MICROORGANISMS FOUND IN FAST AND TRADITIONAL FAST FOODS

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By S. M. H. Easa

Microbiology Department, Faculty of Science, Ain Shams University, Cairo, Egypt

ABSTRACT

Sixty food samples were collected from 60 random restaurants of fast and traditional foods in El Qassium, Saudi Arabia and investigated for the presence of bacteria using different temperatures (10°C, 20°C, 30°C, 40°C and 50°C) incubated for 24-48 hours and analyzed for fungi and yeasts incubated at 25°C. The results revealed that from 45 samples of traditional foods, twenty two species of eighteen genera of bacteria were found plus fourteen species of twelve genera of fungi and three species of three genera of yeasts. While fast food results revealed that from 15 fast food samples collected from 15 restaurants a total of ten species of ten genera of bacteria and eight species of seven genera of fungi were recorded. The species of bacteria isolated in this study are, Acetobacter sp., Achromobacter sp., Bacillus coagulans, B. subtilis, Clostridium perfringens, Erwinia carotovora, Escherichia coli, Flavobacterium sp., Klebsiella pneumoniae, Lactobacillus plantarum, Leuconostoc mesenteroides, Listeria monocytogenes, Microbacterium lacticum, Micrococcus sp., Pseudomonas aeruginosa, Pseudomonas fluorescens, Pseudomonas putrefaciens, Salmonella sp., Staphylococcus aureus, Streptococcus lactis, Streptococcus thermophilus, Campylobacter jejuni, Citrobacter fruendii, Proteus vulgaris and Yersinia sp. The occurrence of some of these bacteria illustrates that fast foods in these restaurants may act as a reservoir of pathogenic bacteria for human. Fungi isolated are Aspergillus glaucus, A. niger, Alternaria sp., Chaetomium candidum, Cladosporium herbarum, Fusarium sp., Monilia sp., Mucor rouxii, Neuropora sp., Penicillium expansum, Penicillium sp., Rhizopus nigricans, Sporotrichum carnis and Thamnidium elegans. Yeasts were represented by **Torulopsis** caroliniana, Saccharomyces rouxii and Zygosaccharomyces sp. Total viable count of bacteria (CFU) was higher in foods containing vegetable salad and fresh vegetables than heated foods (e.g., chicken Shawirma, Beef burger). Some bacteria resist heat and grow at 50°C. Contamination occurred through raw foods, use of polluted irrigation waters, human handling and the use of contaminated containers. The binge-eating of fast food can lead to measurable signs of liver injury, inflammation and inexpensive fat-and calorie-packed foods make us the fat. Food poisoning can be controlled by the adjustment of pH, water activity, temperature control. Prevention of toxins in fast foods must become a cooperative effort on the part of all involved in food production. Prevent multiply the microorganisms by washing and dry hands before preparing any foods and after handling raw foods (meat, poultry, vegetables or fruits), food preparation areas, equipment must be cleaned, kitchen areas, restaurants and foods protected from insects, pests and other animals. Patients should not handle foods in restaurants.

Key words: bacteria, contamination, fast food, fungi, poisoning food, temperatures, traditional fast food, yeasts.

1. INTRODUCTION

Food is a chemically complex matrix, and predicting whether, or how fast, microorganisms will grow in any given food is difficult. Most foods contain sufficient nutrients to support microbial growth. Several factors encourage, prevent, or limit the growth of microorganisms in foods, the most important are water availability, pH, and temperature (Makukutu and Guthrie, 1986; Smith and Fratamico, 1995 and ICMSF, 1996). The busy and hectic life schedule has opened the way for the fast food industry in most parts of the world. The traditional or conventional way of cooking is over and the fast food joints are visible everywhere. Fast food does not only include the traditional fast food items like pizza, burger or French fries but it also includes Chinese as well as Indian. The most typical fast food meals eaten in Germany are similar to those eaten in America namely burger, pizza and fries. Other popular meals are (a sliced sausage with ketchup and mayonnaise), Kebab (the meat is served in flatbread along with lettuce, onion, cucumber, tomatoes) (Ockerman and Stec, 1980; El-Sherif *et al.*, 1991). Although fast food restaurants are often viewed as a representation of modern technology, the concept of "ready-cooked food to go" is as old as cities themselves; unique variations are historical in various cultures. Ancient Roman cities had bread-and-olive stands, flat bread and falafel are ubiquitous in the Middle East. Food habits, pattern and behavior vary widely from culture to culture. Popular Saudi Arabian traditional foods include meat, rice, wheat, vegetables and spices that give these recipes a special flavor.

There are many popular foods in Saudi Arabia like Jarish, Qursan, Saliq, Masapep, Keshta, Mataziz, Freek, Hunayni and Harisah.

In recent years just about all the quick service restaurants have added salad fresh vegetables (Lettuce, Cabbage, Carrot, Cucumber, Onion, ketchup, maymonise). Some foods are cooked prior to consumption others are eaten raw. Products that might be classed with both fresh and processed vegetables are the chopped salad ingredients sold in the grocery store and to the institutional trade. Although essentially fresh, contamination during processing, and changes in microbial growth patterns during storage, may later lead to microflora of these foods quantitatively and qualitatively. The inner tissues of healthy plants and animals are free of microorganisms. However, the surfaces of raw vegetables and meats are contaminated with a variety of microorganisms and this depends on the condition of the raw product, the method of handling, and the time and conditions of storage (Woodward , 1996; Odumeru et al., 1997 and Pelczar et al., 2006).

Microbial food safety is an increasing public health concern worldwide. It is estimated that each year in the United States there are approximately 76 million food borne illnesses. Cases are caused by Campylobacter species, nontyphoidal Salmonella, pathogenic Escherichia coli and all colonize the gastrointestinal tracts of a wide range of wild and domestic animals, especially animals raised for human consumption (Meng and Doyle, 1998). Food contamination with these pathogens can occur at multiple steps along the food chain, including production, processing, distribution, retail marketing and handling or preparation. Numerous epidemiological reports have implicated foods of animal origin as the major vehicles associated with illnesses caused by food-borne pathogens (Petersen and James, 1998).

Contaminated raw or undercooked poultry and

red meats are particularly important in transmitting food borne pathogens. Other sources of human infections include contaminants produced by contact with farm animals and pets. Person-to person transmission has also been described. These microorganisms are carried on hands, wiping cloths and utensils, especially choping boards. The slightest contact can transfer them to food and cause food borne diseases. Examples of zoonotic pathogens that may be transmitted in this way include *Salmonella*, *Campylobacter, Escherichia coli* and eggs of the tape worm, *Taenia solium* (Meng and Doyle, 1998).

The gut is the most important source of bacteria, contributing *Clostridium perfringens*, coliforms, *Salmonella* and *Staphylococcus* to the meat surface. Mesophiles, including pathogens, cannot grow on chilled carcasses, but psychrotrophs of the *Pseudomonas, Achromobacter* grow readily, and eventually spoil the meat (Meng and Doyle, 1998). The conditions in a well wrapped piece of meat encourage the growth of the *Lactobacillus* at the expese of the *Pseudomonas, Achromobacter* group (Petersen and James, 1998).

Pathogenic microorganisms on raw vegetables and fruits suggested that the use of poor quality water for irrigation could increase the incidence of pathogens (E. coli)., enteric Erwinia spp., Aeromonas, Serratia spp. and some gram negative bacteria (Pseudomonas spp., Citroacter freundii) and Clostridium and Xanthomonas. Staphylococcus aureus. Corynebacterium, Listeria spp., Lactobacillus spp., Streptococcus and spp. Micrococcus spp., as gram positive bacteria. The consumption of fast foods, raw milk and raw milk products have been reported to be associated with serious health problems (De Boer and Hahne, 1990; Pacini et al., 1997; Food and Drug Administration (FDA), 2000, 2010 and Pelczar et al., 2006).

Microorganisms in fast and traditional foods are responsible for many human diseases. *e.g.*, *Salmonella* bacteria are common cause of food borne illness, particularly in undercooked chicken and chicken eggs (Woodward, 1996; Kaneko *et al.*, 1999 a; Uyttendaele *et al.*, 1999 and Angelillo *et al.*, 2000).

However, some studies suggested the incidence of *Listeria* sp. in retail foods, ready-cooked chicken, on the hands of food workers, food stuffs, human faeces, sewage and soil from urban sources (MacGowan *et al.*, 1994).

Furthermore, it was reported the prevalence of *Campylobacter* spp., *Staphylococcus* spp., *Escherichia coli, Salmonella* spp., *Yersinia* spp. and *Listeria* on meat, sea foods, vegetable ingredients, chicken shawirmas, raw and cooked foods, raw

chicken, beef burger sandwiches, ready-to eat salad vegetables, commercial Mayonnaise, frozen chicken, poultry products and on the hands of food workers (Kaneko *et al.*, 1999b and Pelczar *et al.*, 2006).

Uzeh *et al.* (2009) reported that microorganisms isolated from salad containing raw vegetables include *Mucor* sp., *Aspergillus fumigatus*, *Trichoderma, Neurospora crassa and Aspergillus niger*.

It was investigated that fast food contains high levels of refined sugar, white flour, trans fat and polyunsaturated fat, salts and numerous food additives, at the same time it is lacking in proteins, vitamins and fibers. Consumption of fast food in the world has been associated with obesity leading to many diseases (Canadian Institute for Health Information, 2007).

Raw materials, including water, ice and milk, may contaminated with dangerous be microorganisms. Toxic chemicals may be formed in damaged and mouldy foods. Care in selection of raw materials and simple measures such as washing and peeling, may reduce the risk. Contaminated water, for example, has been associated with outbreaks of Salmonella, Campylobacter and Escherichia coli, whilst infections with Salmonella, Campylobacter, Mycobacterium (TB), Brucella and Escherichia coli can be acquired through the consumption of contaminated milk or dairy products that are not pasteurized (Meng and Doyle, 1998). When raw milk is left standing for a while, it turns "sour" this is the result of fermentation, where lactic acid bacteria ferment the lactose inside the milk into lactic acid. Prolonged fermentation may render the milk unpleasant to consume. This fermentation process is exploited by the introduction of bacterial cultures Lactobacillus sp., Streptococcus sp., (e.g., Leuconostoc sp. ... etc) to produce a variety of fermented milk products. The reduced pH from lactic acid accumulation denatures protein and caused the milk to undergo a variety of different transformations in appearance and texture (Deak and Beuchat, 1996). Fast foods are sold in a restaurant or store with low quality preparation and served to the customer in a packaged form for take out/take away. In most fast food operations, menu items are generally made from processed ingredients prepared at a central supply facility and then shipped to individual outlets where they are reheated, cooked (usually by microwave or deep frying) or assembled in a short amount of time. Fast foods are often very high in calories, saturated fat and sodium that can make us fat, clog our arteries and send our blood pressure soaring. Food substances that have been

prepared by a fermentative process, or have been exposed to microbial contamination during aging or storage, are likely to contain amines. Alcoholic beverages such as beers can contain biogenic amines, as do some other fermented foods such as sauerkraut and soy bean products (Sanchez, 2009). Amines are also considered as endogenous plant substances that are commonly used for food, where some vegetables and fruits were found to contain high concentrations of various amines. The biogenic amine contents of various foods and feed have been widely studied and found in cheese, fish, meat products, eggs and mushrooms. Biogenic amines may also be considered as carcinogens because of their ability to react with nitrites to form potentially carcinogenic nitrosamines. The toxicity of biogenic amines to chicks in terms of loss of weight and mortality was also reported (Heaton and Jones, 2008).

Meat, produce and soft cheeses (*e.g.*, brie cheese, cottage cheese) have more water content, allowing any bacteria, viruses or molds present to multiply quickly (Bichai *et al.*, 2008).

It was suggested that food must be cooked thoroughly to the correct temperature because proper cooking kills almost all dangerous microorganisms. Studies have shown that cooking food to a temperature of 70°C can help ensure that it is safe for consumption, for example poultry, minced meat products (e.g., hamburger and sausages). Ideally, the center of the food should reach a temperature of 70°C for at least two minutes (Makukutu and Guthrie, 1986). Chilled, ready to eat foods must be kept to temperature below 5°C. Hot foods must be kept at temperatures above 60°C before serving. Cool rapidly and refrigerate left over foods if they are not to be used within 2 hours. Food should be cold before placing in the refrigerator since it may take a while to cool off in the refrigerator and hot food may warm up other foods (Kaneko et al., 1999 b). Some of the highest aerobic counts have been reported for tubers and other vegetables that are in contact with the soil *e.g.*, lettuce, carrots, potatoes, cabbage and flafel with fresh vegetable salads record the highest count of microorganism (Kaneko et al., 1999 a). The high acid and sugar content of fruits often permits yeasts and molds to predominate, while the high carbohydrate content of many vegetables favors the lactic acid bacteria. It was reported that the major source of the organisms on frozen vegetables is contaminated equipment; such as chopers, slicers, conveyor and inspection belts, and filling machines. Some of these units possess surfaces that are difficult to reach for proper

cleaning. Belts may present problems because of the tenacity with which organisms adhere to certain surfaces, and because some fabrics absorb moisture and thus permit a microbial build up within the belt interior. Many organisms do not survive in a low pH environment; for example, *Salmonella and Shigella* die off rapidly in citrus juices (Norberg, 1981).

Contamination of the egg shells occurs after laying nesting material, dirt, and fecal matter. It was reported that the flora of egg shells are dominated by gram positive cocci while the gram negative rods are present in numbers. The eggs may contain organisms from the shell surface by improper washing and storage methods. The most common genera of bacteria found in liquid eggs are gram negative types, including Pseudomonas, Proteus and Escherichia. In commercial egg breaking operations the egg shell is a source of contamination and may contribute large numbers of gram positive cocci to the liquid egg (Administration Urged to Boost Food Safety, 2009). Colonization of the shell contents is characterized by a mixed flora of gram negative bacteria. The most common contaminants are the coliforms, Achromobacter, Pseudomonas, Serratia, Proteus, Alcaligenes and Citrobacter (Finegold and Martin, 1982; Uyttendaele et al., 1999). The major pathogen associated with eggs and egg products is Salmonella.

Some meat products such as flame seared beef patties and cooked beef are processed at lower temperatures. These temperatures are sufficient to destroy pathogens, but the final bacterial counts include some of the more heat resistant vegetative bacteria such as the enterococci. Unless cooked products are packaged hot and immediately frozen, recontamination invariably occurs from equipment, food handlers, raw products or dust (Angelillo *et al.*, 2000).

Human health and mycotoxicoses: It was reported that the ubiquitous fungal strains involved could utilize wide variety of foodstuffs for toxin production. Several mycotoxins have been verified as naturally occurring foods and feeds (Barnett et al., 2000). Most mycotoxicoses of man or animals have been recognized by observation of the toxicity of moldy foods and feeds. The toxigenic storage fungi are primarily Aspergillus, Penicillium, while some like Fusarium may be either field or storage organisms. The mycotoxins presently considered to present the most potential for human health hazard are the toxins of the storage fungi in the genera, Aspergillus, Penicillium and Fusarium are those elaborating mycotoxins which are more important in foods and feeds.

Some of the mycotoxin findings reported represent

extensive survey, others are very limited. Fungi, such as the moulds commonly seen on bread, can also cause illness while viruses such as hepatitis may also be food borne (Wart, 1989). It was reported that when foods such as meat, spaghetti sauce or vegetables are canned, the oxygen can not get in therefore growth of aerobic organisms is controlled and the food is preserved. Some microorganisms will grow only in anaerobic conditions. Botulism is a rare type of food borne illness caused by microorganism that prefers anaerobic conditions. Home canned food that have not been preserved properly are the most common source of this type of food borne illness (Kao and Shih, 1993 and Pacini *et al.*, 1997).

The objectives of this study were to determine the presence of pathogenic bacteria, fungi and yeasts in fast and traditional foods in some restaurants causing human diseases, as well as to investigate the association of microbial contamination with component, type, temperature, season of foods. This study was carried out to give information about the methods of prevention of diseases due to food borne pathogens and how to control it.

2. MATERIALS AND METHODS

Sixty restaurants were tested in the present study. Fast and traditional food samples were obtained from fast and traditional fast food restaurants in El Qassim, Saudi Arabia in the Summer (2007). The Centers for Disease Control Food Borne Diseases Active Surveillance Network (Food net) data indicate that outbreaks and clusters of food-borne infections peak occur during the warmest months of the year (Centers for Disease Control and Prevention, 2001).

2.1. Samples of foods: A) Traditional foods: The traditional fast foods which were used in the research: 1. Jarish: composed of animal fats, crushed wheat, oil, salts and apices. Jarish may be simply boiled and served with a topping of chopped hot pepper and onion or it may be browned in butter or oil and then cooked into a sort of pilaf with a chunks of meat, chopped onion and tomato for the richly flavored dish. 2. Mataziz: composed of flour, meat, cucurbita, onion, oil, spices, tomato and salt. 3. Qursan: included meat, oil, different vegetables onion, tomato, limon and salt. 4. Keshta: included of palm, butter and flour. 5. Mathbib: contained eggs, oil, flour, salt and sugar. 6. Freek: composed of eggs, flour, sugar, small amount of salt. 7. Hunayni: included palm, bread, butter, spices and water. 8. Saliq: contains chicken, rice, milk and spices. The rice first half cooked in meat or chicken both and then with milk for one hour.

9. *Harees:* composed of meat, sugar, butter, wheat, salt and cinnamum zeylani.

B) Fast foods: 1. Chicken shawrmas with salads. 2. Hamburger (beef burger) with salads. 3. Flafel with salads (lettuce, tomato, onion, cucumber).

2.1.1. Collection of samples: Sixty samples were obtained from 60 restaurants at a temperature 35° - 37° C in the summer(2007). The food samples were taken from restaurant in sterile plastic bags in Ice-Box, according to Cheesbrough (1984).

2. 1.2. Preparation of food samples: From each sample 25 g was aseptically weighed and macerated and 225 mls of sterile distilled water was added. Serial dilution was carried out using sterile distilled water as diluents. From each dilution 1 ml was plated using the pour plate methods of Swanson *et al.* (1992).

2.1.3. Isolation of Microorganisms

A) **Isolation of Bacteria:** Samples were cultivated on different media. The inoculated media were cultured at different temperatures. Pure cultures of the microorganisms were identified using the standard procedures of Barrow and Feltham (1993). The tests employed for the identification of isolates were the Gram stain, biochemical test, pigments and colony morphology.

The streak plate method was used for the recovery of the various bacteria species. The Total Viable Count (TVC) of bacteria species was done on different media. The inoculated plates of bacteria were incubated at different temperatures (10°C, 20°C, 30°C, 40°C and 50°C) for 24-48 hours. The colony forming units (CFU) were counted with a Gallenkamp colony counter, the result reported as (CFU) per ml of sample. The same process was repeated in respect of fungi and yeasts, which were incubated at (25-30°C).

2.2. The culture media

2.2.1.Nutrient agar medium: was used for total bacterial count (enumeration of bacteria). The medium contained 3.0 g beef extract and 5.0 g peptone (per liter of distilled water) according to Atlas (1993) and Swanson *et al.* (1992).

2.2.2.MacConkey agar medium: it is a differential and low selectivity medium used to distinguish lactose fermenting (*e.g., Klebsiella and Esherichia coli*) from non lactose fermenting bacteria (*Pseudomonas aeruginosa, Salmonella* species and *Proteus mirabilis* (Oxoid 1992).

2.2.3.Salmonella-Shigella agar medium: was used for isolation of *Salmonella* and *Shigella* species and the cultures were incubated at 35°C for 24-48 hours (Feng *et al.*, 2007).

2.2.4.Violet red bile agar: was used to distinguish

coliform bacteria, and Eosin-Methylene blue Agar (EMB) was used for the isolation of *Escherichia coli* (Oxoid, 1992).

2.2.5.Mannitol salt agar: it is a differential and selective plate medium used to isolate *Staphylococcus aureus*, the medium is available in dehydrated form from Oxoid Ltd. Mannitol is fermented by *Staphylococcus aureus* (yellow in medium) (Finegold and Martin, 1982).

2.2.6. Staphylococcus medium (No.110):

Staphylococcus medium was used for the isolation of *Staphylococcus* spp. and *Micrococcus* spp. (G+bacteria) according to Matthews *et al.* (1997).

2.2.7.Rosef broth: *Campylobacter* strains were grown in stationary cultures in 5 ml of Rosef broth without antibiotics for 48 hours in microaerophilic atmosphere created by using BBL gas peak plus microaerophilic system envelops without the palladium catalyst (Ryan and Ray, 2004).

Clostridium perfringers isolates were grown in a stationary culture in an anaerobic atmosphere and subsequently diluted in sterile Rosef broth or sterile saline to concentrations of 10^6 to 10^8 CFU per ml (Baumgart *et al.* 2007).

1. **MRS Broth (de Man, Rogosa and Sharpe):** *Lactobacilli* bacteria were counted with M.R.S. agar medium according to (Laner and Kandier, 1980). After incubation, colonies developed on the plate were counted. The plates with between 30 and 300 colony recorded as colony forming units [total viable count (CFU/ml)]. Pure cultures of the isolates were obtained by subsequent sub culturing on fresh agar plates.

Identification of microbial isolates:

Isolates of bacteria were identified by the API for enterobacteriaceae only system following the method adapted by Collins *et al.* (1995). This was done based on cultural, morphological and biochemical characteristics of the isolates, the method of bacterial classification, was the Gramstaining stain method, as described by Barrow and Feltham (1993) and identification according to Krieg and Holt (1984) and Sneath *et al.* (1989).

2.3. Isolation and identification of fungi

The purpose of screening was to isolate potent pure cultures from different samples of fast and traditional foods. Test-samples were plate on the surface using the dilution plate method Swanson et al. (1992) using (1) Sabourauds Dextrose agar (SDA) (2)Potato Dextrose agar (PDA). Identification to the genus level was carried out throughout macroscopic and microscopic examination, followed by accurate more identification to the species level according to John and Pitt (1979), Domsch *et al.* (1993) and Robert *et al.* (2008).

2.4. Isolation of yeasts: Yeats were isolated from samples of traditional and fast foods by using the dilution plate method of Swanson *et al.* (1992), media used were (1) Peptone Yeast Malt Agar (PYM), (2) Dextrose-Yeast Broth (DYB), (3) Nutrient agar (NA). Yeast isolates were identified according to Arx (1981); Barnett *et al.* (2000).

Statistical analysis:

Statistical analysis was carried out using statistical program SAS (1988). Duncan's multiple range test was used to separate means.

3. RESULTS

3.1. Microorganisms isolated from traditional food samples

The results of isolation of some species of microorganisms from 60 food samples collected from 60 different restaurants at El Oasim revealed that, a total of the bacteria isolated from traditional foods include Acetobacter species, Achromobacter species, Bacillus coagulans, Bacillus subtilis, Clostridium perfringens, Escherichia coli, Erwinia carotovora, Flavobacterium species, klebsiella pneumoniae, Lactobacilus plantarum, Lactobacllus species, Listeria monocytogenes, Micrococcus species. Microbacterium lacticum, Proteus vulgaris Pseudomanas aeruginos, Ρ. fluorescens, Pseudomonas putrefaciens, Staphylococcus aureus, Streptococcus lactis, Streptococcus thermophilus and Leuconostoc mesenteroides. While from fast foods: bacteria isolated include Campylobacter *jejuni,Citrobacter* fruendii, Bacillus subtilis, Escherichia coli, Listeria monocytogenes, proteus vulgaris, pseudomonas aerginosa, Staphylococcus aureus, Salmonella species and Yersinia species. The bacterial isolates were recorded in Tables (1, 2 and 3).

(2) Bacteria isolated from fast food samples: The results of isolation of bacteria from 15 fast food samples collected from 15 restaurants at El Qassim revealed that ten genera were identified (Table 4).

3.2. Effect of different temperatures on isolated bacteria: The occurrence of isolated bacteria at different temperatures (from traditional foods and fast foods) at 10°C, 20°C, 30°C, 40°C and 50°C are found in (Tables 5 and 6).

3.3. Fungal species from fast food samples: The results of isolation of fungi from 15 fast food samples collected from 15 restaurants revealed that eight species of seven genera were isolated (Table 7) and this agree with Pelczar *et al.* (2006).

foods: In the present study the restaurants were chosen because of people in these localities are more exposed to pathogenic fungi, bacteria, yeasts which may be transmitted to them from foods, soil or hands of workers and also from person to another. The total bacterial count was higher in food samples (Mathabib, Jarish and Qursan) than those from Hunayni and Mataziz, Keshta. While the total bacterial count was lower in food samples Keshta (Freek and Saliq) and the lowest number was in Harees (Table 8).

Table (1) shows the bacteria isolated from traditional foods include Acetobacter species, Achromobacter species, Bacillus coagulans, Bacillus subtilis, Clostridium perfringens, Escherichia coli, Erwinia carotovora, Flavobacterium species, Klebsiella pneumoniae, Lactobacillus plantarum, Lactobacillus species, Listeria monocytogenes, Micrococcus species, Microbacterium lacticum, Proteus vulgaris, Pseudomonas aeruginosa, P. Pseudomonas fluorescens, putrefacien, *Staphylococcus* **Streptococcus** aureus, lactis. thermophilus Streptococcus and Leuconostoc mesenteroides.

3.5. The total bacterial count from fast foodsBacteria of this group of fast food samples may cause disease to human. Isolated bacteria included *Campylobacter jejuni, Citrobacter fruendii, Bacillus subtilis, Escherichia coli, Listeria monocytogenes, Proteus vulgaris, Pseudomonas aeruginosa, Staphylococcus aureus, Salmonella species, and Yersinia* species (Tables 4 and 9).

On the other hand, we can divide the isolated bacteria from fast and traditional food samples into:

3.5.1.Gram negative bacteria: isolated from foods traditional and fast foods are: Campylobacter jejuni, Escherichia coli, Klebsiella pneumoniae, Salmonella species, Yersinia species, Proteus vulgaris, Acetobacter species, Flavobacterium species, Pseudomonas aeruginosa, Citrobacter freundii, Erwinia carotovora, Pseudomonas fluorescens, Pseudomonas Achromobacter putrefaciens, species.

3.5.2.Gram positive cocci: Isolated from traditional foods and fast foods are: *Luconostoc mesenteroides, Micrococcus species, Staphylococcus aureus, Streptococcus Lactis and Streptococcus thermophilus.*

3.5.3.Gram positive non sporing rode: *Listeria monocytogenes* and *Lactobacilli plantarum* were isolated from traditional foods and fast foods.

No.	Name of Sample	Types of bacteria isolated from samples		
1	Jarish	Escherichia coli, Lactobacillus plantarum, Pseudomonas putrefaciens, Streptococcus thermophilus.		
2	Mataziz	Acetobacter species, Erwinia carotovora, Micrococcus species, Pseudomonas aeruginosa, Staphylococcus aureus,		
		Clostridium perfringens		
3	Qursan	Acetobacter species, Escherichia coli, Pseudomonas aeruginosa, Erwinia carotovora, Lactobacillus species.		
4	Keshta	Klebsiella pneumoniae, Micrococcus sp., Staphylococcus aureus		
5	Mathabib	Bacillus subtilis, Flavobacterium species, Klebsiella pneumoniae, Micrococcus sp., Pseudomonas aeruginosa,		
		Microbacterium lacticum, Leuconostoc mesenteroides.		
6	Freek	Bacillus subtilis, Listeria monocytogenes, Proteus vulgaris, Micrococcus species, Shewanella (Pseudomonas)		
		putrefaciens, Staphylococcus aureus.		
7	Hunayni	Bacillus coagulans, Micrococcus species, Pseudomonas fluorescens, Staphylococcus aureus		
8	Saliq	Escherichia coli, Listeria monocytogenes, Lactobacillus plantarum, Micrococcus species, Achromobacter species,		
	-	Staphylococcus aureus, Streptococcus lactis.		
9	Harees	Flavobacterium species, Micrococcus species, Staphylococcus aureus.		

Table (1): Bacterial species isolated from traditional foods.

 Table (2): Fungal species isolated from traditional foods.

No.	Name of Sample	Fungal species
1	Jarish	Aspergillus niger, Alternaria species, Fusarium species, Neurospora species.
2	Mataziz	Penicillium expansum, Chaetomium candidum.
3	Qursan	Aspergillus glaucus, Penicillium expansum, Neurospora species, Chaetonium candidum, Mucor rouxii, Alternaria species.
4	Keshta	Penicillium expansum, Rhizopus nigricans
5	Mathabib	Aspergillus niger, Cladosporium herbarum, Penicillium species.
6	Freek	Aspergillus glaucus, Cladosporium herbarum, Monilia species, Rhizopus nigricans
7	Hunayni	Alternaria species, Thamnidium elegans
8	Saliq	Sporotrichum carnis, Penicillium species.
9	Harees	Alternaria species, Penicillium species.

No.	Name of sample	Yeast species		
1	Jarish	Torulopsis caroliniana.		
2	Mataziz	ND.		
3	Qursan	Torulopsis caroliniana .		
4	Keshta	Saccharomyces rouxii, Torulopsis caroliniana,		
		Zygosaccharomyces species.		
5	Mathabib	Torulopsis caroliniana .		
6	Freek	Torulopsis caroliniana .		
7	Hunayni	Saccharomyces rouxii, Torulopsis caroliniana .		
8	Saliq	Torulopsis caroliniana, Saccharomyces rouxxi.		
9	Harees	Torulopsis caroliniana .		

Table (3): Yeast species isolated from traditional foods.

ND = Not detected.

Table (4): Bacterial species isolated from different fast foods.

Type of sample	Bacteria isolated from samples			
Chicken Shawirmas	Campylobacter jejuni, Escherichia coli, Listeira			
	monocytogenes, Salmonella species.			
Hamburger (beef burger)	Listeria monocytogenes, Salmonella species,			
	Staphylococcus aureus, Yersinia species.			
Flafel with salads (Lettuces, Tomato, Cucumber)	Campylobacter jejuni, Escherichia coli, Listeria			
	monocytogenes, Salmonella species, Pseudomonas			
	aeruginosa, Citrobacter fruendii, Proteus vulgaris,			
	Bacillus subtilis			

Table (5): Occurrence of bacteria at different temperatures isolated from traditional foods obtained from different restaurants.

Na	Bacterial species	Temperature				
No.		10°C	20°C	30°C	40°C	50°C
1	Acetobacter species	-	+	+	-	-
2	Acrhomobacter species	+	+	-	-	-
3	Bacillus coagulans	-	+	+	+	+
4	Bacillus subtilis	+	+	+	+	+
5	Clostridium perfringens	-	-	-	-	+
6	Erwinia carotovora	-	+	+	+	-
7	Escherichia coli	+	+	+	+	-
8	Flavobacterium species	+	+	+	-	-
9	Klebsiella pneumoniae	-	+	+	+	-
10	Lactobacillus plantarum	+	+	+	-	-
11	Lactobacillus species	+	+	+	-	-
12	Leuconostoc mesenteroides	-	+	+	-	-
13	Listeria monocytogenes	+	+	+	+	-
14	Microbacterium lacticum	-	+	+	+	+
15	Micrococcus species.	+	+	+	+	-
16	Proteus vulgaris	-	-	-	-	+
17	Pseudomonas aeruginosa	-	+	+	+	-
18	Pseudomonas fluorescens	+	+	-	-	-
19	Shewanella(Pseudomonas) putrefaciens	+	+	+	+	-
20	Staphylococcus aureus	+	+	+	+	+
21	Streptococcus lactis	+	+	+	-	+
22	Streptococcus thermophilus	-	+	+	+	+

No.	Isolated he starial species	Temperature				
	Isolated bacterial species	10°C	20°C	30°C	40°C	50°C
1	Campylobacter jejuni	-	+	+	+	-
2	Escherichia coli	-	+	+	+	-
3	Listeria monocytogenes	+	+	+	+	-
4	Salmonella species.	-	-	+	+	-
5	Staphylococcus aureus	-	+	+	+	-
6	Pseudomonas aeruginosa	+	+	+	+	-
7	Citrobacter fruendii	-	+	+	+	-
8	Proteus vulgaris	+	+	+	+	-
9	Bacillus subtilis	+	+	+	+	+
10	Yersinia species.	+	+	+	+	-

 Table (6): Occurrence of bacteria at different temperatures isolated from fast foods obtained from different restaurants

Table (7): Occurrence of fungal species isolated from different fast foods at 25°C.

		Fast food				
No.	Isolated fungal species	Chicken shawirma	Beef burger	Flafel with vegetable salads		
1	Aspergillus fumigatus	-	+	+		
2	Aspergillus niger	-	-	+		
3	Mucor species.	+	-	-		
4	Rhizopus nigricans	-	-	+		
5	Trichoderma species.	-	-	+		
6	Alternaria species.	+	-	+		
7	Penicillium species.	-	+	-		
8	Cladosporium species.	-	+	-		

3.5.4.Gram positive nonsporing irregular rods: *Microbacterium* species, isolated from traditional foods only.

3.5.5.Gram-positive rods and cocci (endospores): Isolated from traditional foods and fast foods. It was noticed that the samples of traditional foods included many bacteria not isolated from the samples collected from fast foods such as

Clostridium perfringens, Acetobacter species, Bacillus coagulans, Erwinia carotovora, Flavobacterium species, Klebsiella pneumoniae, Lactobacillus plantarum, Luconostoc mesenteroides, Microbacterium lacticum. Micrococcus species, Pseudomonas fluorescens, Pseudomonas putrefaciens, Streptococcus, lactis, Streptococcus thermophilus and Achromobacter species.

Matching the results recorded in Table (1) with the results recorded in Table (4), it was noticed that many species of pathogenic bacteria were isolated from traditional foods and fast foods causing many diseases for human.

The total count of bacteria was the highest in

Flafel with vegetable salad samples follow by beef burger sample and the bacterial count was lower in chicken shawirma samples (Table 9).

3.6.Isolation of fungi:A-From traditional food

Isolation and identification of fungi were performed and those included Alternaria species, *Aspergillus niger, Aspergillus glaucus, Cheotomium candidium, Cladosporium herbarum, Fusarium* species, *Monilia* species, *Mucor rouxii, Neurospora* species, *Penicillium expansum, Penicillium* species, *Rhizopus nigricans, Sporotrichum carnis and Thamnidium elegans* (Table 2).

B- From fast foods: The fungi isolated from fast food samples included *Alternaria* sp., *Aspergillus fumigatus, Aspergillus niger, Cladosproium herbarum. Mucor* species, *Rhizopus nigricans, Trichoderma* species, *Penicillium* sp. (Table 7).

(5) Isolation of yeasts from traditional foods: In the present study yeasts were isolated from traditional food samples and included *Saccharomyces rouxii, Torulopsis caroliniana and Zygosaccharomyces* sp. as shown in Table (3).

4. DISCUSSION

The effect of microorganisms on human health has been reported. The present study was performed in Saudi Arabia (2007) to give information of the distribution and presence of pathogenic microorganisms in traditional and fast foods from different restaurants. This is important to human and to discuss their role in the food poisoning and also the causation of many human diseases. Studies on the isolation of pathogenic bacteria, fungi and yeasts in this investigation indicated that some Gram negative and Gram positive bacteria were found and recorded in Tables (1 & 4). Bacteria of the greatest importantce to human pathology are the most common causes of human infection and extensively widespread in the environment using fast foods. Our results are in agreement with those of Kay et al. (1994). Their findings are consistent with our results that revealed some pathogenic bacteria, fungi and veasts in fast and traditional foods. Most investigators indicated that bacteria, fungi and yeasts may exert their pathogenic action either through infection of the body, or as a source of toxic substances demonstrated in contaminated foods. The most common infections causing food poisoning and other diseases are those associated with contaminations due to fast and traditional foods (Kay et al., 1994).

More recent studies have focused attention on the food poisoning diseases due to several pathogenic microorganisms. Many human diseases are associated with contamination of fast and traditional fast foods. Many workers reported that raw food, especially meat, poultry and sea food, as well as their juices, can contain dangerous microorganisms, such Campylobacter, Listeria as Salmonella, and Escherichia coli, which may be transferred into other foods during food preparation and storage (Uyttendaele et al., 1999). Our work is in agreement with the above studies. In raw milk, the presence of lactic acid producing bacteria, under suitable conditions ferments the lactose present to lactic acid. The increasing acidity in turn prevents the growth of other organisms, or slows their growth significantly. During pasteurization however, these lactic acid bacteria are mostly destroyed (Christison and Ivany, 2006). These results are in agreement with our results.

Feng *et al.* (2007) reported that some microorganisms are harmful and cause disease while others are benevolent neutral, or even helpful (*e.g.*,

Streptococcus lactis to make butter milk and break down toxins in our environment, while others can sick us (*e.g.*, contaminants in food like *Escherichia coli or Salmonella*), or can kill us for example,

	bacteria samples.	counted in traditiona
No.	Sample	Total viable count (Log 10 cfu/gm)
1	Jarish	5.20
2	Mataziz	5.07
3	Qursan	5.20
4	Keshta	1.02
5	Mathabib	5.38
6	Freek	0.94
7	Hunayni	5.15
8	Saliq	0.94
9	Harees	0.59

Table (8): Total viable counts (cfu/gm) of

cfu: colony forming unit

tomato, cucumber)

samples.			
Type of sample	Total count of bacteria (Log 10 cfu/gm)		
Chicken shawarmas	5.28		
Beef burger	5.53		
Flafel with vegetable salads (lettuce,	5.76		

Table (9): Total viable counts (cfu/gm) of bacteria counted in fast food samples

Proteus cause amoebic dysentery, fungi cause athelete's foot and ringworm, bacteria cause pneumonia, legionmaire's disease, *Streptococcus* throat, tetanus and other diseases.

Some animal disease bacteria can cause human diseases with close animal-man contact. Some of these are Brucella, enteropathgoenic E. coli, Mycobacterium, Corvnebacteria, Leptospira, Coxiella burnetii and Clostridium tetani. Heaton and Jones (2008) suggested that coliforms, E. coli, aureus, C. perfringens enterococci, S. and Salmonella are often present on fresh tissues since slaughter process does not include a bacterial step. The frequency and levels of these bacteria will vary, depending upon farm, climatic, and processing conditions.

Recent studies indicated that *Staphylococcus aureus*, *Clostridium perfringens*, and *Salmonella* frequently are present in low numbers on raw meat surface *Clostridium botulinum* occurs infrequently.

As shown in Table (10) pathogenic bacteria, important diseases and prevention of each disease were recorded to indicate the possible causative

Organism	Where they be found	Important diseases	Prevention
Campylobacter jejuni	Contaminated drinking water and unpasteurized milk, contaminated food, with incorected prepared meat and poultry	Human gastroenteritis in the world, food poisoning, abdominal pain diarrhea, fever, and malaise	Treated with antibiotics in severe cases such as ciprofloxacin, erythromycin, azithromycin or norfloxacin
Escherichia coli	Food or water or with the individuals handling the infant's child, unwashed vegetables or undercooked meat, raw ground beef, raw seed sprouts, raw milk, unpasteurized juice, and foods contaminated, by infected food workers via fecal-oral route. Found in recreational waters and its presence is used to indicate the presence of recent fecal contamination	Most <i>E. coli</i> strains are harmless, but some can cause serious food poisoning in humans, and are occasionally responsible for product recalls, the harmless strains are part of the normal flora of the gut. Produce potentially lethal toxins, food poisoning, diarrhea in humans, rabbis, dogs, cats and horses, urinary tract infections	Cooking food property, preventing cross- contamination instituting barriers such as gloves for food workers, pasteurization of juice or dairy products and proper hand washing requirements treated with antibiotics.
Listeria monocytogenes	Foods as raw milk, pasteurizede fluid milk, cheeses, ice cream, raw vegetables, fermented raw meat sausages, raw and cooked poultry, raw meats (of all types), and raw and smoked fish and refrigerated foods. Isolated also from soil and silage.	It is the causative agent of listeriosis. It is one of the most virulent food borne pathogens with fatality rates exceeding even <i>Salmonella</i> and <i>Clostridium botulinum</i> . Gastrointestinal symptoms such as nausea, vomiting, and diarrhea	For gastrointestinal using antacids or cimetidine. May using also Vancomycin or Ampicillin Alcohol as an effective topical sanitizer or quaternary ammonium added to alcohol.
Pseudomonas aeruginosa	It is found in soil, water, skin flora, in normal atmospheres and also in little oxygen. This bacterium is also found on and in medical equipment including catheters, causing cross infections in hospital and clinics	It infects the pulmonary tract, urinary tract, burns wounds, and also causes other blood infections, gastrointestinal infection and external ear infection	Many antibiotics, for example, ear infections or nail infections, topical gentamicin or colistin may be used.
Salmonella sp.	It can be transmitted by humans to animals and vice versa. Transmitted to humans by eating foods contaminated with animal feces. Contaminated food such as beef, poultry, milk, or eggs, any food including vegetables, food also become contaminated by the hands of an infected food handler who did not wash hands with soap after using the bathroom. The human pathogen of Salmonella abdominals transmission by contact and infected food, water or fly. Contamined foodstuffs.	Diarrhea, fever, or abdominal cramps, food poisoning.	Food be heated for at least ten minutes at 75°C (167°F) so that the center of the food reaches this temperature. It is not destroyed by freezing.
Staphylococcus aureus	Part of the skin floral found in the nose and on skin. It presents in prepared foods left too long at room temperature (e.g., cooked hamburger, salads, dairy products).	It can cause skin infections, pneumoniae, meningitis osteom-yelitis, endocarditis, toxic shock syndrome, bacteremia and septicemia, food poisoning.	Treatment using penicillin, gentamicin
<i>Yersinia</i> sp.	Food products (especially vegetables, milk-derived products and meat)	Gastroenteritis	By oxidizing agents such as hydrogen peroxide and potassium permanganate solutions

Table (10): Pathogenic bacteria, important diseases and prevention of each disease (bacteria isolated from fast and traditional foods)

factor (Richard *et al.*, 2007; FDA Food, Drug Adminsitration, 2010).

These species are most hazardous when they grow without competition as in cooked foods (Talarico et al., 1997; Baumgart et al., 2007). Our results are also in agreement with the previous studies. The hazard potential from foods precooked in commercial establishment is high but the incidence of outbreaks has been low. The Center for Disease Control reported that although more than half of all food borne disease outbreak can be traced to meat and poultry products, there was a serious departure from good practices at the serving level (homes, restaurants, institutions) in nearly all instances. Baumgart et al. (2007) showed that the heating step in the production of cooked cured meats destroyes the typical raw meat flora except the spores. Salt and nitrite in the cure inhibit the growth of survivors and contaminats somewhat selectively. These agree with the results of this study (Table 9).

Ryan and Ray (2004) found that upon prolonged refrigeration, lactic acid bacteria, micrococci, enterobacteria, Bacillus, and yeast may grow and form slime. If the product is in a tight, gas impermeable package, the package may swell. Products of bacterial action sometimes combine with meat pigments to form a green color. Human contact may sometimes introduce a few Escherichia coli or Staphylococcus aureus. These results are in agreement with our results. The food bacteria associated with food fermentation are capable of producing different types of metabolites. They have antimicrobial properties (organic acids e.g., lactic, acetic propionic, aldehydes, ketones, and alcohols (ethanol, diacetyl and acetaldehyde), hydrogen peroxide, reuterine and bacteroides. It was reported that in the presence of the mesophilic lactic acid bacteria (e.g., Lactococcus lactis, some lactobacillus species, and Pediococcus sp.), the growth of psychrotrophic spoilage and pathogenic bacteria was reported to be controlled.

In refrigerated raw milk, meat, egg., and seafood, cells of *Lactobacillus*, *Lactococcus* and *Leuconostoc* species were added to control the growth of psychorotrophic spoilage bacteria such as *Pseudomonas* spp. The inhibitory property could be due to the release of antimicrobial compounds from the cells by the nonmetabolizing lactic acid bacteria.

It was reported that some strains of *Lactobacillus reuteri*, found in the gastrointestinal tract of humans and animals, produce a small molecule, reuterine that is antimicrobial against Gram-positive and Gram negative bacteria. It produces antibacterial action by inactivation some important enzymes, such as ribonucleotide reductase. Many strains of species

from genera *Lactococcus, Streptococcus, Leuconostoc, Pediococcus, Bifidobacterium*, and *Propionibacterium* used in food fermentation have been reported to produce different bacteriocins (Daeschel and Panner, 1992).

Moreover, many of the organisms recorded in this study are in agreement with many researches, several yeast isolates normally present on the surface of fruits and vegetables were reported to prevent spoilage of the products by molds. Some of the inhibitory compounds are small proteins, while some others are enzymes. It was reported that cells of one such yeast isolate were found to adhere tightly with the mold mycelia and produce β -gluconase that degrades the cell wall of the molds and kills them. As many of these yeasts are normally present in fruits and vegetables that are eaten raw, they are not considered pathogenic and thus human may be sick when eat it or when used in salads in fast food or traditional food (Barnett *et al.*, 2000).

Uzeh et al. (2009) showed that decay caused by molds and certain bacteria accounts for much of spoilage of fresh fruits and vegetables. Many of these organisms are true plant pathogens in that they can invade healthy plant tissue. While bacterial rot is caused mainly by genus Erwinia, numerous mold such as Alternaria, Botrytis, Phytophthora are responsible for a variety of market diseases. Our data are in agreement with the results obtained by Bichai et al. (2008) who showed that, the presence of Escherichia coli can be related to the use of polluted irrigation waters during growth, contamination through human handling, the use of contaminated containers, or washing after harvest with polluted water. It was suggested that it could increase the incidence of enteric pathogens (Angelillo et al., 2000). Thus products such as fresh or processed vegetables are the chapped salad ingredients (Lettuce, cabbage, carrots, tometo, cucumber....etc) sold in the grocery store and to the institutional trade (Kaneko et al., 1999a). These otained results are in agreement with those obtained by Ali and Shalaby (1999), who reported that biogenic amines are natural antinutrition factors and are important from a hygienic point of view as they have been implicated as the causative agents in a number of food poisoning episodes, and they are able to initiate various pharmacological reaction. Histamine, putrescine, cadaverine, tyramine, tryptam, Bphenylethylamine, spermine, and spermidine are considered to be the most important biogenic amines occurring in foods.

It was suggested that eating of fast food can lead to measurable signs of liver injury and inflammation. The plentiful availability of relatively inexpensive fat-and calorie-packed foods, has helped to make us the fattest. Fats which are commonly found in fast food have been shown in many tests to have a negative health effect on the body. It was suggested that fast food consumption has been shown to increase calorie intake, promote weight gain, and elevate risk for diabetes. A food may start with a pH which precludes bacterial growth but as a result of the metabolism of other microorganisms (yeasts or molds), pH shifts may occur and permit bacterial growth (Hathcox *et al.*, 1995 and Robert *et al.*, 2008).

The interplay of factors affecting microbial growth in foods such as (water activity, pH, temperature) ultimately determines whether a microorganisms will grow in a given food. Often, the results of such interplay are unpredictable, as poorly understood synergism or antagonism may occur (Smith and Fratamico, 1995). Similar results obtained by Richard et al. (2007) who were reported that some pathogenic bacteria cause sickness for human. These bacteria such as: Listeria monocytogenes have been associated with such foods as raw milk, pasteurize fluid milk, cheeses,, ice cream, raw vegtables, fermented raw-meat sausages, raw and cooked poultry, raw meats (of all types), and raw and smoked fish. Its ability to grow at temperature as low as 0°C permits multiplication in refrigerated foods. In refrigeration temperature 4°C the amount of ferric iron promotes the growth of L. monocytogenes (Dharmarha and Vaishali, 2009; Farber and Peterkin, 2009). Their results are in agreement with our results. Richard et al. (2007) showed that gastrointestinal disease has been reported by eating raw or inadequately cooked meat containing Bacillus spores. B. cereus causes food poisoning B. subtilis, B. coagulans were isolated from traditional food samples (Table 1) and (Table 4) from fast food samples of our study.

Campylobacter jejuni is widely distributed in nature, it infects the intestine, where it can cause ulcerative, inflammatory lesions in the jejunum, ileum, or colon. Diarrhea should be treated using fluids. Prevention is induced by good hygiene avoiding contaminated water, pasteurizing milk and milk products, and thoroughly cooking potentially contaminated food (*e.g.*, poultry). *C. jejuni* was isolated from fast food samples only (Table 4). *Clostridium perfringen* is a part of the normal flora of the vagina and gastrointestinal tract. Its spores are found in soil. Acute food poisoning is caused by the generation of spores in improperly cooked food, resulting in the production of enterotoxin in the small intestine. It was isolated from traditional food samples (Table 1).

Feng et al. (2007) suggested that Escherichia coli is a part of the normal flora in the colon of human and other animals, but can be pathogenic both within and outside the gastrointestinal tract. Enterotoxigenic E. coli (ETEC), is a common cause of "traveler's diarrhea" in developing countries, it infects only humans, with transmission occurring through food and water contaminated with human waste, or by person to person contact, diarrhea can be prevented by taking precaution in food and water consumption, hand washing and disinfection. E. coli was isolated from traditional food samples (Table 1) and from fast food samples (Table 4). These results are in agreement with those obtained by AVI Biopharma (2008) who suggested that Pseudomonas aeruginosa often includes the production of both pyocyanin and fluorescein, as well as its ability to grow at 42°C. P. aeruginosa is capable of growth in diesel and jet fuel, where it is known as a hydrocarbon-utilizing microorganism (or HUM bug), causing microbial corrosion. It creates dark gellish mats sometimes called "algae" because of their appearance. Several studies indicated that P. aeruginosa is the common cause of infections of burn injuries and of the external ear (otitis externa), and is the most frequent colonizer of medical devices (e.g., catheters). However, research indicates that salicylic acid can inhibit pyocyanin production (Prithiviraj et al., 2005). P. aeruginosa is widely distributed in nature (soil, water, plants, animals). P. aeruginosa can grow in distilled water, laboratory hot water baths, hot tubes, wet IV tubing, and other water containing vessels. This explains why the organism is responsible for so many nosocomial infections. This result is in agreement with our results. P. aeruginosa was isolated from traditional food samples and fast food samples (Tables 1 and 4).

To protect against Salmonella infection, it is recommended that food should be heated for at least ten minutes at 75°C (167°F) so that the center of the food reaches this temperature. Salmonella is not destroyed by freezing. It can survive several weeks in a dry environment and several months in water thus, they are frequently found in polluted water, contaminated from the excrement of carrier particularly animals being important (Administration Urged to Boost Food Safety, 2009; FDA/CFSAN-Food Safety, 2009). S. typhi is transmitted generally through food or water contaminated by human faeces. Public food handlers who are carriers can present a serious

public health problem. *Salmonella* sp. was isolated from fast food samples (Table 4). Our results agree with the previous studies.

Staphylococcal gastroenteritis is caused by ingestion of food contaminated with toxin produced by Staphylococcus aureus superantigen. Some strains of S. aureus are capable of producing staphyloxanthin a carotenoid pigment that act as a virulence factor. It has an antioxidant action that helps the microbe to evade killing with reactive oxygen used by the host immune system. It is thought that staphyloxanthin is responsible for S. aureus characteristic golden colour (Clouditz et al., 2006). It was reported that emphasis on basic hand washing techniques are therefore effective in preventing the transmission of S. aureus. The use of disposable gloves by staff reduces skin-to-skin contact that therefore further reduces the risk of transmission.

Recent reports demonstrated that many researches showed that the introduction of *Staphylococcus aureus* into the blood stream can lead to various complications (Tables 1 and 4). Washing of hands and disinfection of fomites are important in the control of nosocomial *Staphylococcus aureus* epidermidis (Cosgrove *et al.*, 2009). *Streptococcus lactis, Streptococcus thermophilus* were isolated from traditional food samples (Table 1).

Yersinia species is endemic distributed worldwide. Infection is transmitted by fleas, the organism can also be transmitted by ingestion of contaminated animal tissues, or via the respiratory route. Y. species was isolated from fast food samples (Table 4). Some members of Yersinia are pathogenic for humans, in particular, Y. pestis is the causative agent of the bubonic plaque. Rodents are the natural reservoirs of Yersinia, less frequently other mammals serve as hosts. Infection may occur either through blood (in the case of Y. pestis) or in an alimentary fashion, occasionally via consumption of food products (especially vegetables, milk-derived products and meat) contaminated with infected urine or feces (Ryan and Ray, 2004). These results are agree with our results. Yersinia may be associated with Crohn's disease, an inflammatory autoimmune condition of the gut by treatment and prevention by streptomycinis the drug of choice, gentamicin and tetracycline are acceptable alternatives.

For individuals in enzootic areas, efforts to minimize exposure to rodents and fleas is important. The above studies are in agreement with our research.

Barnett et al. (2000) reported that many mycotoxins found in various foods such as: Aspergillus toxins: (a) Aflatoxins: Aspergillus and Pencillium sp. have been reported to produce aflatoxins, and the aflatoxins have been isolated from legumes, grains, fruits, meats, spices. cheeses, milk, rice, corn, cotton seeds, others compounds with carcinogenic, hemorrhagic, heptaotoxic, neurotoxic and uterotrophic properties have been isolated from food stuff and identified as metabolites of fungi common to a variety of agricultural commondities (Shank et al., 1972). (b) Ochratoxins: these substances are a group of closely related compounds produced by Aspergillus ochraceus, A. sulphureus, and A. melleus. A. ochraceus group is common in soils and decaying vegetation, grains, wheat, corn, cotton seeds, legumes, peppers, onions and pears (Adams and Moss, 2000). (c) Sterigmatocystin: another common food contaminant is Aspergillus versicolor, A. nidulans, a compound bearing some structural resemblance to the aflatoxins in that, it produces liver and kidney damage and is like the aflatoxing a hepatocarcinogen (Stack and Rodricks, 1973). (d) Other Aspergillus toxins: there are numbers of other metabolites of this genus which have been shown toxic to animals, which are potential food contaminants.

Penicillium toxins: (a) Patulin: among the more important of the large number of mycotoxins produced by the penicillia is the potent antibiotic, Penicillic patulin. acid. rubrotoxin. and tremorgens and cyclopiazonic acid. that compounds elaborate a toxin and have potential carcinogenic agents (Harwig et al., 1973). (b) Rice toxins: Storage fungi proliferate in improperly stored rice. Most are of the genera Aspergillus and Pencillium, and about 10% of the isolates tested are toxigenic. The toxic effects of these substances interactions among them, and their natural occurrence have been reported (Saito et al., 1971). A polyenic compound called citreoviridin and an acidic compound known as citreomycetin. Some of these toxins affect the liver and kidney, some are neurotoxic.

Fusarium toxins: (a) Zearalenone, trichothecenes (diacetoxyscirpenol) and other toxigenic fungi, their potential for human health effects is probably realized by growth of *Fusarium* sp. It was reported that it is probably one of the more common mycotoxin contamination of food and feed (Pelczar *et al.*, 2006). (b) Trichothecenes: There are mold metabolites which have structural features similar to those of the compound known as diacetoxyscirpenol. The acute toxicity of some of mycotoxin trichothecenes cause hemorrhage on the lip and mouth, throat, and entire gastrointestinal tract.

International Commission on Microbiological Specifications for Foods (ICMSF, 1996) reported that there are hundreds of fungal species which have been shown to be toxigenic from animal feeds, peanuts, and seeds, flour, spaghetti, black and red peppers. The following genera showed toxic isolates: Cheotomium, Cladosporium, Alternaria; Curvularia, Gliocladium. Rhizoctonia, Trichoderma, Trichothecium. Scopulariopsis, Pithomyces chartarum produce mold metabolites and known sporidesmins Rhizoctonia as leguminicola produce Slaframine causing diarrhea for animals. Also, suggested that control of fungal toxin production can be controlled by the adjustment of pH, water activity, and temperature control. Temperature does not protect from all toxigenic molds, however, for many will grow at refrigeration temperatures. ICMSF found several toxigenic species capable of growth and toxin production at temperatures down to 10°C. There is evidence that some strains may be more toxigenic at low temperatures than at optimum growth temperatures. Adjustment of water activity, is the best means of controlling growth of microorganisms in foods. e.g., Campylobacter cells when ingested with food or water, it enters the host intestine via the stomach and colonize the distal ileum and colon. The most effective means of eliminating human exposure to mycotoxins in foods is by the prevention of toxin formation. This requires agricultural and industry practices designed to reduce the opportunity for fungal growth from harvest to ultimate commodity use.

In conclusion, the need for good hygienic practices, proper handling, storage and retail of salads in clean environment and at refrigeration temperature can not be over emphasized to ensure good quality and safe salads.

Prevention of mycotoxins must become a cooperative effort on the part of all involved in food production.

We can also conclude that if people have meals regularly and in suitable quantities, there will not be any health problems, relating habits concerning to nutrition according to what the healthy nutritional experts specify, if all the society follow right nutritional habits, healthy foods, they have health.

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محتوى الوجبات السريعة والشعبية من الميكروبات

سعدية محمد حسانين عيسى

قسم الميكروبيولوجي - كلية العلوم - جامعة عين شمس - القاهرة - مصر

ملخص

تسبب الوجبات السريعة الشائعة والشعبية كثيرا من الامراض مثل امراض التسمم الغذائي وأمراض السمنة التي تؤثر على صحة الإنسان. عزلت العديد من الميكروبات (بكتيريا، فطريات، خمائر) كان لها دور هام في إصابة الإنسان بكثير من الأمراض.

إستهدفت الدراسة الحالية معرفة أنواع الميكروبات المتواجدة في الوجبات السريعة عن طريق عزلها وتنقيتها وتعريفها ومعرفة العوامل التي تؤثر على نموها ومن أهمها درجة الحرارة.

استخدم فى هذه الدراسة عدد 60 وجبة سريعة جمعت من 60 مطعم للوجبات السريعة فى أنحاء مختلفة من منطقة القصيم بالمملكة العربية السعودية عام 2007 تم تقسيمها إلى مجموعتين: الأولى عبارة عن وجبات سريعة منتشرة شائعة وهى عبارة عن سندوتشات طعمية، بيف برجر، شاورما دجاج (15 عينة من 15 مطعم)، والثانية: عبارة عن مطاعم الوجبات الشعبية السريعة (45 عينة من 45 مطعم) وهى عبارة عن جريش، مطازيز، كرسان، قشطة، مصابيب، فريك، حنينى، سليق، هريس.

استخدمت درجات حرارة مختلفة 10 °م، 20°م، 30°م، 40°م و50°م لمعرفة انواع الكبتيريا التى تنمو عند درجات الحرارة المرتفعة والممرضة للإنسان وأيضا البكتيريا التي تفسد الأغذية. تم إجراء الاختبارات التقسيمية التأكيدية باستخدام أشرطة API (API 20E, bio Merieus, 2010) لتعريف البكتيريا. تم استخدام الطرق والاختبارات التأكيدية لتعريف الفطريات والخمائر المعزولة من الوجبات السريعة جميعها بعد تحضينها عند درجة حرارة 25°م.

تم فى هذه الدراسة عزل 22 نوع من البكتيريا و14 نوع من الفطريات وثلاثة أنواع من الخمائر من 60 عينة وجبة سريعة. تم تنمية البكتيريا عند درجات الحرارة المختلفة حيث وجد أن بعض البكتيريا لها القدرة على النمو عند 50 م وذلك يسبب فساد للوجبات السريعة.

أثبتت النتائج أن 9 أنواع من البكتيريا محبة للحرارة حيث أمكنها النمو عند درجة حرارة 50 م.

تبين أن هناك اختلاف واضح فى العدد الكلى للبكتيريا فى عينات الوجبات السريعة معتمدة على درجة pH الغذاء، درجة الحرارة، استخدام مواد غذائية خام لم تتعرض للطبخ أو التسخين مثل الخضروات المضافة للسلطة الخضراء (طماطم - خس - بقدونس - خيار - كرنب) والمقبلات التى تضاف لسندوتشات الوجبات السريعة. ولذلك نرى أن أكثر أعداد البكتريا كانت فى السندوتشات التى أضيفت لها خضروات طازجة وهى سندوتشات الطعمية يليها سندوتشات بيف بيرجر ثم أقلها فى العدد سندوتشات شاورما الدجاج. أما بالنسبة لأعداد البكتيريا فى الوجبات الشعبية السريعة فقد كانت أقل من حيث العدد (ماعدا بعض العينات) حيث يتم طبخها جيداً واستخدام درجة حرارة مدة أطول ولا يضاف إليها خضروات طازجة قبل تتاولها. قد يرجع وجود البكتيريا، الفطريات، الخمائر فى الوجبات السريعة مثل بين جر، شاورما، الطعمية والوجبات الت الشعبية الأخرى إلى عدم غسيل الخضروات التى تستخدم فى عمل السلطة الخضراء بماء نظيف حيث تستخدم خام دون تعريضها للحرارة. قد تكون أيدى العاملين مصدر لتلوث السندوتشات أو يكونوا حاملين للأمراض، الأوعية والأجهزة والأدوات التى تستخدم فى تحضير وتجهيز الوجبات السريعة.

أظهرت نتائج البحث أن تناول الوجبات السريعة تسبب كثير من الامراض لما تحتويه من ميكروبات ودهون تسبب السمنة وتؤدى إلى أمراض عديدة. يمكن منع هذه الامراض الناتجة عن الغذاء او الإقلال منها عن طريق: الاهتمام بتطبيق الاشتراطات الصحية فى المطاعم واستمرار التفتيش، توفير أجهزة الكشف ومستلزمات الاختبارات السريعة (Kits) لمفتشى الجهات الرقابية على المطاعم، الاهتمام بتنظيف الخضروات الورقية التى يصنع منها السلطة لضمان خلوها من الأتربة، الطفيليات والفيروسات وتخزينها بطريقة صحية لحين استخدامها، استخدام درجات الحرارة المناسبة فى إنضاج الطعام وحفظه فى أماكن بعيدة عن الحشرات والقوارض وحفظه عند درجات حرارة منخفضة داخل ثلاجات، استخدام مياه نقية نظيفة فى تحضير الوجبات السريعة والتأكد من المواد الخام من حيث مطابقتها للمواصفات مثل اللحوم، البيض، عدم ترك الوجبات السريعة موالية فى درجة حرارة الجو وخاصة التى تحتوى على مواد غذائية حيوانية وبخاصة فى فصل الصيف.

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