Diacylglycerol lipase-α and -β control neurite outgrowth in Neuro-2a cells through distinct molecular mechanisms

Kwang-Mook Jung, Giuseppe Astarita, Dean Thongkham, and Daniele Piomelli

Department of Pharmacology (K.-M.J., G.A., D.T., D.P.) and Department of Biological Chemistry (D.P.), University of California, Irvine, Irvine, CA; Unit of Drug Discovery and Development, Italian Institute of Technology, Genova, Italy (G.A., D.P.).
Running Title: 2-arachidonoyl-sn-glycerol biosynthesis in neurite outgrowth

Corresponding Author: Dr. Daniele Piomelli

Department of Pharmacology, 3101 Gillespie NRF, University of California, Irvine, CA 92697-4625.
Tel: (949) 824-6180
Fax: (949) 824-6305
E-mail: piomelli@uci.edu

Text pages: 30
Table: 0
Figures: 5
References: 42
Supplemental Table: 1
Supplemental Figures: 2
Words in Abstract: 216
Words in Introduction: 618
Words in Discussion: 1142

Abbreviations: ABHD, α-β-hydrolase domain; CB, cannabinoid; DAG, 1,2-diacyl-sn-glycerol; DGL, DAG lipase; DHPG, (S)-3,5-dihydroxyphenylglycine; DMEM, Dulbecco’s Modified Eagle’s Medium; EGFP, enhanced green fluorescence protein; ER, endoplasmic reticulum; ESI, electrospray ionization; FA, fatty acid; FBS, fetal bovine serum; FGF, fibroblast growth factor; GAPDH, glyceraldehyde-3-phosphate dehydrogenase; HDA, heptadecanoic acid; HDG, (1,3)-heptadecanoyl-sn-glycerol; LC-MS, liquid chromatography-mass spectrometry; MAG, monoacyl glycerol; mGlu, metabotropic glutamate; PCR, polymerase chain reaction; PLC, phospholipase C; RA, all-trans-retinoic acid; SDS-PAGE, sodium dodecylsulfate-polyacrylamide gel electrophoresis; shRNA, small-hairpin RNA; SIM, selected ion-monitoring; 2-AG, 2-arachidonoyl-sn-glycerol.
Abstract

The endocannabinoid 2-arachidonoyl-sn-glycerol (2-AG) is produced through hydrolysis of 1,2-diacyl-sn-glycerol (DAG), which is catalyzed by DAG lipase (DGL). Two DGL isoforms have been molecularly cloned, but their respective roles in endocannabinoid signaling have not been fully elucidated. Here we report that DGL-α and DGL-β may contribute to all-trans-retinoic acid (RA)-induced neurite outgrowth in neuroblastoma Neuro-2a cells through distinct mechanisms. RA-induced differentiation of Neuro-2a cells was associated with elevations of cellular 2-AG levels and DGL activity, which were accompanied by temporally separated transcription of DGL-α and DGL-β mRNA. Knock-down of either DGL-α or DGL-β expression attenuated neurite outgrowth, which indicates that both isoforms contribute to neuritogenesis. Immunostaining experiments showed that DGL-β is localized to peri-nuclear lipid droplets, whereas DGL-α is found on plasma membranes. Following RA-induced differentiation, both DGL-α- and DGL-β-GFP were distributed also in neurites, but in distinguishable patterns. Overexpression of either DGL-α or DGL-β increased the number of neurite-bearing cells, but DGL-β caused substantially larger morphological changes than DGL-α did. Finally, the CB₁ antagonist rimonabant (1 μM) inhibited DGL-α-induced neuritogenesis, whereas it had no such effect on DGL-β-induced morphological differentiation. The results indicate that RA-induced DGL expression is required for neurite outgrowth of Neuro-2a cells. The findings further suggest that DGL-α and -β may regulate neurite outgrowth by engaging temporally and spatially distinct molecular pathways.
Introduction

The endocannabinoids are a class of endogenous lipids that bind to and activate cannabinoid receptors. 2-arachidonoyl-sn-glycerol (2-AG) and anandamide have been identified as two major endocannabinoids and have been implicated in a variety of physiological processes (Mackie and Stella, 2006; Chevaleyre et al., 2006). In the adult brain, the endocannabinoids may act as locally restricted retrograde messengers, which are produced at postsynaptic sites, travel across the synaptic cleft, and engage presynaptic CB1-type cannabinoid receptors to regulate the release of glutamate, γ-aminobutyric acid (GABA) and other neurotransmitters (Piomelli, 2003; Freund et al., 2003; Kano et al., 2009).

Evidence suggests an essential role of the endocannabinoid system during early neural development (Harkany et al., 2007; Galve-Roperh et al., 2006). Molecular players of the endocannabinoid system – including CB1 receptors, endocannabinoid molecules and their synthesizing and degrading enzymes – have been identified from the earliest stages of embryonic development to postnatal maturation (Harkany et al., 2008; Aguado et al., 2005; Aguado et al., 2006; Berghuis et al., 2005). CB1 receptors are enriched in axonal growth cones of the pyramidal neurons and GABAergic interneurons of the developing rodent brain, and neurons lacking CB1 receptors displayed impaired axonal path finding and target selection (Berghuis et al., 2007; Mulder et al., 2008). These data identify endocannabinoids as axon guidance cues and implicate CB signaling in the regulation of synaptogenesis and target selection in vivo. In cultured cells, activation of CB1 receptors causes neurite outgrowth by regulating the proteasomal degradation of Rap1GAPII and activating Rap1 and Src-Stat3 signaling (Jordan et al., 2005; He et al., 2005; Bromberg et al., 2008). Moreover, inhibition of 2-AG synthesis attenuates fibroblast growth factor (FGF)-induced neurite outgrowth in primary cerebellar neurons (Williams et al., 2003; Bisogno et al., 2003), suggesting that 2-AG might be specifically involved in the activation of CB1 receptor-dependent neurite outgrowth.
2-AG is produced from the sequential hydrolysis of membrane phosphatidylinositol-4,5-
bisphosphate into 1,2-diacyl-sn-glycerol (DAG), which is catalyzed by phospholipase C-β (PLC-
β), and then of DAG into 2-AG, which is catalyzed by DAG lipase (DGL) (Stella et al., 1997;
Piomelli, 2003). 2-AG is further cleaved by monoacylglycerol lipase or α-β-hydrolase domain
(ABHD)-6 (Piomelli, 2004; Blankman et al., 2007), to be deactivated. Two mammalian isoforms
of DGL have been cloned: DGL-α and DGL-β (Bisogno et al., 2003). DGL-α is expressed in the
hippocampus, striatum, ventral tegmental area and cerebellum of adult mouse brain (Katona et
al., 2006; Yoshida et al., 2006; Uchigashima et al., 2007; Mátyás et al., 2008), where it is
thought to initiate 2-AG-mediated signaling at excitatory synapses (Katona and Freund, 2008;
Tanimura et al., 2010; Gao et al., 2010). By contrast, the functional roles of DGL-β in the adult
brain are largely unknown. Time-dependent expression for both DGL-α and DGL-β has been
observed in neurons during development at developing axonal tracts (Watson et al., 2008;
Berghuis et al., 2007; Buckley et al., 1998; Bisogno et al., 2003), however, the contribution of
DGL and 2-AG signaling to the control of neurite outgrowth has not been investigated on the
molecular level.

In the present study, we asked whether transcriptional control of DGL expression and
subsequent activation of 2-AG-mediated signaling contribute to all-trans-retinoic acid (RA)-
induced neuronal differentiation in neuroblastoma Neuro-2a cells. We found that RA stimulates
expression of both DGL-α and DGL-β, and elevates 2-AG levels, which are required for neurite
outgrowth in Neuro-2a cells. Both α- and β-isoform of DGL, upon exogenous expression in
Neuro-2a cells, increased 2-AG levels and triggered neurite outgrowth. Unexpectedly, we found
segregative signaling characteristics between the two DGL isoforms, e.g., distinguishable
subcellular localization and differential sensitivity to CB1 receptor blockade. Our findings suggest
that transcriptional regulation of DGL-α and DGL-β contributes to RA-induced neurite outgrowth
through temporally and spatially distinct cellular signaling pathways.
Materials and Methods

Chemicals

Hexadecanoic acid, heptadecanoic acid, octadecanoic acid, Δ⁹ octadecenoic (oleic) acid, Δ⁹,12 octadecadienoic acid, Δ⁹,12,15 octadecatrienoic acid, Δ⁸,11,14 eicosatrienoic acid, eicosapentaenoic acid, docosahexaenoic acid, (1,3)-palmitoyl-sn-glycerol, (1,3)-heptadecanoyl-sn-glycerol, (1,3)-stearoyl-sn-glycerol, (1,3)-linolenoyl-sn-glycerol, (1,3)-eicosatrienoyl-sn-glycerol, (1,3)-docosahexaenoyl-sn-glycerol were from Nu-Chek Prep (Elysian, MN). 2-AG, 2-linoleoyl-sn-glycerol, noladin ether, JZL184 and WWL70 were from Cayman Chemicals (Ann Arbor, MI), and 2-oleoyl-sn-glycerol and carbachol from Sigma-Aldrich (St. Louis, MO). (S)-3,5-dihydroxyphenylglycine (DHPG) was obtained from Tocris (Ellisville, MO). Rimonabant (SR141716A) was from RTI International (Research Triangle Park, NC). Solvents were purchased from Burdick and Jackson’s (Muskegon, MI).

Plasmids

We amplified the full-length coding sequence of DGL-β by polymerase chain reaction (PCR), using first-strand mouse brain cDNAs as a template. High Fidelity PCR Master (Roche, Indianapolis, IN) was used for the amplification, following the manufacturer’s protocol. The primers were 5’-DGL-β (5’-GTGGGAGGTGCGCCATGCC-3’) and 3’-DGL-β (5’-CGGTACACTTGAGCCGCCTTGCC-3’). The PCR product was subcloned into a pEF-V5-His vector by TOPO cloning (Invitrogen, Carlsbad, CA). Mouse DGL-α and DGL-α-pEGFP constructs were prepared as previously described (Jung et al., 2007). A construct encoding an enhanced green fluorescence protein (EGFP)-fusion protein to the C-terminal of DGL-β was generated in a pEGFP-N2 vector (Clontech, Mountain View, CA) using a Sal I site. All constructs obtained through PCR amplifications were verified by DNA sequencing. Cloning and screening of small-hairpin RNA (shRNA) constructs targeting mouse DGL-β were performed as
previously described for DGL-α shRNA (Jung et al., 2007). Briefly, we designed the shRNA constructs that contained both a cytomegalovirus promoter-driven GFP and a U6 promoter-driven shRNA expression system using BLOCK-iT RNAi Designer (https://rnaidesigner.invitrogen.com/mae-express/), and synthesized the corresponding oligonucleotides. Oligonucleotides used for DGL-α has been previously described (Jung et al., 2007). The oligonucleotides for DGL-β, which was selected after screening six independent shRNA sequences as described below, was 5’-CACCGCAGTACAGGGATTTCATTCACGAATGAATGAAATCCCTGTACTGC-3’ (top) and 5’-AAAAGCAGTACAGGGATTTCATTCATTCGTGAATGAAATCCCTGTACTGC-3’ (bottom). Sense and antisense oligonucleotides were annealed and ligated into pENTR entry vector (Invitrogen) to generate U6 promoter-shRNA-Pol III terminator cassette. RNAi silencing was tested from six independent constructs by quantitative PCR and western blotting. The selected shRNA expression cassette was amplified by PCR using M13F and M13R primers containing a MluI overhang at both 5’ and 3’ ends. The PCR product was ligated into an adeno-associated virus vector (pAAV-hrGFP; Stratagene) using MluI sites.

**Cell cultures and western blot analyses**

We transfected Neuro-2a cells (American Type Culture Collection, Manassas, VA) using Superfect reagent (Qiagen, Valencia, CA) as recommended by the manufacturer, and incubated them until harvest at 37°C with 5% CO₂ in Dulbecco’s Modified Eagle’s Medium (DMEM) (Gibco-Invitrogen, Carlsbad, CA) supplemented with 10% fetal bovine serum (FBS) and antibiotics (Gibco-Invitrogen). Stable DGL expressing cell lines were generated by transfecting Neuro-2a cells in a 100-mm dish with 10 µg of control pEF6, DGL-α-V5-pEF6 or DGL-β-V5-pEF6. Individual blasticidin-resistant colonies were isolated 14 days after transfection and screened for DGL expression by western blotting using a monoclonal anti-V5 antibody
Stable cell lines were maintained in DMEM with 10% FBS and penicillin/streptomycin in the presence of 10 µg/ml blasticidin (Invitrogen). For protein analyses, we prepared lysates in a buffer containing Tris-HCl (10 mM, pH 7.4), NaCl (150 mM), Triton X-100 (1%), Nonidet P-40 (0.25%), EDTA (2 mM) supplemented with a mixture of protease inhibitors (Roche). Lysates were centrifuged at 14,000 x g for 10 min and protein concentrations from the supernatants were measured using BCA protein assay (Pierce, Rockford, IL). Proteins (20 µg) were separated by 4-20% sodium dodecylsulfate-polyacrylamide gel electrophoresis (SDS-PAGE), transferred to PVDF membranes, and subjected to western blotting. We used monoclonal anti-V5 (1:5,000, Invitrogen), polyclonal anti-DGL-β (1:3,000) (Jung et al., 2005) or monoclonal anti-Actin (1:10,000, Calbiochem, San Diego, CA) as primary antibodies.

Lipid analyses
Lipid analyses were conducted by liquid chromatography-mass spectrometry (LC-MS), as previously described (Jung et al., 2007; Astarita et al., 2008; Astarita et al., 2009). Briefly, cells were rinsed with ice-cold phosphate-buffered saline and scraped into 1 ml of methanol/water (1:1, vol:vol). Protein concentrations were measured with a BCA protein assay kit (Pierce). Lipids were extracted into a chloroform/methanol mixture (2:1, vol:vol; 1.5 ml) containing (1,3)-heptadecanoyl-sn-glycerol (HDG) and heptadecanoic acid (HDA) as internal standards (100 pmol/sample). Organic phases were collected, dried under N₂ and dissolved in methanol/chloroform (3:1, vol:vol) for analyses.

**Monoacylglycerols (MAG)** – We used an Agilent 1100-LC system coupled to a 1946D-MS detector equipped with an electrospray ionization (ESI) interface (Agilent Technologies, Inc., Palo Alto, CA). MAGs were separated on a reversed-phase XDB Eclipse C18 column (50x4.6 mm i.d., 1.8 µm, Zorbax, Agilent Technologies). They were eluted with a gradient of methanol in water (from 85% to 90% methanol in 2.0 min and 90% to 100% in 3.0 min) at a flow rate of 1.5
ml/min. Column temperature was kept at 40°C. MS detection was in the positive ionization mode, capillary voltage was set at 3 kV and fragmentor voltage was 120 V. N₂ was used as drying gas at a flow rate of 13 l/min and a temperature of 350°C. Nebulizer pressure was set at 60 PSI. Commercial MAGs were used as reference standards. For quantification purposes, we monitored the Na⁺ adducts of the molecular ions [M+Na]⁺ in the selected ion-monitoring (SIM) mode, using HDG (mass-to-charge ratio, m/z = 367) as an internal standard. Non-esterified fatty acids (FA) – We used a reversed-phase XDB Eclipse C18 column (50x4.6 mm i.d., 1.8 μm, Zorbax, Agilent Technologies) eluted with a linear gradient from 90% to 100% of A in B for 2.5 min at a flow rate of 1.5 ml/min with column temperature at 40°C. Mobile phase A consisted of methanol containing 0.25% acetic acid and 5 mM ammonium acetate; mobile phase B consisted of water containing 0.25% acetic acid and 5 mM ammonium acetate. ESI was in the negative mode, capillary voltage was set at 4 kV and fragmentor voltage was 100 V. N₂ was used as drying gas at a flow rate of 13 l/min and a temperature of 350°C. Nebulizer pressure was set at 60 PSI. We used commercially available fatty acids as reference standards, monitoring deprotonated molecular ions [M-H]⁻ in the SIM mode. HDA (m/z = 269) was used as an internal standard.

Fluorescence immunostaining and neurite outgrowth assay
We cultured Neuro-2a cells on Labtek chamber slides (Nunc, Roskilde, Denmark), transfected, fixed and immunostained them as described (Jung et al., 2005). Monoclonal anti-V5 (1:2000, Invitrogen), polyclonal anti-V5 (1:500, Covance, Berkeley, CA) and monoclonal anti-β-Tubulin (1:1000, Sigma-Aldrich, St. Louis, MO) were used as primary antibodies. Alexa 488- or Alexa 546-labeled secondary antibodies (1:1000, Molecular Probes, Eugene, OR) were used for detection. BODIPY 493/503 (1:1000, Molecular Probes) was used to stain lipid droplets in Neuro-2a cells treated with 400 μM oleate for 18 hours, and DAPI-containing media (Vector
Laboratories, Burlingame, CA) was used for nucleus staining and mounting. We captured the images using an Eclipse E600 fluorescence microscope (Nikon, Japan) equipped with a digital camera (Diagnostic Instruments, Sterling Heights, MI). For neurite outgrowth assay, images from random fields of stained cells were manually analyzed in a blind fashion. We considered neurite-bearing cells for those displaying more than one process which is at least twice the length of the cell body. For each culture condition, we scored three regions from the slide containing more than 100 cells. Neurite lengths were measured only from the neurite-bearing cells.

**mRNA quantification**

We measured mRNA levels using a quantitative real-time PCR method. We extracted RNA from cultured cells using a TRIzol (Invitrogen)/RNeasy (Qiagen) hybrid method and synthesized first-strand complementary DNA from 2 µg of total RNA. Reverse transcription was carried out using Superscript II RNase H reverse transcriptase (Invitrogen) and oligo(dT)12-18 primers, for 50 min at 42°C. Quantitative PCR was conducted using Mx3000P system (Stratagene). DGL mRNA levels were normalized using glyceraldehyde-3-phosphate dehydrogenase (GAPDH) as an internal standard. The primer/probe sets were as follows: for mouse DGL-α (GI: 54312093), forward, 5′-CCAGGCCTTTGGGCG-3′; reverse, 5′-GCCTACCACAATCAGGCCAT-3′; TaqMan probe, 5′-ACCTGGGCCGTGGAACCAAACA-3′ and for mouse CB1 receptor, forward, 5′-CAACGACGCAATAACACAA-3′; reverse, 5′-ACAGTGCTCTTGATGCAGCTTT-3′; TaqMan probe, 5′-CCAGCATGCACAGGGCCGC-3′ (TIB Molbiol, Adelphia, NJ). We used Taqman gene expression assays for mouse DGL-β (Mm00523381_m1), MGL (Mm00449274_m1) and ABHD6 (Mm00481199_m1) (Applied Biosystems, Foster City, CA).

**In vitro DGL activity assay**
We harvested cells in 50 mM Tris-HCl (pH 7.0, 1 ml/dish) and homogenized them by passing through a 23-gauge needle for 20 times. The homogenates were centrifuged at 800 x g for 5 min at 4°C, and the resulting supernatants were used for the assays. DGL activity was measured at 37°C for 30 min in 50 mM Tris-HCl, pH 7.0, containing 0.1% Triton X-100, 0.1 mg of cellular protein and substrate diheptadecanoylglycerol (50 µM). Reactions were stopped by adding chloroform-methanol (1:1, vol:vol) containing [²H₈]-2-AG (100 pmol/sample). Lipids were extracted and monoheptadecanoylglycerol was quantified as [M+Na]⁺ (m/z = 367) by LC-MS using [²H₈]2-AG (m/z = 409) as an internal standard.

**Statistical analyses**

Results are expressed as means ± SEM. Analyses were conducted with GraphPad Prism v.4.0 (GraphPad Software, San Diego, CA) and differences were considered significant if \( P < 0.05 \) by two-tailed Student's \( t \)-test.
Results

RA-induced differentiation of Neuro-2a cells is associated with temporally distinct changes in DGL-α and DGL-β expression. Treatment of Neuro-2a cells with RA (20 μM) resulted in extensive formation and elongation of neurite-like processes (Fig. 1A and 1B). Carbachol, a pan-agonist for muscarinic acetylcholine receptors, modestly but significantly increased neurite outgrowth after 48 hours of treatment, whereas DHPG (100 μM), a group I mGlu receptor agonist, had no such effect (Fig. 1B). The RA-induced neurite outgrowth was partially, but significantly attenuated by pre-treatment with the selective CB₁ receptor antagonist rimonabant (1 μM, 24 hours) (Fig. 1C), without affecting cell viability (Supplementary Fig. S1), suggesting that an endocannabinoid signal might be involved in RA-induced neurite outgrowth. To test this possibility, we quantified two main endocannabinoids, 2-AG and anandamide, in cell cultures treated with RA. Cellular levels of 2-AG, but not anandamide, were significantly elevated in RA-treated cells, compared to vehicle-treated controls (Fig. 1D). Consistent with this result, we found that RA increased cellular DGL activity, measured in cell homogenates (Fig. 1E).

We further examined whether RA affects the expression of DGL-α and DGL-β. We found that RA caused a time-dependent increase in DGL-α mRNA, which was statistically significant starting from day 1 of treatment (Fig. 2A). We also observed an increase in DGL-β mRNA, which was quantitatively smaller and temporally delayed relative to DGL-α (Fig. 2B). Additionally, significant increases in CB₁ receptor and ABHD6 mRNA were observed after 3 days of RA treatments, while MGL mRNA was not detectable in Neuro-2a cells (Fig. 2C). Similar changes in mRNA levels, which were accompanied by morphological differentiations, were also observed in SH-SY5Y human neuroblastoma cells treated with RA for 7 days; DGL-α (control, 2.25 ± 0.15; RA, 6.65 ± 0.67, P=0.003),
DGL-β (control, 1.33 ± 0.10; RA, 4.497 ± 0.93, \( P=0.027 \)) and CB₁ receptor (control, 0.014 ± 0.002; RA, 0.91 ± 0.07, \( P<0.001 \)) (mRNA ratio to GAPDH x 1,000, n=3 each).

Next, we used RNA interference to test whether DGL expression is required for RA-induced neuronal differentiation. The efficiency and selectivity of shRNA-induced knockdown of DGL isoforms were confirmed by western blot, immunostaining and quantitative PCR methods (Jung et al., 2007; data not shown). We found that cells transfected with either DGL-α- or DGL-β-silencing shRNA, compared to control non-targeting shRNA, displayed significantly less neurite outgrowth after 48 hours of RA treatment (Fig. 2D). Co-transfection with DGL-α- and DGL-β-silencing shRNAs resulted in a marked inhibition of neurite outgrowth, which was significantly greater than that produced by DGL-β-shRNA alone.

We further asked whether 2-AG is a mediator for neurite outgrowth signal. We observed that treatment with a stable analogue of 2-AG, noladin ether (10 \( \mu \)M, 48 hours), resulted in increased neuritogenesis of Neuro-2a cells (Fig. 2E). In addition, inhibition of 2-AG hydrolysis using the ABHD6 inhibitor WWL70 (10 \( \mu \)M) caused a modest but significant increase in neurite outgrowth. Consistent with the lack of MGL expression in Neuro-2a cells (see above), the MGL inhibitor JZL184 (1 \( \mu \)M) had no effect on neuritogenesis (Fig. 2E).

**DGL-β localizes to perinuclear lipid droplets** The temporal difference between RA-induced expression of DGL-α and DGL-β suggests a segregation in cellular function of these two DGL isoforms. We and others have previously reported that DGL-α localizes to the plasma membrane of neuronal cells (Jung et al., 2007; Katona et al., 2006). To determine the subcellular localization of DGL-β, we constructed and expressed a C-terminal GFP-fused DGL-β in Neuro-2a cells. DGL-β-GFP fluorescence was primarily concentrated in an intracellular peri-nuclear compartment (Fig. 3A), unlike DGL-α-GFP.
or control GFP (Jung et al., 2007). Similar results were obtained when the proteins were expressed in HEK-293 cells (Fig. 3B). In double-fluorescence staining experiments, DGL-β-GFP signal did not overlap with that of either nuclear or endoplasmic reticulum (ER) marker proteins (data not shown). Instead, DGL-β immunofluorescence (Fig. 3C, Anti-V5), but not DGL-α immunofluorescence, overlapped with lipid droplet staining (Fig. 2C, BODIPY) in transiently transfected Neuro-2a cells. Alterations in the size and distribution of lipid droplets were also noticeable in DGL-β-overexpressing cells (Fig. 3C). In cells differentiated with RA, DGL-α-GFP and DGL-β-GFP were localized to neurite-like structures, in addition to the plasma membrane and perinuclear lipid droplets distributions, respectively (Fig. 3D). In neurites, expression of DGL-α-GFP was highly concentrated in large clusters at membranes whereas partially punctuated signals from DGL-β-GFP were dispersed throughout the neurites (Fig. 3D).

Overexpression of DGL-α or DGL-β promotes constitutive neurite outgrowth through CB and non-CB mechanisms Next, we investigated the effect of DGL overexpression on cellular 2-AG biosynthesis and neurite outgrowth. Transient overexpression of DGL-β-V5 (Supplementary Fig. S2A) caused a significant increase in cellular 2-AG levels, compared to control, vector-transfected cells (Supplementary Fig. S2B) and caused lipidomic changes that are comparable to those observed after DGL-α overexpression (Supplementary Table 1) (Jung et al., 2007). We found that transient overexpression of either DGL-α or DGL-β results in significant increases of neurite-bearing cells (Fig. 4A and Supplementary Fig. S2C). It is notable that DGL-β expression caused neurite formation in a significantly higher number of cells compared to DGL-α expression (Fig. 4A). We further established Neuro-2a cell lines that stably express DGL-α-V5 (α18) or DGL-β-V5 (β14), as described in Materials and Methods.
Microscopic observations revealed constitutive morphological changes in β14 cells, which included extended neurite outgrowth and filopodia-like protrusions (Fig. 4B: top, light microscopic images; bottom, β-tubulin-immunofluorescence). α18 cells also displayed morphological changes, albeit to a lesser extent than did β14 cells (Fig. 4B). The number of neurite-bearing cells was significantly higher in both α18 and β14 cells, compared with stable vector-transfected P7 cells (Fig. 4C). The mean length of neurite was significantly higher in β14 cells, compared to P7 cells, but not in α18 cells (Fig. 4D). In contrast, average number of neurites from individual neurite-bearing cells showed no differences (data not shown). Finally, we tested whether neurite outgrowth triggered by DGL overexpression is mediated by CB1 receptors. The α18 and β14 cells, along with control P7 cells, were treated with rimonabant (1 μM) for 24 hours. Strikingly, whereas DGL-α-induced neurite outgrowth was almost completely inhibited by rimonabant, CB1 blockade did not affect DGL-β-induced neurite outgrowth (Fig. 5). The results indicate that the neuritogenic effect of DGL-α expression is mediated through CB1 receptor activation, presumably via 2-AG, whereas DGL-β expression operates through a CB1-independent pathway.
Discussion
In the present study, we investigated the role of DGL isoforms, DGL-α and DGL-β, in the control of neurite outgrowth of Neuro-2a cells. Our results indicate that both DGL-α and DGL-β contribute to RA-induced neuritogenesis, but through two distinct mechanisms: DGL-α by initiating 2-AG-mediated endocannabinoid signaling at plasma membranes, and DGL-β by engaging an as-yet-uncharacterized molecular pathway possibly through intracellular lipid droplets.

We found that RA elevates cellular 2-AG levels and DGL activity in Neuro-2a cells. These increases were due, at least in part, to a transcriptional up-regulation of DGL-α and DGL-β, which occurred, however, within distinguishable time frames, DGL-α expression being more rapid and pronounced than DGL-β’s. These molecular events were associated with increased neuronal differentiation, as previously observed in neuroblastoma cells in vitro, in response to RA treatment (Clagett-Dame et al., 2006; Dehmelt et al., 2003; Wu et al., 1995). On the other hand, agonists for either mGlu5 receptor or muscarinic acetylcholine receptors, which are both known to increase 2-AG levels in the adult brain, caused little or no neurite outgrowth compared to RA. This result could be due to differences between developing and mature neurons. This idea corroborated by the fact that the localization of DGL-α changes dramatically during development (Berghuis et al., 2007; Watson et al., 2008; Bisogno et al., 2003). We found that, importantly, RNAi-induced silencing of either DGL-α or DGL-β attenuates neurite outgrowth, suggesting that both DGL isoforms may contribute to RA-induced differentiation. In addition, knock-down of both DGL-α and DGL-β resulted in an additive inhibitory effect on neuritogenesis. A significant difference was found between silencing DGL-α alone or in combination with DGL-β. This suggests that DGL-α may exert its effects, at least in part independently of DGL-β.
Our immunostaining studies of undifferentiated Neuro-2a cells suggest that DGL-β is localized to lipid droplets, whereas DGL-α is mainly found at plasma membranes (Katona et al., 2006; Jung et al., 2007). DGL-α has been previously identified at the plasma membrane of glutamatergic synapses and its contribution to metabotropic glutamate (mGlu) receptor-dependent plasticity has been extensively studied (Maejima et al., 2001; Varma et al., 2001; Katona et al., 2006; Jung et al., 2007). Electrophysiological or pharmacological stimulation of mGlu5 receptors produces 2-AG, mainly from the sequential hydrolysis of phosphatidylinositol-4,5-bisphosphate by PLC-β and DGL-α (Stella et al., 1997; Jung et al., 2005), which then acts on presynaptic CB₁ receptors to regulate neurotransmitter release. Inhibition of phosphatidylinositol-specific PLC activities abrogated the receptor-dependent upregulation of 2-AG biosynthesis, indicating that there is a preference for utilizing phosphoinositides, among various species of membrane phospholipids, as the precursor of 2-AG. This might be, at least in part, due to the assembly of the receptor with signal transducing enzymes, PLC-β and DGL-α, in a macromolecular protein complex which is mediated by direct interactions with scaffolding proteins such as Homer and Shank (Hwang et al., 2005; Katona et al., 2006; Jung et al., 2007). Recent work with genetically modified mice has confirmed such a role of DGL-α in synaptic regulations (Tanimura et al., 2010; Gao et al., 2010). By contrast, the functions of DGL-β are still largely unknown. The localization of this DGL isoform in lipid droplets, which are emerging as important intracellular organelles involved in lipid homeostasis (Digel et al., 2010), suggests that DGL-β may be involved in regulating membrane dynamics during neuritogenesis. This hypothesis deserves further investigation.

We found that, in Neuro-2a cells differentiated by RA, both DGL-α and DGL-β localize to neurite structures in addition to their main site of distribution sites in non-differentiated
cells. This is in accordance with a previous report indicating that most, if not all, developing axonal tracks co-express both DGL isoforms (Bisogno et al., 2003). Moreover, our results indicate that the distributions of DGL isoforms in neurites are distinguishable. Fluorescence signals from DGL-α-GFP were highly concentrated in large clusters at membranes, whereas DGL-β-GFP were dispersed throughout the neurites displaying partially punctuated signals. Distinctive axonal distributions of the two DGL isoforms had been found in embryonic pyramidal cells, where DGL-α is concentrated in axonal varicosities, whereas DGL-β is distributed uniformly along axons. Additionally, in axonal growth cones, DGL-α is targeted to filopodia, whereas DGL-β is concentrated in the axon stem with a clear demarcation from the growth cones (Mulder et al., 2008). The distributions of both DGL isoforms in elongating neurites, in a distinguishable manner, support the idea that both DGL-α and DGL-β contribute to RA-induced neuritogenesis through distinct mechanisms.

Transient or stable overexpression of either DGL-α or DGL-β induces constitutive morphological changes in Neuro-2a cells. We found that DGL-β expression is more effective than DGL-α at inducing neurite outgrowth, which was even more striking after normalization with efficiency of DGL protein expression (data not shown). Importantly, DGL-β expression promotes neurite elongation, whereas DGL-α does not. The results indicate that functional expression of either DGL isoform, which is expected to increase 2-AG levels in two different subcellular locations, is sufficient to initiate neurite outgrowth. On the other hand, our targeted lipidomic analyses revealed that DGL-α and DGL-β, when overexpressed in Neuro-2a cells, cause similar lipidomic changes in cells (Jung et al., 2007). DGL-β expression increased levels of 2-AG and other unsaturated fatty acid-containing MAGs as well as unsaturated free fatty acids that result from increased 2-acylglycerol hydrolysis. Consistent with these findings, it has been reported
that two DGL isoforms display similar biochemical properties in vitro (Bisogno et al., 2003). In addition, a bioinformatics comparison between the two DGL isoforms indicates that DGL-β has extensive structural and sequence homology with DGL-α, e.g. four transmembrane-spanning domains, a lipase-3 motif and a serine lipase motif. But DGL-β lacks a large C-terminal fragment found in DGL-α. The C-terminal “tail” domain, which is followed by the catalytic motif in DGL-α (Bisogno et al., 2003), contains a few hypothetical post-translational modification sites, as well as the Homer binding motif, which is responsible for its association with mGlu receptor-containing multiprotein complex at plasma membranes (Jung et al., 2007). These results indicate that the differential control of expression, subcellular positioning and structural divergence between DGL-α and DGL-β, rather than differences in enzymatic characteristics, lead to the observed distinctions in physiological function. The CB₁ antagonist rimonabant significantly inhibited DGL-α-induced neurite outgrowth, but failed to prevent DGL-β-induced neuritogenesis. An economical interpretation of these results is that DGL-β expression targets an unknown intracellular pathway that is independent from 2-AG signaling at the CB₁ receptors.

Although our results support the role of DGL in axonal growth and guidance, there are, at least apparently, conflicting observations during differentiation of neural stem cell line Cor-1 (Walker et al., 2010). These discrepancies suggest that 2-AG signaling might be linked to neural differentiation in a cell- and developmental stage-dependent manner.

In conclusion, our study indicates that RA induces the functional expression of DGL, which is required for neurite outgrowth of Neuro-2a cells, and that DGL-α and DGL-β may engage differential cellular mechanisms to regulate neuronal differentiation.
Acknowledgements

The contributions of the Agilent Technologies/University of California, Irvine Analytical Discovery Facility, Center for Drug Discovery is gratefully acknowledged.

Authorship Contributions

Participated in research design: Jung, Piomelli
Conducted experiments: Jung, Astarita, Thongkham
Performed data analysis: Jung, Astarita, Thongkham
Wrote the manuscript: Jung, Astarita, Piomelli
References

Aguado T., Monory K., Palazuelos J., Stella N., Cravatt B., Lutz B., Marsicano G.,

Aguado T., Palazuelos J., Monory K., Stella N., Cravatt B., Lutz B., Marsicano G.,
promotes astroglial differentiation by acting on neural progenitor cells. *J. Neurosci.*
**26**, 1551-1561.


2767.

Endocannabinoids regulate interneuron migration and morphogenesis by

K., Marsicano G., Matteoli M., Canty A., Irving A.J., Katona I., Yanagawa Y., Rakic

Bisogno T., Howell F., Williams G., Minassi A., Cascio M.G., Ligresti A., Matias I.,
Schiano-Moriello A., Paul P., Williams E.J., Gangadharan U., Hobbs C., Di Marzo V.
and Doherty P. (2003) Cloning of the first sn1-DAG lipases points to the spatial and
temporal regulation of endocannabinoid signaling in the brain. *J. Cell Biol.* **163**, 463-
468.


Kano M., Ohno-Shosaku T., Hashimoto Y., Uchigashima M. and Watanabe M.


Stella N., Schweitzer P. and Piomelli D. (1997) A second endogenous cannabinoid that


suggests close proximity between production site of an endocannabinoid, 2-arachidonoyl-glycerol, and presynaptic cannabinoid CB1 receptor. J. Neurosci. 26, 4740-51.
Footnotes

This work was supported by grants from the National Institute on Drug Abuse [ARRA R01 DA-012447].
Legends for Figures

Figure 1. RA treatment induces differentiation of Neuro-2a cells, which is associated with increases in cellular 2-AG. Neuro-2a cells were incubated with 20 μM all-trans-retinoic acid (RA) or vehicle (Veh). Morphological differentiation of the cells was noticeable starting from 24 hours of RA treatment (A). Neuro-2a cells were incubated with RA (20 μM), DHPG (100 μM) or carbachol (10 μM) for 24 or 48 hours at 37°C. Cells were fixed, immunostained using anti-β-Tubulin, and subjected to neurite outgrowth assays. Cellular processes which are at least two-fold longer than the corresponding cell body was regarded as neurites (n=6) (B). Neuro-2a cells were pre-treated with either vehicle or rimonabant (1 μM, Rim) for 10 min and then added with 20 μM RA for 24 hours (n=3) (C). Levels of cellular 2-AG and anandamide (D) and in vitro DGL activity (E) were measured at 48 hours of RA treatment (n=4). *P< 0.05, **P< 0.01, ***P< 0.001 by two-tailed t-test.

Figure 2. Functional expressions of both DGL-α and DGL-β are required for RA induced differentiation of Neuro-2a cells. Neuro-2a cells were incubated with 20 μM RA for the indicated times and DGL-α (A) or DGL-β (B) mRNA levels were measured by quantitative real-time PCR. mRNA levels for CB₁ receptor (CB₁R), MGL, ABHD6 and an internal control 18S were quantified after 3 days of RA treatment (n=4) (C). Neuro-2a cells were transfected with the following shRNA-expressing constructs; the non-targeting control (LacZi), DGL-α-targeting (DGL-αi), DGL-β-targeting (DGL-βi) or both DGL-α-targeting and DGL-β-targeting (DGL-αi+DGL-βi). After 24 hours, cells were treated with 20 μM RA for 40 hours at 37°C (n=6) (D). Neuritogenic effects of 2-AG ether (Noladin ether, 10 μM), a MGL inhibitor JZL184 (JZL, 1 μM) and a ABHD6 inhibitor WWL70
(WWL, 10 μM) were determined upon 48 hours of treatments (n=6) (E). *P< 0.05, **P< 0.01, ***P< 0.001 by two-tailed *t*-test.

Figure 3. Recombinant DGL-β-GFP protein localizes to lipid droplets in transfected cells. 48 hours after transfection of control pEGFP or DGL-β-pEGFP vector in Neuro-2a cells (A) or HEK293 cells (B), cells were fixed and the nuclei were stained with DAPI (blue). Representative images under a fluorescence microscope are shown. V5-fused DGL proteins were expressed in Neuro-2a cells and visualized with anti-V5 antibody using Alexa 546 (red, Anti-V5) fluorescence. Co-stained in the cells were lipid droplets, using a fluorescent dye (green, BODIPY) which binds to neutral lipid. Arrows and arrowheads indicate lipid droplets and its co-localization with DGL-β, respectively (C). After 24 hours from transfection with control pEGFP, DGL-α-pEGFP or DGL-β-pEGFP vector, Neuro-2a cells were treated with 20 μM RA for 40 hours at 37°C. Cells were fixed and stained with anti-β-Tubulin antibody using Alexa 546 (red, Anti-Tubulin) fluorescence and DAPI (blue). Arrows indicate neurite localization of DGL-α-GFP or DGL-β-GFP, whereas arrowheads indicate plasma membrane or lipid droplets distribution of DGL-α or DGL-β, respectively (D).

Figure 4. Overexpression of DGL isoforms induces morphological changes in Neuro-2a cells. (A) Neuro-2a cells were transfected with control pEF6 (Vector), DGL-α-V5-pEF6 (DGL-α) or DGL-β-V5-pEF6 (DGL-β). After 72 hours of transfection, cells were fixed and double-immunostained with a rabbit polyclonal anti-V5 (for DGL staining) and a mouse monoclonal anti-β-Tubulin (for β-Tubulin staining) antibodies. For DGL-α or DGL-β, V5-positive cells were selected and the number of cells bearing neurite-like cellular processes were counted. (B) Representative images under a light microscope (top) or a
fluorescence microscope (bottom) of stable transfectants of pEF6 vector (P7), DGL-β-V5 (β14) or DGL-α-V5 (α18). Cells were immunostained using anti-β-Tubulin antibody (green fluorescence) and the nuclei were stained with DAPI (blue). Percentage of cells containing neurite processes (C) and mean lengths of the neurites (D) were measured from the P7, α18 or β14 stable cell lines. *P<0.05, **P<0.01, ***P<0.001 by two-tailed t-test.

Figure 5. Antagonism of CB₁ receptor selectively blocks neurite outgrowth induced by DGL-α expression. The stable DGL-expressing Neuro-2a cells α18 and β14, along with the control P7 cells, were treated for 24 hours with 1 μM rimonabant (Rim), a CB₁ receptor antagonist, in serum-free media. Cells were then fixed, immunostained using anti-β-Tubulin, and subjected to neurite outgrowth assays. ***P<0.001 by two-tailed t-test.
Fig. 1
Fig. 2
Fig. 3
Fig. 4
Fig. 5