



### Proceedings 1

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# General Perspective and Assessment of the Potential of 2 **Utilizing Paraprobiotics in Food Products<sup>+</sup>**

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Abstract: Paraprobiotics are non-viable microbial cells that, when administered in adequate amounts, confer some health benefits to the consumer. Paraprobiotics can be obtained by subjecting probiotics to physical or chemical treatments and inactivation of the microorganism would lead to the release of some compounds like exopolysaccharides, peptidoglycans, surface proteins, and lipoteichoic acids, all of which have a variety of positive health effects. Paraprobiotics also have numerous technological advantages. Therefore, paraprobiotics are promising components and have a great potential for producing functional food products. However, there are limited studies, most of which concentrate on using paraprobiotics in clinical research and using them directly. The objective of this study is to summarize the way of obtaining paraprobiotics, their health benefits, technological advantages, and their potential for utilization in food products.

Keywords: Paraprobiotic; probiotic; functional food; non-thermal technologies; health benefit

### 1. Introduction

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Functional foods refer to the food conferring health advantages beyond their nutritional value and so the consumers' interest is ever increasing. One of the most important features of functional foods is maintaining the total balance of the intestinal system [1]. The probiotics, live microorganisms which, when administered in adequate amounts confer a health benefit on the host, have often been used to produce functional foods, under the favor of antioxidant, antidiabetic, anti-inflammatory, and hypercholesterolemia effects of probiotics [2, 3]. These beneficial effects of probiotics are ensured by interactions between probiotics and gastrointestinal microbiota and also the immunological system. Probiotics have different properties and effects depending on the strain and administered dose so each strain will exert different health benefits [4]. Different types of microorganisms mainly bacteria acknowledge as probiotics and the probiotic species that appear in the literature the most frequently are Lactobacillus, Bifidobacterium, Streptococcus, Bacillus, and *Enterococcus* [5]. Probiotics are used to relieve the symptoms of high blood pressure, lactose intolerance, diarrhea, irritable bowel syndrome, and obesity [6].

There are various foods that contain probiotics, primarily dairy products, which are a good food matrix for the growth of probiotics. However, there are still many challenges with adding probiotics during food processing, the survival of microorganisms, shelf-life stability, and proper delivery to the gut microbiota [7]. The probiotics will be affected by the food's composition, water activity, antibiotic content, processing conditions of temperature, time, pH, storage conditions like oxygen content, and packaging materials [8]. The term "paraprobiotic" has come into prominence to overcome these problems and in

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light of recent research showing that inactivated probiotic microorganisms may also provide such benefits [9]. There are limited studies, most of which concentrate on using paraprobiotics in clinical research and using them directly. More research is necessary because the paraprobiotics' interactions and mechanisms aren't completely understood. Herein, the way of obtaining paraprobiotics, their health benefits, technological advantages, and their potential for utilization in food products are reviewed.

### 2. Paraprobiotics

Until recent years, the idea and consensus were that probiotics should be alive and present in the final product at a minimum concentration of 10<sup>6</sup>-10<sup>7</sup> CFU/g or CFU/ml until reaching the gut microbiota to confer health benefits [10]. However, this admission begins to change in light of the findings concerning non-viable microorganisms and their health effects [11]. Paraprobiotics are non-viable microbial cells that, when administered in adequate amounts, also confer some health benefits for consumers. Paraprobiotics are also referred to as "inactivated probiotics" and "ghost probiotics" [12].

Microorganism viability is lost when microbial cell structures are altered by mechanical damage to the cell envelope, DNA filament breaks, and cell membrane disruption. Furthermore, enzyme inactivation and changing membrane selectivity are other factors affecting microorganism viability [13].

Paraprobiotics are acknowledged as non-culturable and immunologically active. There are several methods for paraprobiotic assessment, including plating and flow cytometry (FC). Even though the plating is easy to implement, there are some drawbacks, such as the lack of information about metabolic activity and cell integrity of paraprobiotics. FC, on the other hand, is a more advanced method for comprehensive paraprobiotic determination. In this regard, flow cytometry can characterize both cell structure and function in real-time [14]. The best inactivation method can be chosen more precisely and easily based on data collected by FC. Beyond that, FC helps to understand the mechanisms by which the health effects of paraprobiotics come out. Furthermore, FC can be used to track the metabolic activity of the paraprobiotics from processing to the end of the product's shelf life [11]. Additionally, there are alternative methods with high sensitivity that depend on the presence of specific nucleic acids including polymerase chain reaction (PCR) and mass spectrometry. On the other hand, a scanning electron microscope can be used for morphological change detections in inactivated cells [15].

### 3. Inactivation Methods to Produce Paraprobiotics

Paraprobiotics can be obtained by subjecting probiotics to physical or chemical treatments. Physical treatments can also be classified as thermal or non-thermal treatments such as high pressure, sonication, ultraviolet rays, ionizing radiation, irradiation, pulsed electrical field, and supercritical CO<sub>2</sub> [14]. The following changes have taken place: destroying the viability of microorganisms by rupturing and/or damaging cell walls, cell membranes, and DNA, as well as inactivating enzymes, lowering intercellular pH, and denaturing or altering the structure of nucleic acids, proteins, and ribosomes [13].

Probiotics have a strain-specific mechanism of action and efficiency, and various approaches have different effects on biological activities and components. The most convenient inactivation technique should be chosen based on the desired health benefits and the targeted microorganism. The chosen approach should inactivate probiotics while maintaining the beneficial effects of microorganisms. Furthermore, the method has the potential to influence the immunomodulatory activity of paraprobiotics [6, 12]. Nonetheless, the thermal process has been used primarily for cost-effectiveness and provides a wide range of time-temperature combinations. Additionally, a variety of processes may be used in combination to carry out the inactivation [13]. The mechanisms of inactivation methods are listed below for each;

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1 2	<i>i</i> .	<i>Thermal treatments;</i> Cell membrane damage, inactivation of enzymes, protein coagulation, ribosome aggregation, nutrient and ion loss, RNA filaments break
3 4	ii.	<i>High pressure;</i> Cell membrane damage, inactivation of enzymes, protein denaturation, changes in ribosome and nucleoids, solute loss, reduction of pH
5	iii.	Sonication; Cell membrane damage and rupture, DNA damage
6	iv.	Ultraviolet rays; Protein denaturation, formation of DNA photoproducts
7	v.	Ionizing radiation; Nucleic acid damage
8	vi.	Pulsed electrical field; Cell membrane disrupture
9 10 11	vii.	<i>Supercritical CO</i> <sup>2</sup> ; Cell membrane damage, inactivation of enzymes, removal of cell and cell membrane vital constituents, disorder in the intracellular balance of electrolytes, direct effects of CO <sub>2</sub> in the metabolism, reduction of pH
12 13 14	viii.	<i>Ohmic heating</i> ; Cell membrane damage, inactivation of enzymes, protein coagulation, ribosome aggregation, nutrient and ion loss, DNA filaments break, electroporation
15 16	ix.	<i>Dehydration</i> ; Cell membrane damage, changes in the structure of proteins, nucleic acids, and ribosomes
17	х.	pH; Cell membrane damage, chemical changes in fundamental components [6].
18		
19	4. Healt	h Benefits of Paraprobiotics
20	The	e results of studies demonstrated that when a microorganism is inactivated, it
21	releases	compounds like exopolysaccharides, peptidoglycans, surface proteins, and
22	lipoteich	noic acids, all of which have a variety of positive health effects. Even if the cells
23	are dead	d, the metabolites continue secretion and provide health benefits [16]. Thus,
24	parapro	biotics can provide some health benefits to consumers like modulating the
25	immune	system and inhibiting pathogens through adhesion to intestinal cells [17]. Also,
26	paraprol	biotics can help the recovery of intestinal injuries, decrease bacterial translocation
27	and mai	ntain the intestinal barrier, alleviate the symptoms of diarrhea, inflammation, lac-
28	tose into	lerance, respiratory diseases, and liver diseases especially alcohol-induced, lower
29	choleste	rol, treatment of dental caries, atopic dermatitis, colitis, intestinal lesions, visceral
30	pain, an	d prevent the onset of aging manifestations, as well as reduce stress and anxiety
31	[6]. Pha	rmaceutical companies have already begun using paraprobiotics to create
32	pharmac	ceutical products. Two examples are Nyaditum resae® and LacteoI <sup>IM</sup> which are
33 24	produce	a by neat-inactivated strains of <i>Mycobacterium manresensis</i> and <i>Lactobacillus</i> ,
34 25	respectiv	very. Also, CytoFlora® is produced as an immunomodulatory supplement, by
30 26	utilizing	the cell walls of different paraproblotic strains including Lactobacillus,
30 37	Біраовас	vertum, and Streptococcus [11].
3/ 28	Е Та-1	alogical Advantages and the Potential of Utilizing Parametricities in Fac 1
30 20	5. Techn	longical Auvantages and the rolential of Utilizing Parapropiotics in Food
37 40	Froducts	annahiatian hava numanana takanlariarlariarlariartara 1915 - (1919) - (1919)
4U 41		apropionics have numerous technological advantages like stability over a wide
41	DE and f	encoerance range, and no interaction with other components in the food matrix

pH and temperature range, and no interaction with other components in the food matrix, which facilitate easy food processing, industrial usage, commercialization, and extending the shelf life of food [18]. The use of paraprobiotics can be promising especially when the processing and shelf-life conditions deleterious for probiotics. Moreover, antibacterial and

	Probiotic strain	Food matrix	Inactivation method	Results	References	
25	Table 1. Applications of para	aprobiotics in foods				
24		Table1;				
23		years have seen an increase in the use of foods to deliver paraprobiotics, as shown in				
22		Although	n clinical studies on paraprobiot	ics are the main focus of r	esearch, recent	
21						
20		throug	hout shelf-life [6].			
19		• The sta	ability and activity of the parap	robiotics in the food matrix	k should assess	
18		The big	ological effects of paraprobiotics	should be carefully evaluat	ed.	
17		method	d should be determined for relev	ant species.		
16		proved	l or optimized standard metho	dologies. The most adequa	ate inactivation	
15		• Even t	hough various inactivation tech	niques have been applied t	here aren't ap-	
14		exert d	ifferent health benefits.			
13		Probio	tic species and strains should sel	ect carefully due to each sp	ecies and strain	
12		that requires fu	rther investigation. In this insta	nce;	1	
11		fects when cons	sumed directly. In this regard, th	e use of foods as carriers is a	a novel concept	
10		Numerou	is studies have recently shown t	hat paraprobiotics have pos	sitive health ef-	
9		metabolic by-p	roducts [12, 19].		-	
8		are highly likel	y to provide not only live bacteri	a but also inactivated micro	organisms and	
7		levels of probio	otics that are stated [13]. For this re	eason, the health benefits of	probiotic foods	
6		to some report	s, some commercially produced	probiotic foods don't actua	ally contain the	
5		otics, of which	the exact numbers of live and d	ead bacteria are unknown	[19]. According	
4		with shelf-life.	Herewith, foods with probiotic	bacteria contain both live a	nd dead probi-	
3		probiotic bacte	ria become inactive during pro	cessing and this loss ever i	increases along	
2		rently [12]. On	the other hand, even if the prec	autionary conditions are m	et, some of the	
1		antifungal ager	nts have no effect on paraprobio	tic bioactivity, so they can b	e used concur-	

Probiotic strain	Food matrix	Inactivation method	Results	References
			Viscosity↑, WHC↑, Syneresis↓, Storage	
		Heat treatment	modulus↓, Loss modulus↓, Stress	
L. acidophilus and B. lactis	Yogurt	(121°C, 15 min)	crossover point↓, Loss tangent↓, Sen-	[20, 21, 22]
			sory properties $\uparrow$ , L* $\leftrightarrow$ , a* $\leftrightarrow$ , b* $\leftrightarrow$ , pH $\downarrow$ ,	
			Acidity↑, Redox potential↑	
	Whey-grape juice	Ohmic heating (8V/cm, 95°C/7min, 60 Hz)	Glucose rate↑, Maximum glucose	
			value↔, Glucose incremental percent-	
L. casei subsp. paracasei 01			age↔, Peak blood glucose time↔, Gly-	[23]
			cemic responses (AUC, AIg, PGV, HP,	
			GB)↔, Glucose postprandial level↓	

26 ↑indicates increment is statistically different; ↓indicates decreasement is statistically different; ↔ indicates increment or decreasement
 27 is not statistically different.

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29 Paraprobiotic yogurt produced with Lactobacillus acidophilus ATCC SD 5221 and 30 Bifidobacterium lactis BB-12 and compared to probiotic yogurt in terms of physicochemical, 31 microstructural, biochemical, rheological, microbiological, and sensory properties during 32 28 days of storage. The results demonstrated that paraprobiotics increased water holding 33 capacity and apparent viscosity while decreasing syneresis due to producing of exopoly-34 saccharides by heat-killed cells. Another crucial point is that the viability of starter cul-35 tures is enhanced by providing nutrients released from the paraprobiotics. Moreover, 36 paraprobiotics affected the rheological characteristics and sensory properties of yogurt. 37 To sum up, the incorporation of paraprobiotics into a food product is a promising alter-38 native to probiotics with some technological and health advantages like no interaction with other ingredients in the food matrix, enhanced shelf-life, resistance to environmental 39 40 changes, and ease of processing and commercialization. Nevertheless, more research is

1 required to validate the findings using different paraprobiotics and food products [20, 21, 2 22]. 3 Lacticaseibacillus casei subsp. casei 01 was inactivated by using ohmic heating and produced a paraprobiotic whey-grape juice drink. According to the results, the paraprobiotic 4 drink displayed reduced glucose postprandial levels like the control probiotic drink. Even 5 6 though more research is needed, it can be claimed that paraprobiotics can be effective in 7 lowering postprandial glycemia [23]. 8 9 6. Conclusions 10 To summarize, the paraprobiotic term has been evolving and gaining attention in recent years. Clinical studies demonstrated that paraprobiotics have health benefits for the 11 12 consumers like probiotics. Therefore, paraprobiotics can be an alternative to probiotics for 13 people with a sensitive immune system or who are immunocompromised/immunodefi-14 cient and avoid probiotic consumption. Also, paraprobiotics can be used when the use of 15 probiotics is a technological challenge. In the case of processing and shelf-life conditions, 16 not convenient probiotics survive. However, there isn't enough information and research 17 in the literature about paraprobiotics just yet. The mechanism of action of paraprobiotics 18 is not fully understood and requires further investigation. In this regard, the following 19 studies should focus on determining valid conditions for emerging inactivation methods, 20 the biological activities and stability of paraprobiotics in vitro and in vivo, and the terms 21 for wide application and easy commercialization of paraprobiotics. Furthermore, it's criti-22 cal to establish a precise definition by subject-matter experts and prevent the misuse of 23 paraprobiotics. 24 25 Author Contributions: Conceptualization, T.T. and GN.Y.; investigation, T.T. and GN.Y; resources, 26

T.T. and GN.Y; writing—original draft preparation, T.T.; writing—review and editing, GN.Y; visualization, M.G.; supervision, M.G.; project administration, M.G. All authors have read and agreed to the published version of the manuscript.

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