistor capable of identifying bacteria based on metabolites released by bacteria. Reproduced with permission.¹⁰⁰ Copyright 2015, Elsevier. B) A fluorescent nanosensor based on SWCNTs for clinically essential bacteria discrimination. Reproduced with permission.¹⁰¹ Copyright 2020, Nature. SWCNTs: single-walled carbon nanotubes.

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learning algorithms, the C-dot-IDE gas sensor exhibits excellent recognition performance, enabling continuous monitoring of bacteria proliferation and differentiation.

3.1.3. Detection of nonvolatile compounds generated by bacteria metabolic processes. The detection of VOCs is prone to be affected by humidity and most of them require sample collection. On the contrary, direct detection of nonvolatile compounds in solutions does not need careful humidity control and complex sample collection techniques. In recent years, some sensor arrays based on detecting nonvolatile metabolites produced by bacterial metabolism have been developed.⁹⁹ For example, Webster *et al.* constructed an AuNP chemiresistor that monitored changes in the resistance of AuNPs in the liquid culture medium (Fig. 9A).¹⁰⁰ In this way, 4 types of bacteria, including *Escherichia coli, Bacillus subtilis, Staphylococcus epidermidis*, and *Pseudomonas aeruginosa* were successfully differentiated based on their metabolic fingerprints.

Nonvolatile metabolites are diverse among bacteria. For instance, endotoxin is a lipopolysaccharide released upon the lysis of Gram-negative bacteria, which is a unique metabolite of Gram-negative bacteria. Siderophores are a class of low molecular weight compounds synthesized by microorganisms under conditions of low iron concentration. They exhibit a specific high affinity for Fe³⁺ and are a common pathway for bacteria to acquire iron from the environment. Nißler et al. developed a set of near-infrared fluorescent nanosensors for remote identification of clinical bacteria (Fig. 9B).¹⁰¹ Nine sensing units, consisting of functionalized single-walled carbon nanotubes (SWCNTs) specific for lipopolysaccharide, siderophores, deoxyribonucleases, and proteases, along with four generic low-selective SWCNTs and one stable nearinfrared fluorescent reference, were integrated into a system biocompatible hydrogel. This successfully differentiated 6 common bacteria in the clinic aided by PCA. Multiplexing was achieved by spectral encoding (900 nm, 1000 nm, 1250 nm), which was capable of differentiating Pseudomonas aeruginosa and Staphylococcus aureus. The nearinfrared fluorescence also endowed this approach with the ability of tissue penetration (>5 mm).

3.2. Identification of bacteria by metabolic labeling

Metabolic labeling is an effective method to study the metabolic processes of living systems, allowing efficient and



Fig. 10 Bacterial recognition through metabolic labeling. A) The RGB-emitting probes for the detection of bacteria by labeling peptidoglycan. Reproduced with permission.¹⁰⁶ Copyright 2020, Royal Society of Chemistry. B) A fluorescent sensor array for identifying bacteria based on metabolic labeling at various acidities and times. Reproduced with permission.¹⁰⁷ Copyright 2022, Elsevier. C) Metabolism-triggered colorimetric sensor array constructed from three types of D-AA functionalized AuNPs for bacterial identification. Reproduced with permission.¹⁰⁸ Copyright 2022, American Chemical Society. D) Identification of antibiotic-resistant bacteria based on metabolic labeling and click reaction. Reproduced with permission.¹⁰⁹ Copyright 2023, American Chemical Society.

non-destructive incorporation of functionalized metabolites biomolecules through into target biosynthetic mechanisms.^{102,103} Peptidoglycan, which is widely present in bacterial cell walls, is a multilayered network structure composed of glycans and short peptides. Functionalized monosaccharides and amino acids can be incorporated into peptidoglycan by metabolic labeling.^{104,105} A sensing strategy combining metabolic labeling with fluorophores was proposed by Hong et al., with bacteria metabolizing three different fluorescent metabolic derivatives to identify different bacterial species or microbiota types (Fig. 10A).¹⁰⁶ Due to disparities in metabolism ability, unique colorimetric fingerprints for different bacterial species and microbiota after metabolic labeling were obtained. By utilizing a portable spectrometer combined with a smartphone for direct reading of fluorescent RGB values, discrimination of different bacterial species and microbiota types was achieved.

D-AAs are persistent almost exclusively in bacterial cell walls, endowing them with unique advantages in bacteria labeling. Unnatural D-AAs can be incorporated into bacterial cell walls by swapping amino acids at the 5th or 4th position of the pentapeptide, mediated by D,D-transpeptidase or L,D-transpeptidase, respectively. Recently, our group proposed a chemical nose approach based on metabolic labeling and click reactions (Fig. 10B).¹⁰⁷ Bacteria first metabolized clickable handle tagged D-AAs at two acidities (pH = 5 and 7) for 20 min and 60 min, respectively, followed by clicking with up-conversion nanoparticles (UCNPs), generating four-dimensional signal outputs. Multidimensional fluorescent response analysis by machine learning successfully classified eight types of bacteria in the training set on strain, genus, and Gram phenotype. The differences in signal response under four incubation conditions reflected variations in the quantity and activity of enzymes involved in metabolic labeling, endowing this strategy with good anti-interference capabilities. This approach can accurately classify unknown samples from the validation set into clusters independent of the training set and can accurately identify their Gram properties.

Culture-based bacterial detection and antibiotic susceptibility testing (AST) are the gold standard methods for bacterial detection, but they are time-consuming. In such cases, clinicians often resort to broad-spectrum antibiotics for treatment at an early stage, which worsens the issue of antimicrobial resistance. Therefore, the development of accurate and rapid methods for bacterial differentiation and AST is urgently needed. Liu's group constructed a colorimetric sensor array for bacterial identification using D-ala, D-2,3-diaminopropionic acid, and D-glutamate-functionalized gold nanoparticles (AuNPs) as probes (Fig. 10C).¹⁰⁸ Bacteria were competitively metabolized with exogenous D-AAs on the surface of AuNPs through the mediation of D,D-transpeptidase, causing the AuNPs to become unstable and aggregate. AuNPs exhibit distinct aggregation phenomena due to the different metabolic capabilities of bacteria towards various D-AAs. By analyzing the unique responses by LDA, eight bacterial species, including an antibiotic-resistant strain, were successfully differentiated. Furthermore, by monitoring the bacterial metabolic activity of D-AAs under different antibiotic treatments, a colorimetric-based rapid AST method was developed, and the whole procedure was completed in 24 h, promoting the clinical practicability of this approach.

For applying array sensor approaches in the clinic, it should be sensitive enough to detect low-concentration bacteria in real clinical samples. Our group constructed a sensor array based on metabolic labeling and the "antibiotic responsive spectrum" for distinguishing antibiotic-sensitive and resistant strains at bacteria concentrations as low as 10⁵ CFU mL⁻¹ (Fig. 10D).¹⁰⁹ This strategy incorporates p-ala with azido groups into bacterial cell walls through bacterial metabolism, followed by clicking with dibenzocyclooctynefunctionalized UCNPs (DBCO-UCNPs). By proceeding with metabolic labeling in the presence of six antibiotics, an "antibiotic responsive spectrum" was obtained as a result of a unique fluorescent fingerprint corresponding to a certain type of bacterium. By combining with various machine learning algorithms, eight bacterial strains were effectively clustered based on species, Gram phenotype, and antibiotic resistance. This metabolic labeling strategy exhibits excellent anti-interference capabilities, accurately identifying bacteria (concentration: 10⁵ CFU mL⁻¹) in artificial urinary tract infection samples and determining their resistance.

3.3. Identification of bacteria by biosynthesis

Bacteria, as efficient biofactories, can utilize various enzymes, proteins, peptides, and electron transfer pathways in vivo to regulate inorganic ions and reduce them into inorganic nanoparticles of different shapes and sizes. Up till now, researchers have employed bacteria to synthesize over 100 types of inorganic nanoparticles.^{110,111} Enzymes are the primary biomolecules responsible for the reduction of inorganic ions. Different microorganisms produce various reductases, including nitrate and nitrite reductases, nicotinamide adenine dinucleotide phosphate-dependent reductases, and sulfate (SO_4^{2-}) and sulfite (SO_3^{2-}) reductases, which co-regulate microbial biosynthesis.112-115 Xianyu's group constructed a novel sensor array based on differences in the properties of microbially biosynthesized AuNPs, such as particle size, surface plasmon resonance (SPR) spectra, and surface potential (Fig. 11).¹¹⁶ By employing PCA, LDA, and random forest (RF) algorithms to explore the relationships between features of microbial-synthesized AuNPs and microbial types, classification models were established. Microorganisms were classified at the levels of kingdom, order, genus, and species. This microbial discrimination result was in accordance with the results obtained by traditional classification approaches.

4. Statistical analysis and machine learning

Machine learning algorithms can rank the significance of input variables, identify trends in the data, and classify the



data, enhancing its visualization and readability.¹¹⁷ Since chemical nose strategies are typically constructed using multiple sensor units, resulting in large volumes of high-dimensional data with poor readability, combining machine learning with sensor arrays can extract more valuable information.^{118,119} In this section, we categorize the machine learning algorithms used for statistical analysis into two types: clustering methods and classification methods.

4.1. Clustering methods

Clustering algorithms, including HCA, PCA, and t-distributed stochastic neighbor embedding clustering (t-SNE), are unsupervised algorithms capable of clustering data. In highdimensional space, HCA typically uses Ward's minimum variance as the clustering criterion and Euclidean distance as the distance metric to cluster data.^{120,121} The HCA dendrogram intuitively expresses the similarity between samples and can effectively convey relationships between similar species. However, since the HCA dendrogram has only one quantitative axis (distance) and the orientation of the dendrogram does not affect data clustering, it may have misinterpretation issues due to changes in the relative position of the linkage axes.

PCA is a commonly used dimensionality reduction algorithm in sensor arrays.⁷⁶ It transforms linear combinations of the initial variables into a set of orthogonal combinations, projecting data from sensor arrays into 2D or 3D space to represent the feature relationships of data.^{120–122} PCA projects the original data onto a new coordinate system, where the first coordinate axis (the first component) is

determined by finding the maximum variance of the dataset along this vector. Subsequent components are orthogonal to previous ones and follow the principle of maximum variance. The score plots after dimensional reduction by PCA are more visualizable than the original data. However, biological data tends to be nonlinear, and PCA, being a linear dimensionality reduction method, fails to capture these nonlinear relationships, potentially leading to information loss. t-SNE, a nonlinear dimension reduction algorithm, is more suitable for analyzing high-dimensional biological information.¹²³ t-SNE creates a probability distribution for each data point by calculating Euclidean distances between data points, with closer points representing higher similarity. It then creates a similarity matrix based on data similarities and can preserve both local and global structures of the data, which is beneficial for discovering more relationships within biological information. All the above algorithms are unsupervised and cannot predict unknown samples. When new data is added to the dataset, the entire clustering must be recalculated.⁷⁶

4.2. Classification methods

The classification method, which usually belongs to supervised algorithms, enables the prediction of unknown samples. In sensor arrays for microbial identification, the classification method utilizes a dataset labeled with microbial species as the training set. It discovers the relationship between microbial species and corresponding data through the training set and establishes a mathematical model.^{15,76} By inputting the data of unknown samples into

In recent years, LDA has been widely applied in sensor arrays. It performs orthogonal transformations on the potentially linear dependence original variables to nonlinear dependence variables, similar to the PCA algorithm. Unlike PCA, LDA is a supervised algorithm that can predict unknown samples by establishing a mathematical model.¹²⁴ It follows the classification principle of minimizing intraclass variance and maximizing interclass variance, meaning that it aims to make projections of data with the same label as close as possible while keeping projections of data with different labels as far apart as possible, in an attempt to find the dimension with the best separation. The LDA score plot can intuitively evaluate the effectiveness of classification and discrimination. However, LDA is not suitable for classification with high-dimensional data but few classes. SVM is an optimization-based algorithm aimed at finding the optimal hyperplane for classifying data. When dealing with linearly separable models, SVM achieves optimal classification by maximizing the margin between classes. For linearly inseparable models, it maps data to a highdimensional space to achieve linear separation.125,126 SVM performs exceptionally well with data of high-dimensional and small sample populations; however, as sample populations increase, its processing efficiency decreases. The evolution of deep learning algorithms is promoted by the rapid development of hardware and software technology of computers. CNN is a type of supervised deep learning algorithm that extracts features from data through convolution and pooling, mapping the input data into a high-dimensional space, and then performing classification or regression of the features through dense layers.¹²⁷ CNNs are characterized by strong feature extraction capability and high robustness, and they have been widely used in the field of biosensing.

5. Outlook and perspectives

Machine learning-assisted sensor arrays are an effective strategy for identifying bacteria, enabling rapid qualitative analysis of various bacteria, which plays an important role in detecting bacteria and inhibiting their spread in real life. We compared the reported sensor arrays in terms of recognition principle, types of interactions between bacteria and the sensor, probe materials, types of pathogens that can be identified, channel numbers employed, reaction time, minimum bacteria concentrations that can be detected (not the limit of detection), and signal readout of the sensor arrays, with details listed in Table 1.

Array-based sensing strategies have achieved certain success in bacterial discrimination, but these methods are still in their infancy and need to be further improved to meet the requirements of practical applications. The sample matrices in real bacterial-containing samples are complex, unknown, and with low concentrations of bacteria. However, current sensing strategies are mostly based on known bacterial strains cultivated in the laboratory, which may fail in practical applications due to the following limitations. First of all, as the surface charge and hydrophobicity vary among bacteria due to environmental changes, noncovalent interactions are easily affected by environmental disturbances, leading to low interference tolerance; the expression of proteins, sugars, and other biomolecules on the bacterial surface may differ as bacteria growth environments vary. Therefore, it is difficult to detect complex real samples solely by recognizing the external properties of bacteria. The sensor arrays based on the detection of the products of bacterial enzymatic catalysis processes typically require sample pretreatment to minimize interference from the sample matrices. Therefore, it is necessary to develop strategies for detecting bacteria based on multi-recognition. Secondly, sensor arrays are constructed based on a library of laboratory strains and can only identify bacteria within the library, which may pose limitations in detecting unknown bacteria in real samples. Therefore, it is necessary to develop a sensing strategy to detect a large number of clinical samples to construct a training set based on actual samples, making it possible to apply it to clinical samples. On the other hand, it is necessary to develop sensor arrays that are capable of clustering bacteria by genus, species, Gram properties, and antibiotic sensitivity, enabling rapid preclassification of bacteria outside the database for targeted detection in the next step. Thirdly, currently developed arrays are greatly influenced by bacteria sensor concentration and struggle to achieve fingerprint For sensors that differentiate bacteria recognition. populations in samples, signal intensity is also related to bacterial concentration. Therefore, appropriate preprocessing of the signal to eliminate interference from concentration is necessary. Additionally, bacterial heterogeneity, especially the heterogeneity of antibiotic-resistant bacteria caused during antibiotic treatment, is a key focus for future monitoring. Fourthly, current data analysis of chemical nose strategies for bacterial identification primarily relies on linear dimensionality reduction methods, such as PCA, which struggle to capture the complex structure of highdimensional data. However, in deep learning, CNN can better process data distribution in high-dimensional spaces through nonlinear dimensionality reduction. Therefore, it is essential to continuously optimize mathematical classification models and develop more robust machine learning algorithms to achieve more reliable identification results.

In summary, to enable the clinical translation of chemical nose strategies, sensor arrays capable of discriminating bacteria at the single bacterium level are helpful to avoid the concentration-dependence issue of the current sensor arrays. Secondly, sufficient strains isolated from clinical samples should be incorporated into the

Table 1 Summary of sensor arrays aided by machine learning for bacteria identification

Recognition mechanism or substance	Type of identification	Probes	Types of pathogens identified	of channels	Reaction time	Bacteria concentration	Signal readout	Ref.
Electrostatic and hydrophobic interactions	Noncovalent interaction	AuNP/AuNCs	10 (3 antibiotic-resistant strains)	3	10 min	OD ₆₀₀ = 0.015	Fluorescence, UV-vis absorption and	33
Electrostatic and hydrophobic interactions	Noncovalent interaction	AuNPs	12 (6 antibiotic-resistant strains)	6	20 min		light-scattering Surface plasmon resonance spectra	34
Electrostatic and hydrophobic interactions	Noncovalent interaction	Lanthanide NPs	5	3	A few minutes		Fluorescence	35
Electrostatic interaction	Noncovalent interaction	AIEgens	8 (2 antibiotic-resistant strains)	5	30 min	$OD_{600} = 0.002$	Fluorescence	37
Electrostatic interaction	Noncovalent interaction	AIEgen/GO	6	7	2 h	$OD_{600} = 1$	Fluorescence	38
Electrostatic interaction	Noncovalent interaction	VIEgens	10 (2 antibiotic-resistant strains)	8	60 min	$10^4 ext{ CFU} ext{mL}^{-1}$	Fluorescence	39
Electrostatic and hydrophobic interactions	Noncovalent interaction	Conjugated polymers and GO	10	3	Within minutes	OD ₆₀₀ = 0.25	Fluorescence	23
Electrostatic interaction	Noncovalent interaction	Conjugated polymers and fluorescent transducers	4	4	30 min	OD ₆₀₀ = 0.25	Fluorescence	44
Electrostatic, hydrophobic, π - π , and hydrogen bonding interactions	Noncovalent interaction	Conjugated polymers	16	12×4	<30 min	OD ₆₀₀ = 0.04	Fluorescence	45
Electrostatic interaction	Noncovalent interaction	Conjugated polymers and	20	6	30 s	OD ₆₀₀ = 0.001	Fluorescence	46
Electrostatic and hydrophobic interactions (according to reference)	Noncovalent interaction	AMPs and conjugated	14	4	30 min	OD ₆₀₀ = 0.01	Fluorescence	53
AMPs – lipid and membrane (according to reference)	Specific recognition	Short antimicrobial	6	4	30 min	$\begin{array}{l} 1\times 10^8 \text{ CFU} \\ \text{mL}^{-1} \end{array}$	Fluorescence	54
Mercaptophenylboronic acid- <i>cis</i> -diol interactions and electrostatic interactions	Covalent and noncovalent interactions	AgNPs	17	8	15 min	OD ₆₀₀ = 0.05	UV-vis absorption	60
BA- <i>cis</i> -diol interactions	Covalent interaction	Au-Fe ₃ O ₄	6	5	20 min	OD ₆₀₀ = 0.1	UV-vis absorption and	62
Glycan-lectin interactions	Specific	Well plate	9	5	1.5 h	OD ₆₀₀ = 1	Fluorescence	68
Glycan-lectin interactions	Specific	Cu:CdS quantum dots	3	6	30 min	$OD_{600} = 0.3$	Fluorescence	69
Van–D-ala-D-ala, polymyxin–phospholipid and BA– <i>cis</i> -diol interactions	Covalent interaction and specific recognition	Carbon dots	6	3	60 min	OD ₆₀₀ = 1	Fluorescence	75
Van–D-ala-D-ala, BA– <i>cis</i> -diol and electrostatic interaction, and metabolic labeling	Covalent and noncovalent interactions	Fe single-atom nanozymes	10	4	30 min	10^4 CFU mL ⁻¹	UV-vis absorption	76
HSA-cell walls, Lf-lactoferrin receptors, Lys-peptidoglycans, and Van-p-ala- p-ala	Specific recognition	AuNCs	6 (2 antibiotic-resistant strains)	4	60 min	OD ₆₀₀ = 1	Fluorescence	78

interactions

Table 1 (continued)

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Recognition mechanism or substance	Type of identification	Probes	Types of pathogens identified	Number of channels	Reaction time	Bacteria concentration	Signal readout	Ref.			
Van–D-ala-D-ala, bacitracin–pyrophosphate groups, and Lys – peptidoglycans	Covalent interaction and specific recognition	AuNCs and Ti_3C_2 MXenes	5	3	30 min	OD ₆₀₀ = 0.2	Fluorescence	83			
Metabolites (H ⁺)	Chemical reaction	Acid–base indicators	11 (2 antibiotic-resistant strains)	3	6 h	$10^7 \text{ CFU} \text{mL}^{-1}$	UV–vis absorption	89			
Metabolites (VOCs)	Chemical reaction	_	6	12	3 h	OD ₆₀₀ = 1	PTR-MS and FGC-PTR-MS	97			
Metabolites (VOCs)	Chemical reaction	Dye-containing agar gel	4	16 × 3	12 h	$10^3 ext{ CFU} ext{mL}^{-1}$	—	90			
Metabolites (VOCs)	Chemical reaction	C-Dot-IDE	4	3	1.3 s, continuous monitoring	$OD_{600} = 0.5$	Capacitance	98			
LPS-binding peptide – LPS, hemin [Fe ³⁺]– siderophores, <i>etc.</i>	Specific and nonspecific recognitions	SWCNTs	6	9	Incubation after 1 h, acquired in 30 s intervals	_	Fluorescence	101			
Metabolic labeling	Covalent interaction	Small molecules	3	3	12 h	$OD_{600} = 0.4$	Fluorescence	106			
Metabolic labeling and click reaction	Covalent interaction	UCNPs	8 (2 antibiotic-resistant strains)	4	100 min	$10^6 \text{ CFU} \text{mL}^{-1}$	Fluorescence	107			
Metabolic labeling and hydrophobic interactions	Covalent interaction (metabolism) and noncovalent interaction (AuNPs aggregation)	AuNPs	8 (2 antibiotic-resistant strains)	3	6 h	OD ₆₀₀ = 0.1	UV-vis absorption	108			
Metabolic labeling and click reaction	Covalent interaction	UCNPs	8 (3 antibiotic-resistant strains)	7	3 h	10^5 CFU mL ⁻¹	Fluorescence	109			
Biosynthesis		AuNPs	17 (1 antibiotic-resistant strains)	3	12 h	OD ₆₀₀ = 0.5	UV–vis absorption, zeta potential, and particle size	116			
Note: Ref. represents reference.											

training set to increase the accuracy of the responses to real clinical samples. Finally, machine learning algorithms suitable for biological samples should be selected and optimized for eliminating signal fluctuations caused by possible interferences and discovering more correlations among strains, thus assisting in the qualitative analysis of bacteria outside the training set. In addition to these three aspects, to achieve the widespread application of chemical nose strategies, efforts should also focus on simplifying the preprocessing procedures and constructing portable, precise, and low-noise instrumentation.

Data availability

No data, models, or code were generated or used in this review paper.

Author contributions

Xin Wang: investigation, writing – original draft, review & editing. Ting Yang: writing – review & editing, funding acquisition, resources, project administration, supervision. Jian-Hua Wang: funding acquisition, resources, supervision.

Conflicts of interest

The authors have no conflicts of interest to declare.

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