

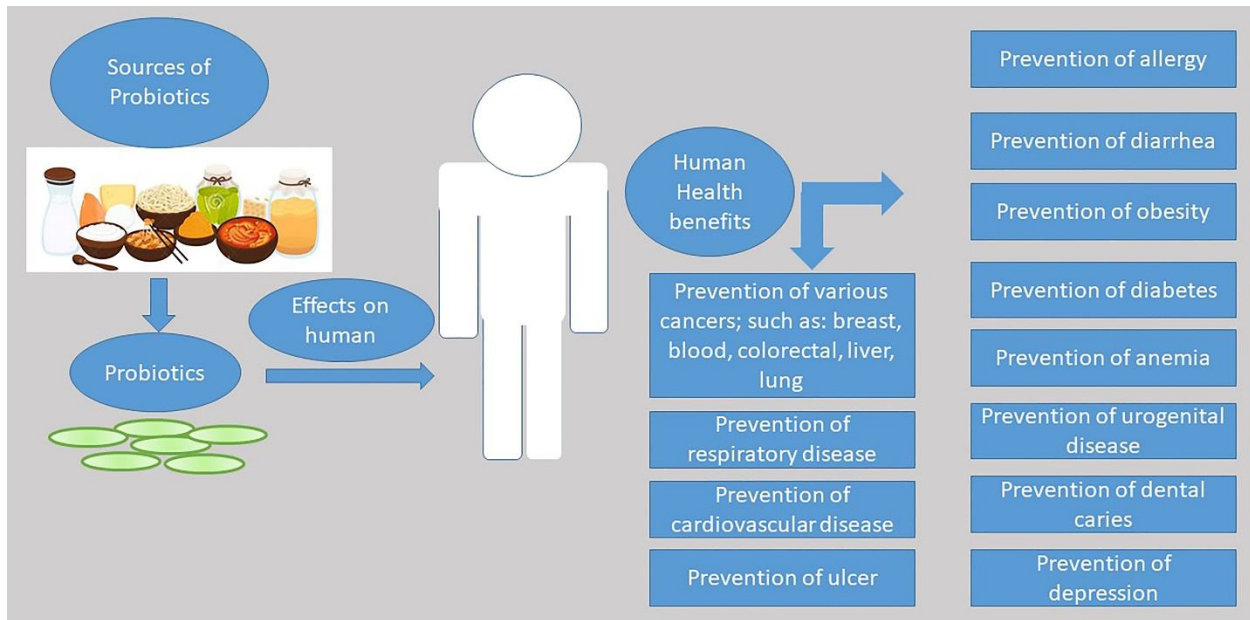
Emerging Role of Probiotics in Advancement of Combating Physical Abnormalities and Disease: A Systematic Perspective Analysis

ABSTRACT

Probiotics are live microorganisms, primarily bacteria that give the host health benefits when given in sufficient amounts. It works in various defensive mechanisms, including releasing antimicrobial compounds, competitive adhesion to the mucosa and epithelium, fortifying the gut epithelial barrier and regulating the immune system. Probiotics have a considerable effect in reducing allergies and gastrointestinal problems like diarrhoea and ulcers. Probiotics are quite beneficial for managing and preventing complex diseases including diabetes, cardiovascular diseases, anaemia, urogenital diseases, dental caries, URTI, cancers and various abnormalities like obesity, depression, etc. Nowadays, Antibiotic resistance is a global challenge associated with high morbidity and mortality. The current lack of efficacious treatments necessitates the creation of innovative therapeutic approaches and substitute antimicrobial therapies. So, natural products or probiotics would be a better option for treating human health. Therefore, we systematically reviewed the sources, including Google Scholar, PubMed, Science Direct, Jstor and Base; studies only about human health were included here. All types of animal diseases and articles before 1990 are excluded. A total of 6,410 articles were identified and 300 articles were taken for further evaluation based on context. However, Probiotics have already become a successful therapeutic intervention strategy among physicians. This investigation covers a comprehensive overview of the current state of probiotic research which comprises a wide range of topics including sourcing, mode of action, probiotic strains, isolating procedures

and the role of probiotics.

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Graphical abstract

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14 **Keywords:** Abnormality, Bacteria, Diseases, Defensive Mechanism, Probiotics.

15

16 1. Introduction

17 Antibiotics have been widely employed in managing bacterial infections because of their

18 capacity to either stop the growth or eradicate living microbes [1]. However, the spread of

19 antibiotic-resistance genes into harmful bacteria has sparked worries about the short-term

20 efficacy of the current antibiotic inventory [2]. Due to the global expansion of antibiotic and

21 antimicrobial resistance, the World Health Organization has classified these concerns as an

22 unanticipated global health danger with wide-ranging effects on the safety of humans, animals,

23 food and the environment across numerous sectors [2]. Thus, Probiotics are considered one of

24 the best substitute treatments [3]. German physicist Werner Kollath first used the term "probiotic"

25 in 1953 [4]. It is a combination of the Latin word "pro" and the Greek word "βίος," which means

26 "for life" [4]. Kollath uses this word to describe the "active substances" that are essential to the
27 healthy growth and well-being of living things [4]. According to the definition given to
28 probiotics in 1965, they are chemicals secreted by microbes that encourage the growth of other
29 microorganisms [5]. Probiotics are defined by the World Health Organization (WHO) as "living
30 microorganisms that, when administered in adequate amounts, confer benefits to the host's
31 health" [6].

32
33 Probiotics which can exist in food, supplements and the intestine are a significant class of
34 advantageous bacteria that are taken or added to diets [7]. These bacteria are capable of
35 colonizing the intestine and competing with dangerous microorganisms. Additionally, these
36 bacteria produce certain enzymes that aid in feed breakdown. Probiotics can enhance immune
37 activities by interacting with various immune cells and altering the composition of gut bacteria
38 when ingested [8][9][10]. Fundamental selection criteria should generally be used in the
39 selection and characterization of novel probiotic strains [11]. The Food and Agriculture
40 Organization (FAO) and the World Health Organization (WHO) of the United Nations released
41 guidelines in 2002 outlining the various factors that need to be considered when choosing
42 probiotic microorganisms [11]. These factors include antimicrobial activity, the ability to adhere
43 to epithelial tissues, resistance to harsh conditions inside the human body and safety for use [11].

44 Furthermore, a probiotic must be highly able to endure under intestinal circumstances, such as
45 acidic pH, enzymes, bile salts, etc. [12]. They have no harmful side effects and are not poisonous
46 or pathogenic [13]. To preserve the intended qualities, they are also appropriately labelled and
47 compatible with the product matrix, processing, and storage circumstances [13]. It has been
48 demonstrated that consuming 10^9 – 10^{10} viable colony-forming units (CFU) daily improves host
49 health [14]. So, probiotics are frequently prescribed to enhance health in nations like Canada,
50 Switzerland and Japan [15]. This manuscript reviews available information and summarizes the

51 current research knowledge about probiotics, their defensive mechanism, sources, isolation
 52 techniques, and the effects of various diseases that impact human health.

53

54 **Table 1.** List of some probiotic bacteria (gram-positive and gram-negative).

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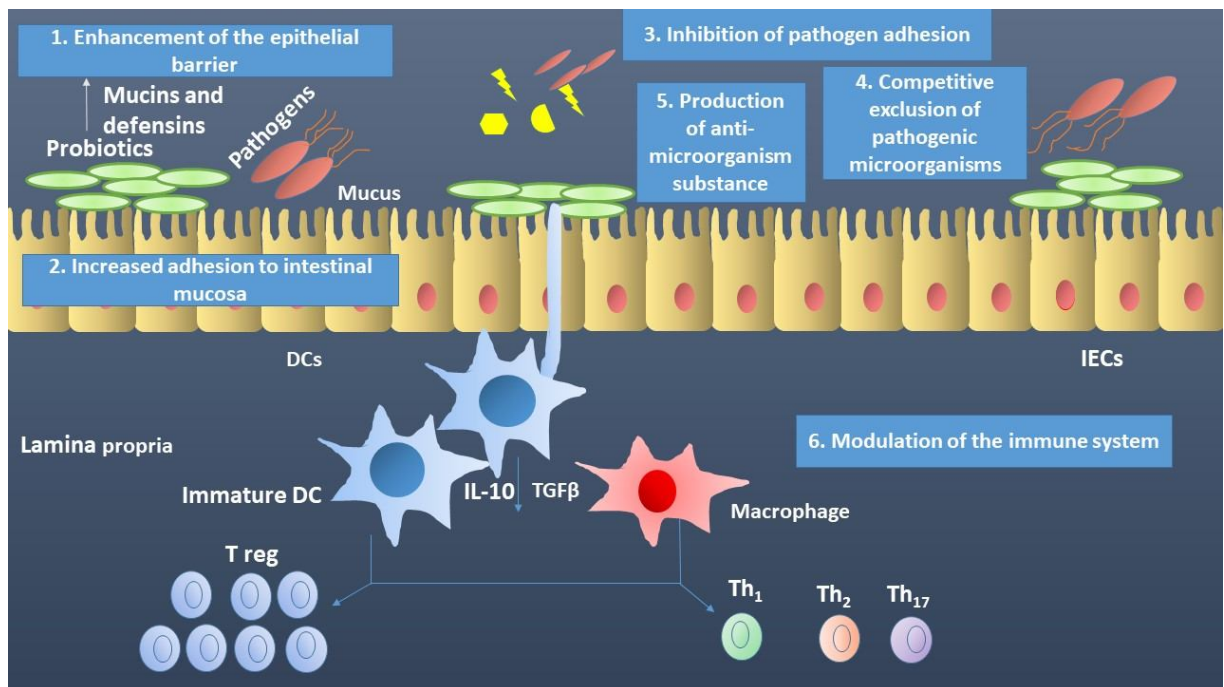
Previous Generation Probiotics		
Bacteria type	Bacteria name	References
Gram-positive	<i>Lactobacillus acidophilus</i> , <i>Lactobacillus casei</i> , <i>Lactobacillus fermentum</i> , <i>Lactobacillus gasseri</i> , <i>Lactobacillus johnsonii</i> , <i>Lactobacillus paracasei</i> , <i>Lactobacillus plantarum</i> , <i>Lactobacillus salivarius</i> , <i>Bifidobacterium adolescentis</i> , <i>Bifidobacterium animalis</i> , <i>Bifidobacterium bifidum</i> , <i>bifidobacterium genera</i> , <i>Leuconostoc</i> , <i>Enterococcus</i> , <i>Streptococcus salivarius</i> , <i>Bifidobacterium longum</i> , <i>Lactobacillus crispatus</i> , <i>Lactocaseibacillus rhamnosus</i> , <i>Lactocaseibacillus casei</i> , <i>Lactiplantibacillus plantarum</i> , <i>Limosilactobacillus reuteri</i> , <i>Levilactobacillus brevis</i> , <i>Ligilactobacillus salivarius</i> , <i>Lactococcus lactis</i> , <i>Pediococcus mesenteroides</i> , <i>Streptococcus thermophiles</i> , <i>Bacillus subtilis</i> , <i>Bacillus coagulans</i> .	[16][17][18][19][20][21][22]
Gram-negative	<i>Escherichia coli</i> , <i>Veillonella sp.</i>	[23][24][25]
New Generation Probiotics		
Bacteria type	Bacteria name	References
Gram-positive	<i>Clostridium butyricum</i> , <i>Anaerobutyricum hallii</i> , <i>Roseburia spp.</i>	[26]
Gram-negative	<i>Akkermansia muciniphila</i> , <i>Faecalibacterium prausnitzii</i> , <i>Bacteroides spp.</i> , <i>Bacteroides uniformis</i> , <i>Bacteroides acidifaciens</i> , <i>Bacteroides thetaiotaomicron</i> , <i>Prevotellacopri</i> , <i>Christensenellaminuta</i> , <i>Parabacteroides goldsteinii</i> .	[27][28][29]

56

57 *1.1 Mode of Action of Probiotics*

58 Though their exact processes are unclear, probiotics are thought to have several positive impacts
59 [13]. Probiotics work against various pathogens in several ways, including releasing
60 antimicrobial compounds, competitive adhesion to the mucosa and epithelium, fortifying the gut
61 epithelial barrier and regulating the immune system in Fig. 1.[13]. Probiotics can increase the
62 intestinal defence system through various strategies, including acting as an antagonist against
63 pathogens, enhancing the intestinal epithelial layer, producing neurotransmitters, increasing
64 innate immunity, and supporting adaptive immunity in Fig. 1.[30][31]. The intestinal barrier is
65 one important defensive mechanism that keeps the organism safe from the environment and
66 preserves epithelial integrity [32]. The mucous layer, antimicrobial peptides, secretory IgA and
67 the epithelial junction adhesion complex are the intestinal barrier's defences[32]. When this
68 barrier function is damaged, food antigens and bacteria can enter the submucosa and cause
69 inflammatory reactions that could lead to digestive problems. Probiotic microorganisms have
70 been thoroughly investigated in terms of their role in maintaining this barrier. Mucin
71 glycoproteins, often called mucins, are important macromolecules that make up epithelial mucus
72 and have been linked to health and illness. Probiotics may increase mucus secretion to enhance
73 barrier performance and pathogen exclusion in Fig. 1. [18].

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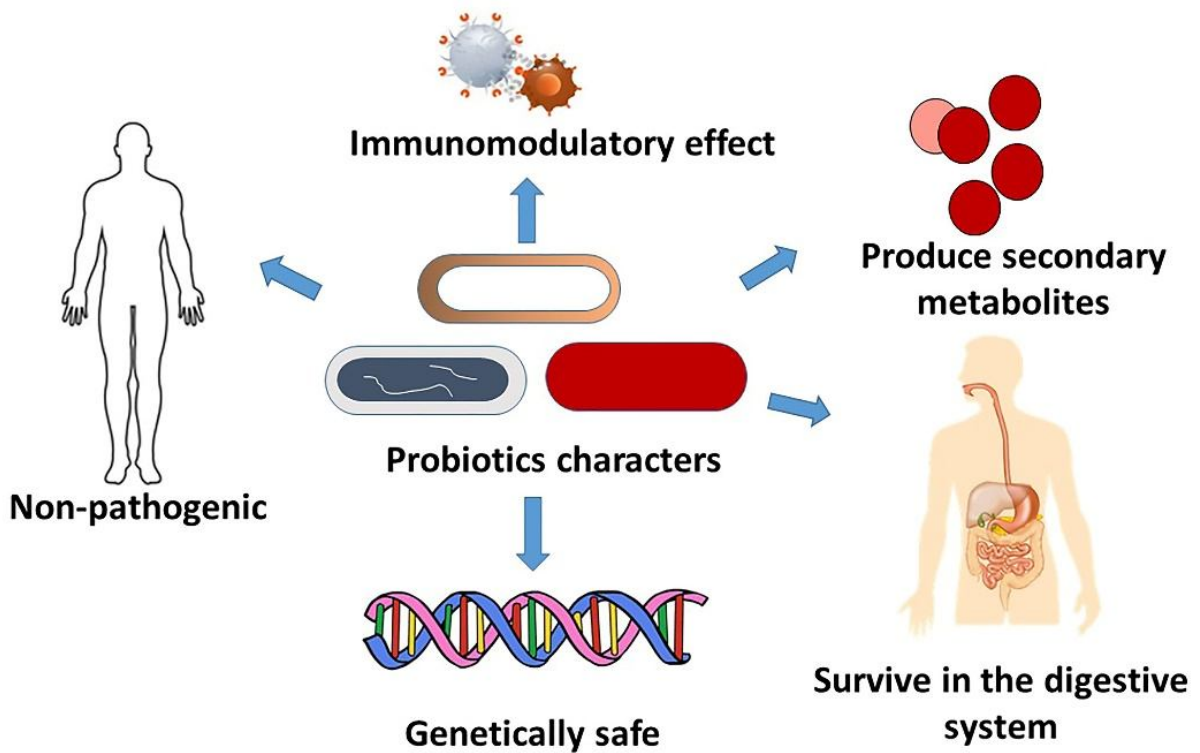
76

77 **Fig. 1.** Major mechanisms of action of probiotics [16] [17] [18].

78

79 It is thought that adhesion to intestinal mucosa is necessary for colonization and plays a crucial
 80 role in the relationship between probiotic strains and their host. The particular interaction has
 81 suggested a potential link between probiotic bacterial surface proteins and pathogens'
 82 competitive removal from mucus [33][34]. Probiotics also control innate and adaptive immune
 83 responses by influencing B and T lymphocytes, dendritic cells (DC) and macrophages [35].
 84 Probiotics interact with intestinal epithelial cells and draw macrophages and mononuclear cells,
 85 increasing the generation of anti-inflammatory cytokines[35]. Additionally, probiotics can use
 86 the gut-brain axis to create neurotransmitters in the gut. The amounts of serotonin, gamma-
 87 aminobutyric acid (GABA) and dopamine can be modulated by some probiotic strains which can
 88 impact mood, behaviour, gut motility and stress-related pathways [36][37][38].

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90

91 **Fig. 2.** Features of probiotics [21] [22] [23] [24].

92

93 *1.2 Sources of probiotics*

94 Probiotic foods are the subject of intense interest due to their positive effects on human health

95 [39]. Thus, there is a corresponding increase in the study of probiotic microbes [39].

96 Consequently, probiotics may be presented in many ecological niches besides humans [40].

97 Probiotics are becoming increasingly common in the food, pharmaceutical and fermentation

98 industries, and customers generally accept them favourably[40]. Besides, probiotic-rich foods

99 include dairy and dairy-related products [41]. The primary supplies could come from human

100 origins such as breast milk or the large or small intestine of humans [41]. Milk is an excellent

101 medium for both good and bad microbes due to its high nutritional content [42]. Moreover, the

102 best source of probiotics for human consumption is the faeces of newborns, kids and adults[43].

103 Their capacity to withstand gastrointestinal transit and colonize the intestines for beneficial

104 purposes is the reason behind this [43]. Additionally, foods of animal origin and foods of plant

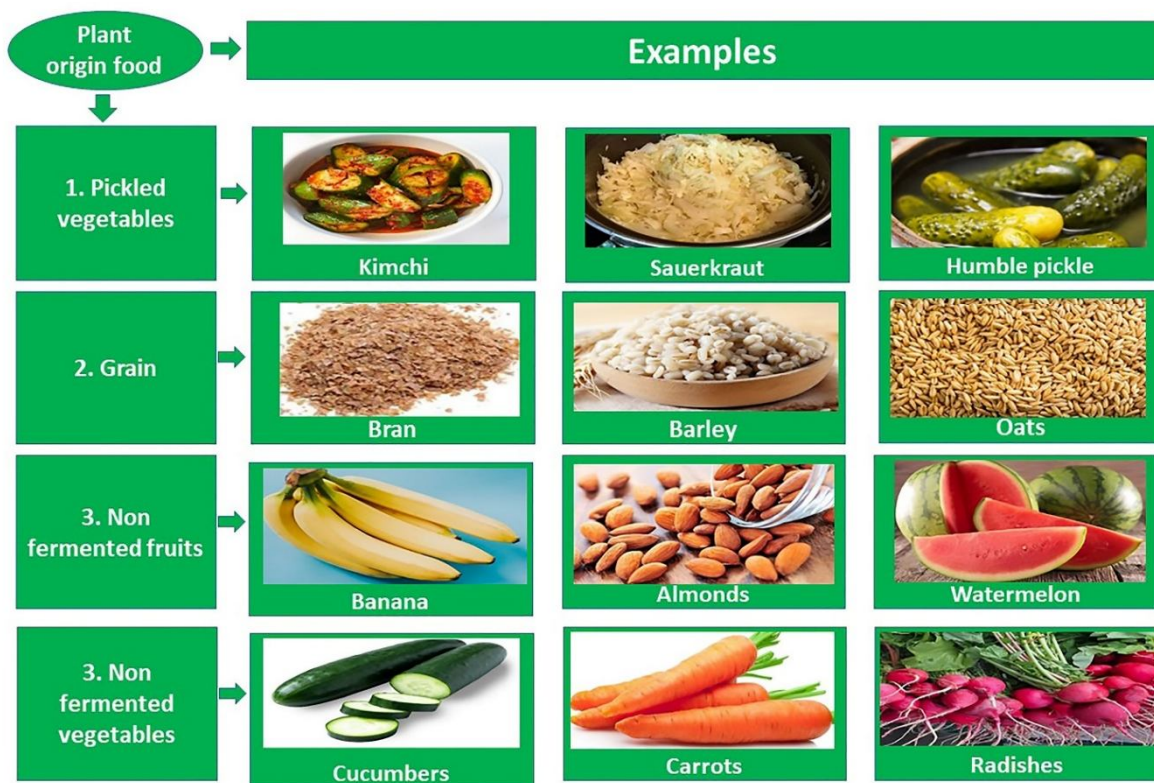
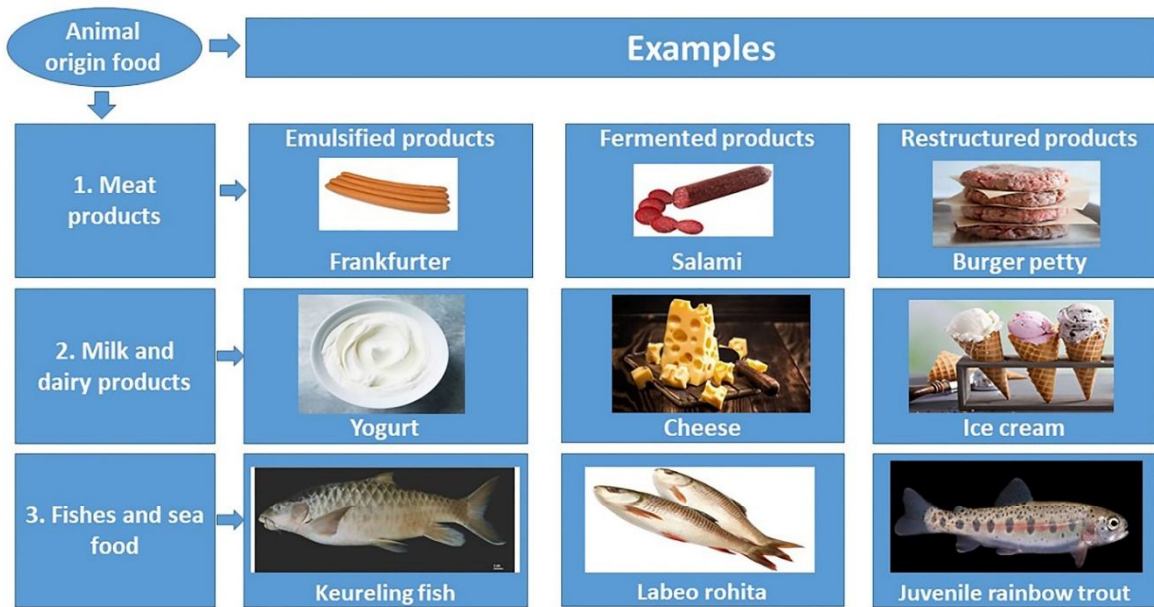
105 origin may provide additional sources for isolating probiotic strains [44]. Probiotics found in

106 animal origin can be divided into three categories such as meat products(emulsified, fermented,
107 and restructured products), milk and dairy products(yogurt, cheese, and ice cream) and fish and
108 seafood(keureling fish, *labeorohita* and juvenile rainbow trout) [45][46][47].

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110

111 Probiotics can also be found in plants of origin. They can be divided into four categories such as
112 pickled vegetables(kimchi, sauerkraut and humble pickle), grain(bran, barley and oats), non-
113 fermented fruits(banana, almonds and watermelon) and non-fermented vegetables (cucumbers,
114 carrots and radishes) [48][49][50]. However, Supplements containing probiotics are also offered
115 [51]. The bacterial strains found in probiotic foods and supplements should be essential for the
116 following processes: lowering and metabolism of cholesterol; colonization of the intestinal,
117 respiratory and urogenital tracts; inhibition of carcinogenesis, either directly or indirectly,
118 through immune system stimulation; lactose metabolism; calcium absorption and vitamin
119 synthesis potential; **the** reductive potential of yeast and vaginal infection; reduction of
120 constipation and diarrheal disorders; mitigation of gastritis and ulcers; reduction of acne, rash
121 face and skin issues; and the synthesis of natural antimicrobials [52][53].



122

123 **Fig. 3.** Probiotic sources: animal and plant-origin foods [54].

124

125 1.3 Isolation of probiotics

126 According to FAO/WHO guidelines, the first step is to isolate possible probiotic strains [55].

127 Researchers have determined that new probiotic bacterial strains are still needed for certain

128 therapeutic applications [55]. The probiotics derived from dairy products differ in several ways
129 from the probiotics isolated from the intestines of people and animals [56]. A prominent
130 distinction between these probiotics is their adherence. The adhesion activity of intestinal
131 isolates is often higher than that of dairy isolates [56]. To extract probiotic strains from dairy
132 products, samples were gathered from various sources, including camel milk, sheep yogurt, cow
133 yogurt, goat milk and native buttermilk [57]. Also, the sample was taken from clinical samples
134 including blood, urine, respiratory, pericardial, abdominal and cerebral fluid (CSF) [58]. These
135 samples were cultured on agar and broth media and incubated for 24.0 to 48.0 hours at 37.0
136 °C[57]. The strains were identified using a variety of tests, such as Gram staining, catalase
137 testing, growth at temperatures between 15.0 °C and 45.0 °C, production of acid and gas from
138 glucose, production of ammonia from arginine, and fermentation of sugars (arabinose,
139 cellobiose, mannitol, mannose, melibiose, raffinose, ribose, salicin, rhamnose and xylose)
140 [57][58][59].

141

142 2. Methodology

143 Different sources were generated for article searches including Google Scholar, PubMed,
144 Science Direct, Jstor, and Base. The information mainly searches in several articles from 2000 to
145 July 2024. The keywords that were used for searches included probiotics, advancements and
146 disease management. The obtained articles were either full text or just the title and abstract.
147 Some criteria followed to search the articles included Probiotics, Sources of probiotics,
148 Mechanism of action of probiotics, Probiotics in human health, Role of probiotics in different
149 diseases (probiotics and allergy, probiotics and diabetes, probiotics and cancer, probiotics and
150 probiotics and cardiovascular diseases, probiotics and diarrhoea, probiotics and anaemia,
151 probiotics and ulcers, probiotics and obesity). The articles that met the requirements for inclusion
152 in this review were either published in peer-reviewed journals, reviewed or authored in English.
153 Only studies about human health were included in the review. The exclusion criteria were as

154 follows: no conference publications or book chapters, no mention of animal diseases that are host
155 specific like foot and mouth disease, no article searches in other languages and yeast or fungus
156 strains not acting as probiotics.

157

158 **3. Role of probiotics in various diseases and abnormalities of the human body**

159 **3.1 Prevention of allergies**

160 Nowadays, Food allergies are becoming very common globally which lowers the quality of life
161 for people who are impacted. People of all ages have been especially affected by food allergies.
162 Nearly 6.0 % of children and 3.0 to 4.0 % of adults suffer from food allergies [60]. Mainly, food
163 allergy is a consequence of several genetic and environmental variables, leading to an absence or
164 reduction in tolerance of particular foods. Sometimes, Food-induced immune responses can
165 result in various symptoms and conditions, such as anaphylaxis and severe allergic reactions
166 [61]. So, avoiding allergens and taking emergency medicine is the only way to manage food
167 allergies without a successful course of treatment [62]. Probiotics help allergies by healing the
168 digestive tract [63]. As a result of this intervention, the immune system stabilizes, the intestinal
169 epithelial barrier is strengthened and inflammation is reduced. Immunological processes cause
170 hypersensitivity reactions which are the outward manifestation of allergies [64]. Probiotics can
171 rearrange antigens, lessening their capacity to elicit an immunological response [64]. This
172 ultimately results in a reduction of intestinal permeability and the class of pro-inflammatory
173 compounds. People who suffer from various allergies often experience these symptoms.
174 Research indicates that *L. rhamnosus* GG, a probiotic bacteria, may lessen the intensity of allergic
175 reactions and ease the symptoms associated with food allergies [63][64]. According to a more
176 contemporary hygiene hypothesis, certain organisms such as *helminths* and microorganisms such
177 as *bifidobacteria*, *lactobacilli* and *saprophytic mycobacteria* are necessary for the healthy
178 development of the human immune system. To support immunological tolerance, these
179 microorganisms are believed to have coevolved with the human immune system as part of our

180 microbiota during human evolution. This involves the production of regulatory T cells (Treg)
181 and the regulation of the balance between Th2 and Th1 which helps avoid the development of
182 autoimmune and allergic diseases [65][66]. A study was conducted that discovered that
183 supplementing with *L. casei* CRL431 and *B. lactis* Bb-12 for twelve (12.0) months did not
184 influence the development of tolerance in 119 infants who were allergic to cow's milk[60].

185

186 3.2 Antimicrobial effect

187 One possible strategy to stop the spread of antibiotic-resistant bacteria is using probiotics [67].
188 Probiotics use several important antimicrobial pathways to accomplish this, including immune
189 system modulation, mucin and tight junction protein production, competitive exclusion, and
190 improved intestinal barrier function [68]. Probiotics produce antimicrobial peptides essential for
191 competitively excluding microorganisms and are a major factor in their antimicrobial
192 effectiveness. The antibacterial activity of *Lactobacillus* strains such as *S. aureus*, *P. aeruginosa*,
193 *Shigella sp.*, *Clostridium difficile*, *Klebsiella sp.*, and *E. coli* has been particularly effective
194 against a range of infections [69]. They produce lactic acid, bacteriocin, and hydrogen peroxide
195 which prevent the attachment of harmful bacteria to the mucosa and enhance the immune
196 response, among other efforts to outcompete and suppress these pathogenic bacteria [70]. The
197 release of chemicals known as bacteriocins is the main way that *Lactobacillus* strains
198 demonstrate antibacterial activity [71]. Antimicrobial peptides known as bacteriocins fight both
199 Gram-positive and Gram-negative bacteria. Interestingly, the bacteria that make these
200 bacteriocins have evolved defence mechanisms against the effects of their antimicrobial peptides
201 [72]. It is also known that certain strains of *Lactobacillus* and *Bifidobacterium* produce
202 bacteriocins. The antimicrobial peptides exhibit multiple modes of action such as blocking the
203 creation of peptidoglycan and pore formation and inhibiting lipid II, an essential constituent of
204 bacterial cell membranes [73]. A study showed that by restoring cell viability and regulating the

205 production of inflammatory cytokines, *B. longum* and *L. reuteri* provide protective benefits in
206 corneal infection induced by *P. aeruginosa*[74].

207

208 **3.3 Prevention of diarrhoea**

209 Numerous clinical investigations have evaluated the probiotics' effectiveness in preventing acute
210 diarrhoeal illness [75]. The prevalent side effect of repeated short- and long-term antibiotic usage
211 is diarrhoea[75]. To stop antibiotic-associated diarrhea (AAD), two meta-analyses utilizing the
212 strains *Lactobacillus acidophilus*, *Lactobacillus rhamnosus* strain GG, *Lactobacillus bulggaricus*
213 and yeast *Saccharomyces boulardii* could be employed [75]. A study was conducted in patients
214 with spinal cord injury (SCI) who frequently took proton pump inhibitors (PPI) in 7.0 days. This
215 study aimed to evaluate the effectiveness of a probiotic containing at least 6.5×10^9 live
216 *Lactobacillus casei*Shirota (LcS) in reducing antibiotic-associated diarrhoea (AAD). When LcS
217 is administered regularly to SCI patients who fall into a specific vulnerable group, it may be able
218 to prevent AAD [76].Another study was conducted to assess *Lactobacillus rhamnosus* GG
219 (LGG) effectiveness in preventing antibiotic-associated diarrhoea (AAD) in adults and
220 children(1499 participants). This finding indicates that in children and adults receiving antibiotic
221 treatment for any cause, *Lactobacillus rhamnosus* GG is useful in avoiding antibiotic-associated
222 diarrhoea[77]. Thus, nosocomial diarrhoea is a major issue in pediatric hospitals and is a global
223 problem. Probiotics (*Streptococcus thermophiles*, *Bifidobacterium bifidum*, and *L*
224 *rhamnosus*starin GG) can be used prophylactically to prevent acute diarrhoea, especially in
225 babies [75].A study was also conducted in children aged six (6.0) months to five (5.0) years with
226 watery diarrhoea in five (5.0) days. The result shows that probiotics are more effective in treating
227 acute watery diarrhoea in hospitalized children. Probiotics may help lessen the intensity and
228 length of diarrhoea when taken in addition to conventional treatment [78].

229

230 **3.4 Prevention of obesity**

231 Human studies that examined the benefits of probiotic use included changes in lipid profiles,
232 namely lower levels of total cholesterol, LDL cholesterol and plasma TG and higher levels of
233 HDL cholesterol [79]. During 4.0 to 6.0 weeks of intake, the majority of research indicates the
234 anti-obesity advantages of some types of probiotics such as *Lactobacillus* (primarily
235 *Lactobacillus casei* strain Shirota (LAB13), *Lactobacillus gasseri*, *Lactobacillus rhamnosus* and
236 *Lactobacillus plantarum*) and *Bifidobacterium* (particularly *Bifidobacterium infantis*,
237 *Bifidobacterium longum*, and *Bifidobacterium breve* B3) [80][81]. Additionally, probiotics with
238 anti-obesogenic properties, including *L. rhamnosus* CGMCC1.3724, help prevent obesity-related
239 issues [82]. A study was conducted to examine how the probiotic *Lactobacillus gasseri* SBT2055
240 (LG2055) affected body weight, other body measurements and abdominal adiposity in persons
241 who tended toward obesity while consuming 200.0 g/day of Fermented milk for 12.0 weeks. The
242 result demonstrated that the probiotic LG2055 decreased the effects on body weight, abdominal
243 obesity and other metrics [83]. Another study was conducted with 45.0 patients with obesity,
244 separated into three groups: probiotics (30.0 g/day), prebiotics (30.0 g/day) and single diets (low-
245 carb and energy-reduced diets with *Bifidobacterium longum*, *Lactobacillus helveticus*,
246 *Lactococcus lactis* and *Streptococcus thermophilus* per day). There is clinical evidence that the
247 prebiotic and probiotic groups showed a significant decrease in fat mass, increased muscle
248 strength, and reduced fasting blood glucose [84]. The other study was carried out with 66 .0 obese
249 patients for three (3.0) months with a BMI in the range of 30.0 to 40.0 kg/m² in obese adults. The
250 result shows that total cholesterol, triglycerides and low-density lipoprotein levels were
251 decreased [85].

252

253 3.5 Prevention of diabetes

254 Patients with type 2 diabetes may have changed gut microbial composition. This implies that
255 altering the gut microbiota by probiotic medication could be a strategy for controlling glucose
256 metabolism. Probiotics may have a hypoglycemic effect, according to numerous studies but the

257 exact processes behind this are still up for debate because the aetiology of type 2 diabetes is so
 258 complicated [86]. Probiotics may help to mitigate several metabolic syndrome components,
 259 many of which are connected to type 2 diabetes [87]. The preservation of a range of mucosal
 260 immune cells may inhibit potential pathogens; pro-inflammatory cytokines (TNF- α and
 261 interleukin-6) may be reduced; delayed glucose absorption in the gut may contribute to these
 262 improvements; and rate-controlling enzymes of gluconeogenesis, such as glucose-6-phosphatase
 263 (G-6-Pase) and phosphoenolpyruvate carboxykinase (PEPCK), may be suppressed. In patients
 264 with type 2 diabetes, unchecked gluconeogenesis results in hyperglycemia [156]. Thus, probiotic
 265 therapy that inhibits G6Pase and PEPCK lowers glycemia by reducing the amount of glucose
 266 released into the bloodstream from the liver [88].

267
 268 **Table 2.** Overview of important studies demonstrating the effect of probiotics on type 2 diabetes
 269 mellitus.

Probiotics	Model types	Time of experiment	Finding	Reference
<i>Lactobacillus brevis</i> 15 and <i>Lactobacillus plantarum</i> 13	24, 2.0-month-old Wistar rat	8.0 weeks	According to this study, the recently identified Bulgarian <i>Lactobacillus</i> strains <i>L. brevis</i> 15 and <i>L. plantarum</i> 13 may be potential probiotics that might help avoid certain metabolic disorders and diabetes-related difficulties.	[89]
Symbiter	53, patients	8.0 weeks	A little improvement in insulin resistance.	[90]
<i>L. rhamnosus</i> BSL and <i>L. rhamnosus</i> R23	Male Sprague-Dawley rats ($n = 24$)	1.0 months	In a streptozotocin-induced diabetic rat model, the liver showed better gene expressions and reduced fasting blood glucose after receiving 30 days' worth of <i>L. rhamnosus</i> BSL and <i>L. rhamnosus</i> R23 dose, reduction of glucose-6-phosphatase to increase insulin sensitivity.	[91]
<i>Bifidobacterium imalis</i> A6	28, type II diabetic patients	4.0 weeks	Patients with type 2 diabetes can greatly lower their fasting blood glucose by taking probiotic camel milk powder twice a day for four (4) weeks in a row.	[92]
Ecologic®Barrier	patients with type 2 diabetes	6.0 months	Multistrain probiotics show promise as an adjunct treatment for diabetes.	[93]

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273 **3.6 Prevention of anemia**

274 Numerous researches have demonstrated a strong correlation between iron deficiency and gut
 275 microbiota. Moreover, studies have demonstrated the positive effect of probiotic bacteria on
 276 improving iron absorption and the reciprocal association between gut microbiota composition
 277 and iron insufficiency [94][95]. Probiotic bacteria such as *Lactobacillus cerevisiae*,
 278 *Lactobacillus cremoris*, *B. pseudocatenulatum*, *Candida famata*, *B. adolescentis*, *Candida*
 279 *glabrata*, *Candida guilliermondii*, *S. cerevisiae*, *Yarrowialipolytica* and *Pichia glucozyma* are
 280 the source of folic acid, a water-soluble B vitamin that is essential for treating and managing
 281 anemia. These microbes are used to improve folic acid absorption through the intestines. *P.*
 282 *denitrificans* and *P. shermanii* have also been discovered to be effective in treating vitamin B12
 283 insufficiency. Foods fermented with lactic acid have a crucial role in boosting the absorption of
 284 iron and are utilized in the treatment of anaemia in patients. They aid in regulating the digestive
 285 tract's pH and stimulate the phytase enzyme which aids in the absorption of nutrients. Moreover,
 286 megaloblastic anaemia has been treated by adding a mix of probiotic bacteria to the diet. These
 287 probiotics mitigate the negative effects of antibiotics and encourage intestinal fermentation [96].

288

289 **Table 3.** Different studies on the treatment of anaemia using probiotics.

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Subject	Number of patients	Study duration	Observation	References
Healthy woman	Study 1- 14; Study 2- 28	4.0 weeks	In the probiotics group: significant increase of iron in serum; In the placebo group: Changes are insignificant.	[97]
Healthy woman	20	1.0 week	Probiotics Group: Significant increase in iron.	[98]
Children with anaemia	52	3.0 years	Nosignificant differences in serum ferritin levels between the	[99]

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3.7 Prevention of urogenital disease

According to recent research, maintaining a lactobacilli-dominated microbiome is crucial for preserving women's quality of life and preventing STDs and premature labour. It was questioned if artificial lactobacilli supplementation may reduce infection rates because women who are prone to vaginal and urinary infections lose lactobacilli organisms. However, uro-pathogens are becoming more resistant to common antibiotics such as trimethoprim/sulfamethoxazole and patients are turning increasingly to natural alternatives like cranberry juice which seems to contain antiadhesive compounds that are effective against uro-pathogens and can help prevent UTIs. Probiotic therapy using lactobacilli has been explored as a non-chemotherapeutic way to maintain and repair a healthy urogenital tract. There is evidence that some strains of lactobacilli can be beneficial when inserted directly into the vagina or when ascending from the rectum after oral ingestion. Probiotic consumption regularly may be beneficial for all healthy women as well as those who experience recurrent urogenital infections to prevent serious infections and superinfections of the vaginal mucosa. Probiotics have been shown to help keep the vaginal flora in balance which protects against UTIs [100]. A study was conducted with 174 premenopausal women with a history of recurrent UTIs at four (4.0) months and examined the efficacy of oral and intravaginal probiotic supplementation as a preventative measure against recurrent urinary tract infections. The result demonstrated the effectiveness in preventing recurrent symptomatic UTI episodes [101]. Another study was conducted with a population of 132 individuals with probiotics, antibiotics and probiotics plus antibiotics being administered to 42, 48 and 42 of the participants in each group at 30.0 days. This study found that probiotics are a safe and efficient way to treat UTIs in Bangladeshi women [102].

316 **3.8 Prevention of dental caries**

317 Dental caries is one of the most prevalent illnesses in the world, causing exorbitant expenses and
318 being regarded as a public health issue in youngsters. Various therapies have been suggested to
319 lower the prevalence of this illness, with probiotic supplementation being one of the main ones.
320 According to recent studies, probiotic supplementation appears to lower the risk of dental caries
321 [103][104][105]. *Streptococcus mutans* (SM), one of the several bacterial species found in saliva,
322 is widely regarded as the pathogen primarily responsible for the early stages of tooth decay,
323 whereas *Lactobacillus* plays a significant part in the disease's progression. The primary traits of
324 *Streptococcus mutans*' pathogenicity include its acidogenicity, capacity to endure in acidic
325 conditions, capacity to form biofilms, and capacity to cling to teeth. The probiotic *Lactobacillus*
326 *paracasei* SD1 was introduced and used in the oral cavity due to its well-known ability to
327 prevent *Streptococcus mutans* from growing, lessen its acid production, and adhere well to oral
328 epithelial cells [106]. A study involved 260 participants of which 130 were in the control group
329 and 130 in the intervention group at 4.0, 8.0 and 12.0 months. The group that took probiotic milk
330 tablets demonstrated statistically significant decreases in carious lesions when compared to the
331 control group [107].

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334 **3.9 Prevention of depression**

335 The gut-brain axis is the term for the reciprocal connections between the brain, intestinal
336 microbiota and the digestive system via immunological, neurological and endocrine pathways.
337 The gut microbiota responds to the physical and physiological strain its host endures. Stress
338 drastically reduces the amount of Firmicutes phylum, particularly in *Bifidobacterium* and
339 *Lactobacillaceae* family bacteria and is a major risk factor for depression. It also modifies the
340 percentage and makeup of the gut microbiota. Stress weakens the brain's relationship with the
341 gut microbiota which can result in metabolic depression. However, Changes in brain activity in

342 patients with depression are known to be caused by higher levels of cortisol in plasma and
343 corticotrophin-releasing factor in the cerebrospinal fluid. Furthermore, catecholamine levels
344 (primarily those of norepinephrine and dopamine) rise when these factors are repeatedly
345 activated. This raises the risk of various physiological problems such as attention deficit
346 disorder, balance and posture, cognitive function, emotions and sleep-wake cycles.
347 *Bifidobacterium* baby restored the stress-induced drop in norepinephrine, indicating that
348 probiotics influenced catecholamine metabolism [108]. A clinical trial consisted of 45 patients
349 with *Bifidobacterium breve* CCFM1025 in 4.0 weeks to examine the psychotropic potential for
350 the treatment of major depressive disorder (MDD). This trial shows that *B. breve* CCFM1025 is
351 a good candidate that reduces depression and related gastrointestinal issues [109].

352

353 ***3.10 Prevention of cardiovascular diseases***

354 Cardiovascular problems commonly affecting the heart include arrhythmia, thromboembolic
355 disease, cardiomyopathy and atherosclerosis. Globally, the prevalence of cardiovascular
356 disorders is rising. Cardiovascular diseases accounted for 18.0 million fatalities in 2015 or over
357 one-third of all deaths with documented causes. This number indicates a noteworthy increase of
358 12.50% when compared to the data from 2005. Predictions from the American Heart Association
359 suggest that by 2030, around 43.90% of the US population will be suffering from cardiovascular
360 disease. In the United States, 92.10 million persons currently suffer from cardiovascular disease.
361 Atherosclerosis is a chronic inflammatory disease characterized by the creation and progression
362 of lipid-fueled atherosclerotic plaques in the artery wall [110]. From spontaneously fermented
363 mustard, researchers obtained the *Lactobacillus plantarum* DMDL 9010 strain (CGMCC No.
364 5172). The potential benefits of this strain in lowering cholesterol were examined using an in
365 vivo model. The study's findings demonstrated that rats given 10⁹ cells of *L. plantarum* DMDL
366 9010 daily experienced significant reductions in their levels of total cholesterol (TC), low-
367 density lipoprotein cholesterol content (LDL-C), and atherosclerosis index (AI) of 23.03%,

368 28.00%, and 34.03%, respectively. After administering *L. plantarum* DMDL 9010, there was no
369 statistically significant alteration in blood triglyceride levels. Rat hepatocyte protection against
370 steatosis was demonstrated by morphological and pathological alterations in the liver following
371 *L. plantarum* DMDL 9010 therapy. Significant reductions in hepatic cholesterol (33.20%) and
372 triglyceride levels (40.86%) were also observed upon administration of substantial dosages of *L.*
373 *plantarum* DMDL 9010. Significant increases were also observed in bile acid (+70.18%) and
374 faecal cholesterol (+31.07%). After receiving *L. plantarum* DMDL 9010 treatment, there was a
375 reduction in blood and total hepatic cholesterol and triglyceride levels and increased bile acid
376 excretion through stool. These outcomes seemed to depend on the dosage of *L. plantarum*
377 DMDL 9010. Thus, Probiotic supplementation may help those who are susceptible to ischemic
378 heart disease escape the devastation that comes with it [111].

379 380 **3.11 Prevention of upper respiratory tract infections**

381 The most prevalent illnesses are upper respiratory tract infections (URTIs), including tonsillitis,
382 pharyngitis, laryngitis, sinusitis, acute respiratory distress known as coronavirus, and the
383 common cold[112][113]. Adults experience colds two to three times a year, and 17.2 billion
384 URTIs are thought to have occurred globally in 2015 [114][115]. Numerous viruses, including
385 the influenza virus, adenovirus, rhinovirus and respiratory syncytial virus are the primary cause
386 of URTIs. The symptoms can include chills, fever, headache, body aches, sore throat, runny or
387 stuffy nose, cough and so on [116][117]. Retaining the immune system in a healthy, normal state
388 reduces the likelihood of recurrent infections. Salivary immunoglobulin A (IgA) and NK cell
389 activity are thought to be crucial in the prevention of URTIs. The immune system can be
390 weakened by several environmental variables such as leading a stressful lifestyle which could
391 raise the risk of URTIs. Evidence shows that certain probiotic strains increase salivary IgA levels
392 and activate NK cells again. Therefore, regular use of a probiotic beverage may help to regulate
393 URTIs and preserve normal immunological function. A probiotic known as *Lactobacillus casei*

394 strain Shirota (LcS) can survive in the digestive system and restore a balanced gut flora.
395 Numerous studies have been conducted on human and animal models to examine the strain's
396 immunomodulatory properties. Research involving healthy individuals with low NK cell activity
397 revealed that consumption of LcS-fermented milk (LcS-FM) increased the activity [118].

398

399 *3.12 Prevention of ulcers*

400 The sores that form on the skin's surface that are painful and inflammatory are called ulcers.
401 Clinical characteristics determine whether the illness is considered mild, substantial, or
402 herpetiform. Many theories have been put out to explain the illness, including immunologic
403 abnormalities, viral infections, and environmental and psychological stress. However, the
404 **disorder's aetiology** is still unknown and up for debate. Probiotics given in sufficient amounts
405 have been shown to improve host health. One way **probiotics help** reduce sickness is by
406 enhancing and modulating the immune system. Probiotics can produce a variety of substances,
407 such as peptides, bacteriocins and organic acids. Consequently, there is a decreased chance of
408 harmful germs colonizing [119]. **Oral ulcers are a common oral ailment that is distinguished by a**
409 **high degree of complexity, diversity and frequency.** Several etiological factors such as allergic
410 reactions, traumatic experiences and infectious agents, can result in oral ulcers. **The hallmarks of**
411 **mouth ulcers are the loss of connective tissue and the development of crater-like depressions**
412 **brought on by persistent disruption or damage to the integrity of the mouth epithelium.** Bacteria
413 that are resistant to infections and benign are known as probiotics. In addition to strengthening
414 immunity and preventing infection by fighting harmful bacteria, they may also improve digestion
415 and make it easier for the host to absorb nutrients. **They influence both immune system**
416 **regulation and tissue healing.** It may be possible to increase the effectiveness of probiotics in
417 aiding the healing process of ulcers by using fucoidan (FD), a prebiotic obtained from marine
418 sources. **Probiotics can create a biofilm in the oral cavity that can successfully compete with**
419 **periodontal and cancer-causing bacteria.** **They also help** prevent diseases by enhancing immune

420 system performance and modulating the host's immunological response. Dental health is
421 significantly improved while using oral probiotics [120][121].

422
423 The development of peptic ulcers in the duodenum or stomach is known as peptic ulcer disease.
424 The main risk factors for gastric and duodenal ulcers are the use of aspirin or nonsteroidal anti-
425 inflammatory medicines (NSAIDs) and the existence of an *H. pylori* infection. The function of
426 the barrier may be improved by probiotics. The primary defence against pathogenic bacteria is
427 the acid and mucus barrier found in the stomach's gastric mucosa. Some probiotics can increase
428 the upregulation of fitting linked proteins and promote the secretion of mucus and mucin which
429 strengthens the mucus secretion process and improves the protective capacity of the stomach
430 mucosal barrier. Furthermore, some probiotic bacteria can create antimicrobial substances such
431 as hydrogen peroxide, lactic acid, short-chain fatty acids (SCFAs) and bacteriocins. Short-chain
432 fatty acids (SCFAs) and lactic acid both show signs of partial dissociation and when these
433 organic acids are in their dissociated stages, *H. pylori* is negatively impacted [122][123].

434

435 ***3.13 Prevention of breast cancer***

436 Breast cancer makes for 29.0 % of all cancer cases identified in women, making it the most
437 frequent cancer among them. An estimated 2.50 million instances of breast cancer were reported
438 in 2016 [124]. It was previously believed that disparities in hormonal components, reproductive
439 patterns, and detection techniques at various stages were the causes of the wide variances in
440 breast cancer incidence [125][126]. Breast cancer mortality has continuously dropped as a result
441 of advancements in breast cancer detection and treatment. It was previously believed that
442 probiotic supplements could lower the incidence of breast cancer in perimenopausal women.
443 Bacteria believed to have anticancer properties include *Lactococcus* and *Streptococcus* which are
444 more common in healthy breast tissue. To see whether they may cause DNA damage, bacteria
445 from malignant females were cultivated on human breast cancer cells. Triple-stranded DNA

446 breaks generated by *Enterobacteriaceae*, *Escherichia coli*, and *Staphylococcus* bacteria were
447 alarmingly visible throughout the proliferation of these malignant cells. Carcinogens can be
448 broken down by the probiotic bacterium *Lactobacillus*. The probiotic *S. thermophilus* strain
449 produces antioxidants that reduce DNA damage [127]. Moreover, Research has indicated that
450 *Lactobacillus crispatus* and *Lactobacillus acidophilus* exhibit antiproliferative properties against
451 breast cancer cells. Furthermore, the transcriptional activity of four distinct cancer-testis antigens
452 can be decreased by lactobacilli. It has been revealed that epigenetic controls have a role in the
453 expression of cancer-testis antigens. Thus, its expression may be epigenetically downregulated
454 by lactobacilli. A new era of clinical research applications could be enabled by lactobacilli-
455 mediated expressional downregulation of cancer-testis antigens, as these antigens have been
456 associated with high-grade malignancies and poor prognosis [128].

457

458 **3.14 Prevention of blood cancer**

459 Most hematopoietic malignancies, often known as blood cancers, start in the bone marrow,
460 which is where blood cells are made. Research indicates that probiotics may be useful in the
461 treatment of cancer. A study found a correlation between chronic myeloid leukaemia and the
462 therapeutic effects of heat-inactivated *Lactobacillus casei* and *Lactobacillus paracasei* the
463 K562 cell line. Bacteria were cultivated in an oxygen-free environment using a particular
464 medium, and their cells were then exposed to a temperature of 100.0 °C to induce cell death.
465 Lyophilization was used to extract moisture from the dead cells and autoclaving was used to
466 sterilize them thereafter. The next step involved creating several heat-inactivated bacterial
467 suspensions at different concentrations (125.0, 250.0, 500.0, 1000.0 and 2000.0 g/ml). Three
468 distinct time points 24.0, 48.0 and 72.0 hours were used to evaluate the in vitro anticancer
469 qualities of heat-killed bacteria using the MTT test. The results showed that *L. casei* and *L.*
470 *paracasei* heat-killed cells have anticancer properties in K562 cells. Moreover, the anticancer
471 activities of heat-killed cells are closely correlated with their concentration. In summary, heat-

472 inactivated *Lactobacillus* has great potential and should be investigated further in the field of
473 treatments for chronic myeloid leukaemia[64].

474

475 ***3.15 Prevention of colorectal cancer***

476 One of the most prevalent cancers in the world and the primary cause of cancer-related mortality
477 is colorectal cancer (CRC) [129]. The development of colorectal cancer and its incidence are
478 intimately correlated with the gut microbiota. The altered gut microbiota can initiate
479 carcinogenesis through modifications to the immune system, epithelial hemostasis, metabolic
480 profile and activity, DNA damage and aberrant cellular and molecular activities in colonocytes.
481 Numerous unfavourable treatment-associated side effects negatively impact the quality of life.
482 Although multiple cutting-edge medical methods (such as immunotherapy, surgery,
483 chemotherapy, and radiation therapy) are available for the treatment of colorectal cancer, the
484 survival rates are poor [130][131]. So, One well-known bioactive option for treating several
485 illnesses and disorders is probiotics [132][133]. A powerful probiotic strain can help prevent
486 tumorigenesis, including colorectal cancer (CRC), by competing for adhesion sites, producing
487 microbicidal agents like bacteriocin, improving intestinal permeability, releasing bioactive
488 metabolites, regulating immune pathways and stimulating cell protective responses [131].
489 Despite several attempts to elucidate the mechanism behind probiotics' anticarcinogenic
490 properties, a definitive explanation of probiotics' anti-CRC action remains to be discovered
491 [134][135].

492

493 The production of carcinogenic compounds such as cresols, ammonia, phenols, aglycones and N-
494 nitroso compounds, is linked to certain bacterial enzymes, including nitrate reductase,
495 azoreductase, β -glucosidase, β -glucuronidase and 7- α -dehydroxylase. These compounds also
496 induce antiapoptotic pathways, which contribute to the development of colorectal cancer
497 [136][137]. Research demonstrated that adding probiotics to the diet dramatically decreased the

498 activity of bacterial enzymes [138][139][140]. The carcinogenic chemicals are released through
499 faeces after binding with peptidoglycan, which is found in the probiotic microorganisms' cell
500 walls. Certain probiotic strains can metabolize carcinogenic substances, particularly amines and
501 N-nitroso substances. Probiotic supplementation has been shown to reduce the incidence of
502 colorectal cancer (CRC) through a variety of mechanisms, including altered metabolic activity of
503 intestinal microbiota (i.e., decreased endogenous generation of carcinogenic chemicals) and the
504 binding and destruction of carcinogens [141].

505

506 ***3.16 Prevention of liver cancer***

507 The second greatest death rate globally is from primary liver cancer. There are various types of
508 malignant **tumor** in this condition which can include fibrolamellar HCC (FLC), pediatric
509 neoplasm hepatoblastoma, mixed hepatocellular cholangiocarcinoma (HCC-CAA [142]),
510 intrahepatic cholangiocarcinoma (iCAA) and HCC. Of all primary liver malignancies, HCC
511 accounts for 75.0 to 85.0 % of cases [142][143]. The more aggressive proliferative variant of
512 HCC is linked to elevated serum levels of α -fetoprotein (AFP), progenitor cell phenotype
513 expression, mutations in **tumor** protein 53 (TP53) and activation of the transforming growth
514 factor β (TGF- β), hepatocyte growth factor receptor (MET), protein kinase B (AKT), and
515 insulin-like growth factor (IGF) 2 pathways [144]. The prognosis for HCC patients is still not
516 good generally, despite advances in treatment such as immune therapy, multi-kinase inhibitors,
517 and intra-arterial therapy [145]. Pathogenesis of HCC is connected to adverse changes in the gut
518 microbiota because the hepatic portal circulation connects the liver and the intestines directly
519 [146]. According to this anatomical link, probiotic bacteria may be able to reverse the dysbiosis
520 linked with HCC by reestablishing the complexity and colonization resistance of the gut flora
521 [147]. The expression of tight junction proteins is upregulated by probiotic metabolites and
522 probiotic cell components, which improve gut epithelial integrity. The integrity of the gut
523 epithelium can be preserved by the probiotic bacteria's cell surface proteins which can reduce gut

524 epithelial cell inflammation and prevent epithelial cell apoptosis. Probiotics also shield the gut
525 epithelium from harmful bacteria by increasing goblet cells' mucus production and antimicrobial
526 peptides [148]. Probiotic bacteria, particularly *Bifidobacterium spp.* and *Lactobacillus spp.*, can
527 lessen fatty liver, lowering the risk of HCC [149].

528

529 **3.17 Prevention of lung cancer**

530 Lung cancer is the second most frequent type of cancer and the primary cause of cancer-related
531 mortality [150] . According to estimates, smoking is the primary cause of 90.0 % of lung cancer
532 occurrences and other known risk factors include tobacco smoke, air pollution and other toxins.
533 The precise mechanics, nevertheless, are not fully understood. Lungs offer a special chance to be
534 exposed to germs and environmental hazards since they are the mucosa site with the biggest
535 surface area in the body and an important interaction with the outside world. Despite being
536 believed to be sterile, many microorganisms reside in the lungs. Introducing new bacteria, their
537 removal by the body and immune system, and their ability to reproduce in the given environment
538 all influence the prevalence of germs. Long-term obstructive pulmonary disease and cystic
539 fibrosis are two lung conditions that seem to have dysregulated lung microbiome
540 [151][152][153][154]. One research topic is the connection between immunological response to
541 medication and gut flora. Recent research indicates that the gut microbiota's composition may
542 influence how effective immunotherapy is [155]. Probiotics may increase the effectiveness of
543 immunotherapy and help to overcome some of its drawbacks by enhancing gut bacteria diversity
544 and fostering a supportive immunological milieu [7, 154-157].

545

546 **4. Future of probiotics**

547 The future of probiotic research is promising and multifaceted with numerous important areas
548 poised for significant advancement. Though antibiotic resistance is very common nowadays, so
549 probiotics can be replaceable treatments for the future. Identifying and utilizing probiotics from

550 diverse and sustainable sources, including soil and marine environments, can provide new strains
551 with unique benefits. This approach also supports biodiversity and environmental sustainability.
552 Innovations in delivery mechanisms such as encapsulation technologies and smart delivery
553 systems, can improve the survival and efficacy of probiotics as they pass through the digestive
554 system, ensuring they reach their target site in the gut. Advances in genetic engineering could
555 enable the creation of genetically modified probiotics with enhanced functionality such as
556 targeted delivery of therapeutic compounds or improved survival and colonization in the gut.
557 Further research into the precise mechanisms of action of different probiotic strains can provide
558 deeper insights into their role in health and disease, facilitating more targeted and effective
559 applications.

560

561 **5. Conclusion**

562 Probiotics are an exciting and rapidly developing field of study that significantly affects human
563 health. These good bacteria play a major role in preserving and improving the balance of the gut
564 microbiota. They are primarily obtained via fermented foods and nutritional supplements. Their
565 potential for therapeutic use is highlighted by their advantageous activities which include
566 enhancing the intestinal barrier, regulating the immunological response and generating vital
567 metabolites. Probiotics have been shown to help treat various illnesses, including mental health
568 problems, systemic diseases like diabetes and gastrointestinal ailments. Although the efficacy of
569 probiotics varies greatly depending on the strain and the individual's health status, the isolation
570 and characterization of particular probiotic strains have permitted focused therapeutic uses. The
571 growing understanding of the gut microbiome's critical role in general health fuels the field's
572 expansion. More studies are necessary to comprehend and maximize the use of probiotics in
573 clinical settings completely.

574

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584

585 **Declaration of Conflict of Interest**

586 The authors declared that this manuscript could influence no conflict of interest.

587

588 **Data Availability**

589 Data is available on request.

590 **CRedit authorstatement**

591 MTK: Conceptualization, Methodology, Formal analysis, Data curation, Writing- original draft,
592 Funding Acquisition. FH: Data curation, Funding Acquisition. NSH: Writing-Original Draft,
593 Data curation, Funding Acquisition. MSH: Visualization, Funding Acquisition. MAA: Software,
594 Funding Acquisition. SA: Visualization, Funding Acquisition. TNE: Visualization, Funding
595 Acquisition. RTT: Supervision, Investigation, Methodology, Software, Data curation, Writing-
596 Review and Editing, Funding Acquisition.

597

598 **Disclaimer (Artificial intelligence)**

599 Author(s) hereby declare that NO generative AI technologies such as Large Language Models
600 (ChatGPT, COPILOT, etc.) and text-to-image generators have been used during the writing or
601 editing of manuscripts.

602

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