


RESEARCH ARTICLE

Open Access



Concurrent regulation of LKB1 and CaMKK2 in the activation of AMPK in castrate-resistant prostate cancer by a well-defined polyherbal mixture with anticancer properties

Amber F. MacDonald¹, Ahmed Bettaieb¹, Dallas R. Donohoe¹, Dina S. Alani¹, Anna Han^{1,5}, Yi Zhao^{1,4} and Jay Whelan^{1,2,3*} 

Abstract

Background: Zyflamend, a blend of herbal extracts, effectively inhibits tumor growth using preclinical models of castrate-resistant prostate cancer mediated in part by 5'-adenosine monophosphate-activated protein kinase (AMPK), a master energy sensor of the cell. Clinically, treatment with Zyflamend and/or metformin (activators of AMPK) had benefits in castrate-resistant prostate cancer patients who no longer responded to treatment. Two predominant upstream kinases are known to activate AMPK: liver kinase B1 (LKB1), a tumor suppressor, and calcium-calmodulin kinase kinase-2 (CaMKK2), a tumor promoter over-expressed in many cancers. The objective was to interrogate how Zyflamend activates AMPK by determining the roles of LKB1 and CaMKK2.

Methods: AMPK activation was determined in CWR22Rv1 cells treated with a variety of inhibitors of LKB1 and CaMKK2 in the presence and absence of Zyflamend, and in LKB1-null HeLa cells that constitutively express CaMKK2, following transfection with wild type LKB1 or catalytically-dead mutants. Upstream regulation by Zyflamend of LKB1 and CaMKK2 was investigated targeting protein kinase C-zeta (PKC ζ) and death-associated protein kinase (DAPK), respectively.

Results: Zyflamend's activation of AMPK appears to be LKB1 dependent, while simultaneously inhibiting CaMKK2 activity. Zyflamend failed to rescue the activation of AMPK in the presence of pharmacological and molecular inhibitors of LKB1, an effect not observed in the presence of inhibitors of CaMKK2. Using LKB1-null and catalytically-dead LKB1-transfected HeLa cells that constitutively express CaMKK2, ionomycin (activator of CaMKK2) increased phosphorylation of AMPK, but Zyflamend only had an effect in cells transfected with wild type LKB1. Zyflamend appears to inhibit CaMKK2 by DAPK-mediated phosphorylation of CaMKK2 at Ser511, an effect prevented by a DAPK inhibitor. Alternatively, Zyflamend mediates LKB1 activation via increased phosphorylation of PKC ζ , where it induced translocation of PKC ζ and LKB1 to their respective active compartments in HeLa cells following treatment. Altering the catalytic activity of LKB1 did not alter this translocation.

(Continued on next page)

* Correspondence: jwhelan@utk.edu

¹Department of Nutrition, University of Tennessee, 1215 West Cumberland Avenue, 229 Jessie Harris Building, Knoxville, TN 37996, USA

²Tennessee Agricultural Experiment Station, University of Tennessee, Knoxville, TN 37996, USA

Full list of author information is available at the end of the article



(Continued from previous page)

Discussion: Zyflamend’s activation of AMPK is mediated by LKB1, possibly via PKCζ, but independent of CaMKK2 by a mechanism that appears to involve DAPK.

Conclusions: Therefore, this is the first evidence that natural products simultaneously and antithetically regulate upstream kinases, known to be involved in cancer, via the activation of AMPK.

Keywords: Zyflamend, AMPK, LKB1, CaMKK2, CWR22Rv1, HeLa, DAPK, PKC-zeta, Prostate cancer, Castrate-resistant

Background

Prostate cancer is the second leading cause of death for men in the United States [1]. While early stages of the disease are treatable, with 5-year survival rates near 100%, prognosis for advanced forms are less promising [2]. Initially, prostate cancer cells rely on androgens for growth, and chemically-mediated deprivation (hormone deprivation therapy) is a common therapy that results in cancer regression [3]. Relapse in the absence of androgens (castrate-resistant prostate cancer) is inevitable for most individuals and is associated with increased expression and activation of the androgen receptor, a major determinant in survival [4, 5]. Due to the poor prognosis of castrate-resistant prostate cancer, concomitant use of natural products to enhance effectiveness is being explored clinically and experimentally [3, 6–10].

Zyflamend (New Chapter, Inc. Brattleboro, VT) is a poly-herbal supplement derived from the extracts of ten different herbs: rosemary, turmeric, holy basil, ginger, green tea, hu zhang, barberry, oregano, Chinese goldthread, and baikal skullcap. Most research using Zyflamend has focused its effects on a variety of cancer models, including oral [11], mammary [12], bone [13], pancreas [14, 15], skin [11, 16], colorectal [15], with an emphasis on prostate [6–9, 17–21], and its beneficial effects appear to be related to the synergy of action of its components [22]. The effects of Zyflamend

and its mechanisms on prostate cancer has been reviewed elsewhere and can be summarized in Fig. 1 [3]. Zyflamend inhibits signaling pathways of inflammation, affects cell survival by enhancing apoptotic and tumor suppressor genes, epigenetically modifies histones, down regulates the androgen receptor and influences the energetics of the cell. The latter pathways are critically important in cancer as rapidly dividing cells rely on the increased synthesis of macromolecules (lipids, proteins, nucleotides, etc) (as reviewed in [23].

5’-adenosine monophosphate-activated protein kinase (AMPK) is a key regulator of energy in the cell and responds to deficits in adenosine triphosphate (ATP). The protein contains a catalytic subunit (α-subunit), and two regulatory subunits, β and γ-subunits. Under conditions of energy stress the following occurs, (i) increased levels of AMP or ADP bind to the γ-subunit causing allosteric activation of the protein (ATP is a competitive inhibitor), (ii) increased affinity for upstream kinases that target phosphorylation at Thr172 of the α-subunit (increasing catalytic activity > 100 fold), and (iii) reduced affinity for phosphatases that are involved in dephosphorylation at Thr172 [24]. When activated, AMPK is instrumental in inhibiting anabolic pathways that consume ATP, such as lipogenesis and protein synthesis, and enhances

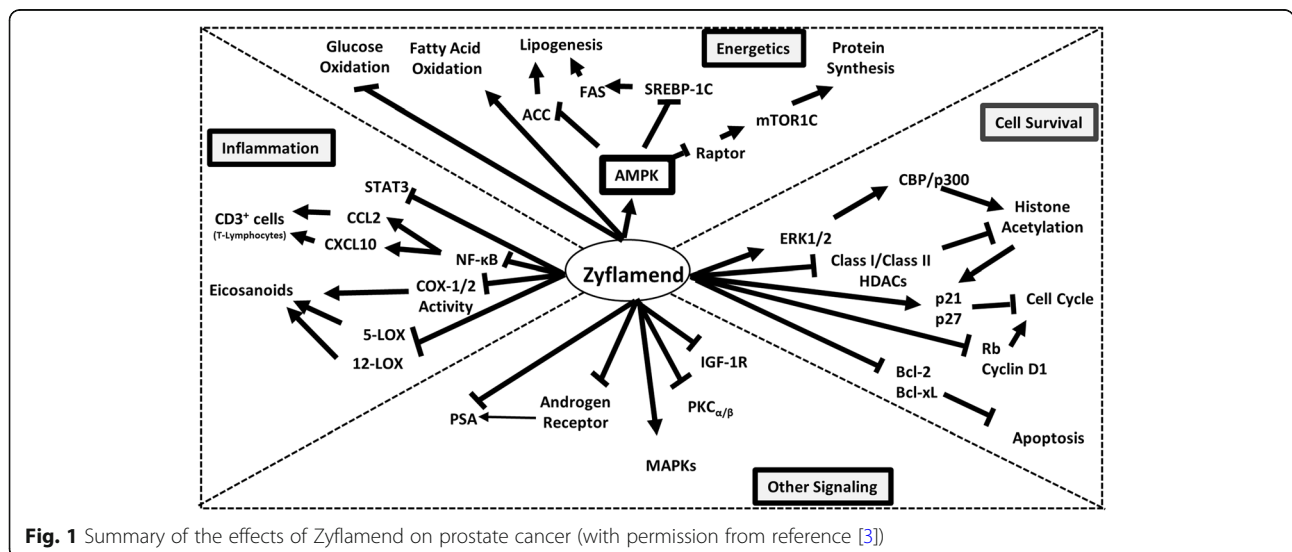


Fig. 1 Summary of the effects of Zyflamend on prostate cancer (with permission from reference [3])

catabolic pathways that generate ATP, such as fatty acid oxidation [23].

Clinically, treatment with Zylamend and/or metformin (activator of AMPK) had benefits in castrate-resistant prostate cancer patients who no longer responded to a variety of treatments (e.g., hormone ablation, immune-, chemo-, and radiation therapy). Recently, it was determined that tumor suppressor properties of Zylamend are associated with the activation of AMPK and its downstream signaling, where siRNA knockdown, pharmacological inhibition and over expression of AMPK confirmed Zylamend's involvement [10]. This involves inhibiting the mammalian target of rapamycin complex-1 (mTORC1) and protein synthesis, lipogenesis by targeting the expression of (i) fatty acid synthase, (ii) the sterol regulatory element-binding transcription factor-1c, and (iii) inhibiting the activity of acetyl CoA carboxylase (ACC). What is not known is how Zylamend upregulates AMPK. Four kinases have been identified that activate AMPK at Thr172, liver kinases B1 (LKB1), calcium-calmodulin kinase kinase-2 (CaMKK2), transforming growth factor- β activated protein kinase-1 (TAK1) and mixed lineage kinase 3 (MLK3) [25–28]. LKB1 and CaMKK2 are important in a number of cancers, including castrate-resistant prostate cancer (as reviewed in [29]), while the involvement of TAK1 and MLK3 has yet to be determined. LKB1 responds to increases in AMP and ADP, while increases in intracellular calcium is needed for activation of CaMKK2 without requiring elevation in AMP or ADP.

Interestingly, while both LKB1 and CaMKK2 are involved in activating AMPK, their effects on cancer appear to be quite different. LKB1 has anticancer properties because its mutation/deletion is associated with a variety of cancers [30]. CaMKK2, on the other hand, is overexpressed in a number of cancers, including castrate-resistant prostate cancer [31, 32]. Therefore, the overall objective of this paper was to interrogate how Zylamend activates AMPK in a model of castrate-resistant prostate cancer and the roles LKB1 and CaMKK2 play in that activation.

The major findings of this research revealed that although LKB1 and CaMMK2 are upstream kinases that can activate AMPK, Zylamend inhibits CaMMK2, a protein overexpressed in many cancers, while simultaneously upregulates LKB1, a reported tumor suppressor. This is the first report linking the simultaneous antagonistic regulation of these two proteins. Upstream regulation of CaMKK2 appears to be mediated by the antitumorigenic death associated protein kinase (DAPK) [33–36], not epigenetically, but via phosphorylation at Ser511 [37]. This is only the second paper to link DAPK activity with the negative regulation of CaMKK2 via phosphorylation at Ser511, and the only one involving cancer cells.

Methods

Zylamend (New Chapter, Inc. Brattleboro, VT), purchased from Earth Fare Supermarket (Knoxville, TN), is composed of extracts from the following herbs (w/w): rosemary (*Rosmarinus officinalis* 19.2%), turmeric (*Curcuma longa* 14.1%), holy basil (*Ocimum sanctum* 12.8%), ginger (*Zingiber officinale* 12.8%), green tea (*Camellia sinensis* 12.8%), hu zhang (*Polygonum cuspidatum* 10.2%), barberry (*Berberis vulgaris* 5.1%), oregano (*Origanum vulgare* 5.1%), Chinese goldthread (*Coptis chinensis* 5.1%), and baikal skullcap (*Scutellaria baicalensis* 2.5%). Detailed description and characterization of the preparation of Zylamend and quality assurance of the mixture has been described previously in detail [7]. This description includes rigorously generated verifiable quality control of its constituents via multiple independent laboratories whose biological effects have been duplicated using different lots, at different times, under different experimental conditions, in different laboratories across the United States.

Dulbecco's Modified Eagle Medium (DMEM), G418, penicillin/streptomycin, puromycin, fetal bovine serum (FBS) and trypsin were purchased from Invitrogen (Carlsbad, CA). Cloning vectors were purchased from Addgene (Cambridge, MA). Antibodies for PKC-zeta (PKC ζ), LKB1, phospho-LKB1, green fluorescent protein (GFP), Histone B, Flag, and Tubulin were from Santa Cruz Biotechnology (Santa Cruz, CA). AMPK and phospho-AMPK were from Cell Signaling Technology (Beverly, MA). The following chemical reagents were purchased: 5-aminoimidazole-4-carboxamide ribonucleotide (AICAR) (AdipoGen Life Sciences, San Diego, CA); 1,2-bis(*o*-aminophenoxy)ethane-*N,N,N',N'*-tetraacetic acid acetoxymethyl ester (BAPTA-AM) and ethylene glycol-bis(β -aminoethyl ether)-*N,N,N',N'*-tetraacetic acid (EGTA) (Thermo Scientific, Rockville, IL); STO-609, radicicol, and PKC ζ Pseudo-substrate Inhibitor (Santa Cruz Biotechnology, Dallas, TX); ionomycin (Sigma-Aldrich, St. Louis, MO); and Death Associated Protein Kinase Inhibitor (DAPKi) (Merck Millipore, Billerica, MA).

Cell culture

CWR22Rv1 cells (American Type Culture Collection, Rockville, MD), a human-derived castrate-resistant prostate cancer cell line, were cultured in RPMI 1640 media, supplemented with 10% FBS. To mimic an androgen-depleted state, the cells were incubated overnight with 0.5% FBS. HeLa cells (ATCC, Rockville, MD), a human-derived cervical cancer cell line that do not express LKB1, and HCT116 cells (ATCC, Rockville, MD), a human derived colorectal cancer cell line, were cultured in DMEM media supplemented with 10% FBS and 25 mM glucose. All cells were incubated under an atmosphere of 5% CO₂, at 37 °C. For activation of AMPK via

LKB1-dependent or CaMKK2-dependent pathways, cells were treated with AICAR (a cell permeable analog of AMP) (1 mM, 1 h) or ionomycin (calcium ionophore) (1 μ M, 1 h), respectively. For experiments using inhibitors of CaMKK2, cells were pre-treated with the selective CaMKK2 inhibitor STO-609 (10 μ M, 30 min) or the calcium chelators BAPTA-AM (30 μ M, 30 min) or EGTA (2 mM, 30 min). For inhibition of LKB1 or DAPK, cells were pre-treated with radicicol (5 μ M, 24 h) or DAPKi (20 μ M, 24 h), respectively. For inhibition of PKC ζ , cells were pre-treated with the selective PKC ζ pseudo-substrate inhibitor (5 μ M, 30 min). For all experiments involving Zyflamend, cells were treated with Zyflamend at 200 μ g/mL for 30 min unless otherwise indicated.

Down regulation of LKB1 by small interfering RNA

CWR22Rv1 cells were seeded in RPMI medium containing 10% FBS and incubated overnight before media was replaced with RNA transfection medium containing 0.5% FBS. Cells were transfected with 20 nmol of siRNA targeting LKB1 (Thermo Scientific/Dharmacon #L-005035-00) and a siRNA non-targeting control (Thermo Scientific/Dharmacon #D-001810-10-05). Western blot analysis confirmed the efficiency of knockdown to be 76% 48 h after transfection, at which time cells were treated with a vehicle or Zyflamend (200 μ g/mL) for 30 min.

Overexpression of LKB1 in HeLa cells

Human WT or catalytically dead (KD) mutants of LKB1 were transfected into HeLa cells using Lipofectamine 3000 (Invitrogen, Carlsbad, CA) following manufacturer's guidelines. Cells were cultured for additional 48 h prior experiments. For total protein lysates, cells were lysed in radio-immunoprecipitation assay buffer (RIPA: 10 M Tris-HCl, pH 7.4, 150 mM NaCl, 0.1% sodium dodecyl sulfate [SDS], 1% Triton X-100, 1% sodium deoxycholate, 5 mM EDTA, 1 mM NaF, 1 mM sodium orthovanadate and protease inhibitors). Lysates were clarified by centrifugation at 13,000 rpm for 10 min, and protein concentrations were determined using a bicinchoninic acid assay kit (Pierce Chemical). Proteins (10 μ g) were separated by SDS-polyacrylamide gel electrophoresis (SDS-PAGE) (8–12%) [38], transferred to polyvinylidene difluoride (PVDF) membranes and immunodetected using the indicated antibodies. Proteins were detected using enhanced chemiluminescence (Amersham Biosciences). Resulting immunoreactive bands were quantified using FluorChem Q Imaging software (Alpha Innotech).

Subcellular fractionation

Following Zyflamend treatment, fractionation was performed in HeLa cells as described previously with modifications [39, 40]. Briefly, cells were washed

with cold buffer A (100 mM sucrose, 1 mM EGTA, 20 mM 3-(N-morpholino)propanesulfonic acid (MOPS), pH 7.4) and resuspended in lysis buffer B (100 mM sucrose, 1 mM EGTA, 20 mM MOPS, 0.1 dithiothreitol (DTT), 5% freshly added percoll, 0.01% digitonin, 1 mM phenylmethylsulfonyl fluoride (PMSF) and cocktail of protease inhibitors, pH 7.4). Membranes were broken using a dounce homogenizer (200 strokes/sample). Debris and unbroken cells were removed by centrifugation (500 g for 10 min) and supernatants were then centrifuged (2500 g, 5 min) to separate nuclei (pellet). Supernatants were centrifuged again (15,000 g, 15 min) to separate mitochondria. Nuclear fraction was resuspended in radioimmunoprecipitation assay (RIPA) buffer containing proteases inhibitors. Cellular distribution and translocation of the indicated proteins were analyzed by SDS-PAGE and Western blot as described above. Purity of nuclear and cytoplasmic fractions was verified using antibodies against histone B and tubulin (Santa Cruz Biotechnology, Dallas, TX), respectively.

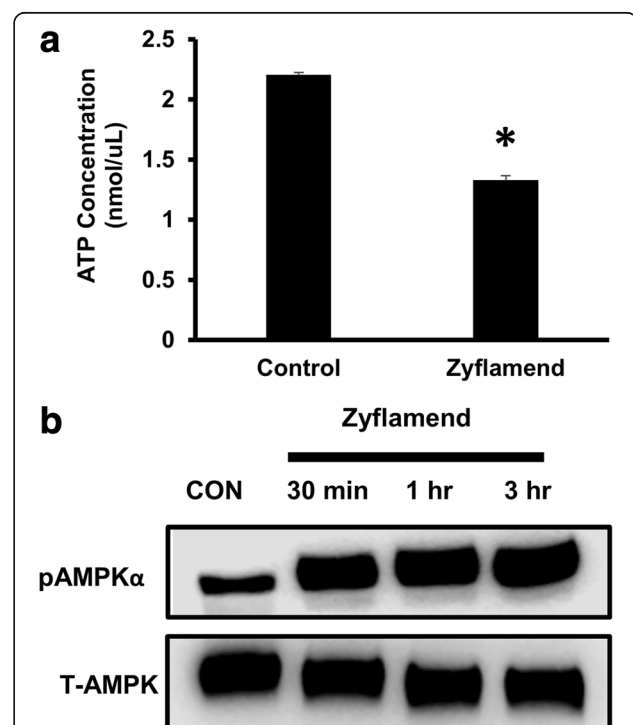


Fig. 2 The effects of Zyflamend on cellular ATP levels and phosphorylation of AMPK α (at Thr172) in CWR22Rv1 cells. **a** ATP levels of CWR22Rv1 cells treated in the presence or absence of Zyflamend (200 μ g/mL, 30 min). **b** The effects of Zyflamend (200 μ g/mL, 30 min – 3 h) on phosphorylation of AMPK α in CWR22Rv1 cells. Data is presented as mean \pm SEM, $n = 4$. Abbreviations: Con, Control

Western blotting

Cells were lysed in RIPA lysis buffer (Thermo Scientific, Rockford, IL). Protein concentration was measured using a Bradford protein assay (Thermo Scientific, Rockford, IL). Equal amount of protein (30 μ g) were separated by 8% SDS-PAGE and transferred to a PVDF membrane by electroblotting. Membranes were blocked by 5% non-fat dry milk (LabScientific, Highlands, NJ) or bovine serum albumin (Santa Cruz Biotechnology, Dallas, TX) in 0.1% Tris-buffered saline-Tween-20 (TBST) for 1 h at room temperature and incubated in TBST containing primary antibodies overnight at 4 $^{\circ}$ C. Membranes were incubated with anti-rabbit or anti-mouse secondary antibody conjugated with horseradish peroxidase (HRP) (Cell Signaling Technology, Danver, MA) for 1 h at room temperature. Protein expression was detected with Super

Signal West Pico Chemiluminescent Substrate (Thermo Scientific, Rockford, IL) and membranes were exposed and analyzed via *Li-Cor Odyssey FC* imaging system (Li-Cor, Lincoln, NE). Antibodies against p-AMPK α (Thr172), AMPK α , p-ACC (Ser79), ACC, p-LKB1 (Ser428), LKB1, p-PKC ζ (Thr410), PKC ζ , and p-CaMKK β (Ser511) were used to detect target protein level at 1:1000. β -Actin or glyceraldehyde 3-phosphate dehydrogenase (GAPDH) (Santa Cruz Biotechnology, Dallas, TX) was used as the loading control.

ATP assay

Cellular ATP concentration was determined using a fluorometric ATP assay kit (BioRad, Milpitas, CA) following the manufacturer's instructions, and fluorescence

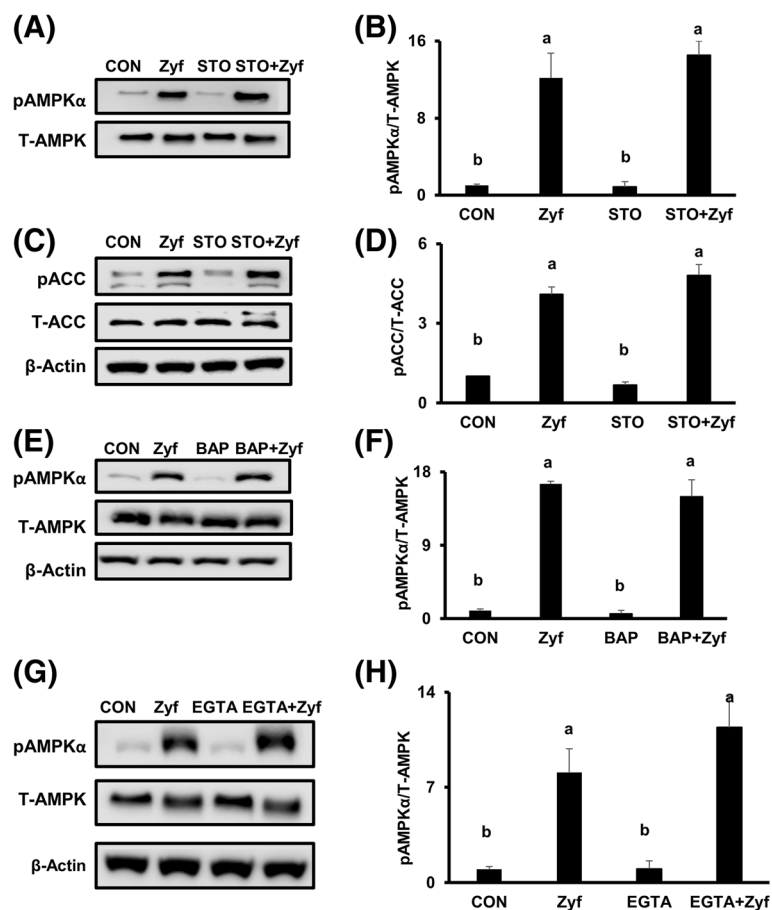


Fig. 3 The effects of CaMKK2 inhibition by STO-609, BAPTA-AM and EGTA on pAMPK α (at Thr172) and pACC (at Ser79), \pm Zylamend in CWR22Rv1 cells. **(A-D)** Western blot of pAMPK α and pACC following treatment of STO-609 (10 μ M, 30 min) \pm Zylamend (200 μ g/mL, 30 min). **(E, F)** Western blot of pAMPK α following treatment of BAPTA-AM (30 μ M, 30 min) \pm Zylamend (200 μ g/mL, 30 min). **(G, H)** Western blot of pAMPK α following treatment of EGTA (2 mM, 30 min) \pm Zylamend (200 μ g/mL, 30 min). Data are presented as mean \pm SEM, $n = 3$. Bars with different letters are statistically different at $p < 0.05$. Abbreviations: BAP, BAPTA-AM (1, 2-bis(o-aminophenoxy)ethane- N,N,N',N' -tetraacetic acid acetoxymethyl ester); CON, Control; EGTA, ethylene glycol-bis (β -aminoethyl ether)- N,N,N',N' -tetraacetic acid; STO, STO-609; Zyf, Zylamend

was read at 525 nm on a Glowmax Multi Detection System (Promega Corporation, Madison, WI).

Statistics

For Western blot, protein was analyzed from 3 independent samples and presented as mean \pm SEM. For ATP concentration, results are presented as mean \pm SEM. For multiple comparisons, data was analyzed using IBM SPSS Statistics 24 and tested by one-way analysis of variance (ANOVA) followed by a Tukey's post-hoc test. Two-group comparisons were analyzed by two-tailed Student's T-test. Results were considered statistically significant at $p < 0.05$.

Results

Effect of Zylamend on cell proliferation, ATP levels and AMPK phosphorylation in CWR22Rv1 cells

Zylamend (200 μ g/mL) inhibited cell proliferation in CWR22Rv1 cells in a concentration and time dependent manner (Additional file 1: Figure S1, replicating previously published results [6, 8]). Similar results were replicated in a variety of immortalized prostate-derived cells lines [8] and in the HCT116 colorectal cell line (Additional file 2: Figure S2). Because Zylamend has been shown to change the energetics of CWR22Rv1 cells [3, 10], levels of ATP were determined. In the presence of Zylamend, ATP levels were reduced by $\sim 40\%$ (Fig. 2a) and AMPK phosphorylation (Thr172) was significantly increased (Fig. 2b). These results were replicated in the HCT116 colorectal cell line to demonstrate that these effects are not specific for prostate cancer cells (Additional file 2: Figure S2).

AMPK activation by CaMKK2 in the presence and absence of Zylamend in CWR22Rv1 cells

Zylamend significantly increased the phosphorylation of AMPK (Thr172) (Fig. 3A and B) and its downstream target ACC (Ser79) (Fig. 3C and D), results unaffected by pretreatment with the CaMKK2 inhibitor STO-609 (Fig. 3A-D, lane 4/bar 4). To confirm that the activation of AMPK by Zylamend is independent of CaMKK2, cells were pre-treated with the calcium chelators BAPTA-AM and EGTA, as CaMKK2 activation is dependent upon intracellular calcium. Pretreatment with BAPTA-AM (Fig. 3E and F) and EGTA (Fig. 3G and H) failed to prevent phosphorylation of AMPK in the presence of Zylamend. Zylamend increased the phosphorylation of CaMKK2 at Ser511 (Fig. 4A), a phosphorylation site that results in its inhibition and reported to be mediated by DAPK [37]. Pretreatment with a DAPK inhibitor attenuated Zylamend's ability to increase phosphorylation of CaMKK2 at Ser511 in a time-dependent manner (Fig. 4A-C), suggesting phosphorylation of CaMKK2 in the presence of Zylamend is mediated by DAPK. Pretreatment of cells with DAPKi failed to prevent phosphorylation of AMPK in the presence of Zylamend, further confirming that the activation of AMPK by Zylamend is independent of CaMKK2 (Fig. 4D and E).

AMPK activation by LKB1 in the presence and absence of Zylamend in CWR22Rv1 cells

Zylamend significantly increased the phosphorylation of LKB1 (Ser428) (Fig. 5A), AMPK (Thr172) (Fig. 5C) and ACC (Ser79) (Fig. 5E). In the presence of radicicol, a

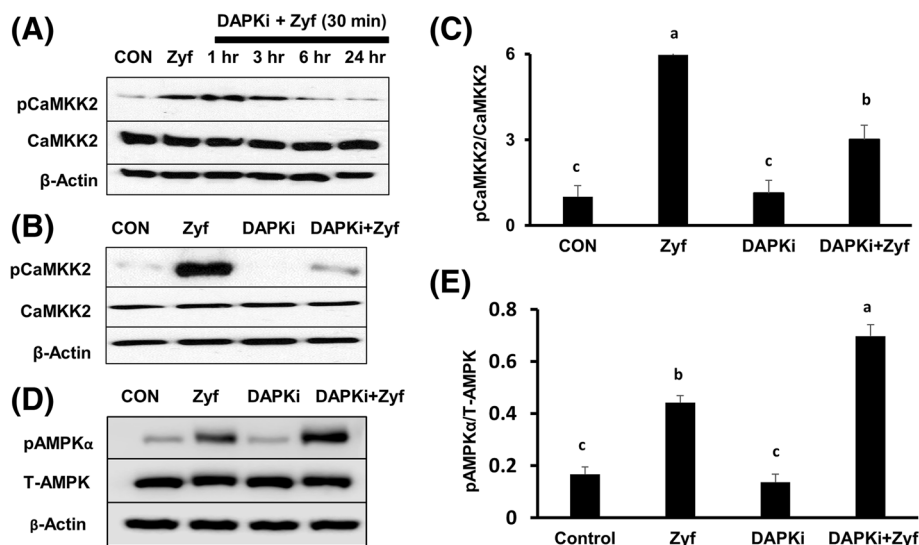


Fig. 4 The effects of Zylamend and an inhibitor of DAPK on the phosphorylation of CaMKK2 (at Ser511), in CWR22Rv1 cells. **A** Western blot of pCaMKK2 \pm Zylamend (200 μ g/mL, 30 min) and following treatment of DAPKi (20 μ M, 1 h – 24 h) followed by the treatment of Zylamend (200 μ g/mL, 30 min). **B, C** Western blot and graph comparison of pCaMKK2 following treatment of DAPKi (20 μ M, 24 h) \pm Zylamend (200 μ g/mL, 30 min). **(D, E)** Western blot and graph comparison of pAMPK following treatment of Zylamend (200 μ g/mL, 30 min) and DAPKi (20 μ M, 24 h) \pm Zylamend (200 μ g/mL, 30 min). Data are presented as mean \pm SEM, $n = 3$. Bars with different letters are statistically different at $p < 0.05$. Abbreviations: CON, Control; DAPKi, DAPK inhibitor; Zyf, Zylamend

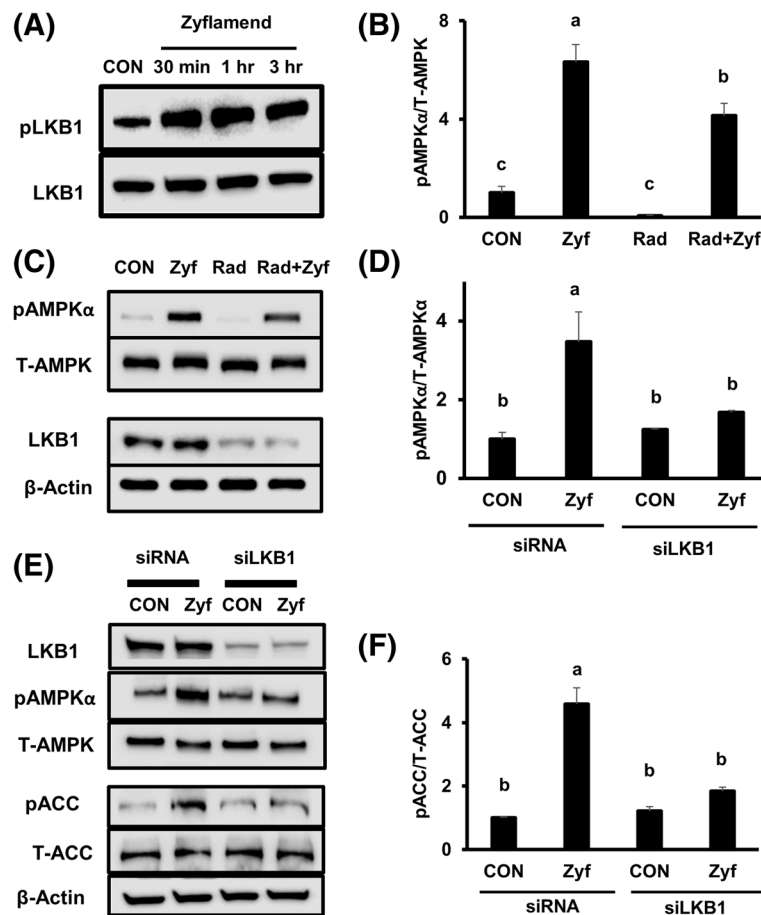


Fig. 5 The effects of Zyflamend on phosphorylation of LKB1 (at Ser428) and AMPK α (at Thr172) following inhibition with radicicol and knockdown of LKB1 in CWR22Rv1 cells. **A** Western blot of pLKB1 \pm Zyflamend (200 μ g/mL, 30 min – 3 h). **B, C** Western blot of pAMPK α following treatment with radicicol (5 μ M, 24 h) \pm Zyflamend (200 μ g/mL, 30 min). **D–F** Western blot of pAMPK α and pACC (at Ser79) following knockdown of LKB1 \pm Zyflamend (200 μ g/mL, 30 min). Data are presented as mean \pm SEM, $n = 3$. Bars with different letters are statistically different at $p < 0.05$. Abbreviations: CON, Control; Rad, Radicicol; Zyf, Zyflamend

non-specific inhibitor of LKB1 that results in reduction of total LKB1 protein levels (Fig. 5C, row 3, column 3), phosphorylation of AMPK failed to be fully restored upon Zyflamend treatment (Fig. 5B and C). Likewise, knockdown of LKB1 by siRNA (Fig. 5d-f) inhibited Zyflamend-induced phosphorylation of AMPK (Fig. 5D and E) and its downstream target ACC (Fig. 5E and F).

Zyflamend-induced AMPK phosphorylation is LKB1 dependent

To confirm that Zyflamend-induced phosphorylation of AMPK is LKB1 dependent and CaMKK2 independent, we treated LKB1-null HeLa cells, that constitutively express CaMKK2, with AICAR (an activator of AMPK that is commonly used as a positive control), ionomycin (activator of CaMKK2) and Zyflamend (Fig. 6a). The same experiment was conducted on HeLa cells that were stably transfected with the wild-type (WT) or the catalytically dead (KD) mutants of the human LKB1. Two different constructs of the

KD mutants were used, Flag-tagged or green fluorescent protein (KD-LKB1 Flag or KD-LKB1 GFP, respectively) (Fig. 6b). In both mutants, lysine 78 was mutated to isoleucine, abolishing auto-phosphorylation and activation of LKB1 [41, 42]. Ionomycin (Fig. 6b, column 3), but not AICAR (column 2) or Zyflamend (column 4), induced phosphorylation of AMPK in LKB1-null HeLa cells (control) (Fig. 6b, columns 1–5). Following transfection with WT-LKB1 (Fig. 6b, columns 6–8), Zyflamend induced phosphorylation of LKB1 and AMPK (column 7), an effect more pronounced with co-treatment of AICAR (column 8). However, no phosphorylation of LKB1 and AMPK was observed in the KD mutants following treatment with Zyflamend and Zyflamend+AICAR (Fig. 6b, columns 9–14).

Zyflamend-mediated LKB1 phosphorylation is linked to PKC ζ

In an effort to determine how Zyflamend may be mediating the phosphorylation of LKB1, PKC ζ was investigated

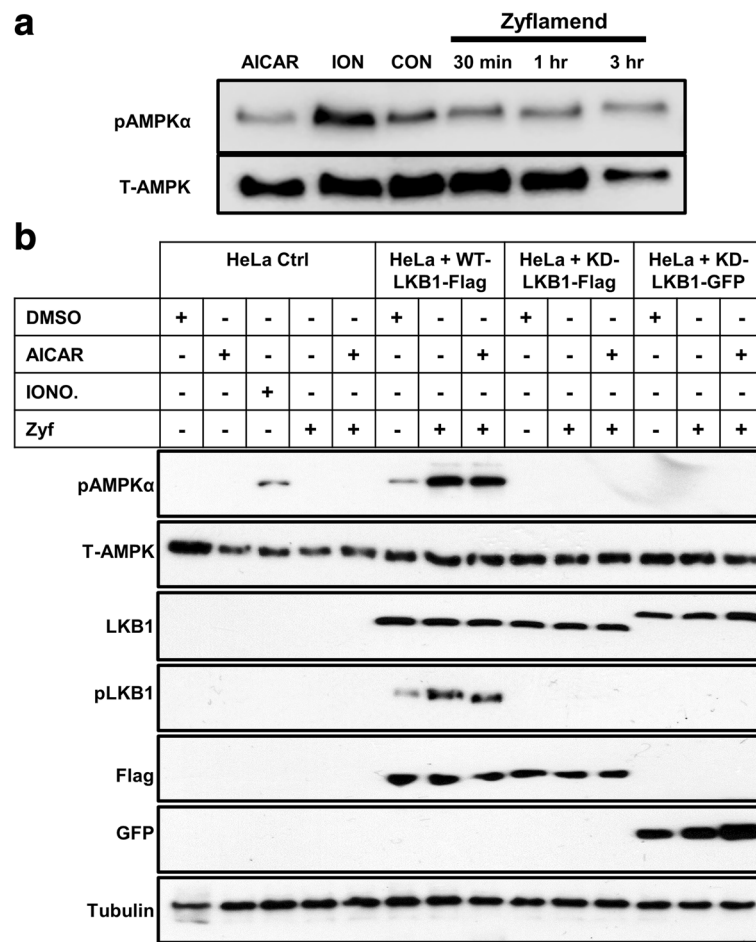


Fig. 6 Effects of Zyflamend on pAMPK (at Thr172) and pLKB1 (at Ser428) in HeLa cells null for LKB1, transfected with wild type (WT) LKB1 or with two catalytically dead (KD) mutants of LKB1. **a** Western blot of pAMPK in HeLa cells \pm AICAR, (1 mM, 1 h), ionomycin (1 μ M, 1 h) or Zyflamend (200 μ g/mL, 30 min – 3 h). **b** Western blot of pAMPK in HeLa cells, WT-LKB1 HeLa cells, and KD-LKB1 HeLa cells \pm AICAR (1 mM, 1 h), ionomycin (1 μ M, 1 h), and/or Zyflamend (200 μ g/mL, 1 h). Representative immunoblots from 3 independent experiments are shown. Abbreviations: Con, Control; DMSO, dimethyl sulfoxide; Ion, ionomycin; Zyf, Zyflamend

as a possible upstream target (Fig. 7). Phosphorylation of PKC ζ and LKB1 increased when CWR22Rv1 cells were treated with Zyflamend (Fig. 7A). However, in the presence of a highly selective PKC ζ pseudo-substrate inhibitor, phosphorylation of LKB1 could not be restored to control levels following Zyflamend treatment (Fig. 7B and C). Using our informative HeLa cell constructs, null for LKB1 and transfected with WT-LKB1 or KD-LKB1, we further investigated the relationship between PKC ζ and LKB1 (Fig. 7D). In HeLa cells devoid of LKB1 (control cells), PKC ζ is located in the cytosol (Fig. 7D, row 1, column 3), but appears to translocate to the nucleus upon treatment with Zyflamend (Fig. 7D, row 1, column 2). In cells transfected with WT-LKB1, PKC ζ is located in the cytosol (Fig. 7D, row 1, column 7) and LKB1 is located in the nucleus (Fig. 7D, row 2, column 5). Following Zyflamend treatment, their locations switch, where PKC ζ translocates to the nucleus (Fig. 7D, row 1, column

6) and LKB1 is found in the cytosol (Fig. 7D, row 2, column 8). This translocation following Zyflamend treatment appears to be independent of a catalytically active protein, as the same results were observed with the KD-LKB1 mutant (Fig. 7D, columns 9–12).

Discussion

Zyflamend is a unique blend of ten herbal extracts with tumor suppressor properties whose biochemical and physiological effects have been replicated in different laboratories, at different times, using different lots, with similar doses/concentrations [6–8, 10, 11, 13–17, 19, 21, 43]. The quality control of this preparation has been summarized elsewhere [7] and is most likely responsible for the reproducibility of results. Importantly, the effects of Zyflamend have been reported in a variety of cell lines, not just prostate cancer cells (CWR22, CWR22R, CWR22Rv1, PC3, LNCaP, RWPE-1, RAW 264.7, H1299, KBM-5, U266, MeW0, A365,

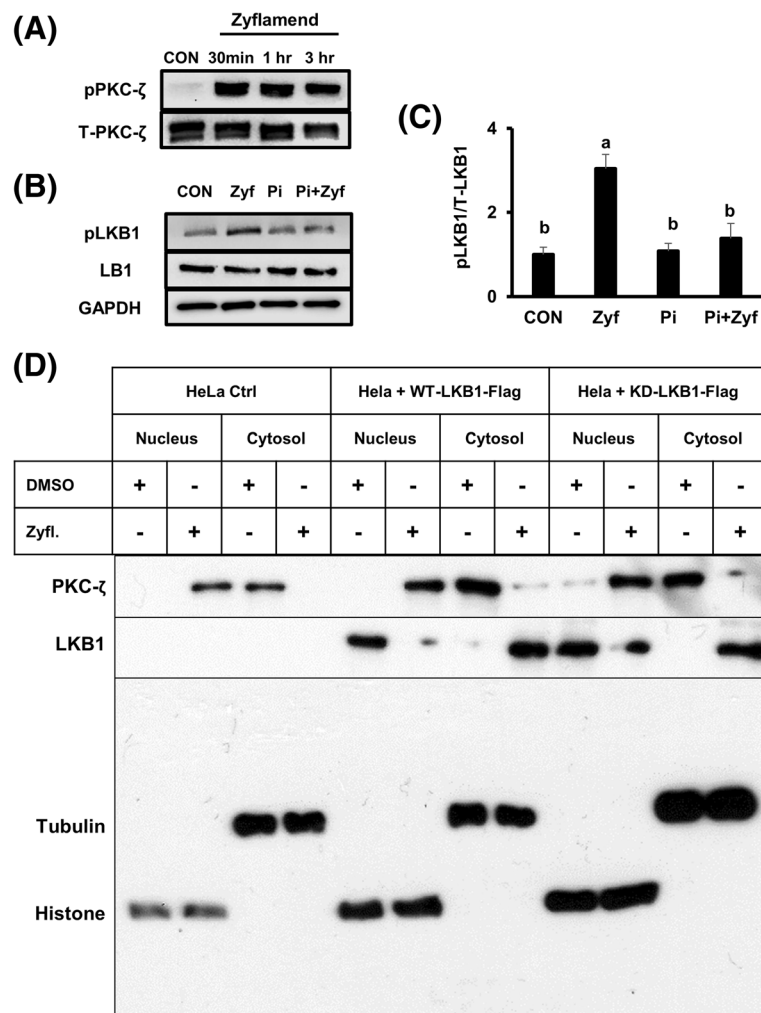


Fig. 7 The effects of Zyflamend on phosphorylation of PKC ζ (at Thr410) and cellular location of PKC ζ and LKB1 in HeLa cells null for LKB1, transfected with wild type (WT) LKB1 or with a catalytically dead (KD) mutant of LKB1. **A** Western blot of phosphorylation of PKC ζ \pm Zyflamend (200 μ g/mL, 30 min – 3 h). **B, C** Western blot of pLKB1 (at Ser428) following treatment with Zyflamend, PKC ζ inhibitor (5 μ M, 30 min) or pretreatment with the PKC ζ inhibitor plus Zyflamend (200 μ g/mL, 30 min). **D** Western blot for PKC ζ and LKB1 following subcellular fractionation (cytosol and nucleus) in HeLa cells null for LKB1, transfected with WT LKB1 or with a KD mutant of LKB1 treated with or without Zyflamend (200 μ g/mL, 1 h). Representative immunoblots from 3 independent experiments are shown. Abbreviations: CON, Control; Pi, PKC ζ inhibitor; Zyfl, Zyflamend

MSK-Leuk1, HaCaT, HCT116, THP-1, HEK293) [6–8, 10–19, 21, 43]. While there is clinical evidence for the beneficial effects of Zyflamend on prostate cancer [9, 18, 20, 44], it is not possible to tease out the contributions and/or interactions of each constituent due to the high number of possible combinations (as many as 1024). However, recent studies have demonstrated that combining components from this blend enhances their cellular and molecular effects by orders of magnitude as compared to isolated components, indicating highly synergistic interactions when combined [22]. The ability of this mixture to function at human equivalent doses and in clinical trials is most likely due to this synergy. This combination has been shown to be clinically effective in prostate cancer patients by reducing levels of prostate

specific antigen (PSA), a biomarker used to monitor prostate cancer progression. Case studies from M.D. Anderson Cancer Center report dramatic reductions in PSA levels following treatment with Zyflamend and/or metformin (an activator of AMPK) in patients whose prostate cancer no longer responds to a variety of standard therapies [9], an effect also observed in our recent clinical trial with prostate cancer patients undergoing radical prostatectomy [44].

Castrate-resistant prostate cancer is the focus of this research and the research in our laboratory. To study mechanisms of action, we use human prostate cancer cells derived from the CWR22 lineage [6–8, 10, 45, 46]. Similar to the progression of human prostate cancer, these cells are originally androgen dependent and can transform to a

castrate-resistant line in vivo (i.e., CRW/22R) following hormone ablation [46–48]. Unlike some other prostate cancer cell lines (i.e., PC3 cells), the CWR22Rv1 cells express a constitutively active androgen receptor and PSA, characteristics shared by castrate-resistant prostate cancer in humans.

The effectiveness of Zyflamend on prostate cancer rests, in part, with its ability to upregulate AMPK, an effect observed in other cell types and tissues (i.e., HCT116, PC3, adipose) suggesting a general effect [10, 44]. However, the mechanism as to how Zyflamend upregulates AMPK is unknown, although many of its constituents have been shown to independently activate AMPK by modifying mitochondrial ATP production (as reviewed in [49]). This is the first paper to delineate the coordination of potential upstream pathways involved in the activation of AMPK, viz., LKB1 and CaMKK2, and to do so using natural products.

The role of AMPK in prostate cancer is controversial in the sense that upstream kinases responsible for its activation appear to have contradictory effects on cancer [29]. CaMKK2 is a known tumor promotor whose expression is linked to the upregulation of the androgen receptor, a key step in castrate-resistant prostate cancer [4, 31]. Interestingly, Zyflamend down regulates the androgen receptor and its nuclear localization [6]. DAPK is a differentially methylated gene where most of its effects on cancer have focused on its epigenetic regulation [33–36]. Uniquely, a catalytic downstream target of DAPK is CaMKK2 where it phosphorylates CaMKK2 at Ser511 [37], a site adjacent to the Ca²⁺-calmodulin regulatory domain, preventing autophosphorylation and inhibiting catalytic activity [37].

In contrast, LKB1 exhibits tumor suppressor properties, where loss of LKB1 is involved in a variety of

cancers [24, 30]. LKB1-mediated activation of AMPK is dependent upon increases in the AMP(ADP):ATP ratios. Zyflamend significantly decreases ATP in the cells. LKB1 contains a nuclear localization domain and is typically (but not always exclusively) found in the nucleus. Following activation, LKB1 co-localizes with STE20-related adaptor (known as STRAD) protein and scaffolding mouse 25 (known as MO25) protein and translocates to the cytosol where it exerts its kinase activity on a number of downstream targets, including AMPK [50]. Nuclear export, in part, appears to involve phosphorylation at Ser428 by PKCζ [51].

A key finding from this research is that Zyflamend antithetically regulates two parallel pathways important in the phosphorylation of AMPK that is potentially important in castrate-resistant prostate cancer. These effects are summarized in Fig. 8. Zyflamend-mediated activation of AMPK appears to be LKB1 dependent, while coordinately and negatively regulating CaMKK2 activity. This was observed using LKB1-null and KD-LKB1 transfected HeLa cells that constitutively express CaMKK2. The addition of ionomycin (activator of CaMKK2) robustly increased phosphorylation of AMPK, but Zyflamend (with and without AICAR, an AMP analog) had no effect. Our results suggest that Zyflamend inhibits CaMKK2 following DAPK-mediated phosphorylation at Ser511, as this effect is prevented by the presence of a DAPK inhibitor.

On the other hand, Zyflamend robustly increased the phosphorylation of AMPK only in HeLa cells transfected with WT LKB1. Using the various constructs of the HeLa cells, we confirmed nuclear localization of LKB1, with translocation to the cytosol following Zyflamend treatment (Fig. 7). Zyflamend increased phosphorylation

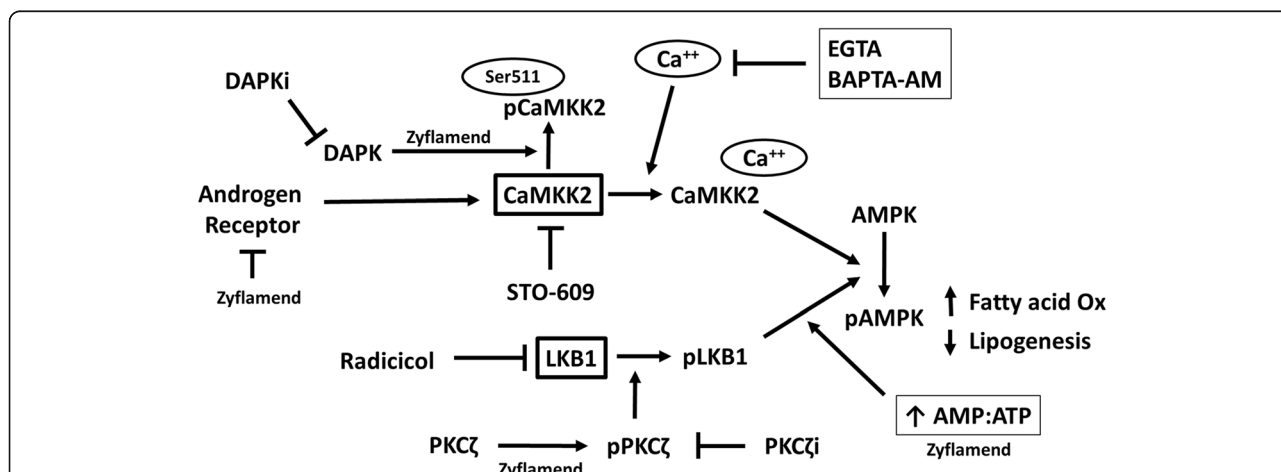


Fig. 8 Summary of the effects of Zyflamend on AMPK regulation by signaling pathways of LKB1 and CaMKK2. Zyflamend has been shown to inhibit castrate-resistant prostate cancer, in part, through the activation of AMPK. This activation is mediated by the increase in AMP:ATP ratio and the activation of the tumor suppressor protein LKB1 following phosphorylation by PKCζ. Simultaneously, Zyflamend inhibits the tumor promotor CaMKK2 via phosphorylation at Ser511 by DAPK

of PKC ζ , a known activator of LKB1, and inhibition of PKC ζ reduced LKB1 phosphorylation in the presence of Zylamend. Importantly, translocation of PKC ζ from the cytosol to the nucleus occurred concomitantly.

These results help explain why inhibitors of CaMKK2 (STO-609, BAPTA-AM, EGTA, DAPKi) failed to prevent the activation of AMPK in the presence of Zylamend in CWR22Rv1 cells. This was due to the simultaneous activation of LKB1, and this was confirmed when activation of AMPK (in the presence of Zylamend) was not completely rescued following inhibition (radicol) and knockdown (siRNA) of LKB1.

Conclusions

In summary, Zylamend has been shown to inhibit castrate-resistant prostate cancer, in part, through the activation of AMPK, results confirmed using chemical and molecular interventions [10]. In conjunction with reducing ATP levels, this activation is mediated by the tumor suppressor protein LKB1, via activation of PKC ζ . Simultaneously, Zylamend inhibits CaMKK2, a tumor promoter that is over-expressed in many cancers, including castrate-resistant prostate cancer. This inhibition appears to be uniquely mediated by DAPK. More studies are warranted to interrogate the relationship of Zylamend with DAPK signaling. In conclusion, this is the first evidence that multiple upstream pathways involved in the activation of AMPK, an important signaling molecule in cancer, can be simultaneously regulated using a well-defined natural product.

Additional files

Additional file 1: Figure S1. Effect of Zylamend on the proliferation of a castrate resistant prostate cancer cells in vitro. CWR22Rv1 cells were treated with Zylamend (0–200 $\mu\text{g}/\text{ml}$) from 0 to 96 h and cell proliferation was monitored using the MTT assay. (PDF 71 kb)

Additional file 2: Figure S2. Effect of Zylamend on the proliferation of a colorectal cancer cell line in vitro and the subsequent phosphorylation of AMPK α at Thr172. (A) HCT116 cells were treated with Zylamend (0–200 $\mu\text{g}/\text{ml}$) from 0 to 72 h and cell proliferation was monitored using the MTT assay. (B) Phosphorylation of AMPK α at Thr172 was determined following Zylamend treatment (200 $\mu\text{g}/\text{ml}$ for 3 h). (PDF 254 kb)

Abbreviations

ACC: Acetyl CoA carboxylase; AICAR: 5-aminoimidazole-4-carboxamide ribonucleotide; AMPK: 5'-adenosine monophosphate-activated protein kinase; ANOVA: Analysis of variance; BAPTA-AM: 1,2-bis(o-aminophenoxy)ethane-*N,N,N',N'*-tetraacetic acid acetoxymethyl ester; CaMKK2: Calcium-calmodulin kinase kinase-2; CR-PCa: Castrate-resistant prostate cancer; DAPK: Death-associated protein kinase inhibitor; DTT: Dithiothreitol; EGTA: Ethylene glycol-bis(β -aminoethyl ether)-*N,N,N',N'*-tetraacetic acid; GAPDH: Glyceraldehyde 3-phosphate dehydrogenase; GFP: Green fluorescent protein; LKB1: Liver kinases B1; MLK3: Mixed lineage kinase 3; MO25: Scaffolding mouse 25 protein; MOPS: 3-(*N*-morpholino)propanesulfonic acid; PKC ζ : Protein kinase C-zeta; PMSF: Phenylmethylsulfonyl fluoride; RIPA: Radioimmunoprecipitation assay; STRAD: STE20-related adaptor protein; TAK1: Transforming growth factor- β activated protein kinase-1

Acknowledgements

GFP-LKB1 was a gift from Junying Yuan (Addgene plasmid # 21147), while pcDNA3-FLAG-LKB1 was a gift from Lewis Cantley (Addgene plasmid # 8590).

Funding

This work was supported, in part, by a USDA HATCH grant (#TEN00441) through the Tennessee Agricultural Experiment Station, (JW), University of Tennessee, Knoxville, TN 37996, and by a grant from the National Institutes for Health (R00DK100736) (AB). The University of Tennessee Institute of Agriculture, the University of Tennessee Agricultural Experiment Station, the National Institutes for Health, the manufacturer of Zylamend (New Chapter, Brattleboro, VT) or anyone affiliated with them, have not been consulted or associated with the research in this paper, or involved in the study design; in the collection, analysis, or interpretation of data; in the writing or review of the manuscript; or in the decision to submit the paper for publication.

Availability of data and materials

The data supporting the conclusions of this article are included within the manuscript. The raw data and materials of current study are available from the corresponding author upon reasonable request.

Authors' contributions

All of the authors have read and approved the manuscript. AFM conducted the research, analyzed the data, help write the paper, read and approved the final manuscript. AB helped in the design of the experiments, provided essential technical expertise/supplies/materials, conducted research, read, commented on, approved the final manuscript and provide some financial support through his NIH grant. DRD helped in the design of the experiments, provided essential technical expertise, supplies/materials, conducted research, read, commented on and approved the final manuscript. DSA provided essential technical expertise to the research, read, and approved the final manuscript. AH provided essential technical expertise to the research, read, commented on and approved the final manuscript. YZ provided essential technical expertise to the research, generated preliminary data for this research, read, commented on and approved the final manuscript. JW was primarily responsible for the design of the research, oversight and supervision of the research, writing of the manuscript, had final responsibility for the content and funded the project through his USDA Hatch grant.

Author's information

The impact of food and food components (including natural products) on delaying or inhibiting the promotion and progression of cancer and modifying the ability of relapse following the use standard therapies is the focus and line of research of the primary author (JW). If you are unfortunate enough to get an advanced form of prostate cancer (castrate-resistant prostate cancer), what can be done to enhance the effectiveness of standard therapies? This research focuses on castrate-resistant prostate cancer, a potentially deadly form of the disease, using an experimental design consistent with this condition to explore underlying mechanisms.

Ethics approval and consent to participate

Manuscripts reporting studies involving human participants, human data or human tissue must include a statement on ethics approval and consent (even where the need for approval was waived) and include the name of the ethics committee that approved the study and the committee's reference number if appropriate.

This research did not use any preclinical or animals models and did not involve human subjects. Therefore, this section is not applicable.

Consent for publication

This manuscript does not contain any individual person's data (adult or child) in any form (including individual details, images or videos). Therefore, this section is not applicable.

Competing interests

The authors have no conflicts of interest. They have not received any reimbursement fees or funding from the manufacturers or distributors of Zylamend as outlined by the publisher. None of the authors hold stock or shares that can result in financial gain in anyway, nor have applied for patents or reimbursements or fees that would constitute a conflict of

interest. The University of Tennessee Agricultural Experiment Station, the National Institutes for Health, the manufacturer of Zyflamend (New Chapter, Brattleboro, VT) or anyone affiliated with them, have not been consulted or associated with this study, or involved in the study design; in the collection, analysis, or interpretation of data; in the writing or review of the manuscript; or in the decision to submit the paper for publication. No financial support or product has been provided by New Chapter (Brattleboro, VT), manufacturer of Zyflamend, for this study. This research has been reviewed and accepted by all authors. It has not been published or submitted for publication elsewhere.

Publisher's Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Author details

¹Department of Nutrition, University of Tennessee, 1215 West Cumberland Avenue, 229 Jessie Harris Building, Knoxville, TN 37996, USA. ²Tennessee Agricultural Experiment Station, University of Tennessee, Knoxville, TN 37996, USA. ³Department of Nutrition, Laboratory for Cancer Research, University of Tennessee, 1215 West Cumberland Avenue, Room 229 Jessie Harris Building, Knoxville, TN 37996-1920, USA. ⁴Present addresses: Kellogg Eye Center, University of Michigan, 1000 Wall St, Ann Arbor, MI 48105, USA. ⁵Present addresses: Department of Cancer Biology, Thomas Jefferson University, 233 S.10th Street, Philadelphia, PA 19107, USA.

Received: 1 February 2018 Accepted: 11 June 2018

Published online: 18 June 2018

References

- Siegel RL, Miller KD, Jemal A. Cancer statistics, 2017. *CA-Cancer J Clin*. 2017; 67(1):7–30.
- Miller KD, Siegel RL, Lin CC, Mariotto AB, Kramer JL, Rowland JH, Stein KD, Alteri R, Jemal A. Cancer treatment and survivorship statistics, 2016. *CA-Cancer J Clin*. 2016;66(4):271–89.
- Whelan J, Zhao Y, Huang EC, MacDonald A, Donohoe D. In: Hardman R, Harikumar KB, editors. *Zyflamend and prostate cancer therapy*. In *Complementary and Alternative Medicines in Prostate Cancer: A Comprehensive Approach*. Boca, Raton, FL: CRC Press; 2017. p. 197–220.
- Conteduca V, Wetterskog D, Sharabiani MTA, Grande E, Fernandez-Perez MP, Jayaram A, Salvi S, Castellano D, Romanel A, Lolli C, et al. Androgen receptor gene status in plasma DNA associates with worse outcome on enzalutamide or abiraterone for castration-resistant prostate cancer: a multi-institution correlative biomarker study. *Ann Oncol*. 2017;28:1508–16.
- Taplin ME, Balk SP. Androgen receptor: a key molecule in the progression of prostate cancer to hormone independence. *J Cell Biochem*. 2004;91(3):483–90.
- Huang EC, Chen G, Baek SJ, McEntee MF, Collier JJ, Minkin S, Biggerstaff JP, Whelan J. Zyflamend reduces the expression of androgen receptor in a model of castrate-resistant prostate cancer. *Nutr Cancer*. 2011;63(8):1287–96.
- Huang EC, McEntee MF, Whelan J. Zyflamend, a combination of herbal extracts, attenuates tumor growth in murine xenograph models of prostate cancer. *Nutr Cancer*. 2012;64(5):749–60.
- Huang EC, Zhao YCG, Baek SJ, McEntee MF, Minkin S, Biggerstaff JP, Whelan J. Zyflamend, a polyherbal mixture, down regulates class I and class II histone deacetylases and increases p21 levels in castrate-resistant prostate cancer cells. *BMC Complement Altern Med*. 2014;14:68.
- Bilen MA, Lin SH, Tang DG, Parikh K, Lee MH, Yeung SC, Tu SM. Maintenance therapy containing metformin and/or Zyflamend for advanced prostate Cancer: a case series. *Case Rep Oncol Med*. 2015;2015:471861.
- Zhao Y, Donohoe D, Huang EC, Whelan J. Zyflamend, a polyherbal mixture, inhibits lipogenesis and mTORC1 signalling via activation of AMPK. *J Funct Foods*. 2015;18:147–58.
- Mohebbati A, Guttenplan JB, Kochhar A, Zhao ZL, Kosinska W, Subbaramaiah K, Dannenberg AJ. Carnosol, a constituent of Zyflamend, inhibits aryl hydrocarbon receptor-mediated activation of CYP1A1 and CYP1B1 transcription and mutagenesis. *Cancer Prev Res*. 2012;5(4):593–602.
- Subbaramaiah K, Sue E, Bhardwaj P, Du B, Hudis CA, Giri D, Kopelovich L, Zhou XK, Dannenberg AJ. Dietary polyphenols suppress elevated levels of proinflammatory mediators and aromatase in the mammary gland of obese mice. *Cancer Prev Res*. 2013;6(9):886–97.
- Sandur SK, Ahn KS, Ichikawa H, Sethi G, Shishodia S, Newman RA, Aggarwal BB. Zyflamend, a polyherbal preparation, inhibits invasion, suppresses osteoclastogenesis, and potentiates apoptosis through down-regulation of NF-kappa B activation and NF-kappa B-regulated gene products. *Nutr Cancer*. 2007;57(1):78–87.
- Kunnumakkara AB, Sung B, Ravindran J, Diagaradjane P, Deorukhkar A, Dey S, Koca C, Tong Z, Gelovani JG, Guha S, et al. Zyflamend suppresses growth and sensitizes human pancreatic tumors to gemcitabine in an orthotopic mouse model through modulation of multiple targets. *Int J Cancer*. 2012; 131(3):E292–303.
- Kim JH, Park B, Gupta SC, Kannappan R, Sung B, Aggarwal BB. Zyflamend sensitizes tumor cells to TRAIL-induced apoptosis through up-regulation of death receptors and down-regulation of survival proteins: role of ROS-dependent CCAAT/enhancer-binding protein-homologous protein pathway. *Antioxid Redox Signal*. 2012;16(5):413–27.
- Ekmekcioglu S, Chattopadhyay C, Akar U, Gabisi A Jr, Newman RA, Grimm EA. Zyflamend mediates therapeutic induction of autophagy to apoptosis in melanoma cells. *Nutr Cancer*. 2011;63(6):940–9.
- Bemis DL, Capodice JL, Anastasiadis AG, Katz AE, Buttyan R. Zyflamend, a unique herbal preparation with nonselective COX inhibitory activity, induces apoptosis of prostate cancer cells that lack COX-2 expression. *Nutr Cancer*. 2005;52(2):202–12.
- Rafailov S, Cammack S, Stone BA, Katz AE. The role of Zyflamend, an herbal anti-inflammatory, as a potential chemopreventive agent against prostate cancer: a case report. *Integr Cancer Ther*. 2007;6(1):74–6.
- Yang P, Cartwright C, Chan D, Vijjeswarapu M, Ding J, Newman RA. Zyflamend-mediated inhibition of human prostate cancer PC3 cell proliferation: effects on 12-LOX and Rb protein phosphorylation. *Cancer Biol Ther*. 2007;6(2):228–36.
- Capodice JL, Gorroochurn P, Cammack AS, Eric G, McKiernan JM, Benson MC, Stone BA, Katz AE. Zyflamend in men with high-grade prostatic intraepithelial neoplasia: results of a phase I clinical trial. *J Soc Integr Oncol*. 2009;7(2):43–51.
- Yan J, Xie B, Capodice JL, Katz AE. Zyflamend inhibits the expression and function of androgen receptor and acts synergistically with bicalutamide to inhibit prostate cancer cell growth. *Prostate*. 2012;72(3):244–52.
- Zhao Y, Collier JJ, Huang EC, Whelan J. Turmeric and Chinese goldthread synergistically inhibit prostate cancer cell proliferation and NF-kB signaling. *Funct Foods Health Dis*. 2014;4(7):312–39.
- Jeon SM. Regulation and function of AMPK in physiology and diseases. *Exp Mol Med*. 2016;48(7):e245.
- Cheng J, Zhang T, Ji H, Tao K, Guo J, Wei W. Functional characterization of AMP-activated protein kinase signaling in tumorigenesis. *Biochim Biophys Acta*. 2016;1866(2):232–51.
- Hawley SA, Davison M, Woods A, Davies SP, Beri RK, Carling D, Hardie DG. Characterization of the AMP-activated protein kinase kinase from rat liver and identification of threonine 172 as the major site at which it phosphorylates AMP-activated protein kinase. *J Biol Chem*. 1996;271(44):27879–87.
- Hawley SA, Pan DA, Mustard KJ, Ross L, Bain J, Edelman AM, Frenguelli BG, Hardie DG. Calmodulin-dependent protein kinase kinase-beta is an alternative upstream kinase for AMP-activated protein kinase. *Cell Metab*. 2005;2(1):9–19.
- Herrero-Martin G, Hoyer-Hansen M, Garcia-Garcia C, Fumarola C, Farkas T, Lopez-Rivas A, Jaattela M. TAK1 activates AMPK-dependent cytoprotective autophagy in TRAIL-treated epithelial cells. *EMBO J*. 2009;28(6):677–85.
- Luo L, Jiang S, Huang D, Lu N, Luo Z. MLK3 phosphorylates AMPK independently of LKB1. *PLoS One*. 2015;10(4):e0123927.
- Khan AS, Frigo DE. A spatiotemporal hypothesis for the regulation, role, and targeting of AMPK in prostate cancer. *Nat Rev Urol*. 2017;14(3):164–80.
- Gan RY, Li HB. Recent progress on liver kinase B1 (LKB1): expression, regulation, downstream signaling and cancer suppressive function. *Int J Mol Sci*. 2014;15(9):16698–718.
- Karacosta LG, Foster BA, Azabdaftari G, Feliciano DM, Edelman AM. A regulatory feedback loop between Ca²⁺/calmodulin-dependent protein kinase kinase 2 (CaMKK2) and the androgen receptor in prostate cancer progression. *J Biol Chem*. 2012;287(29):24832–43.
- Lin F, Marcelo KL, Rajapakshe K, Coarfa C, Dean A, Wilganowski N, Robinson H, Sevcik E, Bissig KD, Goldie LC, et al. The camKK2/camKIV relay is an essential regulator of hepatic cancer. *Hepatology*. 2015;62(2):505–20.
- Yuan W, Chen J, Shu Y, Liu S, Wu L, Ji J, Liu Z, Tang Q, Zhou Z, Cheng Y, et al. Correlation of DAPK1 methylation and the risk of gastrointestinal cancer: a systematic review and meta-analysis. *PLoS One*. 2017;12(9):e0184959.

34. Li L, Guo L, Wang Q, Liu X, Zeng Y, Wen Q, Zhang S, Kwok HF, Lin Y, Liu J. DAPK1 as an independent prognostic marker in liver cancer. *PeerJ*. 2017;5:e35568.
35. Xie JY, Chen PC, Zhang JL, Gao ZS, Neves H, Zhang SD, Wen Q, Chen WD, Kwok HF, Lin Y. The prognostic significance of DAPK1 in bladder cancer. *PLoS One*. 2017;12(4):e0175290.
36. Cai F, Xiao X, Niu X, Zhong Y. Association between promoter methylation of DAPK gene and HNSCC: a meta-analysis. *PLoS One*. 2017;12(3):e0173194.
37. Schumacher AM, Schavocky JP, Velentza AV, Mirzoeva S, Watterson DM. A calmodulin-regulated protein kinase linked to neuron survival is a substrate for the calmodulin-regulated death-associated protein kinase. *Biochemistry*. 2004;43(25):8116–24.
38. Laemmli UK. Cleavage of structural proteins during the assembly of the head of bacteriophage T4. *Nature*. 1970;227(5259):680–5.
39. Bettaieb A, Averill-Bates DA. Thermotolerance induced at a mild temperature of 40 degrees C protects cells against heat shock-induced apoptosis. *J Cell Physiol*. 2005;205(1):47–57.
40. Samali A, Cai J, Zhivotovsky B, Jones DP, Orrenius S. Presence of a pre-apoptotic complex of pro-caspase-3, Hsp60 and Hsp10 in the mitochondrial fraction of jurkat cells. *EMBO J*. 1999;18(8):2040–8.
41. Shaw RJ, Kosmatka M, Bardeesy N, Hurley RL, Witters LA, DePinho RA, Cantley LC. The tumor suppressor LKB1 kinase directly activates AMP-activated kinase and regulates apoptosis in response to energy stress. *Proc Natl Acad Sci U S A*. 2004;101(10):3329–35.
42. Karuman P, Gozani O, Odze RD, Zhou XC, Zhu H, Shaw R, Brien TP, Bozzuto CD, Ooi D, Cantley LC, et al. The Peutz-Jegher gene product LKB1 is a mediator of p53-dependent cell death. *Mol Cell*. 2001;7(6):1307–19.
43. Yang P, Sun Z, Chan D, Cartwright CA, Vijjeswarapu M, Ding J, Chen X, Newman RA. Zylamend reduces LTB4 formation and prevents oral carcinogenesis in a 7,12-dimethylbenz[alpha]anthracene (DMBA)-induced hamster cheek pouch model. *Carcinogenesis*. 2008;29(11):2182–9.
44. Tague ED, Bourdon AK, MacDonald A, Lookadoo MS, Kim ED, White WM, Terry PD, Campagna SR, Voy BH, Whelan J. Metabolomics approach in the study of the well-defined Polyherbal preparation Zylamend. *J Med Food*. 2018;21:306–316.
45. Pretlow TG, Wolman SR, Micale MA, Pelley RJ, Kursh ED, Resnick MI, Bodner DR, Jacobberger JW, Delmoro CM, Giaconia JM, et al. Xenografts of primary human prostatic carcinoma. *J Natl Cancer Inst*. 1993;85:394–8.
46. Nagabhushan M, Miller CM, Pretlow TP, Giaconia JM, Edgehouse NL, Schwartz S, Kung HJ, de Vere White RW, Gumerlock PH, Resnick MI, et al. CWR22: the first human prostate cancer xenograft with strongly androgen-dependent and relapsed strains both in vivo and in soft agar. *Cancer Res*. 1996;56(13):3042–6.
47. Tepper CG, Boucher DL, Ryan PE, Ma AH, Xia L, Lee LF, Pretlow TG, Kung HJ. Characterization of a novel androgen receptor mutation in a relapsed CWR22 prostate cancer xenograft and cell line. *Cancer Res*. 2002;62(22):6606–14.
48. Sramkoski RM, Pretlow TG, Giaconia JM, Pretlow TP, Schwartz S, Sy MS, Marengo SR, Rhim JS, Zhang D, Jacobberger JW. A new human prostate carcinoma cell line, 22Rv1. *In Vitro Cell Dev Biol Anim*. 1999;35(7):403–9.
49. Hardie DG. AMPK: a target for drugs and natural products with effects on both diabetes and cancer. *Diabetes*. 2013;62(7):2164–72.
50. Korsse SE, Peppelenbosch MP, van WW. Targeting LKB1 signaling in cancer. *Biochim Biophys Acta*. 2013;1835(2):194–210.
51. Song P, Xie Z, Wu Y, Xu J, Dong Y, Zou MH. Prot10, lines 222 and 224. Ein kinase Czeta-dependent LKB1 serine 428 phosphorylation increases LKB1 nucleus export and apoptosis in endothelial cells. *J Biol Chem*. 2008;283(18):12446–55.

Ready to submit your research? Choose BMC and benefit from:

- fast, convenient online submission
- thorough peer review by experienced researchers in your field
- rapid publication on acceptance
- support for research data, including large and complex data types
- gold Open Access which fosters wider collaboration and increased citations
- maximum visibility for your research: over 100M website views per year

At BMC, research is always in progress.

Learn more [biomedcentral.com/submissions](https://www.biomedcentral.com/submissions)

