

***In vitro* and *in vivo* biological screening of kefiran polysaccharide produced by *Lactobacillus kefiranofaciens*.**

Elsayed EA^{1,2*}, Farooq M¹, Dailin D³, El-Enshasy HA^{3,4}, Othman NZ⁴, Malek R⁴, Danial E^{5,2}, Wadaan M¹

¹Department of Zoology, Faculty of Science, King Saud University, Riyadh, Saudi Arabia

²Department of Natural and Microbial Products, Pharmaceutical Industries Division, National Research Centre, Egypt

³Institute of Bio Product Development, University Teknologi Malaysia, Malaysia

⁴Department of Bioprocess Development, City for Scientific Research and Technology Applications (CSAT), New Burg Al Arab, Alexandria, Egypt

⁵Department of Biochemistry, Science Faculty of Girls, King Abdulaziz University, Saudi Arabia

Abstract

Kefiran is a functional fermented milk product traditionally used for its beneficial probiotic properties. It exhibits antimicrobial, antioxidant, anti-inflammatory anticancer and different health promoting characteristics. Although kefiran showed potential effects against many cancer cell lines, little information is present in the literature on its effect against cervical and hepatocellular carcinoma as well as on zebrafish embryos. The study aimed at investigating the cytotoxicity (in human cervical and hepatocellular carcinoma cell lines) and developmental toxicity (in zebrafish embryos) of kefiran produced by the fermentation of *Lactobacillus kefiranofaciens*. Cervical and hepatocellular cancer cells were exposed to serial concentrations of kefiran to evaluate its cytotoxic activities. Further biological effects of kefiran on the mortality and developmental abnormalities of zebrafish embryos were investigated. Results showed that kefiran significantly affected the viability of both tested cancer cell lines in a dose-dependent manner with IC₅₀ values of 358.8 ± 1.65 and 413.5 ± 1.05 µg/ml for HeLa and HepG2 cells, respectively. Furthermore, kefiran adversely affected the morphological characteristics of the cells. Kefiran extract was much safer for zebrafish embryos and no mortality was observed up to 100 µg/ml, whereas the LC₅₀ value (≥ 279.76 µg/ml) was also very high. Moreover, no developmental toxicity was observed up to 100 µg/ml concentration. Conclusively, microbially-produced kefiran showed anticancer properties in two tested human cancer cells, while its safer profiles in animals (zebrafish embryos) poses it as potential anticancer agent which does not affect normal tissue growth.

Keywords: Anticancer, HeLa, HepG2, Kefiran, Polysaccharides, Zebrafish embryos.

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Introduction

Kefiran is a water-soluble exopolysaccharide produced by *L. kefiranofaciens*, and constitutes the matrix of the kefir grains, a fermented milk product traditionally consumed in eastern European countries [1,2]. Kefir constitutes a microbial consortium of lactic acid bacteria (*Lactobacillus*, *Lactococcus*, *Streptococcus* and *Leuconostoc*) and acetic acid bacteria (*Acetobacter*) symbiotically growing with different yeast species (*Saccharomyces*, *Candida* and *Kluyveromyces*) [3]. Kefir has many potential applications as a functional food component and exhibited numerous probiotic characteristics. Also, kefir helps in the stabilization of hypertension, decreases the levels of serum cholesterol [4,5]. Kefiran is a heteropolysaccharide having a ratio of 1:1 of glucose and

galactose and is mainly produced by the lactic acid bacteria and yeasts present in the kefir grains [6-8]. In addition, kefiran has been recognized by the US Food and Drug Administration as GRAS product (Generally Regarded as Safe), and found many potential applications in food as well as in pharmaceutical industries [2]. Moreover, kefiran has been reported to possess antibacterial, antifungal, antioxidant and anti-inflammatory properties, and has been used to lower serum cholesterol level and to modulate the immune system [9-12]. Recently, kefiran showed a gastro protective effect on ulcers induced in irradiated rats [13], and was reported to inhibit the growth of several cancer types; i.e. Ehrlich carcinoma, Lewis lung carcinoma and breast carcinoma [14,15].

Zebrafish (*Danio rerio*) are proven *in vivo* model which is best suited for drug discovery and screening of small molecules [16,17]. The embryos of zebrafish are transparent and development occurs externally, enabling an easy and thorough assessment of drug effects on internal organs in live organisms. It is known that zebrafish are used at various stages of the drug discovery process as a cost-effective alternative to some mammalian models [18-20]. Zebrafish embryogenesis is very rapid, with the entire body plan established by 24 hours post-fertilization (hpf). The zebrafish embryo is also an attractive model for studying neurogenesis as it is a vertebrate with the conserved organization of common tissues including the brain and the spinal cord. The neurogenesis starts around 10 hours post fertilization (hpf), synaptogenesis and the first behaviours around 18 hpf [21].

Little information is present in the literature on the biological activities of microbially produced kefirin on human cervical and hepatocellular carcinoma as well as in zebrafish embryos. Therefore, the present study was designed to evaluate the possible cytotoxic properties of kefirin produced by *L. kefiranofaciens* *in vitro* in human cervical cancer cells (HeLa) and human hepatocellular carcinoma cells (HepG2) and *in vivo* developmental toxicity in zebrafish embryos.

Materials and Methods

Materials

Unless otherwise stated, all chemicals, reagents and disposables were of cell culture grade and were purchased from Sigma-Aldrich Chemical Company, St. Louis, USA.

Preparation of kefiran polysaccharide

L. kefiranofaciens ATCC 8007 cells were used to produce the kefiran polysaccharide according to our previously published work [1,2]. Cells were grown in 250 ml Erlenmeyer shake-flasks for 72 h at 30°C and 200 rpm on a rotary shaker (Innova 4080, New Brunswick Scientific, NJ, USA). The optimized cultivation medium contained (g/L): lactose, 50.0; yeast extract, 12.0; KH₂PO₄, 0.25; sodium acetate, 5.0; Triammonium citrate, 2.0; MgSO₄.7H₂O, 0.2; MnSO₄.5H₂O, 0.05. The pH of the medium was adjusted to 5.5. Lactose was sterilized separately at 100°C for 20 min and was added to the cultivation medium before inoculation.

The method Piermaria et al. [22] was adapted to recover and determine the extracellularly produced kefirin. Briefly, the culture supernatant was incubated overnight with an equal volume of cold absolute ethanol at 4°C in order to precipitate kefirin. The mixture was then centrifuged at 9000 rpm for 15 min, and then the obtained precipitate was dissolved in hot distilled water and back-precipitated with ethanol. The last step was repeated three times to obtain kefirin in a pure form. The finally precipitated pure kefirin was collected, dried at 65°C for 48 h, and then stored for further steps.

The purified kefirin was dissolved in DMSO to prepare the stock solution (1 mg/ml), which was used to prepare a series of

different working solutions (0-1000 µg/ml) using DMEM medium. The working solutions were aseptically filtered using 0.22 µm sterile syringe filters (Millipore, USA).

Cell lines and cultivation conditions

Human cervical cancer (HeLa) and human hepatocellular carcinoma (HepG2) cells were obtained from Sigma-Aldrich Chemical Company, St. Louis, USA. Cells were grown on DMEM medium containing foetal bovine serum (10%), penicillin/streptomycin solution (100x, 1%) and NaHCO₃ (3.6 g/L). According to standard cell culture protocols, cells were routinely sub-cultured and in a humidified CO₂ incubator (ShelLab, USA) at 5% CO₂, 37°C and 95% humidity. Viable cell concentration as well as cell viability were assessed using the Trypan blue exclusion method [23,24].

Cytotoxicity assay

Standard MTT (3-(4, 5-Dimethylthiazol-2-yl)-2, 5-diphenyl) cytotoxicity assay [25] was used to determine the percentage of cell viability. Cells were firstly treated with trypsin, washed and then resuspended. Only cells having viability scores higher than 95% were used to perform the cytotoxicity assays. 96 well culture plates were inoculated with cells to reach a concentration of 10⁴ cells/100 µl/well, and were then allowed to grow for 24 h. The consumed medium was then aspirated and replaced with fresh preparations containing different concentrations of kefirin working solutions, and the cells were then grown for another 24 h. DMSA only at a final concentration (≤ 0.5%) served as the control. Firstly, the plates were examined for morphological changes using an inverted contrast microscope (Nikon Eclipse T500, Japan, and 10x). Cells were then treated with MTT (10 µl, 5 mg/ml in PBS) for 4 h, and the resulting formazan crystals were dissolved with 200 µl of DMSO. The absorbance was read at 550 nm using a micro plate reader (Thermo Scientific, USA). Cell viability was calculated as a percentage of the control value. The concentration resulting in 50% inhibition of cell growth was referred to the IC₅₀ value, and was determined from the linear regression of the calibration curve.

Zebrafish embryos

Wild type zebrafish (AB/Tuebingen TAB-14) and transgenic TG (Fli;1:EGFP) y1 [16] were obtained from the zebrafish International Resource Centre (ZIRC University of Oregon, Oregon, USA) and maintained in the animal facility at Bio products research chair, Department of Zoology at King Saud University. The adult tropical zebrafish were kept under the standard laboratory conditions of 28.5°C on a 14-h light/ 10-h dark photoperiod in fresh water (FW) which consists of reverses osmosis water supplemented with a commercially available salt solution (0.6% Instant Ocean) according to the standard guidelines that are described in the literature [17]. All experiments were carried out in accordance with the National and International animal use guidelines and were in accordance with the ethical guidelines of the College of Science, King Saud University.

Animal treatment

The wild type (AB Tubingen) and transgenic Tg (fli-1:EGFP) zebrafish embryos were obtained by natural pairwise mating and raised up to the shield stage (6 hours post fertilization). The extracted kefiran was re-suspended in the cell culture grade DMSO (D8418 Sigma LLC., St. Louis, USA) to prepare a stock concentration of 20 mM. The synchronized embryos (all embryos were at the same stage of development) were treated with a serial dilution of kefiran in order to assess both the toxicity and developmental abnormalities. Approximately thirty (30) embryos were placed in sterile 60 mm Petri dishes that contained 10 ml of the embryo medium (5.0 mM NaCl, 0.17 mM KCl, 0.33 mM CaCl, 0.33 mM MgSO₄) at the desired concentration of kefiran. The 1% (v/v) DMSO treated embryos served as controls. The embryos were then incubated overnight at 28°C in an air incubator. From the second day onward, the embryo medium containing the kefiran was changed daily. Three biological replicate trials (each clutch of the embryo was from different adult pair of fish) were carried out for each experiment [20].

Microscopic examination

All images were acquired using fluorescent stereo microscope Olympus ZX12 with DP72 camera using CellSens standard software (Olympus Capital Holdings Asia Pte Ltd. Singapore).

Statistical analysis

Data were analysed using SPSS 9.0, and the obtained results are represented as mean ± SD of three experiments. One-way ANOVA analysis of variance and student t-Test was used to compare between different experimental groups and data were considered statistically significant for P values less than 0.05.

Results

Evaluation of in vitro cytotoxicity of kefiran polysaccharides

The kefiran polysaccharides produced by *L. kefiranofaciens* were evaluated for its *in vitro* cytotoxic properties against HeLa and HepG2 cells using standard MTT assay. Figure 1 shows the cytotoxic effect of kefiran, which was dose- and cell type-dependent. The IC₅₀ values for HeLa and HepG2 cells were 358.8 ± 1.65 and 413.5 ± 1.05 µg/ml, respectively. Increasing kefiran concentration significantly increased the cytotoxicity of kefiran to HeLa and HepG2 cells (p<0.001). The highest kefiran concentration (1 mg/ml) significantly decreased cell viability by about 72.25 and 81.85% recording 27.75 ± 1.31 and 18.15 ± 0.88% for HeLa and HepG2 cells, respectively. Moreover, the results showed that HeLa cells were not significantly affected when being tested with 15.6 or 31.3 and 125 µg/ml of kefiran. On the other hand, HepG2 cells showed no significant changes when treated with kefiran concentrations ranging from 0.0 to 125 µg/ml.

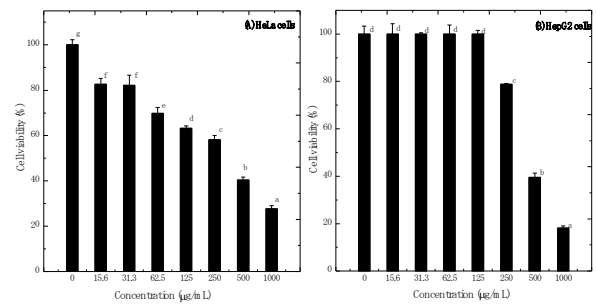


Figure 1. Effect of different concentrations of kefiran microbially produced by *L. kefiranofaciens* on the viability of HeLa and HepG2 cells. Data are presented in Mean ± SD. Different letters with the same cell type represent a significant difference (p<0.05).

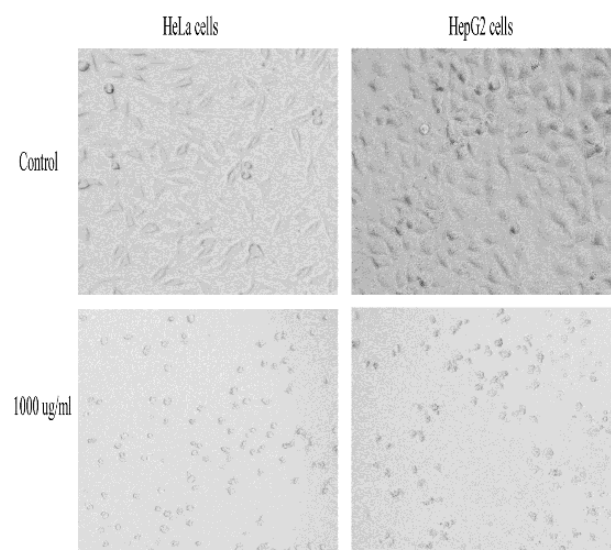


Figure 2. Effect of kefiran on morphological characteristics of HeLa and HepG2 cells after 24 h. Images were captured using inverted contrast microscope at 10x magnification.

Figure 2 represents the effect of variable concentrations of kefiran on the morphological characteristics of both HeLa and HepG2 cells compared to control untreated cells. Increasing kefiran concentration drastically affected the morphological characteristics of both cell lines, with the maximal effect observed upon using the highest kefiran concentration (1 mg/ml). Upon increasing kefiran concentration, cells started to shrink, lost their adherence capacity and finally started to float in the cultivation flask. The maximal kefiran concentration resulted in the complete rounding-up of the cells with the formation of vacuoles leading finally to their death.

In vivo cytotoxic effects of kefiran polysaccharide in zebrafish embryos

In order to determine the *in vivo* developmental toxicity of kefiran in zebrafish embryos, the embryos were exposed to serial dilution of kefiran ranging from 1.0 to 1000 µg/ml. The zebrafish embryos responded dose dependently as depicted in

Figure 3. Results showed that mortality percentage increased with the increase of kefir concentration, where the maximum concentration of kefir (1 mg/ml) resulted in about 80% mortality, and the LD50 concentration was $\leq 279.76 \mu\text{g/ml}$.

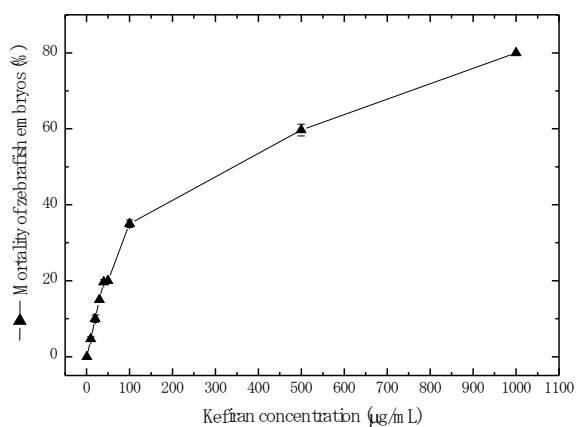


Figure 3. Effect of different concentrations of kefir microbially produced by *L. kefirifaciens* on the mortality of zebrafish embryos.

Figures 4A-4D represent images taken during the assessment of developmental abnormalities in zebrafish embryos induced by kefir (at 10-30 $\mu\text{g/ml}$) in comparison with mock treated embryos at the same stage. The zebrafish embryos hatched normally at 48-52 hpf and started to swim normally directly after being hatched. The brain has developed normally in all treated embryos (Figure 4B) and all the brain structures (fore brain FB, mid brain MB and hind brain HB) along with mid brain hind brain boundary MHB developed normally and there was no obvious abnormality in brain formation compared to control (Figure 4B).

The zebrafish transgenic line TG (fli-1; EGFP), expressing the enhanced green fluorescent protein (EGFP) in endothelial cells under the promoter of fli1 [16], is routinely used to screen antiangiogenesis molecules. This transgenic zebrafish line was also used in the current work to validate whether kefir disrupted the formation of angiogenic blood vessels.

Figure 4 shows representative live images of 3 dpf transgenic zebrafish embryos control (C) and treated with kefir (D). In control-as well as kefir-treated embryos, it can be clearly observed that the intersegmental blood vessels (ISV) sprouted from the dorsal aorta (DA) and extended along with the boundaries of somites and connected to the dorsal longitudinal anastomotic vessel (DLAV, red arrow).

Similarly, the second angiogenic blood vessels, which form the sub intestinal vein (SIV white arrows) and are composed of an arcade of 10 to 12 vessels arranged as a basket like structure on the yolk, developed normally in treated embryos at 72 hpf. Therefore, it can be concluded that treating zebrafish embryos at the tested concentration did not induce any abnormalities during embryonic development.

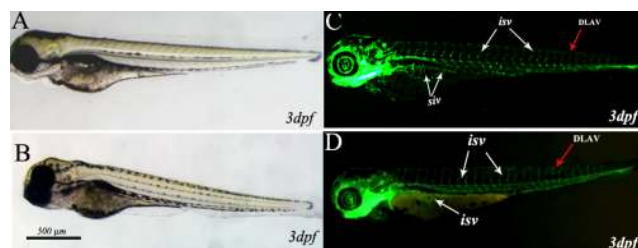


Figure 4. Effect of kefir on morphological and developmental characteristics of wild and transgenic zebrafish embryos. A and C: Controls wild and transgenic, respectively. B and D: Treated wild and transgenic, respectively. Images were acquired using a fluorescent stereo microscope (Olympus ZX12).

Discussion

Kefir is a functional fermented milk product with a wide range of probiotic properties. Regardless of its nutritional value, kefir exhibits different biological activities [9,10,26]. Recently, many studies were conducted to evaluate the cytotoxic activities of kefir against different cancer cell types and animal models. However, little information can be traced in the literature concerning studies on the evaluation of kefir biological activities *in vitro* and *in vivo* on cervical as well as hepatocellular cancers and zebrafish embryos, respectively.

Kefir exhibited a significant cytotoxic activity against both HeLa and HepG2 cell lines. Kefir reduced the cell viability of HeLa and HepG2 cells up to 72.25 and 81.85%, respectively, upon applying the highest kefir concentration (1 mg/ml). Additionally, the IC50 values recorded for HeLa and HepG2 cells were 358.8 ± 1.65 and $413.5 \pm 1.05 \mu\text{g/ml}$, respectively. Polysaccharides produced by microorganisms have been long used for their potential biological properties [27,28]. The obtained results are in good agreement with those reported earlier reporting anticancer activities of kefir against Ehrlich, lung and breast carcinoma [15,29-31]. Recently, Maalouf et al. [32] obtained about 88.1 and 86.7% reduction in cell viabilities of CEM and Jurkat cells after 24 h of treatment with 60 $\mu\text{g}/\mu\text{L}$ of kefir. However, they were not able to determine the IC50 values for kefir because of the high viscosity of kefir. The anticancer effect of kefir can be attributed to the effect of the exopolysaccharide, i.e. kefiran, lentinan, viilian. β -1, 3-Glucans with 1, 6-glucopyranoside branching have been reported to produce effector activities in tumour synergic cell cytotoxicity [33,34]. Furthermore, our results showed that both tested cell lines responded differently to the kefir polysaccharide. This can be explained by the fact that different types of cancer cells differ in their specificity and selectivity arising from differences in cell morphology and membrane structures [24,35,36].

The *in vivo* toxicity profile of kefir on zebrafish embryos could be correlated with the *in vitro* cytotoxicity, with an LD50 value of 279.76 $\mu\text{g/ml}$. This could be considered as a very high concentration of a compound to affect the zebrafish embryos. Generally, the toxicity profile of active compounds lies within the range of 1-30 $\mu\text{g/ml}$. The transgenic zebrafish has been used for the evaluation of the antiangiogenic properties of

newly synthesized compounds [20]. We also tested the effect of kefiran on zebrafish blood vessels formation. As shown in Figure 4, kefiran did not induce any abnormality in zebrafish angiogenic blood vessels formation and development. Moreover, we could not observe any developmental abnormality or teratogenic profile of Kefiran in zebrafish embryos which mean that it turned out to be very safe in animals. Furthermore, the results concerning the effect of kefiran on the development of zebrafish embryos showed that kefiran has no obvious effect. Our results coincide with those reported by Kang et al. [37,38]. They evaluated the antioxidant activity of polysaccharides purified from *Acanthopanax koreanum* Nakai and aloe vera gel in zebrafish model, and found no effect on the survival rate of zebrafish embryos. Moreover, they reported a protective effect when the embryos were pre-treated with the polysaccharide prior to oxidative stress induction. Recently, Guven et al. [39] investigated the effect of kefir on spinal cord injury ischemia in rats. They concluded that kefir has a neuroprotective and anti-oxidant effects spinal cord ischemia and injury, where lactic acid bacteria produce bioactive peptides, which capture the reactive oxygen species and thus inhibit the formation of malondialdehyde, responsible for cell membrane damage mediated by lipid peroxidation.

Additionally, our results can be explained on the basis that the routine protocol depends on sub lethal concentrations to evaluate the developmental abnormalities in zebrafish embryos [19,20]. The sub lethal concentration of kefiran (10-100 µg/ml) did not induce any developmental abnormality which could be considered as safer in zebrafish embryos. The higher toxicity profile, which was observed with kefiran used in this study, could be attributed to the fermentation process as it was separated and extracted from a fermentation culture broth obtained. This in turn can explain the slightly higher data regarding cell toxicity and mortality of zebrafish embryos, in comparison to pure kefiran compounds used in previous studies.

Conclusion

The present investigation provides sufficient evidences that kefiran polysaccharide produced by *L. kefiranofaciens* showed a statistically significant level of anti-proliferative effect on human cervical cancer cells (HeLa) and human liver hepatocarcinoma cells (HepG2). Moreover, the anticancer profile was not only dose dependent, but also cancer cell type specific. Furthermore, kefiran did not induce any developmental toxicity in zebrafish embryos, at sub lethal concentrations. The anticancer profile specifically only in human cancer cells and safer behaviour of kefiran in zebrafish embryos warrants its potential anticancer properties. Furthermore, considering the lack of information on the effect of kefiran on the development of zebrafish embryos, these results can provide a preliminary platform for future research.

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***Correspondence to**

Elsayed AE,
Department of Zoology
King Saud University
Riyadh
Saudi Arabia