Effect of Heat Killed Probiotic and Wheat Bran on Gastrointestinal Symptoms in Hemodialysis Children

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Abstract

Introduction: Gastrointestinal (GIT) disturbances are recorded in patients with end stage renal diseases (ESRD) frequently, killed probiotic was reported in to decrease GIT problems many other diseases; So, the aim of our study is to evaluate whether a combination of heat killed probiotic and wheat bran affect the presence and severity of gastrointestinal symptoms (GIS) in hemodialysis children.

Subjects and methods: Our study was Open label uncontrolled clinical trial conducted on 35 pediatric patients on regular hemodialysis for end stage renal disease. every patient received one sachet of killed probiotic and wheat bran twice daily for two months. Assessment of GIT symptoms were done one month before administration of the drug, then follow them for two months during intervention then reassess the symptoms.

Result: We found that combined heat killed probiotic + wheat bran administration to hemodialysis children produced significant differences in urea, Hb and PO4 levels compared to baseline levels, in addition children produced highly significant decrease in all gastrointestinal symptoms (P<0.001) after 1 month, after 2 months in comparison to baseline.

Conclusion: We concluded that combination of heat killed probiotic and wheat bran decreases presence and severity of gastrointestinal symptoms (GIS) in hemodialysis children; thus we hypothesised that, our results could represent an important preliminary work that need further justification.

Keywords: Heat killed probiotic; Wheat bran; Gastrointestinal; Hemodialysis children; Lactobacillus rhamnosus; Uremic toxins; Bifidobacteria

Introduction

Patients with renal disease usually have impaired intestinal microbiome. It is suggested that almost two thirds of individuals with uremia have abnormalities in the gastrointestinal mucosa and a disequilibrium in the intestinal ecosystem [1]. The incidence of GI symptoms can largely be attributed to the underlying conditions, such as increased level of uremic toxin, the effect of dialysis, lifestyle change, or the medications required for treatment [2], the most common gastrointestinal symptoms in patients with renal failure include nausea, vomiting, abdominal pain, constipation, and diarrhea [3].

The term of probiotic was derived from Greek, meaning “for life”. The joint Food and Agriculture Organization (FAO) and World Health Organization (WHO) Expert Consultation on evaluation of health and nutritional properties of probiotics in food including powder milk with live lactic acid bacteria defined probiotics as: “live microorganisms that, when administered in adequate amounts, confer a health benefit on the host” [4]. Prebiotics have the ability to support the growth of Probiotics [5]. It is believed that diets high in phytate contribute to Fe deficiency [6]. In contrast, some naturally occurring non-digestible oligosaccharides in wheat, known as prebiotics, have been suggested to have an enhancing effect on Fe absorption [7].

Studies have demonstrated that heat-killed probiotics are safer than live probiotics for purposes such as eliminating antibiotic resistant genes, preventing production of recombinant strains, controlling the microbial load during probiotic supplementation, and application in children and immunosuppressed patients [8,9]. Equal effectiveness of viable and heat-killed Lactobacillus rhamnosus GG [10] and greater efficacy of heat-killed Lactobacillus acidophilus LB than viable non-specified strain of L. acidophilus [11], for the treatment diarrhea were noted. It was further denoted that application of inactivated cells may be a safer approach than
using live microbes in premature infants immediately after birth or in immuno-suppressed individuals. Li Net al. [12].

The aim of our study is to evaluate whether a combination of heat killed probiotic and wheat bran affect the presence and severity of gastrointestinal symptoms (GIS) in hemodialysis children.

Patients and Methods

Patients

Our study was an open label uncontrolled clinical trial conducted on 35 pediatric patients aged 10-18 years. There were 17 males and 18 females, they are all on regular hemodialysis for end stage renal disease (ESRD), in Pediatric Nephrology Unit, Zagazig University Hospitals. The study was conducted after taking an informed consent from patient and /or their parents, 35 out of 54 on patients on hemodialysis accept to participate in our study, we included Pediatric patients clinically stable on regular hemodialysis with arteriovenous fistula, duration of dialysis was 7.37±2.921 years and receiving thrice-weekly hemodialysis for at least 3 months before start the study. In this study we excluded Patients with usual intake of Probiotics, omega-3 fatty acids, Patients with immunosuppressive and/or non-steroidal anti-inflammatory drugs and Patients with medical illnesses that affect nutritional and inflammatory status (cancer, decompensated heart failure, chronic liver diseases, intestinal malabsorption, active infections, and acquired immunodeficiency syndrome).

Methodology

All studied infants were subjected to the following at the beginning of the study; Full detailed history taking including; Personal history (age, sex, address), Compliant, Present history and GIT symptoms (Anorexia, Nausea, Vomiting, Heart burn, Gastric discomfort, Gastric distention, Constipation, Diarrhea, Complication related to other systems and Drugs given). We also take past history; History of causative diseases, Duration of dialysis and Family history. Thorough clinical examination was done including the following; General examination, Anthropometric measurements and local examination. Routine laboratory investigation was done to all patients including.

Daily intake of Probiotics, patients received standard dose of (Lacteal forte) twice daily starting with the first day of the study till complete 2 months and receive also one wheat bran capsule twice daily as a source of insoluble fiber. Lacteal forte was used in the form of sachets product by the company of the tenth of Ramadan/Egypt with a license from the French company Axcan Pharma - France. each sachet contains Lactobacillus LB, corresponding to lactobacillus delbruekii and lactobacillus fermentum one sachet containing 10 billion of lyophilized heat-killed L. acidophilus LB, twice daily plus one capsule of wheat bran starting with the first day of the study till two months later; The main action of Lactobacillus LB consist in stimulating and regulating the intestine’s host defense mechanisms.

Every patient received also Wheat bran as a concentrated source of insoluble fiber may result in synergistic effects on gastrointestinal function, in the form of capsules each capsule contains Bran (treated) 400mg product by the company of Arab Company for pharmaceuticals and medicinal plants (MEPACO).

Assessment of GIT symptoms were done before administration of the drug, then follow them for two months during intervention with reassessment of the symptoms (Table 1).

Table 1: Questionnaire design of GIT symptoms.

| Symptoms            | None | Mild | Moderate | Sever | Very Sever |
|---------------------|------|------|----------|-------|------------|
| Anorexia            | 1    | 2    | 3        | 4     | 5          |
| Nausea              | 1    | 2    | 3        | 4     | 5          |
| Vomiting            | 1    | 2    | 3        | 4     | 5          |
| Gastric discomfort  | 1    | 2    | 3        | 4     | 5          |
| Abdominal distention| 1    | 2    | 3        | 4     | 5          |
| Constipation        | 1    | 2    | 3        | 4     | 5          |
| Diarrhea            | 1    | 2    | 3        | 4     | 5          |

None=absence of symptom
Mild=presence of symptom, but no effect of daily living
Moderate=obvious symptom, with slight effect on daily living
Sever =sever symptom, with great effect on daily living
Very sever =outstanding symptom hardly endurable [12]

Statistical analysis

Data were analyzed by Statistical Package of Social Science (SPSS) version 17, we used Wilcoxon Signed-Rank Test as nonparametric alternative to paired t-test is used to compare two sets of data that come from the same participants and we used Paired-samples t-test (dependent t-test) to compare the means between two related groups on the same continuous, dependent variable. We used Friedman test to test differences between groups when the dependent variable being measured is ordinal (One group is measured on three or more different occasions). P value was considered significant if less than 0.05.
**Results**

**Clinic-demographic data**

We found that combined heat killed probiotic +wheat bran administration to hemodialysis children produced significant differences in urea, Hb, CRP and Po4 levels compared to baseline levels. However, other clinic-laboratory parameters showed no significant differences following combined heat killed probiotic+ wheat bran administration compared to baseline levels (Table 2).

Table 2: Comparison between clinic-laboratory data among the studied children (before and after combined heat killed probiotic and wheat bran administration).

| Variables               | Before (probiotic + wheat bran) | After (probiotic + wheat bran) | Test                        | P-value |
|-------------------------|---------------------------------|--------------------------------|-----------------------------|---------|
| Weight (Kg) Mean±SD Median(Range) | 32.6±6.26 32(21-43) | 32.1±6.01 31(21-43) | Wilcoxon signed rank test (Z)=1.357 | 0.175  |
| Albumin (mmol/L) Mean±SD Median(Range) | 4.13±0.260 4.1(3.7-4.6) | 4.14±0.19 | Wilcoxon signed rank test (Z)=0.205 | 0.837  |
| Urea (mg/dl) Mean±SD Median(Range) | 53.68±10.97 53.4(7.3-66) | 60.8±10.40 63(42-77) | Wilcoxon signed rank test (Z)=2.52 | 0.012  |
| Creatinine (mg/dl) Mean±SD Median(Range) | 7.41±2.26 7.2(3.4-13.7) | 7.52±1.46 7.6(4.5-11) | Wilcoxon signed rank test (Z)=0.805 | 0.42   |
| Hb (g/dl) Mean±SD Median(Range) | 9.46±0.65 9.5(7.9-10.6) | 10.54±0.67 10.4(9.2-12.5) | Paired t-test=-6.37 | <0.001  |
| Ca (mg/dl) Mean±SD Median(Range) | 8.5±0.80 8.5(6.9-10.6) | 8.7±0.69 8.9(7.3-9.6) | Paired t-test=-1.02 | 0.316  |
| Po4 (mg/dl) Mean±SD Median(Range) | 5.2±1.22 5.2(3.7-8.7) | 5.71±1.18 5.4(3.7-8.2) | Paired t-test=-2.04 | 0.05   |
| Na (mg/dl) Mean±MD Median(Range) | 137.6±12.72 137(132-143) | 137.45±2.59 137(132-140) | Paired t-test=1 | 0.324  |
| K (mg/dl) Mean±SD Median(Range) | 6.11±1.10 5.9(4.5-8.30) | 6.04±1.29 6.2(3.7-8) | Paired t-test=0.261 | 0.796  |
| Median(Range) | 5.9(4.5-8.30) | 6.2(3.7-8) | Paired t-test=-0.261 | 0.796  |
| PT(seconds) Mean±SD Median(Range) | 12.1±0.66 12(11-13.6) | 11.99±0.59 12.2(11.5-13) | Paired t-test=1.41 | 0.17   |
| PTT(seconds) Mean±SD Median(Range) | 37.01±5.21 37(19-43.6) | 37.9±6.12 38(19-58) | Paired t-test=-1.32 | 0.2    |
| C-reactive protein (CRP)(mg/dl) Mean±SD Median(Range) | 6.57±3.98 5.9(0.9-15.5) | 1.8±1.2 1.7(0.1-4.9) | Wilcoxon signed rank test (Z)= -4.74 | <0.001  |
| Iron (mg/mL) Mean±SD Median(Range) | 93.17±39.37 78(50-180) | 95.57±43.67 78(45-189) | Paired t-test=-0.232 | 0.818  |

**Effect of using heat killed probiotic and wheat**

Combined heat killed probiotic +wheat bran administration to hemodialysis children produced highly significant decrease in all gastrointestinal symptoms (P<0.001) after 1 month, after 2 months in comparison to baseline (Table 3 & Figure 1)

Figure 1: Line graph shows comparison between total GIT symptoms scoring (before, 1 month and 2 months after heat killed probiotic+wheat.
Table 3: Comparison between gastrointestinal symptoms in the studied children (before and after 1 month, after 2 months from combined heat killed probiotic+wheat bran administration).

| Variables                          | Baseline n=35 | Month 1 n=35 | Month 2 n=35 | Friedman test | P-value |
|------------------------------------|---------------|--------------|--------------|---------------|---------|
| Anorexia Median (IQR)              | 4(3-4)        | 2(2-3)       | 2(1-2)       | 63.43         | <0.001  |
| Nausea Median (IQR)                | 2(2-3)        | 2(1-2)       | 1(1-2)       | 50.54         | <0.001  |
| Vomiting Median (IQR)              | 3(2-3)        | 2(1-2)       | 2(1-2)       | 53.01         | <0.001  |
| Abdominal discomfort Median (IQR)  | 3(3-4)        | 2(1-3)       | 1(1-2)       | 55.87         | <0.001  |
| Abdominal distension Median (IQR)  | 3(2-3)        | 2(1-3)       | 1(1-1)       | 55.22         | <0.001  |
| Heart burn Median (IQR)            | 3(2-4)        | 2(1-2)       | 1(1-2)       | 53.91         | <0.001  |
| Constipation Median (IQR)          | 3(2-4)        | 2(1-2)       | 1(1-1)       | 54.84         | <0.001  |
| Diarrhea Median (IQR)              | 1(1-2)        | 1(1-1)       | 1(1-1)       | 16.75         | <0.001  |
| Total scoring Median (IQR)         | 22(21-24)     | 15(14-17)    | 10(10-11)    | 62.64         | <0.001  |

Discussion

Uremic patients show greatly increased counts of pathogenic microorganisms in the intestine, and the absorption of proteins (and other nutrients) is hampered, so that more substrate enters the intestine; thus, more generation and absorption of uremic toxins occur, which worsens the uremic and nutritional status and accentuates gastrointestinal symptoms (GIS) [13]. Moreover, intestinal bacterial overgrowth enhances bacterial translocation and production of gut-derived endotoxins, which may increase systemic inflammation [14]. In addition, in CKD patients there is an increase in pH that promotes the growth of aerobic bacteria in the gastrointestinal tract and the subsequent production of uremic toxins. Bifidobacteria (used as probiotics) ferment carbohydrates and produce acetic and lactic acids to acidify the intestine. Hence, these bacteria prevent the growth of aerobic microorganisms, and they normalize the altered intestinal microbiome in CKD patients [1].

Previous studies have evaluated the probiotic Lactobacillus rhamnosus GG (LGG) and its effects on inflammation in human intestinal epithelial cells, specifically the Caco-2 cell line. One study demonstrated that pretreatment with both live and heat-killed LGG downregulates the production of IL-8 when an inflammatory response is generated in this cell line following cell stimulation by tumor necrosis factor-a (TNF-a) [15]. However, when an inflammatory response was not stimulated by any type of pathogenic ligand, high doses of live LGG significantly increased the production of inflammatory markers in intestinal epithelial cells, whereas heat-killed LGG only caused a slight increase [15].

This study was Open label uncontrolled clinical trial conducted on 35 children on regular hemodialysis for ESRD, we aimed to evaluate whether a combination of heat killed probiotic and wheat bran affect the presence and severity of gastrointestinal symptoms (GIS) in hemodialysis is children or not. The symptoms that were included in our study are anorexia, nausea, vomiting, heart burn, gastric discomfort, abdominal distention, constipation and diarrhea. It is one of the few in this new field of research concerning probiotic and wheat bran and their effects on renal diseases.

As regard weight we found that there was no statistically significant difference between children after and before probiotic+wheat bran, this finding agrees with a national, open, randomized, controlled intervention trial; Weight gain was marginally [16].

In our study we found that there was statistically significant change of urea while there was no change in creatinine after 2 months, this agrees with Alatriste who found statistically significant change of urea before and after using probiotic and no difference as regard creatinine [1]. RENADYL a symbiotic dietary supplement was studied in randomized clinical trials in CKD 3 and 4 and ESRD patients. Other oral therapies for removal of uremic toxins include the use of l-arginin and charcoal sorbents, these trials show that Levels of urea, uric acid, creatinine, CRP, and the lesser known toxic metabolite arising from protein putrefaction due to gut dysbiosis in CKD, can be reduced using some specific probiotic strains with improved QOL. Other methods of toxin removal by keto acid supplementation have disadvantages in comparison to probiotic [17].

This study showed that, there was statistically significant decrease of GIT symptoms (reduction of Anorexia, Nausea, Vomiting, Abdominal distension, Heart burn, Constipation, Diarrhea and Abdominal discomfort) between children after taken combined probiotic+wheat bran administration and before, matching our results Several clinical trials have been performed to assess the efficacy of probiotic in the prevention and the treatment of gastrointestinal diseases such as diarrhea and food allergies. In antibiotic associated diarrhea main studies have reported the efficacy of oral probiotic administration for reduction of duration and frequency of the disease [18]. There are several studies on frequency of defecation and gut transit time during gastroenteritis provide evidence that probiotic supplementation may have a beneficial effect on gut motility, also Gill reported a significant reduction in total and segmental colonic transit time in subjects receiving milk fermented with B. animalis compared with subjects given heat-treated bifidobacteria [19].

Xiao S [11]. studied the efficacy of live and killed probiotics in adult patients with chronic diarrhea and found killed probiotics
were more effective than live probiotics in decreasing abdominal pain, abdominal distention and improving fecal consistency.

Conclusion

We concluded that combination of heat killed probiotic and wheat bran decreases presence and severity of gastrointestinal symptoms (GIS) in hemodialysis children, thus our results could represent an important preliminary work that justifies.

Limitation of the Study

We have some limitations to our study which were mainly due to small number of available children on regular hemodialysis, so we recommend to repeat the study with a control group, and divide the testing group into three groups one receive killed probiotic, one receive wheat bran and 3rd receive combination to test the difference, in addition we recommend to test if we need to repeat the course and weather there is rebound of GIS after stoppage of medication.

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