

Antiproliferative Activity of Kenyan *Trametes versicolor* Aqueous Extract on Selected Cancer and Normal Cell Lines

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Abstract

Cancer is a major public health burden in both developed and developing countries. The current conventional cancer therapies like chemotherapy are expensive and inaccessible to many cancer patients. Commercial and wild edible mushrooms are becoming more important for their nutritional value and are becoming an alternative source of immune modulation and anticancer agents. Although Previous studies with *Trametes versicolor* mushroom from various parts of the world have demonstrated antiproliferative activity on various cancer cell lines, the antiproliferative activity of the recently identified Kenyan *T. versicolor* mushroom have not been studied. This study examined the *in vitro* antiproliferative activity of an aqueous extract of the Kenyan *T. versicolor* mushroom on breast cancer (4T1), prostate cancer (DU145), hepatocellular carcinoma (HCC), rat normal intestinal epithelial cells (IEC-6) and African green monkey normal kidney (vero) cell lines using MTT assay. The results demonstrated that the *T. versicolor* extract at 1.37 µg/ml to 1000 µg/ml dose-dependently inhibited the proliferation of DU145 and 4T1 cell lines with IC₅₀ values: DU145 (71.2 µg/ml) and 4T1 (188.5 µg/ml). The extract however did not exert any significant antiproliferative effect on HCC, IEC-6 and Vero cell lines (IC₅₀>1000 µg/ml) when compared with a chemotherapeutic anticancer drug, tamoxifen (p<0.05), confirming the tumor-selective cytotoxicity on cancer cell lines and its safety on normal cell lines. In all cell lines, the extract showed a significant difference in inhibition of cell proliferation between the untreated cells and the highest concentration (1000 µg/ml) (p<0.05). Presence of phytochemicals such as saponins, tannins, steroids, terpenoids and flavonoids in the *T. versicolor* extract used might be the probable reason for its antiproliferative activity.

Keywords: *Trametes versicolor*; Antiproliferation; Cancer; Cell line; Inhibition; Viability

Introduction

Cancer is a disease characterized by irregular proliferation of cells with a manifestation of malfunctions in immunity as malignant cells manage to escape recognition and elimination by the immune system [1]. Cancer cells display a broad spectrum of genetic alterations that may include gene rearrangements, point mutations, and gene amplifications, leading to disturbances in molecular pathways regulating cell growth, survival and metastasis [2]. Anyone can develop cancer and the risk of being diagnosed with cancer increases with age. About 77% of all cancers are diagnosed in people 55 years of age and older [3].

Chemotherapy, surgery, immunotherapy and radiotherapy remain to be the most effective conventional methods in cancer treatment. However, they are expensive and can cause serious side effects as these do not often show adequate differential effect between tumor and normal cells [2], hence the need for affordable but effective ways of cancer management.

More than 50% of all modern drugs in clinical use are of natural product origins, many of which have antiproliferative ability [4]. According to World Health Organization estimates, more than 80% of people in developing countries depend on traditional medicine for their primary health needs including cancer management [5]. A survey done in the year 2008 showed that more than 60% of cancer patients use herbs as therapy [6]. An attempt has been made to explore the potential of newly discovered anticancer compounds from medicinal mushrooms as a lead for anticancer drug development [5]. Mushrooms have key medicinal uses including anticancer activity [7], immunomodulating effects and antiproliferative activity [8]. *Trametes versicolor* mushroom, also called "Turkey tail" has been widely studied

and its antiproliferative activity on different cancer cell lines has been well documented [9-11].

In Japan, polysaccharide-Krestin (PSK) extracted from *T. versicolor* mushroom is prescribed to cancer patients routinely, both during and after radiation and chemotherapy [12-15]. In the United States of America, whole, freeze-dried *T. versicolor* is commonly prescribed to breast cancer patients [16].

In some studies done in China to test for the antiproliferative activity of *T. versicolor* extract, results showed that the extract could inhibit the proliferation of four breast cancer cells (T-47D, Bcap37, ZR75-30, MCF-7), B-cell lymphoma (Raji), human promyelocytic leukemia (HL-60, NB-4) and liver cancer cell line (7703) [17,18]. However, the extracts of *T. versicolor* mushroom have been reported to be non-toxic to several normal cells [19-21] and in cancer patients on clinical trials [22-24]. Differences in the ecological zones where mushroom exist and the influence of some environmental factors may account for the overall composition and activity of compounds present in mushrooms [25,26]. This study therefore investigated the antiproliferative potential

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of the recently identified Kenyan *T. versicolor* mushroom against breast cancer (4T1), prostate cancer (DU145), hepatocellular carcinoma (HCC), rat normal intestinal epithelial cells (IEC-6) and African green monkey normal kidney (vero) cell lines. We report significant anti proliferative activity of *T. versicolor* aqueous extract on DU145 and 4T1 cell lines, with insignificant antiproliferative activity on HCC, IEC-6 and vero cells.

Material and Methods

Reagents

The cells used were Breast cancer cell line (4T1), hepatocellular carcinoma cell line (HCC), human prostate cancer cell line (DU145), African green monkey kidney cell line (Vero) and rat normal intestinal epithelial cell line (IEC-6) which were obtained from the center for traditional medicine and drug research (CTMDR) at Kenya medical research institute (KEMRI). The RPMI-1640 medium and fetal calf serum were purchased from Gibco Laboratories (Grand Island, NY, USA). The MTT (3-(4,5-dimethylthiazolyl)-2, 5-diphenyl-tetrazolium bromide), dimethyl sulfoxide (DMSO) and other chemicals used were purchased from Sigma Chemical Co. (St. Louis, MO, USA).

Collection and preparation of mushrooms

Trametes versicolor were botanically identified by a botanist and collected from Arabuko Sokoke forest at the Kenyan coast and evaluated at KEMRI laboratories. Voucher specimens were deposited at the University of Nairobi herbarium. The mushroom *T. versicolor* were cut into small pieces and dried at room temperature. The dried materials were ground in a food grinder (mincer) to a fine powder, packed in bags, and stored at room temperature until use.

Aqueous extraction of the mushrooms

Aqueous extraction of *T. versicolor* was done according to Piero [27] and Kigondu [28] with modifications. Briefly, 100 g of the previously prepared powder of the mushroom *T. versicolor* was measured using an electrical beam balance and put into a 500 ml conical flask. Double distilled water was added until the sample was completely submerged. The mixture was then transferred into a water bath at 80°C for 3 hours and cooled to room temperature. The mixture was then filtered using Whatman no.1 filter paper into a clean sterile 1000 ml conical flask and the solvent removed using a freeze drying machine. The resulting extract was weighed and stored in an airtight bottle at -20°C until use.

Antiproliferative activity

The antiproliferative activity of the mushroom extract was evaluated using MTT (3-(4,5-dimethylthiazolyl)-2, 5-diphenyl-tetrazolium bromide) assay (Sigma, USA). The assay detects the reduction of MTT by mitochondrial dehydrogenase to blue formazan product, which reflects the normal function of mitochondria and cell viability [18]. Formazan, which is an insoluble yellow colored product can be measured spectrophotometrically [29,30] and the activity of mitochondrial dehydrogenase enzyme to produce formazan is directly and inversely proportional to the level of cell viability and inhibition respectively [31].

Cells were maintained as monolayer cultures in RPMI-1640 medium supplemented with 10% fetal bovine serum and incubated at 37°C in a humidified incubator at 5% CO₂. Five types of cells were seeded in 96-well plates with the final volume 100 µl containing 24,000 cells per well, with the cell density determined using trypan blue exclusion assay. The fourth, eighth and twelfth columns of the 96 well

plates contained 100 µl RPMI-1640 with no cells as they were treated as blanks for each cell line. The plates were incubated at 37°C for 48 hours. The *T. versicolor* extract in PBS was proportionally diluted with RPMI-1640, and 50 µl of each solution was added to triplicate wells in the concentrations: 1000 µg/ml, 333.33 µg/ml, 111.11 µg/ml, 37.04 µg/ml, 12.35 µg/ml, 4.12 µg/ml and 1.37 µg/ml with the last row of the 96 well plate left untreated to serve as a cell control. After 48 hours incubation in 5% CO₂ humidified environment at 37°C, a volume of 10 µl of PBS containing 5 mg/ml MTT was then added into each well including all the controls and plates further incubated for 3 hours. The medium was removed and 100 µl DMSO added into each well. After the plates were shaken mildly, the absorbance of the samples was measured at 560 nm with a Multiskan Spectrum Microplate Spectrophotometer. Determination of cell proliferation was achieved using the formula developed by Patel et al. [30] and Awasare et al. [32], as follows;

$$\text{Proliferation rate} = \frac{At - Ab}{Ac - Ab}$$

$$\text{Percentage viability} = \frac{At - Ab}{Ac - Ab} \times 100$$

$$\text{Percentage inhibition} = 100 - \left(\frac{At - Ab}{Ac - Ab} \times 100 \right)$$

Where At=Absorbance value of test compound

Ab=Absorbance value of Blank

Ac=Absorbance value of negative control (untreated Cells)

Statistical analysis

Results were expressed as the mean ± standard error of the mean (SEM) for all the different concentrations of the mushroom extracts. Statistical differences between the concentration means were assessed by One-way ANOVA and the means subjected to Tukey's post-hoc test for pairwise comparison with values of p<0.05 considered as statistically significant. Generated dose response curves were used to determine IC₅₀ values. Student's unpaired t-test was then used to compare the IC₅₀ values of *T. versicolor* to that of Tamoxifen. Data analysis was done using Excel spread sheets and Minitab statistical software version 17.0.

Qualitative phytochemical analysis of *Trametes versicolor* extract

A qualitative phytochemical analysis of the crude mushroom extract was performed using standard protocols described by Odebiyi and Sofowora [33].

Test for alkaloids: Weight of 0.5 g of the extract was stirred with 5 ml of 1% aqueous hydrochloric acid for two minutes on a steam water bath. The mixture was filtered and few drops of Dragendorff's reagent was added. The sample was then observed for color change or turbidity to draw inference.

Test for saponins: The persistent frothing test for saponin described by Odebiyi and Sofowora [33] was used. A volume of 30 ml distilled water was added to 1 g of the mushroom extract. The mixture was vigorously shaken and heated on a steam water bath. The sample was observed for the formation of stable froth to draw inference.

Test for phlobatannins: Weight of 0.2 g of the mushroom extract was dissolved in 10 ml of distilled water and filtered. The filtrate was then

boiled with 2% hydrochloric acid solution and observed for deposition of red precipitate which indicates the presence of phlobatannin.

Test for tannins: The method of Trease and Evans [34] was adopted where 0.5 g of the mushroom extract was dissolved in 5 ml of distilled water, then, boiled gently and cooled. One ml of the solution was dispensed in a test tube and 3 drops of 0.1% ferric chloride solution were added and observed for brownish green or blue black colouration which indicates the presence of tannins.

Test for terpenoids: The Salkowski test was used. A weight of 5 g of the extract was dissolved in 5 ml distilled water. The mixture was then added in 2 ml of chloroform, and 3 ml concentrated sulphuric acid was carefully added to form a layer. The solution was then observed for reddish brown colouration which confirms the presence of terpenoids.

Test for steroids: A volume of 2 ml acetic anhydride was added to 0.5 g of the mushroom extract and filtered. Two ml of Sulphuric acid was added to the filtrate and observed for colour change from violet to blue or green, which indicates the presence of steroids.

Test for flavonoids: A volume of 5 ml diluted ammonia solution was added to 0.5 g of the mushroom extract dissolved in 5 ml distilled water. This was then followed by the addition of a concentrated sulphuric acid. The solution was observed for yellow colouration that disappears on standing to confirm the presence of flavonoids.

Test for anthraquinones (Borntrager's test): A weight of 0.5 g of the mushroom extract was shaken with 10 ml of benzene, filtered and 5 ml of 10% ammonia solution added to the filtrate. The mixture was then shaken and observed for the presence of pink red or violet colour in the ammonia layer which indicates the presence of free anthraquinones.

Phytochemical extract	<i>T. versicolor</i> aqueous
Saponin	+
Tannin	+
Steroid	+
Alkaloid	-
Terpenoid	+
Flavonoids	+
Anthraquinone	-
Phlobatannin	-

The (+) sign denotes presence of phytochemicals while (-) sign denotes absence of phytochemicals.

Table 1: Qualitative phytochemical screening of *T. versicolor* aqueous extract.

Concentration (µg/ml)	% Viability					% Inhibition				
	4T1	DU145	HCC	IEC-6	Vero	4T1	DU145	HCC	IEC-6	Vero
Untreated cells	100	100	100	100	100	0	0	0	0	0
1.37	96	84	95	92	97	4	16	5	8	3
4.12	94	74	90	89	94	6	26	10	11	6
12.35	83	68	89	86	92	17	32	11	14	8
37.04	70	52	85	70	88	30	48	15	30	12
111.11	60	41	81	65	86	40	59	19	35	14
333.33	31	35	65	60	74	69	65	35	40	26
1000.00	22	9	59	56	57	78	91	41	44	43

Results are expressed as the Mean of three independent experiments of three wells each which were calculated from the OD values obtained from a spectrophotometer.

Table 2: Percentage viability and percentage inhibition for effect of aqueous extract of *T. versicolor* mushroom on 4T1, DU145, HCC, IEC-6 and vero cell lines.

Cell line	Drug	IC ₅₀ (µg/ml) ± SEM
4T1	1	188.53 ± 4.81 ^a
4T1	2	163.33 ± 8.82 ^a
DU145	1	71.16 ± 3.48 ^b
DU145	2	63.33 ± 3.48 ^{bc}
IEC-6	2	42.20 ± 5.64 ^{cd}
HCC	2	41.37 ± 5.49 ^{cd}
Vero	2	36.46 ± 2.69 ^d
IEC-6	1	-
HCC	1	-
Vero	1	-

1= *T. versicolor*, 2=Tamoxifen (Reference drug).

Results are expressed as the Mean ± SEM of three independent experiments of three wells each. Values that do not share a superscript are significantly different (P>0.05). i.e: IC₅₀ values on 4T1 cells treated with drug 1 and drug 2 are statistically not different. Drug 1 (*T. versicolor*) did not produce IC₅₀ values on IEC-6, HCC and vero cell lines (IC₅₀ >1000 µg/ml).

Table 3: Statistical comparison on the IC₅₀ values between *T. versicolor* extract and Tamoxifen (reference drug) on the 5 cell lines.

Results

Qualitative phytochemical analysis of *T. versicolor* aqueous extract

Results from the phytochemical analysis of the aqueous extracts of the Kenyan *T. versicolor* mushroom revealed the presence of saponins, tannins, steroids, terpenoids and flavonoids while anthraquinones, alkaloids and phlobatannins were absent (Table 1).

Antiproliferative potential of *T. versicolor* aqueous extract

This study evaluated the antiproliferative potential of *T. versicolor* aqueous extracts on five cell lines namely 4T1, DU145, HCC, IEC-6 and Vero cell lines. The *T. versicolor* aqueous extract at 1.37 µg/ml to 1000 µg/ml exhibited dose-dependent inhibitory effects on the proliferation of DU145 and 4T1 cells with more than 90% and 70% suppression respectively at the highest concentration (Table 2). However, the extract induced a low level of suppression on the proliferation of HCC, normal IEC-6 and normal Vero cells showing values of 41.19%, 43.85% and 42.64% respectively at the highest concentration (Table 2). Table 3 shows the concentrations producing 50% growth inhibition (IC₅₀) of the *T. versicolor* extract on the five cell lines of which DU145 proliferation was most potently suppressed with the lowest IC₅₀ value (71.16 µg/ml) followed by 4T1 cells (IC₅₀ =188.53 µg/ml) after incubation with the *T. versicolor* extract. The *T. versicolor* extract did not record any IC₅₀ values against HCC, vero and IEC-6 cell lines even at the highest concentration (1000 µg/ml). The IC₅₀ values of *T. versicolor* against all cell lines used were statistically compared to those of the reference drug (Tamoxifen) where results revealed that there was no statistical difference between the IC₅₀ values of *T. versicolor* extract and Tamoxifen against DU145 and 4T1 cells (Table 3).

All cell lines treated with the aqueous extract of *T. versicolor* had a general reduction in their viability with the extract showing a general increase in the percentage inhibition on all cell lines used (Table 2). The mushroom extract demonstrated a dose dependent inhibition of cell proliferation in all cell lines used. As the concentration of the mushroom extract increased from 1.37 µg/ml to 1000 µg/ml, the percentage viability decreased, showing the lowest viability at 1000 µg/ml and the highest in untreated cells with HCC and DU145 cell lines showing the highest and lowest viability respectively (Table 2). A general reduction on the proliferation rate against all cell lines used was recorded. As the concentration of the mushroom extract increased,

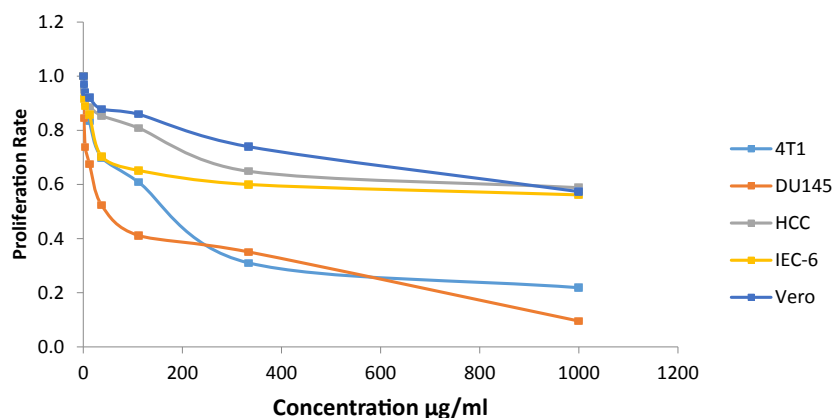


Figure 1: Dose response curve showing the effect of aqueous extract of *T. versicolor* on the proliferation rate on 4T1, DU145, HCC, IEC-6 and Vero cell lines.

Concentration (µg/ml) <i>T. versicolor</i>	Proliferation rate Cell line				
	4T1	DU145	HCC	IEC-6	Vero
Untreated cells	1.00 ± 0.02 ^a	1.00 ± 0.01 ^a	1.00 ± 0.01 ^a	1.00 ± 0.01 ^a	1.00 ± 0.01 ^a
1.37	0.96 ± 0.01 ^a	0.84 ± 0.00 ^{ab}	0.95 ± 0.00 ^{ab}	0.92 ± 0.01 ^b	0.97 ± 0.01 ^{ab}
4.12	0.94 ± 0.02 ^a	0.74 ± 0.04 ^{bc}	0.90 ± 0.01 ^{abc}	0.89 ± 0.00 ^b	0.94 ± 0.01 ^{abc}
12.35	0.83 ± 0.02 ^{ab}	0.68 ± 0.01 ^{cd}	0.89 ± 0.02 ^{bc}	0.86 ± 0.00 ^b	0.92 ± 0.01 ^{bcd}
37.04	0.70 ± 0.01 ^{bc}	0.52 ± 0.02 ^{de}	0.85 ± 0.02 ^{bc}	0.70 ± 0.01 ^c	0.88 ± 0.03 ^{cd}
111.11	0.61 ± 0.04 ^c	0.41 ± 0.05 ^{ef}	0.81 ± 0.01 ^c	0.65 ± 0.03 ^{cd}	0.86 ± 0.01 ^d
333.33	0.31 ± 0.01 ^d	0.35 ± 0.01 ^f	0.65 ± 0.01 ^d	0.60 ± 0.06 ^{de}	0.74 ± 0.00 ^e
1000.00	0.22 ± 0.01 ^e	0.10 ± 0.01 ^g	0.59 ± 0.02 ^d	0.56 ± 0.02 ^e	0.57 ± 0.02 ^f

Proliferation rates in the same cell line were compared to each other among the different concentrations of the extract. Results are expressed as the Mean ± SEM of three independent experiments of three wells each. Values followed by the same superscript in the same cell line are not significantly different ($P > 0.05$). In each cell line, the proliferation rate on the untreated cells is statistically different to that on the highest concentration (1000 µg/ml). i.e: Values followed by superscript 'a' and 'e' on 4T1 cell line are statistically different.

Table 4: Effect of aqueous extract of *T. versicolor* mushroom on the proliferation rate on 4T1, DU145, HCC, IEC-6 and Vero cell lines.

the proliferation rate decreased with DU145 and 4T1 cells showing the lowest proliferation rates at the highest concentration (1000 µg/ml). In all cell lines, the proliferation rate was lowest at 1000 µg/ml and the highest in untreated cells (Figure 1). There was a significant difference between proliferation rate at 1000 µg/ml and untreated cells in all cell lines. ($P < 0.05$) (Table 4).

Discussion

The aqueous extract of the Kenyan *T. versicolor* mushroom used in this study exhibited antiproliferative activity on two cancer cells with high safety levels on normal cells. Previous studies have shown that aqueous extracts of *T. versicolor* mushroom contain natural polysaccharides which have been proven to have some anticancer activity [35]. This study evaluated the antiproliferative activity of aqueous extracts of Kenyan *T. versicolor* mushroom on normal and cancer cell lines namely, 4T1, DU145, HCC, IEC-6 and Vero.

The findings of this study revealed that aqueous extracts of the Kenyan *T. versicolor* mushroom significantly inhibited the proliferation of breast cancer cells (4T1) and human prostate cancer cells (DU145) in a dose dependent manner. Such antiproliferative activity of the *T. versicolor* extract was reflected by its high percentage inhibition (Table 2) and the relatively low IC_{50} values which were significantly not different to those of Tamoxifen, the reference drug used (Table 3). These findings are consistent with previous studies where ethanol-water extract of *T. versicolor* was reported to have significant inhibitory effects on human breast cancer cells, prostate cancer, human leukemia, and lymphoma cells [18]. In another study done by Cheong-Yip et al. [36], ethanol-water extract of *T. versicolor* exhibited significant antiproliferative

activity on three human breast cancer cell lines (MDA-MB-231, MCF-7 and T-47D) in a dose-dependent manner *in vitro*. Carolyn achieved similar findings et al. [37] in a clinical trial on women with breast cancer where oral preparations of *T. versicolor* administered to the subjects showed significant antiproliferative activity.

However, the aqueous extracts of *T. versicolor* in the present study showed a very low inhibition on the proliferation of human liver cancer cells (HCC), rat normal intestinal epithelial cells (IEC-6) and African green monkey normal kidney cells (Vero). This is shown by their high IC_{50} values where even the highest concentration of the mushroom extract used could not inhibit at least 50% of the cells (Table 3). In a study by Xuannwei et al. [17] on polysaccharides isolated from *T. versicolor* mushroom, the results recorded no antiproliferative activity on three human liver cancer cells (HepG2, 7721, PLC) and human normal liver cells (WRL) which corresponds with the results of this study as *T. versicolor* extract failed to inhibit the proliferation of human cancer liver cells (HCC). Xuannwei et al. [17] in their study recorded that the polysaccharide extract of *T. versicolor* showed antiproliferative activity on four breast cancer cell lines (T-47D, Bcap37, ZR7-30, MCF-7) while it inhibited the proliferation of only one liver cancer cell line (7703) out of four liver cancer cell lines used. This therefore might be suggestive that *T. versicolor* extract acts on liver cancer cells selectively.

The results of this study suggest that the aqueous extract of the Kenyan *T. versicolor* mushroom is more actively in the inhibition of breast and prostate cancer cells than human liver cancer cells and normal cell lines. This was reflected by the low IC_{50} values of 71.16 µg/ml and 188.53 µg/ml on prostate cancer (DU145) and breast cancer cells

(4T1) respectively compared to liver cancer cells (HCC), rat normal intestinal epithelial cells (IEC-6) and African green monkey normal kidney cells (Vero) both of which had an IC_{50} value >1000 $\mu\text{g/ml}$ (Table 3). These results tend to agree with previous reports that *T. versicolor* extracts can selectively suppress the proliferation of various cancer cells with no antiproliferative activity in normal cell lines suggesting its high level of safety [17,18,36]. In all cell lines used in this study, there was a significant difference in the proliferation rate between the untreated cells and the highest concentration (1000 $\mu\text{g/ml}$) of the *T. versicolor* extract used (Table 4). This tendency however suggests that the extract used in this study has a dose-dependent activity on the cell lines used.

The antiproliferative activity of the *T. versicolor* aqueous extract used in this study can be associated with the phytochemicals found to be present in the extract. Saponins, flavonoids and Tannins, which were present in the extract have been documented from previous research to possess some anticancer activity on different cancer cell lines [38-40]. Flavonoids have demonstrated antiproliferative activity in prostate, renal, colon, non-Hodgkin's lymphoma, gastric carcinomas and neuroblastoma cancer cells [41-43]. Tannin isolated from *Cuphea hyssopifolia*, *Phyllanthus niruri* and *Cistus ladanifer* among other medicinal plants has been documented to have antiproliferative effects against breast, human promyelocytic leukemia HL-60, ovarian and pancreatic cancer cell lines [44-46].

The findings of this study indicates that the aqueous extract of the Kenyan *T. versicolor* mushroom are consistent with previous results on *T. versicolor* mushrooms collected from other parts of the world. However, it is probable that the phytochemicals found to be occurring in the aqueous extract of the Kenyan *T. versicolor* used in this study are responsible for the antiproliferative nature of the extract. These results suggest the need for a quantitative analysis and study of the phytochemicals of the Kenyan *T. versicolor* mushroom. *In vitro* and *in vivo* anti-proliferative activity and evaluation of cell death mechanisms using purified active compounds extracted from the *T. versicolor* mushroom is also recommended.

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