Isolation of Bacteriocin from Lactobacillus Species and its Application as a Preservative in Dairy Products

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ABSTRACT

The supernatant fluid tested inhibitory assignment no longer most powerful in opposition to some lactic acid microorganism but additionally, in opposition to some pathogenic and meals-spoilage species, at the side of Clostridium, Listeria and Enterococcus. It used to be as soon as purified to homogeneity by the use of a single four-step process: a crude supernatant fluid purchased from early stationary-part subculture in MRS medium was once as soon as subjected to ammonium sulphate fractionation, CM-Sephadex cation-exchange chromatography, Phenyl-Sepharose hydrophobic chromatography and reverse-phase HPLC chromatography. The bacteriocin used to be produced constitutively at some factor of exponential development. It used to be as soon as bactericidal to sensitive cells and the bactericidal effect used to be now not produced with the help of cell lysis. The amino acid composition of the bacteriocin was once determined and no modified amino acid was once as soon as located among the residues recognized.

INTRODUCTION

Lactic acid microorganism (LAB) are Gram-optimistic, non-spore forming, catalase-horrible microorganism which are devoid of cytochromes and are of non-aerobic nevertheless are aero-tolerant, fastidious, acid tolerant and strictly fermentative; lactic acid is the important end-manufactured from sugar fermentation. Lactic acid microorganism (LAB) includes a range of bacterial customary within the Phylum fumicutes. The general Enterococcus, Lactobacillus, Lactococcus, Leuconostoc, Milissoccocus, Oenococcus, Pediococcus, Streptococcus, Tetragenococcus, Vagococcus and Weissella are famous as lactic acid micro-organism. They may be mostly normally essentially the most generally used bacteria as starter cultures for the trade processing of fermented dairy, meat, vegetable and cereal merchandise. Reduction of pH and conversion of sugars to healthful acids is the principal maintaining moves that these microorganism furnish to fermented meals [1-10].

These ordinary isolates of lactic acid microorganism from spontaneous fermentations can be used as special starter cultures or as adjunct traces, after phenotypic and genotypic characterization, they most commonly signify a potential supply of in general new antimicrobial metabolites. Additionally, the applying of lactic acid microorganism and their antimicrobial metabolites within the prevention of food spoilage and the extension of the shelf lifetime of meals that's equipped to eat, modern-day-tasting, nutrient and diet rich, minimally processed and bio preserved are the foremost challenges for the present meals organization [10-20]. The utilization of bacteriocin-producing lactic acid bacteria as defending traces or bacteriocins in variety of purified or centered compounds as bio-preservatives to manage undesirable microorganism remains a predominant core of recognition of researches involving ingredients defend and satisfactory [21-30].

The antimicrobial undertaking of starter cultures and probiotic microorganism has been attributed to the production of metabolites and same to natural and organic acids (lactic and acetic acid), hydrogen peroxide,
ethanol, diacetyl, acetaldehyde, other low molecular mass compounds with antimicrobial recreation and bacteriocins [31-40].

**REVIEW OF LITERATURE**

First of all, lots of the great development in bacteriocin be trained stemmed from investigations of the colicins, the prototype bacteriocins produced via slightly countless members of the adored ones Enterobacteriaceae, and this resulted in large expertise of the genetic groundwork, self-discipline constitution, mode of formation, and killing action of those molecules. Nonetheless, there has now been an potent quantity of be trained assignment based upon the bacteriocin-like ambitions of Gram-constructive microorganism, notably lactic acid microorganism (LAB) [41-50].

The important clear documentation of the personality of an antibiotic agent produced by way of E.Coli was once provided via utilizing making use of Gratia, who validated in 1925 that pressure V (virulent in experimental infections), produced in liquid media, a dialyzable and warmness-steady substance (later referred to as colicin V) that inhibited the development of E. Coli even with a immoderate dilution. Later a series of individual colicins had been discovered over an interval of time [51-53].

The extra natural time period “bacteriocin” used to be coined by way of Jacob et al. Bacteriocins had been principally outlined as protein inhibitors or ‘antibiotics’ of the colicin variety, i.e., molecules characterized by means of lethality after biosynthesis, predominant intraspecies killing venture, and adsorption to distinctive receptors on the dermis of bacteriocin sensitive cells [54].

An analysis of the bacteriocins of Gram-positive microorganism opened with the comment that a lot of the definitive investigations inside the strength of mind of bacteriocins had headquartered on these of Gram-terrible microorganism nonetheless expected in be trained emphasis on bacteriocins of Gram-optimistic lactic acid microorganism [55-60]. Apparently a lot of the renewed curiosity in these add-ons is an instantaneous response to the perceived expertise beneficial application of these shops, every for the renovation of meals or the prevention and therapy of bacterial infections.

**Components protection**

Despite the fact that nisin is the one bacteriocin in america licensed as a compatible away meals additive, there could also be an quality deal of curiosity in exclusive bacteriocins which have equal residences and show off colossal spectrum inhibitory activity. Bacteriocins produced via fermentation can be purified and brought to foods as pure chemical components to inhibit meals pathogens and spoilage organisms simplest after obtaining approval as an immediate constituents additive with the support of the FDA [61-63]. Many bacteriocins can resist excessive temperature utilized in meals processing and might keep clever over a large pH type. Bacteriocins can also be digested through many enzymes inside the human gastrointestinal tract equal to exceptional proteins inside the healthful eating regimen and not emerge as an trouble for worthy gut microflora. Bacteriocins are non-poisonous, odorless, colorless, and tasteless.

**Bactericidal action**

The low-molecular weight bacteriocins of Gram-confident microorganism traditionally appear to be membrane lively. The lantibiotic subgroup of bacteriocins tends to fluctuate from the reverse companies within the voltage dependence of their membrane insertion. Poration complexes had been proposed to be fashioned between one, two, or virtually obviously much more species of amphipathic peptides, major to ion leakage, lack of proton intent force, and eventually cellphone death [64-66].

**Antimicrobial spectrum**

The low molecular weight bacteriocins of Gram-confident microorganism exhibit bactericidal pastime which is directed basically towards exclusive precise Gram-positive microorganism. For illustration, the lantibiotic nisin has been established to be amazing in opposition to many traces of Gram-constructive microorganism, along with staphylococci, streptococci, bacilli, clostridia, and mycobacteria [50].
Detection of bacteriocins produced through lactic acid microorganism (LAB) typical approaches

Agar diffusion system

A original approach for screening bacteriocin mission entails utilizing agar media contained in petri plates. Piddock has reviewed a wide variety of those approaches for detection and dimension of bacteriocin mission. There are really quite a lot of models, most of them derivatives of the “spot-on-outside” procedure, to be equipped to contain an agar overlay [67-70]. On this approach, the bacteriocin from the producer subculture is noticed on the indicator cultures and these plates are incubated even as earlier than examination for zones of inhibition during the growth of producing cultures. Kekessy and Piguet described a system in which the manufacturing and indicator traces had been each and every grown on unique gold common media. The manufacturing tradition used to be spot-inoculated onto a variety plate, and after growth, the agar mass used to be once aseptically dislodged with a spatula from the petri dish bottom and transferred to the lid of the dish [71]. A gentle agar overlay seeded with the indicator was once once then poured over the inverted agar. Following re-incubation, bacteriocin-positive cultures displayed a halo of clearing within the backyard across the average button of progress. This assay minimizes the penalties of acids and bacteriophages, considering that the bacteriocin-producing and indicator strains are bodily separated through an agar layer [72,73].

Mass spectrometry

Mass spectrometry has been tailored for the quick detection of pediocin, nisin, brochocins A and B, and enterocins A and B from subculture supernatants by the use of Rose et al. The system is referred to as matrix-assisted laser desorption/ionization time-of-flight mass spectroscopy (MALDI-TOF MS) and was as quickly as on the devised for the examination of colossal molecules, similar to biopolymers [74-76].

Flow cytometry

In a float cellphone, Mugochi et al. Developed a fast and sensitive system for detection of bacteriocins in fermentation broth. Low concentrations of potassium ions had been measured, so that launched potassium ions from a bacteriocin-touchy indicator stress instantly correlated to concentrations of crude bacteriocin present in fermentation broth injected into the cellular. This system in evaluation excellent to an usual agar exceptional diffusion assay [77,78].

PCR detection systems

The PCR method has been used to rapidly check lactobacilli that may produce just right-characterised bacteriocins, on the other hand of counting on the utilization of problematic biochemical methods in order to even be more almost always than no longer required for the identification and characterization of such bacteriocins. PCR programs had been used to observe genes in fee for bacteriocin development and regulation in bacterial cultures. Rodriguez et al. demonstrated the amplification of a seventy 5 bp gene fragment of the lactocin S structural gene in seven bacteriocinogenic traces of lactobacilli remotened from fermented sausages [79]. In however one different work, Garde et al. detected the genes important for the synthesis of lactocin 481 and nisin utilizing PCR methods with detailed probes on an isolate of L. Lactis sub-sp. Lactis. To be able to decrease the time priceless for detection, procedures to simplify or even cast off the necessity for put up-amplification gel assays were furnished. A method for PCR quantification has been devised and is by and large known as “genuine-time” PCR when you consider that use of a fluorescent label makes it possible for the character to close to view the expand within the large style of motive DNA as it can be amplified [80-84]. The precise-time PCR process is based on detection and quantification using probably the most a quantity of kinds of fluorescent reporter molecules. This signal raises in direct percentage to the quantity of PCR product inside the response. A thermocycler organized with a fluorescence detection system measures the fluorescence in peculiarly designed tubes that include the response add-ons. The amplification final result in an attribute sigmoid formed curve which represents three phases of PCR: the lag part (little product accumulation), the exponential part (fast product accumulation) and the plateau section (no additional product is amplified) [85-89]. With the help of recording the amount of fluorescence emission in each and every cycle, it can be viable to detect the PCR for the duration of the exponential segment the area the primary large develop inside the fluorescence instantly correlates with the preliminary number of goal template and the PCR cycle at which this fluorescence is measured is most commonly called a threshold cycle (Ct) [90].

Components purposes

Three procedures are mostly used in the utility of bacteriocins for biopreservation of foods:
(1) Inoculation of food with LAB that produce bacteriocin within the merchandise. The competencies of the LAB to support and produce bacteriocin within the merchandise is essential for its robust use.
(2) Addition of purified or semi-purified bacteriocins as food preservatives.
(3) Use of a product beforehand fermented with a bacteriocin-producing stress as an ingredient in ingredients processing.

**Biopreservation of dairy merchandise**

L. Monocytogenes has been the documented cause of an enormous form of outbreaks involving dairy merchandise, corresponding to pasteurized milk and cheese. Nisin has been demonstrated mighty inside the direction of L. Monocytogenes in dairy merchandise [91-94]. A important trouble in cheese creation is the Clostridium-associated butyric acid fermentation. Nisin is traditionally dropped at pasteurized, processed cheese spreads to prevent the outgrowth of clostridial spores, similar to Clostridium tyrobutyricum [95,96]. Lacticin 3147, a giant-spectrum, two-part bacteriocin produced by way of Lactis subsp. Lactis DPC 3147, is used to control cheddar cheese satisfactory by way of lowering nonstarter LAB populations in the course of ripening. The lacticin 3147 producing transconjugant has moreover been used as a protecting culture to inhibit Listeria on the epidermis of a mildew-ripened cheese [97-100]. Aside from using nisin, a bacteriocin (pediocin) produced via Pediococcus acidilactici %. Zero has tested to inhibit Gram-confident spoilage organisms like lactobacilli in salad dressings.

**CONCLUSION**

Bacteriocins are protein-like antibiotics. The macromolecular bacteriocins exert their action by way of adsorption to specific receptors located on the outside surface of the sensitive pressure, followed by way of foremost organic and morphological alterations of both the bacterial telephone and the bacteriocin particle. This results within the killing of the sensitive strain without creation of extra bacteriocin particles, one of the vital foremost aspects which make them evidently exclusive to bacteriophages.

Essentially the most intensively studied companies of bacteriocin are the colicins that are produced via E.Coli and carefully related enterobacteria. They are most commonly plasmid-encoded proteins. Although many bacteriocins had been remoted and characterized, only some have confirmed industrial talents in meals application. Nisin is the only purified bacteriocin authorized for meals use in the US. It has been used as a meal preservative in more than 50 international locations, as a rule in cheese, canned vegetables, more than a few pasteurized dairy, liquid egg merchandise, and salad dressings.

Bacteriocins that have been used as meals preservatives have slightly slender endeavor spectra and are customarily new not active in opposition to Gram-bad bacteria. The simultaneous software of bacteriocins and non-thermal processing applied sciences, such as HP and PEF, to reinforce shelf life of foods is attractive since foods produced utilizing these non-thermal technologies most of the time have higher sensory and dietary qualities in comparison with merchandise produced making use of traditional thermal processing system.

**REFERENCES**


