**Research Article**

**R-Limonene Enhances Differentiation and 2-Deoxy-D-Glucose Uptake in 3T3-L1 Preadipocytes by Activating the Akt Signaling Pathway**

Ilavenil Soundharrajan,1 Da Hye Kim, 2 Srigopalram Srisesharam,1 Palaniselvem Kuppusamy,1 and Ki Choon Choi 1

1Grassland and Forage Division, National Institute of Animal Science, Rural Development Administration, Cheonan 31000, Republic of Korea

2Laboratory of Animal Physiology, Graduate School of Agricultural Science, Tohoku University, Aoba, Sendai 980-8577, Japan

Correspondence should be addressed to Ki Choon Choi; choiwh@korea.kr

Ilavenil Soundharrajan and Da Hye Kim contributed equally to this work.

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Adipocyte is an important place for lipid storage. Defects in lipid storage in adipocytes can lead to lipodystrophy and lipid accumulation in muscle, liver, and other organs. It is the condition of mixed dyslipidemia which may favor the development of insulin resistance via lipotoxic mechanisms. Our objective of the study was to investigate the potential role of R-limonene (LM) on differentiation, lipid storage, and 2-deoxy-D-glucose (2DG) uptake in 3T3-L1 preadipocytes. Genes and proteins associated with differentiation, lipid accumulation, 2DG uptake, and its signaling pathways in the adipocytes were analyzed using qPCR and western blot methods. LM treatment increased differentiation, lipid accumulation, and the expression of adipogenic and lipogenic markers such as C/EBP-\(\alpha\), C/EBP-\(\beta\), PPARY, SREBP-1, RXR, FAS, and adiponectin. However, the LM concentration at 10\(\mu\)M decreased \((p < 0.05)\) adipogenesis and lipogenesis via regulating key transcriptional factors. LM treatment increased activation of Akt by increasing its phosphorylation, but p44/42 activation was not altered. MK-2206, an Akt specific inhibitor, reduced the activation of Akt phosphorylation whereas LM treatment aborted the MK-2206 mediated inhibition of Akt activation. LM enhanced glucose uptake in differentiated adipocytes. Overall data suggested that LM treatment favored lipid storage and glucose uptake in adipocytes via activation of key transcriptional factors through activation of Akt phosphorylation in 3T3-L1 adipocytes.

1. Introduction

Differentiation of adipocytes is an essential process for the lipid storage. Inhibiting adipocyte differentiation alone is not an appropriate way to prevent obesity because adipocytes provide a safe place for lipid accumulation. Defects in adipogenesis can lead to stimulating lipodystrophy. In this condition, lipids are accumulated in muscle, liver, and other organs which develop the insulin resistance [1, 2]. Animal and human experiments have proved that type 2 diabetes mellitus is closely associated with adipogenesis and its factors [3].

Adipogenesis is a complex process that is regulated by a cascade of transcription factor and other regulatory proteins. Numerous positive and negative factors have been involved in adipocyte differentiation [4–7]. Among these, C/EBP-\(\beta\) (CCAAT/enhancer-binding protein-\(\beta\)) and C/EBP-\(\delta\) act together and induce the expression of PPAR-\(\gamma\) 2 (peroxisome proliferator-activated protein-\(\beta\)) and C/EBP-\(\alpha\) (CCAAT/enhancer-binding protein-\(\alpha\)). These factors are known to regulate genes and proteins associated with adipogenesis and lipogenesis [8].

The Akt kinases play a critical role in adipogenesis and glucose transport [9]. Lack of or inhibition of Akt activation in fibroblast displays an inability to differentiate preadipocyte into adipocytes. In addition, activation of Akt can promote the differentiation of preadipocytes into mature adipocytes.
An early study indicated that strengthens the role of dietary monoterpenes in human life anti-breast cancer and anti-colorectal carcinoma effects. It could provide significant health beneficial effects especially body weight per one day. Consumption of monoterpenes limonene at a range of 0.27 milligrams per kilogram of chewing gum. Human consumes both synthetic and natural asafragrance additive in candy, ice cream, orange juice, and is relatively high. LM with lemons-like odor has been used pathway [26]. Dietary intake of monoterpenes by the human and extracellular signal-regulated kinases (ERK) signaling respectively [27].

Limonene can enhance glucose uptake in 3T3-L1 adipocytes. In addition, limonene has been shown to be helpful in relieving heartburn and gastroesophageal reflux disorder; [20, 21]; in addition, limonene is antidiabetic properties. In addition, they can stimulate glucose uptake herbivore deterring, pollinator attracting properties, antioxidant, antipollagistic, antitumor, antiviral, and antiinvasive properties. In addition, they can stimulate glucose uptake in C2C12 cells [16–19]. Among monoterpenes, limonene (Figure 1) and α-pinene exhibit potent antitumor properties [20, 21]; in addition, limonene has been shown to be helpful in relieving heartburn and gastroesophageal reflux disorder; it supports normal peristalsis [22].

Limonene can enhance glucose uptake in 3T3-L1 adipocytes via glucose transporters [23]. It possesses antidiabetic effects by preventing dyslipidemia [24]. It also prevents insulin resistance [25], LM enhanced mitochondrial biogenesis, and elevated protein levels of hormonal sensitive lipase (HSL), perilipin (PLIN), AMP-activated protein kinase (pAMPK), Phospho acetyl CoA carboxylase (pACC), Acy coenzyme A oxidase (ACO), cytochrome c oxidase subunit 4 (COX4), carnitine palmitoyltransferase 1 (CPT1), and cytochrome C (CYT-C) at the concentration of 25 and 50μM of limonene, which are involved in lipolysis and lipid catabolism via activation of β-adrenergic receptor-3 (β3-AR) and extracellular signal-regulated kinases (ERK) signaling pathway [26]. Dietary intake of monoterpenes by the human is relatively high. LM with lemons-like odor has been used as a fragrance additive in candy, ice cream, orange juice, and chewing gum. Human consumes both synthetic and natural limonene at a range of 0.27 milligrams per kilogram of body weight per one day. Consumption of monoterpenes could provide significant health beneficial effects especially anti-breast cancer and anti-colorectal carcinoma effects. It strengthens the role of dietary monoterpenes in human life [22]. An early study indicated that D-limonene exhibited antiobesity activity in adipocytes and high-fat diet induced obesity at the concentration of 50μM and 0.5% limonene, respectively [27].

2. Materials and Methods

2.1. Cell Culture and Chemicals. The 3T3-L1 preadipocytes cell line was obtained from the American Type Culture Collection [Rockville, MD, USA]. Dulbecco modified Eagle medium [DMEM] and fetal bovine serum (FBS) were procured from Gibco-BRL [Gaithersburg, MD, USA]. Kits for mRNA extraction, cDNA synthesis, and qPCR were purchased from Bio-Rad [Hercules, CA, USA]. R-limonene (Sigma Aldrich, #183164) and rosiglitazone were obtained from Sigma Aldrich (St. Louis, MO, USA). MK-2206 was obtained from Selleckchem (Houston, TX, USA). Monoclonal antibodies (PPARY, RXR, C/EBP-α, SREBP-1, aP2, FAS, adiponectin, ACC, TAKT, pAKT (Serine 473), p44/42, pp44/42 (Thr 202/ Tyr204), AMPK-α, pAMPK-α (Thr172), and GAPDH) used in the study were obtained from Cell Signaling Technology (Danvers, MA, USA).

2.2. Cytotoxicity of Limonene (LM). Ez-cytox assay kit (iTS-Bio, Korea) was used to determine the cytotoxic effects of LM. Briefly, 3T3-L1 preadipocytes (ATCC, USA) were seeded into 96-well cell culture plates at the density of 1 × 10⁴ cells/well and incubated at 37°C with 5% CO₂ for 24 hours. These cells were then treated with different concentrations of cyclic terpene limonene (LM) and incubated for 1 day, 2 days, and 10 days; every 48 h fresh media was replaced with different concentration of LM until the end of experimental periods. Ten microliters of water-soluble tetrazolium (WST) reagent was added to each well and incubated at 37°C with 5% CO₂ for 1-2 hours. Absorbance (color intensity) of each well was measured at a wavelength of 450 nm using a Spectra Count microplate reader [28].

2.3. Differentiation Induction. 3T3-L1 pre adipocytes were seeded into 6- and 12-well cell culture plates at a density of 3 × 10⁴ and 1.5 × 10⁶ cells/well, respectively, and incubated at 37°C with 5% CO₂. Every 48 h, the culture medium was replenished with fresh medium. When cells reached 100%, further incubation was performed for another two days. Growth medium was then replaced with a differentiation induction medium (DMI) (0.5 mM 3-isobutyl-1-methylxanthine, 1μM dexamethasone, and 1μg/mL insulin) containing LM, agonist, and antagonist [28].

2.4. Quantification of Lipid Using Oil Red O Staining. Differentiated 3T3-L1 adipocytes in 6-well plates were fixed with 2 mL of 10% formalin for 1 h and then rinsed with 40%
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Figure 2: Cytotoxic effect of LM on 3T3-L1 preadipocytes. (a) LM effect on 3T3-L1 preadipocytes viability after 24 h treatment; (b) LM effect on 3T3-L1 preadipocytes viability after 48 h treatment; (c) LM effect on 3T3-L1 preadipocytes viability after 10 days of treatment.

isopropanol. After rinsing, 3 mL of Oil Red O staining solution was added to each well. Plates were incubated at room temperature for 15 min and washed three times with distilled water. Stained cells were photographed with an inverted microscope [CKX41, Olympus Corporation, Tokyo Japan]. Additionally, Oil Red O stain was eluted with 100% isopropanol and measured at 490 nm [28].

2.5. Glucose Uptake Assay. Eight-day differentiated cells were incubated with serum-free media for 12 h. Then, serum-starved cells were treated with LM (5 μM), insulin (10 μM), and MK-2206, an antagonist for Akt (8 nM) individually for 2 h at 37°C with 5% CO2 followed by addition of 10 mM 2DG and incubated for another 20 min. Glucose uptake was then performed using glucose uptake assay kit according to the manufacturer’s protocol (glucose uptake colorimetric assay kit, Abcam, UK).

2.6. Quantification of Gene Expression Using Quantitative RT-PCR. Total RNA was extracted from experimental cells using RNasey lipid mini kit (Qiagen, MD, USA) and quantified by Spectramax i3 ( Molecular devices, CA, USA). Total RNA (500 ng) was then reverse transcribed using iScript cDNA synthesis kit (Hercules, CA, USA). Gene transcripts in the experimental samples were quantified with SYBR Green-based qPCR using specific primers for C/EBP-α (F-gcagaggagatacagaggag, R- acagctcctaatctcccaaca); C/EBP-β (F-gtggctggactgagcctgca, R- aacaccccccagaggaaca); PPAR γ2 (F-ggtgttcaccagatggcctgctgtaa); Adiponectin (F-cctctctttcatcctgca, R- tccctccctctctctctgac); FAS (F-cctctccataagttgtaa, R-atccgagtgactcagcaa); SREBP-1 (F-gaatggaggagacgcctta, R-tatcctcaaaggtggactg); β-actin (F-cctcctctacgcct, R- atgtcagcagcatttcc) on a CFX 96 Real Time PCR detection system (Hercules, CA, USA).

All gene expressions were normalized against housekeeping gene β-actin [28].

2.7. Protein Extraction and Immunoblotting. Proteins were extracted from experimental adipocytes with RIPA lysis buffer containing protease and phosphatase inhibitors cocktail (Roche, Switzerland and Sigma Aldrich, USA). Protein quantification was performed with the Pierce BCA protein assay kit (Thermo Fisher Scientific, MA, USA). Immunoblotting was performed with Cell Signaling Technology monoclonal antibodies [29]. The intensity of immunoreacted bands was quantified with ImageJ software version 1.49 (32 bit), (Wayne Rasband, National Institute of Health, USA).

2.8. Statistical Analysis. Experimental data were subjected to one-way ANOVA with post hoc test and multivariate comparisons with Duncan test using the statistical package of social science [SPSS-Version 16.0, SPSS, Inc., USA]. Statistical significance was considered when p value was less than 0.05.

3. Results

3.1. Cytotoxic Effects of LM on 3T3-L1 Preadipocytes. Preadipocytes were treated with different concentrations (10 μM to 300 μM) of LM and incubated for 1 day, 2 days, and 10 days. LM treatment failed to affect the cell viability significantly at a concentration of 10 μM or less. Furthermore, the increase of LM concentration slightly reduced cell viability compared to the control. However, there was no significant difference in cell viability between control and LM treatment at any concentration tested (Figure 2).

3.2. Effects of LM on 3T3-L1 Preadipocyte Differentiation. Based on microscopic observation, adipocytes treated with
differentiation media started to increase the accumulation of lipid droplets in adipocytes (Figure 3). Such lipid accumulation and differentiation were accelerated \((p < 0.05)\) after the addition of LM at 2.5 \(\mu M\) and 5 \(\mu M\) compared to those in control adipocytes. Lipid accumulation in maturing adipocytes after treatment with LM at 10 \(\mu M\) was reduced \((p < 0.05)\) compared to that in control adipocytes. After extracting Oil Red O dye from experimental adipocytes, higher absorbance values in adipocytes treated with LM at 2.5 and 5 \(\mu M\) but lower absorbance value in adipocytes treated with LM at 10 \(\mu M\) compared to control adipocytes were observed, indicating that the LM could regulate the lipid accumulation in adipocytes in a dose dependent-manner. Maximum lipid accumulation was noted in adipocytes on the 5th and 10th day after treatment with 5 \(\mu M\) of LM without any alteration in cell morphology. Therefore, we used 5 \(\mu M\) of LM for further experiment.

3.3. Effect of LM on mRNA Expression in 3T3-L1 Adipocytes. We next investigated how LM regulated the expression of PPAR\(\gamma\) and C/EBPs in adipocytes. Expression levels of C/EBP-\(\alpha\), C/EBP-\(\beta\), and PPAR\(\gamma\) were significantly \((p < 0.05)\) upregulated in adipocytes treated with LM at 5 \(\mu M\) compared to those in control adipocytes. The expression levels of adiponectin, FAS, and SREBP-1 mRNAs were also significantly increased by LM treatment compared to those in control adipocytes (Figure 4(a)). LM treatment at a concentration of 10 \(\mu M\) significantly \((p < 0.05)\) downregulated PPAR\(\gamma\), C/EBP-\(\alpha\), C/EBP-\(\beta\), SREBP-1, FAS, and adiponectin mRNA expression levels in differentiated adipocytes compared to control (data not shown).

3.4. Effect of LM on Adipogenic and Lipogenic Proteins Expression in 3T3-L1 Adipocytes. Expression levels of protein expressions related to adipocyte differentiation in adipocytes on the 10th day after treatment with LM were analyzed. Treatment with LM at 5 \(\mu M\) significantly \((p < 0.05)\) increased protein levels of PPAR\(\gamma\), RXR, C/EBP-\(\alpha\), SREBP-1, and their downstream targets such as ap2, FAS, ACC, and adiponectin, compared to control adipocytes (Figure 4(b)).

3.5. Comparative Effects of LM and Rosiglitazone (RGZ) on Differentiation and PPAR\(\gamma\) Expression. We also compared the effects of LM with those of PPAR-\(\gamma\) agonist (RGZ) in adipocytes. Confluent preadipocytes were treated with RGZ or LM \((0.1 \mu M \text{ or } 5 \mu M, \text{ respectively})\) for 48h. Results showed that RGZ treatment significantly increased lipid accumulation and PPAR-\(\gamma\) protein expression. Similarly, LM treatment significantly increased lipid accumulation and PPAR-\(\gamma\) protein expression compared to control adipocytes (Figures 5(a)–5(c)).
3.6. Effects of LM on Signaling Pathways. Next, signaling pathways involved in the regulation of adipocyte differentiation and lipid accumulation in control and LM treated adipocytes were investigated. LM treatment induced activation of Akt by increasing its phosphorylation at serine 473 but decreased AMPK-α phosphorylation at Thr172 in adipocytes on the 10th day compared to the control adipocytes. However, p44/42 pathway remained the same in the control and experimental adipocytes (Figure 5(d)).

3.7. A Competitive Study between LM and MK-2206 on Akt Phosphorylation. MK-2206, an antagonist of Akt, was used in this study to further evaluate the role of Akt involved in the effect of LM on adipocyte differentiation. Adipocytes treated with MK-2206 for 48h inhibited lipid accumulation and downregulated PPAR-γ expression compared to the control adipocytes. Cotreatment with both LM and MK-2206 (Akt inhibitor) significantly \( (p < 0.05) \) increased lipid accumulation and upregulated PPAR-γ expression in...
Figure 5: (a)-(d) A comparative study between R-limonene (LM) and rosiglitazone (RGZ) on lipid accumulation and PPAR-γ expression in adipocytes. Adipocytes differentiated with RGZ (0.1 μM) and LM (5 μM) separately in DMI. (a) Microscopic visualization of differentiated adipocytes on day 10; (b) percentage of lipids in the experimental adipocytes on day 10; (c) PPAR-γ protein expression in the experimental adipocytes on day 10. Experiments were performed in triplicate and repeated three times with the same results. Bars display mean ± SEM and statistical analysis was performed by one-way ANOVA. *p < 0.05 indicates a statistically significant difference compared to control and LM treatments; #p < 0.05 indicates a statistically significant difference compared to control adipocytes. (d) Impact of LM on signaling pathways associated with an adipocyte differentiation. Adipocytes were differentiated with differentiation induction medium (DMI) in the presence/absence of LM for 10 days. Experiments were performed in triplicate and repeated three times with the same results. Bars display mean ± SEM and statistical analysis was performed by one-way ANOVA. *p < 0.05 indicates a statistically significant difference compared to control adipocytes.

differentiated adipocytes, indicating that LM treatment could abolish MK-2206 mediated downregulation of PPAR-γ and optical density of Oil Red O stain. These data confirmed that LM treatment could increase differentiation and lipid accumulation via Akt activation by increasing its phosphorylation at serine 473 (Figures 6(a) and 6(b)).

3.8. LM on Glucose Uptake in Differentiated 3T3-L1 Adipocytes. Differentiated adipocytes were then treated with LM, insulin, MK-2206, and MK-2206+LM separately and incubated for 2 h in a CO₂ incubator after serum starvation. Insulin treatment increased 2DG uptake in 3T3-L1. Treatment with MK-2206, an antagonist for Akt, reduced glucose uptakes. LM treatment significantly enhanced 2DG uptake in 3T3-L1 adipocytes. MK-2206 + LM treatment also increased glucose uptakes as compared to control adipocytes. Overall data suggest that the LM could stimulate 2DG uptake in differentiated adipocytes through the Akt signaling pathway in differentiated adipocytes (Figure 6(c)).
4. Discussion

Adipogenesis is a sequential process accompanied by the dramatic increase in the expression of adipocyte genes [7]. The dysfunctional adipose tissue is characterized by reduced adipogenesis, increased cellular senescence, and inflammation. Such dysfunction impairs lipid storage and dysregulates production of adipokines and cytokines, leading to metabolic disorder, adipose tissue inflammation, insulin resistance, and type-2 diabetes [30]. An adequate adipogenesis is essential
to sequester lipids in adipose tissues to prevent ectopic fat deposition and insulin resistance development. Increases in lipid storage capacity are believed to play a vital role in metabolic regulation [31]. In the present study, LM treatment induced differentiation and lipid accumulation in adipocytes at 5th and 10th day after treatment. However, cells treated with LM at the concentration higher than 10 μM inhibited lipid accumulation with slight morphology changes compared to control.

Furthermore, we analyzed effects of LM on expression levels of C/EBP-β, PPARγ, and C/EBP-α known to be essential factors that work sequentially and cooperatively to induce differentiation and lipid accumulation in adipocytes [7]. In the present study, LM induced C/EBP-β, C/EBP-α, PPARγ2, and RXR expression. Crosstalks among PPAR-γ, RXR, and C/EBPα at common regulatory sites are important in the regulation of genes involved in adipocyte differentiation, for lipogenic function requires PPAR-γ/RXR heterodimers while the vital function is mediated by PPAR-γ/RXR α or PPAR-γ / RXRγ heterodimers [32]. C/EBP-β is mainly involved in adipocyte differentiation. It activates key transcriptional factors such as PPARγ and C/EBP-α [33]. Upregulation of these genes was consistent with lipid accumulation results based on Oil Red O staining.

Adiponectin is exclusively secreted in fully differentiated adipocytes. It regulates many metabolic processes via enhancing insulin sensitivity in muscle and liver or by activating fatty acid oxidation in different tissues [34]. In the present study, LM treatment increased mRNA of adiponectin and their protein expression levels in differentiated adipocytes compared to control, suggesting that LM might promote the glucose uptake by cells.

PPARγ and C/EBP-α can regulate their downstream targets such as aP2, FAS, LPL, leptin, and adipQ. These downstream targets can trigger the synthesis of fatty acids and triglycerides synthesis [5]. Adipocytes can dramatically induce lipogenesis after differentiation induction, leading to insulin sensitivity. FAS, aP2, and ACC are known to be increased 10- to 100-fold during differentiation [35]. In the present study, LM treatment increased FAS, ACC, and adiponectin levels in differentiated adipocytes as compared to control, indicating that LM may trigger insulin sensitivity via activating adipocyte differentiation and fatty acid metabolism by regulating key transcriptional factors and their downstream targets. Adipocyte binding protein (aP2) is a key mediator involved in the transport of fatty acids and their metabolism regulation. It is regulated by PPARγ during differentiation and lipid accumulation [36]. LM significantly upregulated aP2 expression in differentiated adipocytes as compared to the control. SREBPs can enhance genes associated with cholesterol and fatty acid biosynthesis and uptake [37]. Furthermore, SREBP-1c induces PPARγ expression in adipocytes. In collaboration with C/EBP-α, SREBP-1c can activate adipocyte markers [38]. Furthermore, activation of AMPK directly phosphorylates precursor of SREBP-1c at ser 372 residue which controls proteolytic maturation and translocation of mature SREBP-1c into the nucleus which leads to reducing the expression of ACC and FAS [39–41]. The present study is indicating that LM treatment could increase mature SREBP-1c expression via AMPK-α downregulation and increased ACC and FAS expression levels which are the key enzymes involved in the lipogenesis.

The insulin signaling pathway plays a crucial role in preadipocyte differentiation [29]. Akt is particularly important in adipocyte differentiation and insulin metabolic functions [42]. Many researchers have reported that Akt regulates PPARγ and adipocyte differentiation [12, 43]. In the present study, insulin, DEX, and IBMX mixture increased Akt phosphorylation at ser 473. Furthermore, activation of Akt by increasing its phosphorylation was accelerated by LM treatment. These data suggest that activation of Akt can activate genes associated with adipocyte differentiation. It is known that Akt phosphorylation can promote adipocyte differentiation via upregulating PPAR γ [44]. In addition, cotreatment with Akt inhibitor and LM significantly increased lipid accumulation and differentiation, indicating that LM treatment could abrogate inhibition of Akt phosphorylation induced by Akt inhibitor and increase phosphorylation of Akt. Taken together, these results suggest that LM might accelerate adipogenesis and lipogenesis via Akt signaling pathway in adipocytes. In the meantime, the Erk1/2 pathway remained the same in control and LM treated cells.

Finally, we examined the effect of LM on glucose uptake properties of cells because LM upregulated Akt phosphorylation at serine 473 and adiponectin level in 3T3-L1 adipocytes. It is known that insulin signaling pathway is activated under nutrient availability. GLUT-4 translocation is required for insulin-dependent PI3K/AKT activation [45]. Akt kinase, a Ser/Thr kinase, is activated by insulin and certain growth factors. Several reports have shown that Akt function is downstream of the PI3 kinase. Akt kinase critically contributes to glucose uptake and metabolism through PI3K kinase in the insulin signaling pathway [14]. In this study, adipocyte treated with LM or insulin increased glucose uptake in 3T3-L1 adipocytes. However, MK-2206, a specific inhibitor for Akt activation, caused a reduction in glucose uptake. These results confirmed that LM could enhance glucose uptake in adipocytes via activating Akt signaling pathway.

5. Conclusions

In summary, the present study showed that limonene (LM) could induce differentiation and glucose uptake in 3T3-L1 preadipocytes. LM regulated adipogenesis and lipogenesis via induction of key transcriptional factors such as PPARγ, C/EBP-α, and C/EBP-β as well as their downstream targets by activation of Akt signaling pathway. Our overall data suggest that limonene is a promising compound found in the natural products that can favor lipid storage in adipocytes and glucose uptake via the Akt signaling pathway.

Data Availability

The data used to support the findings of this study are available from the corresponding author upon request.
Conflicts of Interest
The authors have no conflicts of interest to disclose.

Authors’ Contributions
Ilavenil Soundharrajan and Ki Choon Choi conceived and designed the experiments; Ilavenil Soundharrajan, Da Hye Kim, Srigopalram Srisesharam, and Palaniselvam Kuppusamy performed the experiments; Srigopalram Srisesharam and Palaniselvum Kuppusamy helped in writing the paper; Ki Choon Choi contributed reagents/materials/analysis tools. Ilavenil Soundharrajan and Da Hye Kim contributed equally to this work.

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