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LPA as a Paracrine Mediator of Adipocyte Growth and Function

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Abstract

Adipogenesis corresponds to the recruitment of new adipocytes in adipose tissue and results from proliferation/differentiation of preadipocytes. Production of paracrine and autocrine factors by adipocytes play an important role in adipogenesis. We recently demonstrated the existence of an adipocyte production of lysophosphatidic acid (LPA) both *in vitro* and *in situ*. This production is modulated by catecholamines via α 2-adrenergic receptors. Adipocyte-LPA present in conditioned media increases the growth of a preadipose cell line in culture. This is associated with an activation of Mitogen Activated Protein Kinases, and of the Focal Adhesion Kinase. Because of the close proximity of preadipocytes and adipocytes within adipose tissue, adipocyte-LPA could play an important role in autocrine/paracrine control of adipogenesis.

Introduction

In obesity, adipose tissue overgrowth is associated with adipocyte hypertrophy and adipocyte hyperplasia. Hypertrophy results from excessive triglyceride accumulation in adipocytes. Hyperplasia results from recruitment of new adipocytes involving proliferation and differentiation of preadipocytes: adipogenesis. Identifying factors regulating adipogenesis, is a major point in understanding normal and pathologic growth of adipose tissue. Many hormones, growth factors and cytokines, such as insulin, catecholamines, IGF-I, growth hormone, glucocorticoids and thyroid hormone, are known to promote proliferation and/or differentiation of preadipocytes (1-3). Production of paracrine and autocrine factors by adipocytes themselves could also play an important role. Adipocytes produce several peptidic (TNF α , adipin, angiotensinogen or leptin) and lipidic factors (fatty acids, prostaglandins, monobutyrin) all involved directly or not in adipogenesis (2, 4-6).

Among the different factors controlling adipose tissue growth, catecholamines (epinephrine and norepinephrine) play an important role. These mediators act through different receptor subtypes: β 1, β 2, β 3, α 1 and α 2 coupled to several intracellular transduction pathways (7). Our group is interested in understanding the contribution of the different

adrenergic receptors in the control of adipogenesis. We previously demonstrated the involvement of catecholamines in the control of preadipocyte proliferation, a key event in adipogenesis. This trophic activity involves the specific activation of α 2-adrenergic receptors and the activation of Mitogen Activated Protein Kinases, Focal Adhesion Kinase and the small GTPase p21rhoA (8-10). Recent studies suggest that in addition to those intracellular transduction pathways, an autocrine/paracrine loop involving lysophosphatidic acid (LPA) could also play an important role in α 2-adrenergic control of adipogenesis (11).

Adipocyte production of lysophosphatidic acid

Experiments which allowed to propose the existence of an α 2-adrenergic-dependent production of LPA by adipocytes were based on utilization of conditioned media (CM). Adipocytes were isolated from human adipose tissue or obtained in culture after differentiation of the murine adipose cell line α 2AF2 (8). Both type of adipocyte express abundant functional α 2-adrenergic receptors. CM were prepared from adipocytes exposed or not to α 2-adrenergic agonists, and tested for their ability to exert trophic activities on α 2-adrenergic insensitive preadipocytes (3T3F442A cell line). Control CM (CCM) induced a rapid (within minutes) reorganization of preadipocyte actin cytoskeleton (cell spreading) followed (after 48 h) by an increased in preadipocyte number. Those effects were significantly increased with CM prepared from α 2-adrenergic-stimulated adipocytes (α 2CM). This additional effect was completely suppressed by pretreatment of α 2CM with a lysophospholipase (phospholipase B: PLB). This suggests the existence of an α 2-adrenergic-dependent production of a trophic lysophospholipid by adipocytes. Among different lysophospholipids tested, only 1-oleoyl-LPA and sphingosine-1-phosphate (S1P) were able to reproduce the effects of α 2CM. S1P being insensitive to phospholipase B, we proposed that LPA was responsible for the trophic activities of α 2CM. This was supported by presence of [32 P]LPA in α 2CM prepared from [32 P]Pi prelabelled adipocytes. In order to estimate the amount of LPA produced by adipocytes, phospholipids present in α 2CM were chemically derivatized with a fluorescent molecule (1-pyrenyl diazomethane), separated by thin layer chromatography, and fluorescence was quantified. Concentration of LPA in α 2CM was estimated to 280-360 nM (0.7-0.9 nmol/ 10^6 cells). This amount of LPA is close to that produced by platelets (12) and ovarian cancer cells (13).

LPA is relatively abundant in plasma (bound to albumin) produced by aggregating platelets (12, 14). It is also found in other biological fluids such ascites (15), cerebro spinal liquid (16), aqueous humor (17). Eventhough LPA can originate from various metabolic pathways (12), several groups have proposed that its production mainly results from phospholipase A2 (PLA2--catalysed deacylation of phosphatidic acid (13, 19, 20). In order to determine the existence of adipocyte production of LPA *in situ*, a microdialysis probe was implanted in human subcutaneous adipose tissue and perfused with or without an α 2-

adrenergic agonist. The presence of LPA in microdialysates was tested using a bioassay. Microdialysates obtained in the presence of the $\alpha 2$ -adrenergic agonist exhibited stronger LPA-like activity as compared to control microdialysates. This revealed the presence of LPA in extracellular fluid of human adipose tissue.

LPA effects on preadipocyte growth

LPA is known as a potent bioactive phospholipid able to regulate a number of cellular events including proliferation and differentiation. Most of the cellular responses generated by LPA involve specific G-protein coupled cell surface receptors. Four G-protein coupled receptors can potentially transduce LPA signals in mammals: vzg1/edg2, edg4, PSP24 and edg1 (21-25). Those receptors can be coupled to Gi/Go heterotrimeric proteins, leading to activation of the mitogen activated protein kinases (MAPK) or inhibition of adenylyl cyclase. LPA receptors can also be coupled to G12 or G13 heterotrimeric proteins leading to activation of the Focal Adhesion Kinase (FAK) and actin cytoskeleton reorganization via the small GTPase p21rhoA (18). In preadipocytes, 1-oleoyl-LPA (the most active form of LPA) generated both short term and long term effects. Twenty minutes treatment of serum deprived 3T3F442A preadipocytes led to a dose dependent (EC50 of 1-5 nM) reorganization of actin cytoskeleton resulting in cell spreading. Similar response was obtained with an other bioactive phospholipid, sphingosine-1-P (S1P). If 1-oleoyl-LPA treatment was maintained for 48 h it led to a dose-dependent increase in cell number (EC50 of 100-110 nM). It is noticeable that those EC50s are compatible with estimated LPA concentration (280-360 nM, see above) in adipocyte conditioned media. Chronic exposition (twice a day for 48 h) of 3T3F442A preadipocytes to 10 μ M 1-oleoyl-LPA, strongly desensitized the spreading effect of 1-oleoyl-LPA (12 -fold increase in EC50 and 36% reduction of the maximal effect). This desensitization was specific to 1-oleoyl-LPA since in the same conditions S1P spreading effect was not altered. We further analysed the effects of 1-oleoyl-LPA on the activity of intracellular kinases involved in both cell cytoskeleton and cell proliferation. 1-oleoyl-LPA led to a rapid and dose-dependent activation of Mitogen Activated Protein Kinases ERK1 and ERK2, as well as of the Focal Adhesion Kinase p125FAK (Figure 2). Pertussis toxin pretreatment blocked 1-oleoyl-LPA activation of ERK 1 and ERK2, not that of p125FAK. Conversely, the p21rhoA-inhibitor C3-exoenzyme blocked the activation of p125FAK, not that of ERK1 and ERK2. Similarly, 1-oleoyl-LPA-induced preadipocyte spreading was completely blocked by treatment with C3-exoenzyme, not by pertussis toxin. Therefore trophic activity of LPA in preadipocytes involves the activation of two independent but complementary pathways: the ras/MAPK pathway via heterotrimeric G proteins of Gi/Go subtype, and the p21rhoA/FAK/cytoskeleton pathway via pertussis toxin insensitive G proteins likely belonging to G12/G13 subtypes. This strongly support the involvement of G-protein coupled receptors in trophic activity of LPA in preadipocytes.

Conclusion

Our data demonstrate that LPA is produced by adipocytes under α 2-adrenergic stimulation, and exerts trophic activities in preadipocytes. Because of the close coexistence of adipocytes and preadipocytes in intact adipose tissue, adipocyte-LPA could play an important role in paracrine/autocrine control of adipogenesis by catecholamines (Figure 3). We are currently trying to depict the precise mechanisms involved in adipocyte production of LPA (involvement of different phospholipase A2 subtypes) as well as in the trophic activity of LPA in preadipocytes (characterization and specific contribution of Edg-receptor subtypes). Adipose tissue is present in a lot of different anatomical locations in the body in close contact with several other tissues (muscle, heart, mammary glands, colon, ...). Because of the pleiotropic trophic activity of LPA, adipocyte-LPA could also play an important role in normal and/or pathologic growth of other tissues.

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Legends of the figures

Figure 1: Influence of human adipocyte conditioned media on 3T3F442A preadipocyte spreading. Conditioned media (CM) were prepared from isolated human adipocytes incubated for 16h at 37°C in Krebs-Ringer albumin buffer supplemented or not (CCM) with 1 μ M of the selective α 2-adrenergic agonist UK14304 (UK) alone (α 2CM) or in the presence (α 2CM+RX) of 10 μ M of the selective α 2-adrenergic antagonist RX821002 (RX). Spreading activity was measured by determining the proportion of refringent cells after 20 min exposure of retracted 3T3F442A preadipocytes to conditioned medium pretreated (+) or not (-) with 3U/ml phospholipase B (PLB) for 37°C.

Figure 2: LPA activation of the Mitogen Activated Protein Kinases (ERK1-ERK2) and of the Focal Adhesion Kinase (FAK) in 3T3F442A preadipocytes. Serum-deprived 3T3F442A preadipocytes were exposed to 1 μ M 1-oleoyl-LPA for various period of time. Cells proteins were then solubilized, immunoprecipitated with an anti-phosphotyrosine antibody, separated on SDS-PAGE, transferred on nitrocellulose, and blotted with anti-ERK1/ERK2 (A) or anti-FAK (B) antibodies. * corresponds to the biphosphosphorylated (tyrosin and threonin) and active form of ERK1 and ERK2.

Figure 3: Paracrine control of preadipocyte growth by adipocyte LPA.