Liraglutide prevents cognitive decline in a rat model of streptozotocin-induced diabetes independently from its peripheral metabolic effects

Caterina Palleria a, Antonio Leo a, Francesco Andreozzi b, Rita Citraro a, Michelangelo Iannone c, Rosangela Spiga b, Giorgio Sesti b, Andrew Constanti d, Giovambattista De Sarro a, Franco Arturi b, Emilio Russo a,∗

a Science of Health Department, School of Medicine, University “Magna Graecia” of Catanzaro, Italy
b Department of Medical and Surgical Sciences, University Magna Graecia of Catanzaro, 88100, Viale Europa, Catanzaro, Italy
c Institute of Neurological Sciences, Pharmacology Section, Roccetta di Borgia, Catanzaro, Italy
d Department of Pharmacology, UCL School of Pharmacy, 29/39 Brunswick Square, London, UK

HIGHLIGHTS

• Liraglutide might have anxiolytic and pro-depressant effects.
• Liraglutide demonstrated neuroprotective effects and improved learning and memory.
• Liraglutide protected from hippocampal neurodegeneration likely through mTOR pathway.
• Liraglutide reduced hippocampal neuronal death.
• Liraglutide treatment might be a relevant option for cognitive decline.

ABSTRACT

Diabetes has been identified as a risk factor for cognitive dysfunctions. Glucagone like peptide 1 (GLP-1) receptor agonists have neuroprotective effects in preclinical animal models. We evaluated the effects of GLP-1 receptor agonist, liraglutide (LIR), on cognitive decline associated with diabetes. Furthermore, we studied LIR effects against hippocampal neurodegeneration induced by streptozotocin (STZ), a well-validated animal model of diabetes and neurodegeneration associated with cognitive decline. The study evaluated the effects of STZ-induced diabetes in rats, which were treated with or without LIR. LIR treatment significantly improved learning and memory in the Morris water maze test, reduced hippocampal neuronal death, and prevented the increased phosphorylation of AKT and p70S6K. These findings suggest that LIR might have therapeutic potential in the treatment of cognitive decline associated with diabetes.

Abbreviations: AD, Alzheimer’s disease; APP/PS1, amyloid precursor protein/presenilin 1; Aβ, β-amyloid; CNS, central nervous system; CTRL, control; EPM, elevated plus maze; FST, forced swimming test; GLP-1R, glucagon-like peptide-1 receptor; i.c.v., intracerebroventricular; i.p., intraperitoneally; IT, initial latency; IT, immobility time; LIR, liraglutide; mTORC, mammalian target of rapamycin complex 1; MWM, Morris water maze; NOS, nitric oxide synthase; OF, open field arena; p70S6K, ribosomal protein S6 kinase; PD, Parkinson’s disease; s.c., subcutaneously; STL, step-through latency; STZ, streptozotocin; T2DM, type 2 diabetes mellitus; TNF-α, tumor necrosis factor-alpha.

* Corresponding author at: Chair of Pharmacology, Department of Science of Health, School of Medicine, University of Catanzaro, Via T. Campanella, 115, 88100 Catanzaro, Italy.
E-mail address: erusso@unicz.it (E. Russo).

http://dx.doi.org/10.1016/j.bbr.2017.01.004
0166-4328/© 2017 Elsevier B.V. All rights reserved.
1. Introduction

Type 2 diabetes mellitus (T2DM) is a public health problem currently increasing in prevalence [1]. T2DM is a complex endocrine and metabolic disorder also accompanied by a negative impact on the central nervous system (CNS), particularly in the elderly [2], leading to diabetic encephalopathy and concomitant increased incidence of cognitive problems; the latter are particularly associated with atrophy of the hippocampal formation that is known to be also involved in learning and memory processing [2]. Patients with T2DM show volumetric abnormalities in the hippocampus and amygdala, similar to those found in patients with major depression [3]. Furthermore, neurocognitive deficits in working memory, attention and executive function have been reported in both individuals with diabetes and mood disorders [2]. The precise pathophysiology of cognitive dysfunction and neuronal damage in T2DM is not completely understood, but it is likely that impaired glucose control (hyperglycemia, hypoglycemia), vascular damage, and insulin resistance play significant roles [3]. None of the known antidiabetic drugs currently used for T2DM has a proven effect on cognitive decline in patients, even if, there are some data on the efficacy of metformin and other antidiabetic drugs in animal models of diabetes and Alzheimer’s disease (AD) [4–6] however, the glucagon-like peptide-1 receptor (GLP-1R) agonists, a new class of antidiabetic drugs now used therapeutically in diabetic patients, also exert powerful neuroprotective properties in animal models of neurodegenerative diseases, cerebral ischemia and traumatic brain injury [7,8], therefore might be valuable therapeutic tools for prevention of neurological comorbidity in T2DM. GLP-1 is an endogenous incretin (insulinotropic) peptide hormone secreted from the gastrointestinal tract, that plays a key physiological role in glucose homeostasis by enhancing pancreatic insulin secretion and by suppressing glucagon release and hepatic glucose output [9]. GLP-1 receptor (GLP-1R) agonists such as exenatide, liraglutide (LIR) and lixisenatide have been approved for treatment of T2DM [10]; in December 2014, liraglutide 3.0 mg was approved by the Food and Drug Administration and in March 2015 by the European Medicines Agency for obesity treatment. In addition to its peripheral metabolic effects, GLP-1 acts as a growth factor in the brain, inducing neurite growth and protecting against oxidative injury [11]. It is known that GLP-1Rs are expressed by pyramidal neurons in the hippocampus, hypothalamus, neocortex and Purkinje cells in the cerebellum and are involved in cell differentiation and neuroprotection as demonstrated both in vitro and in vivo [12]. GLP-1 and GLP-1 analogs also increase neuronal progenitor proliferation in the brain, as well as long-term potentiation and paired-pulse facilitation in the hippocampus [13,14]; accordingly, GLP-1R knock-out mice have an impairment of memory formation [14].

The neuroprotective effects of GLP-1 have been widely demonstrated in cultured neurons [15]. Human neuroblastoma cell lines over-expressing GLP-1Rs are protected from oxidative stress-induced cell death [16]. Excitotoxic L-glutamate-induced death of cultured rat hippocampal neurons is also decreased by GLP-1 [17]. The dual actions of GLP-1 in pancreatic β-cells and in neurons have recently generated therapeutic interest, considering that T2DM is a risk factor also for AD. Thus, GLP-1 has beneficial effects not only for the treatment of diabetes but also produces significant neuroprotection in animal models of cerebral ischemia and AD [18,19]. Indeed, GLP-1 reduces endogenous levels of β-amyloid (Aβ) in the rodent brain [20] and intracerebroventricular (i.c.v.) administration of GLP-1 enhances associative and spatial learning [13]. Furthermore, most GLP-1 agonists show remarkable neuroprotective effects in different neurodegenerative disorders such as AD, Parkinson’s disease (PD) and stroke by reducing β-amyloid plaques, preventing loss of synapses and memory impairments, and reducing oxidative stress and chronic inflammatory responses in the brain. Recent studies have established that exenatide, a stable GLP-1R agonist, enhances neuronal progenitor proliferation in the brain of diabetic mice [21] and reduces endogenous levels of Aβ in transgenic AD mice [21]. Similarly, LIR has neuroprotective effects in rats [22]. It also has neuroprotective effects in an amyloid precursor protein/presenilin 1 (APP/PS1) mouse model of AD [23], increases neurogenesis, improves cognitive function and reduces amyloid plaque deposition in a mouse model of AD [19,24] and is now being tested in clinical trials for the treatment of AD patients [25]. LIR additionally attenuates the neuronal damage following cerebral ischemia in rats by preventing apoptosis and decreasing oxidative stress [26]. Recently, lixisenatide was tested in the APPswe/PS1DE9 mouse model of AD in comparison with LIR, showing that lixisenatide was equally effective but more potent [27]. Therefore, GLP-1 agonists apparently have a potential role to facilitate neuronal network repair in cortical tissue and consequently could have beneficial effects in patients with neurodegenerative disorders. Based on this background, we tested the effects of LIR on cognitive decline associated with diabetes mellitus in the established streptozotocin (STZ) rat model; furthermore, to exclude any influence of LIR effects on blood glucose levels, we studied the effects of LIR against the hippocampal neurodegeneration induced by i.c.v. injection of STZ, a well-validated model of neurodegeneration associated with cognitive impairment, not accompanied by diabetes [28]. Finally, in order to clarify the neuroprotective mechanism of action of LIR, we studied its effects on the mammalian
دریافت فوری متن کامل مقاله

| امکان دانلود نسخه تمام متن مقالات انگلیسی |
| امکان دانلود نسخه ترجمه شده مقالات |
| پذیرش سفارش ترجمه تخصصی |
| امکان جستجو در آرشیو جامعی از صدها موضوع و هزاران مقاله |
| امکان دانلود رایگان ۲ صفحه اول هر مقاله |
| امکان پرداخت اینترنتی با کلیه کارت های عضو شتاب |
| دانلود فوری مقاله پس از پرداخت آنلاین |
| پشتیبانی کامل خرید با بهره مندی از سیستم هوشمند رهگیری سفارشات |