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(71) **Applicants:** **CIDQO 2012, S.L.** [ES/ES]; Polígono Industrial Can Verdalet, Carred D, Nau 91, E-08490 Tordera - Barcelona (ES). **UNIVERSITAT DE BARCELONA** [ES/ES]; Centro de Patentes UB, Baldiri Reixac, 4, Torre D, E-08028 Barcelona (ES).

(72) **Inventors:** **VÁZQUEZ CRUZ, Santiago**; Avenida Can Corts, 15, 3-2, E-08940 Cornellá - Barcelona (ES). **LEI-VA MARTÍNEZ, Rosana**; Avenida Madrid, 51-53, 1º 5 A, E-08028 Barcelona (ES). **VALVERDE MURILLO, Elena**; Abat Escarré, 12, E-43711 Banyeres del Penedès - Tarragona (ES).

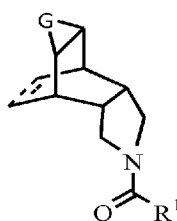
(74) **Agent:** **CARBONELL VALLÈS, Enric**; ABG PATENTES, S.L., Avda. de Burgos, 16D, Edificio EUROMOR, E-28036 Madrid (ES).

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(54) Title: NEW AZA- TETRACYCLO DERIVATIVES



(I)

(57) **Abstract:** The present invention relates to compounds of formula (I); which are capable of inhibiting the 11-beta-hydroxysteroid dehydrogenase type 1 (11- beta-HSD1), to processes for preparing these compounds, to pharmaceutical compositions comprising an effective amount of these compounds, to the use of the compounds in medicine in particular for the treatment of pathological conditions or diseases that can improve by inhibition of 11-beta-HSD1, to the use of said compound for the manufacture of a medicament for the treatment of pathological conditions or diseases that can improve by inhibition of 11-beta-HSD1 and to methods for the treatment or prevention of a disease or conditions susceptible of improvement by inhibition of 11β-hydroxysteroid dehydrogenase type 1 (11β-HD1) in a subject in need thereof comprising the administration to said subject of a therapeutically effective amount of said compounds.



NEW AZA- TETRACYCLO DERIVATIVES

FIELD OF THE INVENTION

5 The invention relates to new aza-tetracyclo derivatives capable of inhibiting the 11-beta-hydroxysteroid dehydrogenase type 1 (11-beta-HSD1).

 Other objectives of the present invention are to provide a procedure for preparing these compounds; pharmaceutical compositions comprising an effective amount of these compounds; the use of the compounds for manufacturing a medicament
10 for the treatment of pathological conditions or diseases that can improve by inhibition of 11-beta-HSD1, such as obesity, insulin resistance, metabolic syndrome, cognitive disfunction associated to Alzheimer's disease, HIV-associated dementia, Creutzfeld-Jakob disease, Parkinson's disease, Huntington's disease, vascular inflammation, atherosclerosis, diabetes type 2, cardiovascular disease, glaucoma, osteoporosis,
15 myocardial infarction and for alcohol abuse treatment.

BACKGROUND OF THE INVENTION

 Glucocorticoids (GCs) are well-known ubiquitous hormones playing a key role in modulating immune and inflammatory responses, regulating energy metabolism and
20 cardiovascular homeostasis and are a key element of stress response. GCs act through the intracellular glucocorticoid receptors and, in some tissues, through mineralocorticoid receptors, both being nuclear transcription factors. The action of GCs on the target tissues depends not only on circulating steroid concentrations and the cellular expression of the receptors, but also on the intracellular enzymes which
25 critically determine up to what point the glucocorticoids will have active access to the receptors. Opposing the action of insulin, GCs promote gluconeogenesis, inhibit β -cell insulin secretion and peripheral glucose uptake. Moreover, GCs contribute to lipolysis, with subsequent fatty acid mobilization, and proteolysis. Thus, overall, GCs prompt catabolic states (e.g., see Dallman, M. F. et al. "Feast and famine: critical role of
30 glucocorticoids with insulin in daily energy flow" in *Front. Neuroendocrinol.* **1993**, *14*, 303-347, and Sapolsky, R. M. et al. "How do glucocorticoids influence stress

responses? Integrating permissive, suppressive, stimulatory, and preparative actions” in *Endocr. Rev.* **2000**, 21, 55-89).

The active human GC cortisol, is synthesized by the adrenal cortex with a strong circadian rhythm. Its secretion is controlled by the hypothalamic-pituitary-adrenal (HPA) axis and induced in stressful conditions. Serum cortisol readily passes cell membranes and exerts its intracellular functions by binding to GC receptors (GRs), which are ligand-activated nuclear receptors. GRs regulate, directly or indirectly, the expression of a plethora of genes involved in various physiological processes (e.g., see Rosen, E. D. “Adipocyte differentiation from the inside out” in *Nat. Rev. Mol. Cell. Bio.* **2006**, 7, 697-706). Furthermore, GCs bind to mineralocorticoid receptors, which regulate blood pressure and sodium balance in the kidney (see Arriza, J. L. et al. “Cloning of human mineralocorticoid receptor complementary DNA: structural and functional kinship with the glucocorticoid receptor” in *Science* **1987**, 237, 268-275).

Along with the HPA axis, GC’s homeostasis is controlled by the 11 β -hydroxysteroid dehydrogenase enzyme. Two isoforms of this enzyme, 11 β -HSD type 1 (11 β -HSD1) and type 2 (11 β -HSD2), catalyse the interconversion between active 11-hydroxy forms (cortisol in humans, and corticosterone in rodents) and inert 11-ketosteroids (cortisone in humans, and 11-dehydrocorticosterone in rodents). The 11 β -HSD1 enzyme, mainly expressed in liver, adipose tissue, bone, pancreas, vasculature and brain, reconverts inactive glucocorticoids into active ones, thus playing an important role in modulating cellular agonist concentration and, therefore, in activating corticosteroid receptors in the target tissues.

METABOLIC DISEASE SYNDROME AND DIABETES

Type 2 diabetes mellitus (T2DM) is characterized by increased cellular insulin resistance with an initial phase of enhanced insulin secretion followed by pancreatic β -cell failure. Regarding obesity, a person with a body mass index of 30 or more is considered obese, entailing health risks. T2DM and obesity are considered as first-world epidemics and are nowadays great healthcare challenges in the developed countries. According to the World Health Organization (WHO), in 2014, 9% of the worldwide adult population had diabetes (see WHO web page. Diabetes Fact Sheet, <http://www.who.int/mediacentre/factsheets/fs312/en/> accessed on 21st March 2016).

That same year, the percentage of total health expenditure in T2DM reached US \$ 612 billion. The expected increase on the diabetic population by 2035 is about 205 million, which will elevate the total health cost still higher (see, European Commission web page.

Major and chronic diseases: Diabetes.

- 5 http://ec.europa.eu/health/major_chronic_diseases/diseases/diabetes/index_en.htm#fragment7 accessed on 21st March 2016). Concerning obesity, in 2014 more than 1.9 billion adults were overweight, and of these, 600 million were obese (WHO web page. Obesity and overweight Fact Sheet. <http://www.who.int/mediacentre/factsheets/fs311/en/> accessed on 21st March 2016).

- 10 From another standpoint, the metabolic syndrome is a cluster of metabolic abnormalities including resistance to insulin, obesity, hyperglycemia, and hypertension that represents a major risk factor for type-2 diabetes and cardiovascular disease. It is diagnosed in a given patient if visceral obesity and two or more risk factors are present, including elevated fasting plasma glucose, elevated triglycerides, elevated blood
15 pressure and reduced high-density lipoprotein (see Huang, P. L. “A comprehensive definition for metabolic syndrome” in *Dis. Model Mech.* **2009**, 2, 231-237”). Among the mechanisms involved in its pathogenesis, glucocorticoid synthesis and metabolism have been proposed to play a key role (e.g., see Wamil, M. et al. “Inhibition of 11 β -hydroxysteroid dehydrogenase type 1 as a promising therapeutic target” in *Drug*
20 *Discover. Today* **2007**, 12, 504-520).

- Currently, the predominant therapeutic strategy for preventing or managing T2DM and metabolic syndrome is the individual treatment of the set of risk factors instead of treating the disorder as a whole. The increase in the prevalence of T2DM and metabolic syndrome over the last few decades in the developed countries has propelled
25 a growing need for the development of novel therapies to treat metabolic syndrome and its associated disorders.

- Evidence has accumulated that enzyme activity of 11 β -hydroxysteroid dehydrogenase type 1 (11 β -HSD1), which regenerates active glucocorticoids from inactive forms and plays a central role in regulating intracellular glucocorticoid
30 concentration, is commonly elevated in fat depots from obese individuals. Notably,

increased tissue 11 β -HSD1 expression and activity have been demonstrated in metabolic tissues including liver and adipose in human obesity and metabolic syndrome.

The impact of tissue-specific GC activation by 11 β -HSD1 in metabolic homeostasis has been examined using several transgenic mouse models. Experiments in mice with adipose tissue-targeted 11 β -HSD1 overexpression developed a phenotype similar to the one of Cushing's syndrome, including insulin resistance, visceral obesity, dyslipidaemia and hypertension, without an increase in the GC plasmatic levels (see, Masuzaki, H. et al. "A transgenic model of visceral obesity and the metabolic syndrome" in *Science* **2001**, 294, 2166–2170, and Masuzaki, H. et al. "Transgenic amplification of glucocorticoid action in adipose tissue causes high blood pressure in mice" in *J. Clin. Invest.* **2003**, 112, 83-90). On the other hand, overexpression of 11 β -HSD1 in liver tissue led to the appearance of metabolic syndrome but without impaired glucose tolerance and obesity (see Paterson, J. M. et al. "Metabolic syndrome without obesity: hepatic overexpression of 11beta-hydroxysteroid dehydrogenase type 1 in transgenic mice" in *Proc. Natl. Acad. Sci USA* **2004**, 101, 7088-7093). Conversely, 11 β -HSD1 knockout (11 β -HSD1 $-/-$) mice resist visceral fat accumulation and insulin resistance even on a high-fat diet and have a 'cardioprotective' phenotype (see Kotelevtsev, Y. et al. "11beta-hydroxysteroid dehydrogenase type 1 knockout mice show attenuated glucocorticoid-inducible responses and resist hyperglycemia on obesity or stress" in *Proc. Natl. Acad. Sci.* **1997**, 94, 14924-14929, Morton, N. M. et al. "Novel adipose tissue-mediated resistance to diet-induced visceral obesity in 11beta-hydroxysteroid dehydrogenase type-1 deficient mice" in *Diabetes* **2004**, 53, 931-938, and Morton, N. M. et al. "Improved lipid and lipoprotein profile, hepatic insulin sensitivity, and glucose tolerance in 11beta-hydroxysteroid dehydrogenase type 1 null mice" in *J. Biol. Chem.* **2001**, 276, 41293-41300), whose effects are also seen with 11 β -HSD1 inhibitors in rodents (see, Wamil, M. and Seckl, J. R. "Inhibition of 11 β -hydroxysteroid dehydrogenase type 1 as a promising therapeutic target" in *Drug Discover. Today* **2007**, 12, 504-520).

Several studies have appeared over the recent years proving the relationship between tissue-specific alterations in 11 β -HSD1 expression in human T2DM, obesity

and metabolic syndrome (see, Morgan, S. A. et al. "11beta-hydroxysteroid dehydrogenase type 1 inhibitors for the treatment of type 2 diabetes" in *Expert Opin. Investig. Drugs* **2010**, *19*, 1067-1076). For instance, the expression of 11 β -HSD1 in human adipose tissue is positively correlated with the degree of obesity (e.g., see
5 Valsamakis, G. et al. "11beta-hydroxysteroid dehydrogenase type 1 activity in lean and obese males with type 2 diabetes mellitus" in *J. Clin. Endocrinol. Metab.* **2004**, *89*, 4755-4761 and Wang, M. "Tissue-specific glucocorticoid excess in the metabolic syndrome: 11 β -HSD1 as a therapeutic target" in *Drug Develop. Res.* **2006**, *67*, 567-569).

10 Hence, modulation of 11 β -HSD enzyme, and in particular inhibition of 11 β -HSD1 in adipose tissue and liver, are of special interest as pharmacotherapeutic strategies in the medical treatment of the metabolic syndrome and associated disorders (see Masuzaki, H. et al. "Tissue-specific glucocorticoid reactivating enzyme, 11 β -hydroxysteroid dehydrogenase type 1 (11 β -HSD1) - a promising drug target for the
15 treatment of metabolic syndrome" in *Curr. Drug Targets Immune Endocr. Metabol. Disord.* **2003**, *3*, 255-262, Morton, N. M. "Obesity and corticosteroids: 11beta-hydroxysteroid type 1 as a cause and therapeutic target in metabolic disease" in *Mol. Cell. Endocrinol.* **2010**, *316*, 154-164, Anderson, A. et al. "11 β -HSD1 inhibitors for the treatment of type 2 diabetes and cardiovascular disease" in *Drugs* **2013**, *73*, 1385-1393,
20 and Wang, M. "Inhibitors of 11 β -Hydroxysteroid Dehydrogenase Type 1 in Antidiabetic Therapy" in *Diabetes – Perspectives in Drug Therapy*, Schwanstecher, M. (ed.), Handbook of Experimental Pharmacology 203, **2011**, 127-146, Springer-Verlag).

Numerous inhibitors of 11 β -HSD enzymes have been identified. Naturally occurring inhibitors include the liquorice derivative glycyrrhetic acid,
25 flavanone/hydroxyl-flavanone, bile acids, and progesterone metabolites; however, most of these compounds inhibit both 11 β -HSD1 and 11 β -HSD2, and as a result their therapeutic benefit is limited due to ensuing hypertension and hypokalemia (see Gathercole, L. L. "11 β -Hydroxysteroid dehydrogenase 1: translational and therapeutic aspects" in *Endocrine Rev.* **2013**, *34*, 525-555). In humans, the first published reports of
30 the metabolic consequences of 11 β -HSD1 inhibition were derived from the nonselective compound, carbenoxolone. In healthy individuals, carbenoxolone improves whole body

insulin sensitivity, and in patients with type 2 diabetes it decreases glucose production rates, principally through a decrease in glycogenolysis with no apparent effect on gluconeogenesis. In addition, it decreases total circulating cholesterol (see Andrews, R. C. et al. "Effects of the 11-hydroxysteroid dehydrogenase inhibitor carbenoxole on
5 insulin sensitivity in men with type 2 diabetes" in *J. Clin. Endocrinol. Metab.* **2003**, 88, 285-291).

Highly potent selective 11 β -HSD1 inhibitors have been developed by several major pharmaceutical companies such as Abbott, Amgen, Astellas, Bristol-Myers-Squibb, Daiichi Sankyo, Merck, Novo Nordisk, AstraZeneca, Sterix, Shionogi,
10 Biovitrum, Janssen, Pfizer, Takeda, Lilly, Taisho, Wyeth, Ono, Incyte, Roche and Novartis (see, Fotsch, C.; Askew, B. C.; Chen, J. J. "11 β -Hydroxysteroid dehydrogenase-1 as a therapeutic target for metabolic diseases" in *Expert Opin. Ther. Patents* **2005**, 15, 289-303, and Boyle, C. D.; Kowalski, T. J. "11 β -hydroxysteroid dehydrogenase type 1 inhibitors: a review of recent patents" in *Expert Opin. Ther.*
15 *Patents* **2009**, 19, 801-825).

In clinical trials, 11 β -HSD1 inhibitors have been well tolerated and have improved glycaemic control, lipid profile and blood pressure, and induced modest weight loss (see, Anderson, A.; Walker, B. R. "11 β -HSD1 inhibitors for the Treatment of Type 2 Diabetes and Cardiovascular Disease", in *Drugs* **2013**, 73, 1385-1393). Proof
20 of concept for the 11 β -HSD1 inhibitor class was demonstrated with INCB13739. In patients with T2DM failing metformin monotherapy, INCB13739 treatment achieved significant reductions in haemoglobin A1c and fasting plasma glucose, and when present improves hyperlipidaemia and hypertriglyceridaemia (see, Hollis, G.; Huber, R. "11 β -Hydroxysteroid dehydrogenase type 1 inhibition in type 2 diabetes mellitus" in
25 *Diabetes, Obes. Metab.* **2011**, 13, 1-6). More recently, novel 11 β -HSD1 inhibitors have been progressed into clinical trials (e. g., see Scott, J. S. "Medicinal chemistry of inhibitors of 11 β -hydroxysteroid dehydrogenase type 1 (11 β -HSD1)" in *J. Med. Chem.* **2014**, 57, 4466-4486). For example, in a phase 1b trial, RO5093151 was effective and safe in reducing liver-fat content in overweight, insulin-resistant patients with liver
30 steatosis (see Stefan, N. "Inhibition of 11 β -HSD1 with RO5093151 for non-alcoholic fatty liver disease: a multicentre, randomised, double-blind, placebo-controlled trial" in

Lancet Diabetes Endocrinol. **2014**, 2, 406-416). Another inhibitor, RO5027838, in a phase 2 study in metformin-treated patients with T2DM over 28 days showed inhibitory effects on 11 β -HSD1 activity. This molecule showed a trend for improved haemoglobin A1c levels and consistent reduction of body weight exceeding that observed with placebo (see Heisel, T. et al. "Safety, efficacy and weight effect of two 11 β -HSD1 inhibitors in metformin-treated patients with type 2 diabetes" in *Diabetes Obes. Meta.* **2014**, 16, 1070-1077). However, most of 11 β -HSD1 inhibitor development programs have been discontinued, predominantly due to insufficient efficacy. Therefore novel 11 β -HSD1 inhibitors are still required.

10 VASCULAR INFLAMMATION AND ATHEROSCLEROSIS

Several lines of evidence points towards 11 β -HSD1 inhibition as an attractive target in atherosclerosis and vascular inflammation (see, e.g. Hadoke, P. W. F. et al. "Modulation of 11 β -hydroxysteroid dehydrogenase as a strategy to reduce vascular inflammation" in *Curr. Atheroscler. Rep.* **2013**, 15:320). Inhibition of 11 β -HSD1 with compound MK-544 slowed plaque progression in a murine model of atherosclerosis, the key clinical sequela of metabolic syndrome. Mice with a targeted deletion of apolipoprotein E exhibited 84% less accumulation of aortic total cholesterol, as well as lower serum cholesterol and triglycerides, when treated with an 11 β -HSD1 inhibitor. These data provide the first evidence that pharmacologic inhibition of intracellular GC activation can effectively treat atherosclerosis, the key clinical consequence of metabolic syndrome, in addition to its salutary effect on multiple aspects of the metabolic syndrome itself (see, Hermanowski-Vosatka A. et al. "11 β -HSD1 inhibition ameliorates metabolic syndrome and prevents progression of atherosclerosis in mice" in *J. Exp. Med.* **2005**, 202, 517-527). More recently, a similar reduction in intra-aortic cholesterol has been shown with a distinct inhibitor (L-750) (Luo, M. J. et al. "11 β -HSD1 inhibition reduces atherosclerosis in mice by altering proinflammatory gene expression in the vasculature" in *Physiol. Genomics* **2013**, 45, 47-57). Additionally, it has been found that levels of 11 β -HSD1 mRNA are increased in human atherosclerotic vessels compared either with nearby intact (atheroma-free) vasculature or with vessels from control patients without coronary artery disease (Atalar, F. et al. "11 β -hydroxysteroid dehydrogenase type 1 gene expression is increased in ascending aorta

tissue of metabolic syndrome patients with coronary artery disease” in *Genet. Mol. Res.* **2012**, *11*, 3122-3132; Ayari, H. et al. “Auto-amplification of cortisol actions in human carotid atheroma is linked to arterial remodelling and stroke” in *Fundam. Clin. Pharmacol.* **2014**, *28*, 53-64).

5 COGNITIVE DYSFUNCTION AND RELATED DISORDERS SUCH AS VASCULAR DEMENTIA

In the CNS, 11 β -HSD1 is highly expressed in regions important for cognition such as hypothalamus, hippocampus, frontal cortex, and cerebellum (see, e.g., Moisan et al., “11beta-hydroxysteroid dehydrogenase bioactivity and messenger RNA
10 expression in rat forebrain: localization in hypothalamus, hippocampus, and cortex” in *Endocrinology* **1990**, *127*, 1450-1455). Growing evidence suggests excessive glucocorticoid activity may contribute to Alzheimer’s disease (AD) and age-associated memory impairment. It is known that glucocorticoids locally generated by 11beta-HSD1 play a major role in age-related cognitive impairments (MacLulich, A. M. J. et al., “11 β -hydroxysteroid dehydrogenase type 1, brain atrophy and cognitive decline” in
15 *Neurobiol. Aging* **2012**, *33*, 207.e1-207.e8) and that 11beta-HSD1 deficiency prevents memory deficits with aging (see, e.g., Yau, J. L. W. et al. “Lack of tissue glucocorticoid reactivation in 11 β -hydroxysteroid dehydrogenase type 1 knockout mice ameliorates age-related learning impairments” in *Proc. Natl. Acad. Sci. U.S.A.* **2001**, *98*,
20 4716–4721). Also, it has been found that aging associates with elevated hippocampal and neocortical 11beta-HSD1 and impaired spatial learning while deficiency of 11beta-HSD1 in knockout mice prevents the emergence of cognitive decline with age (see, e.g., Yau, J. L. W. et al. “Local amplification of glucocorticoids in the aging brain and impaired spatial memory” in *Frontiers Aging Neurosci.* **2012**, *4*, 24). Thus, mice with
25 lifelong deficiency of 11beta-HSD1 resist age-dependent spatial memory impairments even though plasma corticosterone levels are elevated to the same level as wildtype mice (see, e.g., Yau, J. L. W. et al. “Lack of tissue glucocorticoid reactivation in 11 β -hydroxysteroid dehydrogenase type 1 knockout mice ameliorates age-related learning impairments” in *Proc. Natl. Acad. Sci. U.S.A.* **2001**, *98*, 4716–4721). This occurs in
30 various cognitive tasks of spatial memory, such as the Morris water maze and the Y-maze (see, e.g., Yau, J. L. W. et al. “Enhanced hippocampal long-term potentiation and

spatial learning in aged 11beta-hydroxysteroid dehydrogenase type 1 knockout mice” in *J. Neurosci.* **2007**, 27, 10487-10496). Conversely transgenic mice with modest forebrain-specific overexpression of 11beta-HSD1 show accelerated memory deficits with ageing (see, e.g., Holmes, M. C. et al., “11beta-Hydroxysteroid dehydrogenase type 1 expression is increased in the aged mouse hippocampus and parietal cortex and causes memory impairments” in *J. Neurosci.* **2010**, 19, 6916-6920).

Furthermore, short-term pharmacological inhibition of 11beta-HSD1 in already aged mice reverses spatial memory impairments (see, e.g., Sooy, K. et al. “Partial deficiency or short-term inhibition of 11beta-hydroxysteroid dehydrogenase type 1 improves cognitive function in aging mice” in *J. Neurosci.* **2010**, 30, 13867-13872; Yau, J. L. W. et al. “Intrahippocampal glucocorticoids generated by 11beta-HSD1 affect memory in aged mice” in *Neurobiol. Aging* **2015**, 36, 334-343; Wheelan, N. et al. “Short-term inhibition of 11beta-hydroxysteroid dehydrogenase type 1 reversibly improves spatial memory but persistently impairs contextual fear memory in aged mice” in *Neuropharmacol.* **2015**, 91, 71-76. Also, in a well-characterized murine AD model, the Tg2576 mouse, which bears a mutated human amyloid precursor protein (APP) gene, a short-term treatment (four weeks) with a known 11beta-HSD1 inhibitor improves cognition and a long-term (44 weeks) prevents cognitive decline (see, e.g., Sooy, K. et al. “Cognitive and disease-modifying effects of 11beta-hydroxysteroid dehydrogenase type 1 inhibition in male Tg2576 mice, a model of Alzheimer’s disease” in *Endocrinology* **2015**, 156, 4592-4603). Finally, Amgen has also disclosed that aged rats treated with another selective 11beta-HSD1 inhibitor, AMG221, exhibited superior improvement in the novel object recognition test, when compared to rats that had been administered anti-AD drug galantamine (WO2012051139A1). Finally, in humans, four weeks of the non-selective 11beta-hydroxysteroid dehydrogenase inhibitor carbenoxolone improved verbal fluency in healthy elderly men and type 2 diabetics who have a selective impairment in verbal memory (Sandeep, T. C. et al. “11beta-hydroxysteroid dehydrogenase inhibition improves cognitive function in healthy elderly men and type 2 diabetics” in *Proc. Natl. Acad. Sci. USA* **2004**, 101, 6734-6739).

Also, increased cortisol levels after an overnight dexamethasone-suppression test have been reported in vascular dementia and depression, suggesting a beneficial role of 11b-HSD1 inhibitors in these diseases (see, e.g., Martocchia, A. et al. “Recent advances

in the role of cortisol and metabolic syndrome in age-related degenerative diseases” in *Aging Clin. Exp. Res.* **2016**, 28, 17–23).

Thus, 11 β -HSDI inhibitors are of potential therapeutic utility in the treatment of diseases which are characterised by cognitive impairment such as Alzheimer’ disease
5 and related neurodegenerative diseases.

GLAUCOMA

Glucocorticoid target receptors and the enzymes regulating glucocorticoid activity at these receptors are present in mammalian ocular tissues, which regulate aqueous humor formation and outflow. Alteration in the number or affinity of receptors
10 or in the activity of regulatory enzymes may alter the susceptibility of certain individuals to the effects of glucocorticoids on intraocular pressure. Glucocorticoids increase the risk of glaucoma by increasing intraocular pressure when they are exogenously administered and in certain conditions of increased production such as Cushing's syndrome (see, Stokes, J. et al. “Distribution of glucocorticoid and
15 mineralocorticoid receptors and 11 β -hydroxysteroid dehydrogenases in human and rat ocular tissues” in *Invest. Ophthalmol. Vis. Sci.* **2000**, 41, 1629–1638). It has been found that the intraocular pressure can be decreased significantly in humans after few days of oral treatment with carbenoxolone, a known 11 β -HSD1 inhibitor (see, Rauz, S. et al. “Expression and putative role of 11 β -hydroxysteroid dehydrogenase isozymes
20 within the human eye” in *Invest. Ophthalmol. Vis. Science* **2001**, 42, 2037-2042). Also, carbenoxolone lowers intraocular pressure in patients with ocular hypertension mediated through inhibition of 11 β -HSD1 in the ciliary epithelium (Rauz, S. et al. “Inhibition of 11 β -hydroxysteroid dehydrogenase type 1 lowers intraocular pressure in patients with ocular hypertension” in *Q. J. Med.* **2003**, 96, 481-490). Thus, selective and
25 topical inhibitors of 11 β -HSD1 could provide a treatment for patients with glaucoma.

OSTEOPOROSIS

Osteoporosis is characterized by decreased bone mineral density which leads to bone fragility and increased fracture risk due to reduced bone strength. This is due to the imbalance of the bone remodeling process. Glucocorticoids are crucial for normal
30 bone development but are detrimental in excess. Rapid onset of bone loss and fragility

fractures are common complications of long term systemic glucocorticoids treatment and glucocorticoid induced osteoporosis affects up to 50 % of patients taking long term glucocorticoid therapy (e.g., see Raqisz, L. G. "Pathogenesis of osteoporosis: concepts, conflicts, and prospects" in *J. Clin. Invest.* **2005**, *115*, 3318-3325). Glucocorticoid action in bone depends upon the activity of 11 β -HSD1 enzyme. Regulations of 11 β -HSD1 activity may protect the bone against bone loss due to excess glucocorticoids. Treatment with glycyrrhizic acid, a potent 11 β -HSD inhibitor, led to significant reduction in bone resorption marker pyridinoline and improved the bone structure and biomechanical strength of rats receiving long-term glucocorticoids therapy (Ramli, E. S. M. et al. "Glycyrrhizic acid (GCA) as 11 β -hydroxysteroid dehydrogenase inhibitor exerts protective effect against glucocorticoid-induced osteoporosis" in *J. Bone Miner. Metab.* **2013**, *31*, 262-273). In another study, BVT-2733, an 11 β -HSD1 selective inhibitor, protected osteoblasts against endogenous glucocorticoid induced dysfunction (see Wu, L. et al. "11 β -hydroxysteroid dehydrogenase type 1 selective inhibitor BVT.2733 protects osteoblasts against endogenous glucocorticoid induced dysfunction" in *Endocrine J.* **2013**, *60*, 1047-1058). Also, KR-67500, a potent 11 β -HSD1 inhibitor, enhances the osteolastogenesis and inhibits the osteoclastogenesis (see Park, J. S. et al. "A novel 11 β -HSD1 inhibitor improves diabetes and osteoblast differentiation" in *J. Mol. Endocrinol.* **2014**, *52*, 191-202). Therefore, 11 β -HSD1 inhibitors have the potential to be used as new therapeutic agents against osteoporosis.

ANGIOGENESIS AND MYOCARDIAL INFARCTION:

Angiogenesis restores blood flow to healing tissues, a process that is inhibited by high doses of glucocorticoids. Endogenous glucocorticoids, including those generated locally by 11 β -HSD1, exert tonic inhibition of angiogenesis. 11 β -HSD1 knockout mice showed enhanced angiogenesis *in vitro* and *in vivo* within sponges, wounds, and infarcted myocardium (e.g., see McSweeney, S. et al. "Improved heart function enhanced inflammatory cell recruitment and angiogenesis in 11 β -HSD1-deficient mice post-MI" in *Cardiovasc. Res.* **2010**, *88*, 159-167). Thus, inhibition of 11 β -HSD1 may offer a therapeutic approach to improve healing of ischemic or injured tissue, such as in ischemic heart diseases and impaired wound healing (see, Small, G.R. et al. "Preventing local regeneration of glucocorticoids by 11 β -hydroxysteroid dehydrogenase type 1

enhances angiogenesis” in *Proc. Natl. Acad. Sci. U. S. A.* **2005**, *102*, 12165–12170). Recently it has been found that UE2316, a selective 11 β -HSD1 inhibitor, increases angiogenesis and improves cardiac function after myocardial infarction in mice (see, McGregor, K. et al. “Immediate pharmacological inhibition of local glucocorticoid
5 generation increases angiogenesis and improves cardiac function after myocardial infarction” in *Heart* **2014**, *100* (supp 3), A118).

TREATMENT OF ALCOHOL ABUSE:

Few phamacotherapies for alcohol abuse, the most prevalent abused substance in the United Stantes and Europe, are currently available, and these have shown only
10 limited efficacy and compliance. Thus, the development of more effective medications for alcohol abuse is a significant unmet medical need (e.g., see Litten, R. Z. et al. “Medications development to treat alcohol dependence: a vision for the next decade” in *Addict Biol.* **2012**, *17*, 513-527). Alcohol intake activates glucocorticoids. As glucocorticoid effects are modulated in part by the activity of 11 β -HSD, its inhibition
15 may be beneficial for the treatment of alcohol abuse. In fact, the 11-HSD inhibitor carbenoxolone reduces both baseline and excessive drinking in rodents (see Sanna, P. P. et al. “11b-hydroxysteroid dehydrogenase inhibition as a new potential therapeutic target for alcohol abuse” in *Transl. Psychiatry* **2016**, *6*, e760).

The patent literature describe numerous examples of 11-beta-hydroxysteroid
20 dehydrogenase type 1 (11-beta-HSD1) inhibitors. The following are a few examples: WO 2015/077286, WO 2013/025664, WO 2011/161128, WO 2011/068927, WO 2012/061708, WO 2011/049520, WO 2011/002910, WO 2011/011123, WO 2010/139673, WO 2010/141424, WO 2010/089303, WO 2010/091067, WO 2010/022850, WO 2010/010157, WO 2010/010150, WO 2009/135581, WO
25 2009/132986, WO 2009/130496, WO 2009/131669, WO 2009/117109, WO 2009/108332, WO 2009/102460, WO 2009/102428, WO 2009/094169, WO 2009/090239, WO 2009/088997, WO 2009/075835, WO 2009/061498, WO 2009/059666, WO 2008/157752, WO 2008/156601, WO 2008/106128, WO 2008/088540, WO 2008/074384, WO 2008/069313, WO 2008/052638, WO
30 2009/056881, WO 2008/046758, WO 2008/024497, WO 2008/012532, WO

2008/011453, WO 2008/003611, WO 2007/137066, WO 2007/130898, WO
2007/127688, WO 2007/103719, WO 2007/101270, WO 2007/092435, WO
2007/089683, WO 2007/084314, WO 2007/068330, WO 2007/067504, WO
2007/057768, WO 2007/038138, WO 2007/025880, WO 2007/003521, WO
5 2006/134467, WO 2006/132436, WO 2006/055752, WO 2006/053024, WO
2006/048750, WO 2006/034804, WO 2006/020598, US 2006-0025445, WO
2006/000371, WO 2006/002361, WO 2006/012226, WO 2006/002349, WO
2006/012173, WO 2006/012227, WO 2006/002350, WO 2005/110992, WO
2005/108359, WO 2009/026422, WO 2005/097759, US 8,178,546, WO 2005/060963,
10 WO 2005/046685, WO 2005/047250, WO 2004/112784, WO 2004/112779, WO
2004/112781, WO 2004/112782, WO 2004/112783, WO 2004/112785, WO
2004/113310, WO 2004/058730, WO 2004/041264, WO 2004/033427, WO
2004/011410, WO 2003/104208, WO 2003/075660, WO 2003/065983, WO
2003/043999, WO 2003/044000 and WO 01/90091.

15

US 3,475,450 describes bridged tetracyclic indole derivatives as hypoglycemic agents wherein the bridged tetracyclic core structure is substituted on the ring nitrogen by an arylsulfonylamide group.

20 WO 2006/129178 describes bridged aza (tri an tetra)cyclic derivatives as cannabinoid receptor CB1 ligand for treating metabolic, CNS, and inflammatory diseases.

Rey-Carrizo M., et al, "*J. Med. Chem.*" 2014, V. 57, N. 13, pp 5738-5747 describes bridged tetracyclic derivatives as inhibitors of the M2 proton channel for
25 treating influenza virus.

Therefore, there is a need in the art to develop further inhibitors 11-beta-hydroxysteroid dehydrogenase type 1.

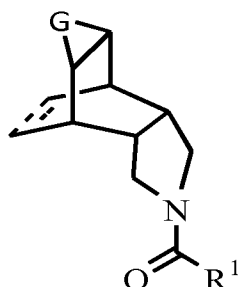
30 SUMMARY OF THE INVENTION

The inventors have surprisingly found that compounds of formula (I) are inhibitors of the 11-beta-hydroxysteroid dehydrogenase type 1 (11-beta-HSD1).

DETAILED DESCRIPTION OF THE INVENTION

5

In a first aspect, the invention relates to compounds of formula (I):



wherein

R¹ is selected from the group consisting of:

- 10 a) C₃₋₈ cycloalkyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl;
- b) C₃₋₈ cycloalkenyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl;
- c) C₆₋₁₀ aromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy;
- 15 d) C₅₋₁₀ heteroaromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NHR³, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₅₋₁₀ heteroaryl (optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy), phenyl (optionally substituted with one or two groups selected from halogen atoms, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy) and C₅₋₇ heterocyclyl (optionally substituted with a group selected from -COR⁴, -COOR⁴, -CN, -CONHR⁴, -CONH₂ and phenyl which is optionally substituted with one or two groups selected from halogen atoms, CONH₂, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy), wherein the C₅₋₇ heterocyclyl group
- 20 comprises a nitrogen atom forming part of the heterocycle and is attached to the C₅₋₁₀ heteroaromatic group through said nitrogen atom;
- 25

- e) C₃₋₈ heterocyclyl optionally substituted with a group selected from C₁₋₄ alkyl, C₃₋₈ cycloalkyl and -COR⁵; and
- f) -C(CH₃)₂-R²;

G is a rest selected from the group consisting of -CH₂-, -CH₂-CH₂- and -CH=CH-;

- 5 R² is selected from the group consisting of hydrogen atom and C₁₋₃ alkyl

R³ is selected from the group consisting of C₁₋₃ alkyl and C₁₋₃ hydroxyalkyl

R⁴ is selected from the group consisting of C₁₋₆ alkyl and phenyl

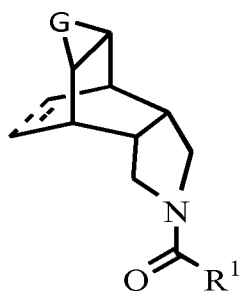
R⁵ is a group C₁₋₆ alkyl

the bond represented by --- is either a single bond or a double bond;

- 10 and pharmaceutically stable salts thereof for use in the treatment or prevention of a disease or conditions susceptible of improvement by inhibition of 11 β -hydroxysteroid dehydrogenase type 1 (11 β -HD1), preferably a disease or condition selected from the group consisting of obesity, insulin resistance, metabolic syndrome, cognitive disfunction associated to Alzheimer's disease, HIV-associated dementia, Creutzfeld-
- 15 Jakob disease, Parkinson's disease, Huntington's disease, vascular inflammation, atherosclerosis, diabetes type 2, cardiovascular disease, glaucoma, osteoporosis, myocardial infarction and alcohol abuse treatment.

The compounds of formula (I) defined in the first aspect are new.

Thus, in a second aspect, the invention relates to new compounds of formula (I):



20

wherein

R¹ is selected from the group consisting of:

- a) C₃₋₈ cycloalkyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl;
- b) C₃₋₈ cycloalkenyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl;
- 5 c) C₆₋₁₀ aromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy;
- d) C₅₋₁₀ heteroaromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NHR³, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₅₋₁₀ heteroaryl (optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy), phenyl (optionally substituted with one or two groups selected from halogen atoms, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy) and C₅₋₇ heterocyclyl (optionally substituted with a group selected from -COR⁴, -COOR⁴, -CN, -CONHR⁴, -CONH₂ and phenyl which is optionally substituted with one or two groups selected from halogen atoms, CONH₂, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy), wherein the C₅₋₇ heterocyclyl group comprises a nitrogen atom forming part of the heterocycle and is attached to the C₅₋₁₀ heteroaromatic group through said nitrogen atom;
- 10 e) C₃₋₈ heterocyclyl optionally substituted with a group selected from C₁₋₄ alkyl, C₃₋₈ cycloalkyl and -COR⁵; and
- 15 f) -C(CH₃)₂-R²;
- 20

G is a rest selected from the group consisting of -CH₂-, -CH₂-CH₂- and -CH=CH-;

R² is selected from the group consisting of hydrogen atom and C₁₋₃ alkyl

R³ is selected from the group consisting of C₁₋₃ alkyl and C₁₋₃ hydroxyalkyl

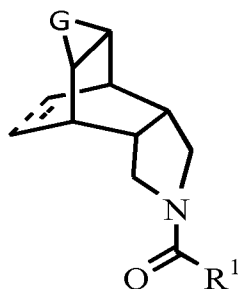
R⁴ is selected from the group consisting of C₁₋₆ alkyl and phenyl

25 R⁵ is a group C₁₋₆ alkyl

the bond represented by --- is either a single bond or a double bond;

and pharmaceutically stable salts thereof.

In a third aspect, the invention relates to the use of a compound of formula (I):



wherein

R¹ is selected from the group consisting of:

- 5 a) C₃₋₈ cycloalkyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl;
- b) C₃₋₈ cycloalkenyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl;
- c) C₆₋₁₀ aromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy;
- 10 d) C₅₋₁₀ heteroaromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NHR³, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₅₋₁₀ heteroaryl (optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy), phenyl (optionally substituted with one or two groups selected from halogen atoms, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy) and C₅₋₇ heterocyclyl (optionally substituted with a group selected from -COR⁴, -COOR⁴, -CN, -CONHR⁴, -CONH₂ and phenyl which is optionally substituted with one or two groups selected from halogen atoms, CONH₂, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy), wherein the C₅₋₇ heterocyclyl group comprises a nitrogen atom forming part of the heterocycle and is attached to the C₅₋₁₀ heteroaromatic group through said nitrogen atom;
- 15 e) C₃₋₈ heterocyclyl optionally substituted with a group selected from C₁₋₄ alkyl, C₃₋₈ cycloalkyl and -COR⁵; and
- f) -C(CH₃)₂-R²;
- 20 G is a rest selected from the group consisting of -CH₂-, -CH₂-CH₂- and -CH=CH-;

R^2 is selected from the group consisting of hydrogen atom and C_{1-3} alkyl

R^3 is selected from the group consisting of C_{1-3} alkyl and C_{1-3} hydroxyalkyl

R^4 is selected from the group consisting of C_{1-6} alkyl and phenyl

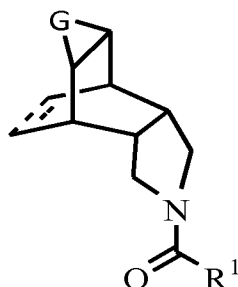
R^5 is a group C_{1-6} alkyl

- 5 the bond represented by --- is either a single bond or a double bond;

and pharmaceutically stable salts thereof for the manufacture of a medicament for the treatment or prevention of a disease or conditions susceptible of improvement by inhibition of 11β -hydroxysteroid dehydrogenase type 1 (11β -HD1), in particular a disease or condition selected from the group consisting of obesity, insulin resistance, metabolic syndrome, cognitive dysfunction associated to Alzheimer's disease, HIV-associated dementia, Creutzfeld-Jakob disease, Parkinson's disease, Huntington's disease, vascular inflammation, atherosclerosis, diabetes type 2, cardiovascular disease, glaucoma, osteoporosis, myocardial infarction and alcohol abuse treatment.

- 15 In a fourth aspect, the invention relates to the method for the treatment or prevention of a disease or conditions susceptible of improvement by inhibition of 11β -hydroxysteroid dehydrogenase type 1 (11β -HD1), in particular a disease or condition selected from the group consisting of obesity, insulin resistance, metabolic syndrome, cognitive dysfunction associated to Alzheimer's disease, HIV-associated dementia, Creutzfeld-Jakob disease, Parkinson's disease, Huntington's disease, vascular inflammation, atherosclerosis, diabetes type 2, cardiovascular disease, glaucoma, osteoporosis, myocardial infarction and alcohol abuse treatment, in a subject in need thereof comprising the administration to said subject of a therapeutically effective amount of a compound of formula (I):

19



wherein

R¹ is selected from the group consisting of:

- 5 a) C₃₋₈ cycloalkyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl;
- b) C₃₋₈ cycloalkenyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl;
- c) C₆₋₁₀ aromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy;
- 10 d) C₅₋₁₀ heteroaromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NHR³, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₅₋₁₀ heteroaryl (optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy), phenyl (optionally substituted with one or two groups selected from halogen atoms, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy) and C₅₋₇ heterocyclyl (optionally substituted with a group selected from -COR⁴, -COOR⁴, -CN, -CONHR⁴, -CONH₂ and phenyl which is optionally substituted with one or two groups selected from halogen atoms, CONH₂, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy), wherein the C₅₋₇ heterocyclyl group comprises a nitrogen atom forming part of the heterocycle and is attached to the C₅₋₁₀ heteroaromatic group through said nitrogen atom;
- 15 e) C₃₋₈ heterocyclyl optionally substituted with a group selected from C₁₋₄ alkyl, C₃₋₈ cycloalkyl and -COR⁵; and
- 20 f) -C(CH₃)₂-R²;

G is a rest selected from the group consisting of -CH₂-, -CH₂-CH₂- and -CH=CH-;

- 25 R² is selected from the group consisting of hydrogen atom and C₁₋₃ alkyl

R^3 is selected from the group consisting of C_{1-3} alkyl and C_{1-3} hydroxyalkyl

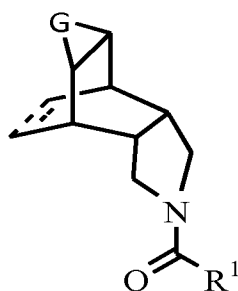
R^4 is selected from the group consisting of C_{1-6} alkyl and phenyl

R^5 is a group C_{1-6} alkyl

the bond represented by --- is either a single bond or a double bond;

5 and pharmaceutically stable salts thereof.

In a fifth aspect, the invention relates to a pharmaceutical composition comprising a compound of formula (I):



10

wherein

R^1 is selected from the group consisting of:

- a) C_{3-8} cycloalkyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C_{1-4} alkyl;
- 15 b) C_{3-8} cycloalkenyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C_{1-4} alkyl;
- c) C_{6-10} aromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF_3 , NH_2 , OH, C_{1-4} alkyl and C_{1-4} alkoxy;
- 20 d) C_{5-10} heteroaromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF_3 , NHR^3 , C_{1-4} alkyl, C_{1-4} alkoxy, C_{5-10} heteroaryl (optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF_3 , NH_2 , OH, C_{1-4} alkyl and C_{1-4} alkoxy), phenyl (optionally substituted with one or two groups selected from halogen atoms, CN, OH, CF_3 , C_{1-4} alkyl and C_{1-4} alkoxy) and C_{5-7} heterocyclyl (optionally substituted with a

- group selected from $-\text{COR}^4$, $-\text{COOR}^4$, $-\text{CN}$, $-\text{CONHR}^4$, $-\text{CONH}_2$ and phenyl which is optionally substituted with one or two groups selected from halogen atoms, CONH_2 , CN , OH , CF_3 , C_{1-4} alkyl and C_{1-4} alkoxy), wherein the C_{5-7} heterocyclyl group comprises a nitrogen atom forming part of the heterocycle and is attached to the C_{5-10} heteroaromatic group through said nitrogen atom;
- 5 e) C_{3-8} heterocyclyl optionally substituted with a group selected from C_{1-4} alkyl, C_{3-8} cycloalkyl and $-\text{COR}^5$; and
- f) $-\text{C}(\text{CH}_3)_2-\text{R}^2$;

G is a rest selected from the group consisting of $-\text{CH}_2-$, $-\text{CH}_2-\text{CH}_2-$ and $-\text{CH}=\text{CH}-$;

- 10 R^2 is selected from the group consisting of hydrogen atom and C_{1-3} alkyl

R^3 is selected from the group consisting of C_{1-3} alkyl and C_{1-3} hydroxyalkyl

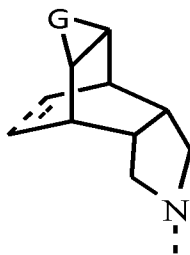
R^4 is selected from the group consisting of C_{1-6} alkyl and phenyl

R^5 is a group C_{1-6} alkyl

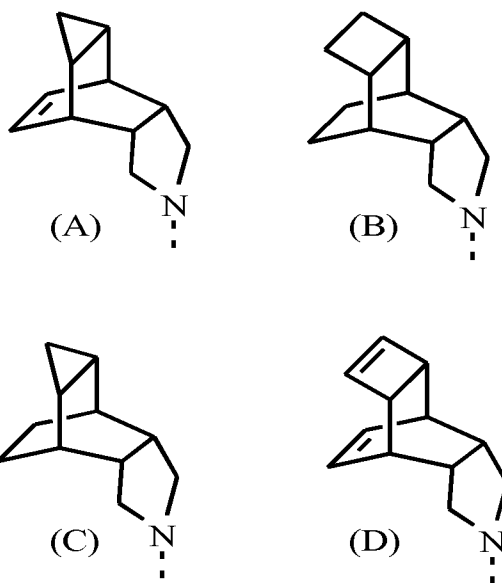
the bond represented by --- is either a single bond or a double bond;

- 15 and pharmaceutically stable salts thereof and a pharmaceutically acceptable excipient.

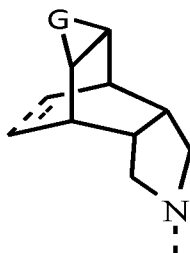
In one embodiment of the first, second, third, fourth or fifth aspects of the present invention in the compounds of formula (I) the rest



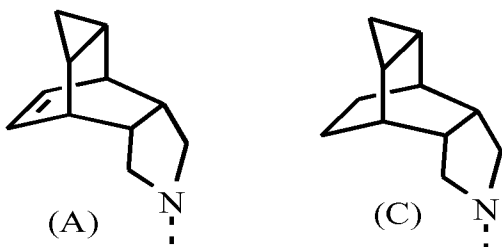
- 20 is selected from the group consisting of rests (A) to (D):



In another embodiment of the first, second, third, fourth or fifth aspects of the present invention in the compounds of formula (I) the rest



5 is selected from the group consisting of rests (A) and (C):



In another embodiment of the first, second, third, fourth or fifth aspects of the present invention in the compounds of formula (I) G is a rest $-\text{CH}_2-$.

In another embodiment of the first, second, third, fourth or fifth aspects of the present invention in the compounds of formula (I) the bond represented by $\equiv$ is a double bond.

In another embodiment of the first, second, third, fourth or fifth aspects of the present invention in the compounds of formula (I) R^1 is a C_{3-8} cycloalkyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C_{1-4} alkyl.

In another embodiment of the first, second, third, fourth or fifth aspects of the present invention in the compounds of formula (I) R^1 is a C_{3-8} cycloalkenyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C_{1-4} alkyl.

5 In another embodiment of the first, second, third, fourth or fifth aspects of the present invention in the compounds of formula (I) R^1 is a C_{6-10} aromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF_3 , NH_2 , OH, C_{1-4} alkyl and C_{1-4} alkoxy.

10 In another embodiment of the first, second, third, fourth or fifth aspects of the present invention in the compounds of formula (I) R^1 is a C_{5-10} heteroaromatic group optionally substituted with 1 to 2 groups selected from halogen atoms, CN, CF_3 , NHR^3 , C_{1-4} alkyl, C_{1-4} alkoxy, C_{5-10} heteroaryl (optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF_3 , NH_2 , OH, C_{1-4} alkyl and C_{1-4} alkoxy), phenyl (optionally substituted with a group selected from halogen atoms, CN, OH, CF_3 , C_{1-4} alkyl and C_{1-4} alkoxy) and C_{5-7} heterocyclyl (optionally substituted with a group selected from $-COR^4$, $-COOR^4$, $-CN$, $-CONHR^4$, $-CONH_2$ and phenyl which is optionally substituted with one or two groups selected from halogen atoms, $CONH_2$, CN, OH, CF_3 , C_{1-4} alkyl and C_{1-4} alkoxy), wherein the C_{5-7} heterocyclyl group comprises a nitrogen atom forming part of the heterocycle and is attached to the C_{5-10} heteroaromatic group through said nitrogen atom, wherein R^3 is selected from the group
15 consisting of C_{1-3} alkyl and C_{1-3} hydroxyalkyl and R^4 is selected from the group consisting of C_{1-6} alkyl and phenyl.
20

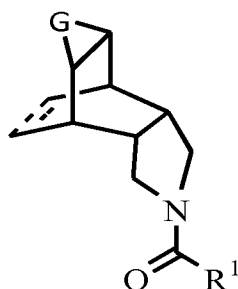
In another embodiment of the first, second, third, fourth or fifth aspects of the present invention in the compounds of formula (I) R^1 is a C_{3-8} heterocyclyl optionally substituted with a group selected from C_{1-4} alkyl, C_{3-8} cycloalkyl and COR^5 wherein R^5
25 is a group C_{1-6} alkyl.

In another embodiment of the first, second, third, fourth or fifth aspects of the present invention in the compounds of formula (I) R^1 is a group $-C(CH_3)_2-R^2$, wherein R^2 is selected from the group consisting of hydrogen atom and C_{1-3} alkyl.

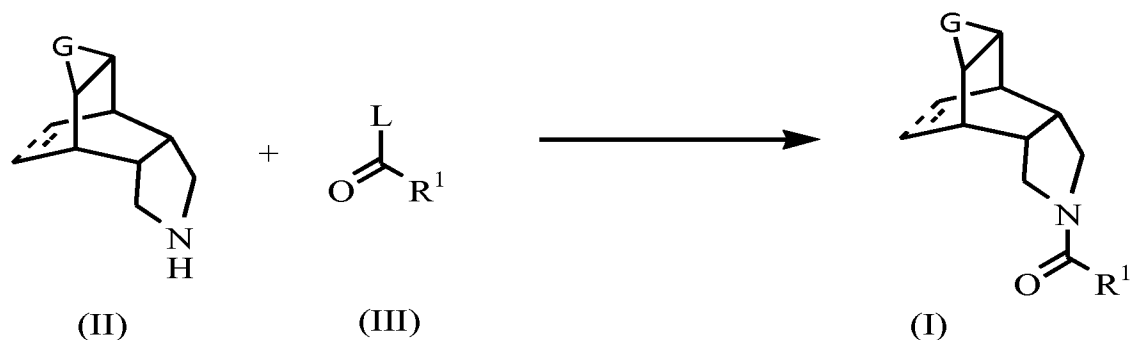
30 In another embodiment of the first, second, third, fourth or fifth aspects of the present invention in the compounds of formula (I) R^5 is a group methyl.

Process for producing the compounds of formula (I)

In a sixth aspect, the invention relates to a process for producing a compound of formula (I):



as defined in the first or second aspects of the invention by causing a polycyclic amine of formula (II) or a salt thereof to react with an acid derivative of formula (III).

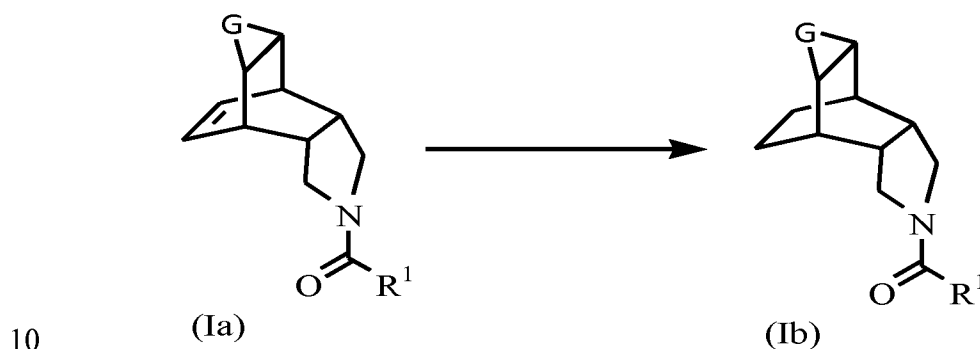


The group L in the compound of formula (III) is selected from the group consisting of halogen atoms, C₁₋₄-alkoxy, OH, and a group of formula –O-CO-R¹, i.e. the compounds of formula (III) are carboxylic acid halides, carboxylic acid esters, carboxylic acids or carboxylic acid anhydrides.

The reaction of compounds of formula (II) with compounds of formula (III) can be carried out following different methods known by any person skilled in the field of organic synthesis. In a preferred embodiment the group L is OH (i.e. the compounds of formula (III) are carboxylic acids) and the reaction is carried out in a solvent such as ethyl acetate in the presence of 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide (EDC), hydroxybenzotriazole (HOBt) and triethylamine (NEt₃), preferably at room temperature. Alternatively, the group L is Cl (i.e. the compounds of formula (III) are carboxylic acid chlorides) and the reaction is carried out in a solvent such as acetone or dichloromethane in the presence of a base such as triethylamine (NEt₃), preferably at room temperature. In turn, the carboxylic acid chlorides (compounds of formula (III) when the group L is Cl) can be obtained from the corresponding carboxylic acid using a

large variety of reagents such as thionyl chloride, sulfuryl chloride, oxalyl chloride, etc. In another embodiment, the group L is a group of formula $-O-CO-R^1$ (i.e. the compounds of formula (III) are carboxylic acid anhydrides) and the reaction is carried out in a solvent such as dichloromethane in the presence of a base such as triethylamine (NEt₃).

The compounds of the invention wherein the bond represented by --- is a single bond (compounds of formula (Ib)) may be also obtained by hydrogenation of the compounds of the invention wherein the bond represented by = is a double bond (compounds of formula (Ia)) as shown in the scheme below:



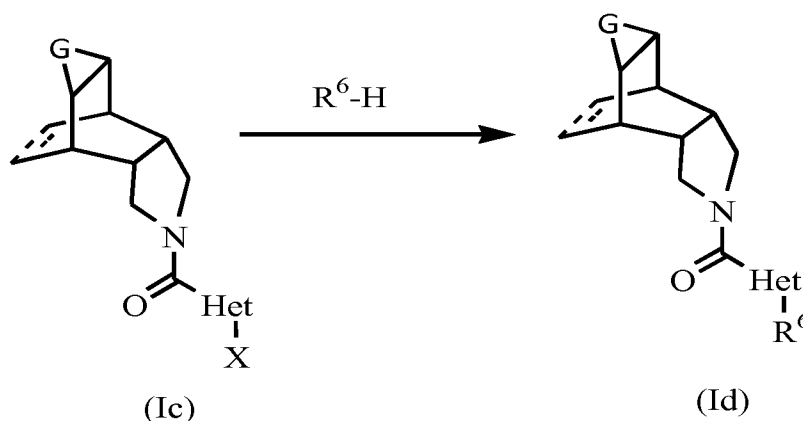
In an embodiment the hydrogenation is carried out in a solvent such as ethanol (or methanol) using hydrogen gas in the presence of a metal catalyst such as Pd/C, preferably at room temperature and at 1 atmosphere of pressure. Alternatively, the reduction can be carried out using other methods known by the skilled person, such as diimide or hydrogen transfer hydrogenation (HCO_2H , Et_3N , Pd).

The compounds of formula (II) are either commercially available or may be obtained by the process detailed in Rey-Carrizo, M et al., J Med Chem. 2014 Jul 10;57(13):5738-47.

Similarly the compounds of formula (III) are either commercially available or
20 may be obtained by processes known to the expert in the field.

When, in the compounds of formula (I), the group R¹ is a C₅₋₁₀ heteroaromatic group substituted with groups selected from NHR³ and C₅₋₇ heterocyclyl (optionally substituted with a group selected from -COR⁴, -COOR⁴, -CN, -CONHR⁴, -CONH₂ and phenyl which is optionally substituted with one or two groups selected from halogen atoms, CONH₂, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy), wherein the C₅₋₇ heterocyclyl group comprises a nitrogen atom forming part of the heterocycle and is attached to the C₅₋₁₀ heteroaromatic group through said nitrogen atom; said compounds

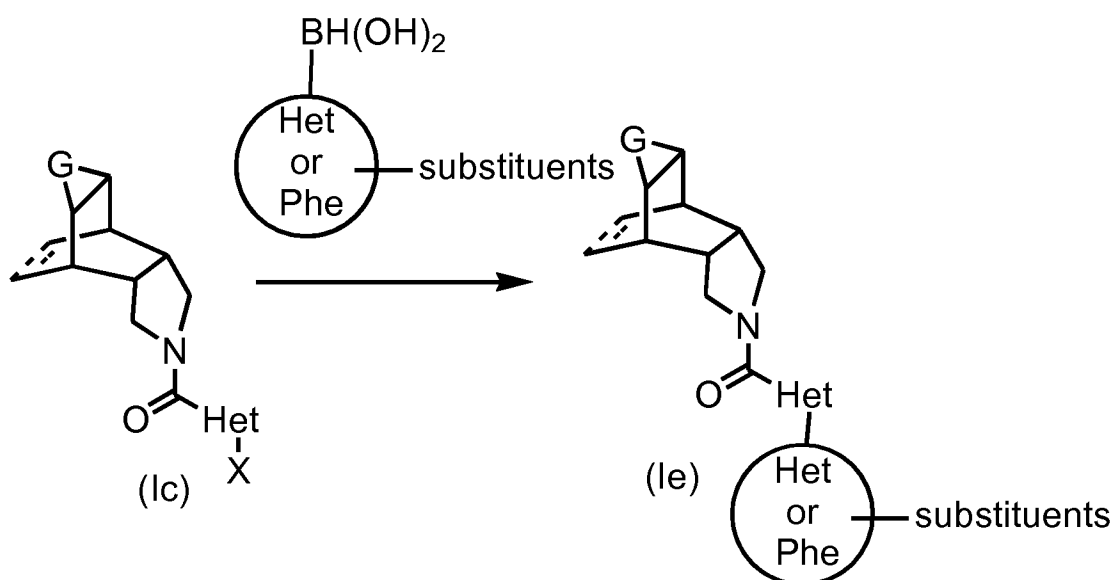
of formula (Id) may be obtained by reacting a compound having the corresponding C₅₋₁₀ heteroaromatic group substituted by a halogen atom (Ic) with a compound of formula R⁶-H wherein R⁶ represents a group selected from NHR³ and C₅₋₇ heterocyclyl (optionally substituted with a group selected from -COR⁴, -COOR⁴, -CN, -CONHR⁴, -CONH₂ and phenyl which is optionally substituted with one or two groups selected from halogen atoms, CONH₂, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy), wherein the C₅₋₇ heterocyclyl group comprises a nitrogen atom forming part of the heterocycle and is attached to the C₅₋₁₀ heteroaromatic group through said nitrogen atom as shown in scheme below:



10

When, in the compounds of formula (I), the group R¹ is a C₅₋₁₀ heteroaromatic group substituted by a group selected from C₅₋₁₀ heteroaryl (which is optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy) or by a phenyl group which is optionally substituted with one or two groups selected from halogen atoms, CONH₂, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy; said compounds of formula (Ie) may be obtained by reacting a compound having the corresponding C₅₋₁₀ heteroaromatic group substituted by a halogen atom (Ic) with a heteroaryl boronic acid or a phenyl boronic acid in the presence of a catalyst such as Pd(PPh₃)₄ and a base, such as K₂CO₃, wherein the phenyl is optionally substituted with one or two groups selected from halogen atoms, CONH₂, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy and the heteroaryl boronic acid is optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy as shown in scheme below:

20



Medical uses of the compounds of the invention

The compounds of the invention can be used in the treatment or prevention of a disease or condition susceptible of improvement by inhibition of 11β -hydroxysteroid dehydrogenase type 1 (11β -HD1), in particular a disease or condition selected from the group consisting of obesity, insulin resistance, metabolic syndrome, cognitive dysfunction associated to Alzheimer's disease, HIV-associated dementia, Creutzfeld-Jakob disease, Parkinson's disease, Huntington's disease, vascular inflammation, atherosclerosis, diabetes type 2, cardiovascular disease, glaucoma, osteoporosis, myocardial infarction and alcohol abuse treatment.

Definitions

The term "alkyl", as used herein, refers to; is used to designate linear or branched hydrocarbon fully saturated radicals ($\text{C}_n\text{H}_{2n+1}$). The indication C_{x-y} preceding the term "alkyl" is used to indicate that the alkyl may have from x to y carbon atoms. Examples of alkyls include methyl, ethyl, n-propyl, i-propyl, n-butyl, sec-butyl, tert-butyl, n-pentyl, 1-methyl-butyl, 2-methyl-butyl, isopentyl, 1-ethylpropyl, 1,1-dimethylpropyl, 1,2-dimethylpropyl, n-hexyl, 1-ethylbutyl, 2-ethylbutyl, 1,1-dimethylbutyl, 1,2-dimethylbutyl, 1,3-dimethylbutyl, 2,2-dimethylbutyl, 2,3-dimethylbutyl, 2-methylpentyl and 3-methylpentyl radicals. In a preferred embodiment said alkyl groups have 1 to 4 carbon atoms or 1 to 3 carbon atoms.

The term “alkoxy”, as used herein, refers to radicals which contain a linear or branched alkyl group linked to an oxygen atom ($C_nH_{2n+1}-O-$). The indication C_{x-y} preceding the term “alkoxy” is used to indicate that the alkoxy may have from x to y carbon atoms. Preferred alkoxy radicals include methoxy, ethoxy, n-propoxy, i-propoxy, n-butoxy, sec-butoxy and t-butoxy.

The term “hydroxyalkyl”, as used herein, refers to radicals which contain a linear or branched alkyl group in which one hydrogen atom has been replaced by a hydroxyl ($-OH$) group. The indication C_{x-y} preceding the term “hydroxyalkyl” is used to indicate that the hydroxyalkyl may have from x to y carbon atoms. Preferred hydroxyalkyl radicals include hydroxymethyl, hydroxyethyl, n-hydroxypropyl, i-hydroxypropyl, n-hydroxybutyl, sec-hydroxybutyl and t-hydroxybutyl.

The term “cycloalkyl”, as used herein, refers to hydrocarbon fully saturated cyclic groups atoms (C_nH_{2n-1}). The indication C_{x-y} preceding the term “cycloalkyl” is used to indicate that the cycloalkyl may have from x to y carbon atoms Said cycloalkyl groups may have a single cyclic ring or multiple condensed rings. Such cycloalkyl groups include, by way of example, single ring structures such as cyclopropyl, cyclobutyl, cyclopentyl, cyclooctyl, and the like, or multiple ring structures such as adamantanyl, bicyclo[2.2.1]heptane, 1,3,3-trimethylbicyclo[2.2.1]hept-2-yl, (2,3,3-trimethylbicyclo [2.2.1]hept-2-yl), or carbocyclic groups fused with an aryl group, for example indane, and the like.

The term “cycloalkenyl”, as used herein, refers to hydrocarbon partially unsaturated cyclic groups atoms comprising only single or double bonds between the carbon atoms of the cycle. A hydrocarbon cyclic group is considered to be partially unsaturated when it has a number of unsaturations which is lower than the maximum possible number of unsaturations admitted by said hydrocarbon cycle. The indication C_{x-y} preceding the term “cycloalkenyl” is used to indicate that the cycloalkenyl may have from x to y carbon atoms. Said cycloalkenyl groups may have a single cyclic ring or multiple condensed rings. Such cycloalkenyl groups include, by way of example, single ring structures such as cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclooctenyl, and the like, or multiple ring structures.

The term “halogen atom”, as used herein, is used to designate an atom selected from the group consisting of chlorine, fluorine, bromine or iodine atoms typically a

fluorine, chlorine or bromine atom. The term halo when used as a prefix has the same meaning.

The term “aromatic group”, as used herein, refers to a cyclic aromatic group wherein the cycle is made exclusively of carbon atoms. According to the IUPAC a cyclically conjugated group is said to be aromatic when it has a stability (due to delocalization) significantly greater than that of a hypothetical localized structure. The indication C_{x-y} preceding the term “aromatic group” is used to indicate that said group may have from x to y carbon atoms. Examples of aromatic groups are benzene, naphthalene, anthracene and phenanthrene.

The terms “heterocyclyl” and “heterocycle”, as used herein, refer to non-aromatic, saturated or unsaturated ring systems comprising carbon atoms and one or more heteroatoms selected from the group consisting of oxygen, nitrogen and sulphur. The indication C_{x-y} preceding the term “heterocyclyl” is used to indicate that the heterocyclyl may have from x to y carbon atoms. Said heterocyclyl groups may have a single cyclic ring or multiple condensed rings. Such heterocyclyl groups include, by way of example, single ring structures such as azetidine, oxetane, thietane, diazetidine, dioxetane, tetrahydrofurane, pyran, dihydropyran, tetrahydropyran, pyrrolidine, thiolane, imidazolidine, pyrazolidine, oxazolidine, isoxazolidine, thiazolidine, isothiazolidine, dioxolane, dithiolane, piperidine, oxane, thiane, piperazine, morpholine, thiomorpholine, dioxane, dithiane, azepane, oxepane and the like or multiple ring structures.

The term “optionally substituted”, as used herein, refers to the fact that a group or rest may have one or more of its hydrogen atoms replaced (substituted) by an atom or group different from hydrogen.

The term “heteroaromatic group” or heteroaryl, as used herein indistinctively, refers to a cyclic aromatic group wherein the cycle is made of carbon atoms and at least one atom selected from the group consisting of oxygen, nitrogen and sulphur. According to the IUPAC a cyclically conjugated group is said to be aromatic when it has a stability (due to delocalization) significantly greater than that of a hypothetical localized structure. The indication C_{x-y} preceding the term “heteroaromatic group” is used to indicate that said group may have from x to y carbon atoms. Examples of heteroaromatic groups are pyridine, pyrrol, furan, thiophene, pyrazine, pyrimidine,

pyridazine, imidazole, pyrazole, oxazole, thiophene, benzimidazole, quinolone, isoquinoline, quinoxaline, quinazoline, cinnoline, phthalazine, imidazopyridine, pyrazolopyridine, indazole, benzooxazole, benzofuran, isobenzofuran, benzothiophene, indol and isoindol.

5 The term “acetyl”, as used herein, refers to the group $\text{CH}_3\text{-C(=O)-}$;

 The term “pharmaceutically acceptable” relates to molecular entities and compositions being physiologically tolerable and normally not causing an allergic reaction or similar adverse reaction, such as gastric discomfort, dizziness and the like, when they are administered to a human being. Preferably, as used in this description, the
10 term “pharmaceutically acceptable” means approved by a governmental regulatory agency or listed in the US pharmacopoeia or another generally recognized pharmacopoeia for use in animals, and more particularly in humans. When applied to salts the term “pharmaceutically acceptable” encompasses salts with a pharmaceutically acceptable acid or base. Pharmaceutically acceptable acids include inorganic acids, such
15 as hydrochloric, sulphuric, phosphoric, diphosphoric, hydrobromic, hydroiodic and nitrate acids, and organic acids, such as citric, maleic, malic, mandelic, ascorbic, oxalic, succinic, tartaric, acetic, methanesulfonic, ethanesulfonic, benzenesulfonic and p-toluenesulfonic acids. Pharmaceutically acceptable bases include hydroxides of alkali metals (e.g. sodium or potassium), alkaline-earth metals (for example, calcium or
20 magnesium) and organic bases (for example, alkylamines, arylalkylamines and heterocyclic amines).

 Other preferred salts according to the invention are quaternary ammonium compounds in which an equivalent of an anion (X^-) is associated with the positive charge of the N atom. X^- may be an anion of diverse mineral acids such as for example,
25 chloride, bromide, iodide, sulfate, nitrate, phosphate, or an anion of an organic acid, such as acetate, maleate, fumarate, citrate, oxalate, succinate, tartrate, malate, mandelate, trifluoroacetate, methanesulfonate and p-toluenesulfonate. X^- is preferably an anion selected from chloride, bromide, iodide, sulfate, nitrate, acetate, maleate, oxalate, succinate and trifluoroacetate. More preferably X^- is chloride, bromide, trifluoroacetate or
30 methanesulfonate.

The term “prevention”, as used herein, refers to the administration of the compounds of the invention in an initial or early stage of a disease, or to also prevent its onset.

The term “treatment” is used to designate the administration of the compounds of the invention to control disorder progression before or after the clinical signs had appeared. By control of the disorder progression it is meant to designate beneficial or desired clinical results including, but not limited to, reduction of symptoms, reduction of the length of the disorder, stabilization pathological state (specifically avoidance of further deterioration), delay in the disorder’s progression, improvement of the pathological state and remission (both partial and total). In a particular embodiment of the invention the compounds of the invention are used to control the disorder progression once at least one of the disorder’s clinical signs has appeared.

The term “medicament”, as used herein, refers to a pharmaceutical composition comprising a compound of formula (I). The medicament may be administered by any suitable route. It is prepared by conventional means with pharmaceutically acceptable excipients.

The term “subject”, as used herein, refers to any animal or human that is suffering from one of the diseases disclosed above. Preferably, the subject is a mammal. The term “mammal”, as used herein, refers to any mammalian species, including but not being limited to domestic and farm animals (cows, horses, pigs, sheep, goats, dogs, cats or rodents), primates, and humans. Preferably, the mammal is selected from a human being. In the context of the present invention, the mammal is suffering from a disease selected from the group consisting of obesity, insulin resistance, metabolic syndrome, cognitive dysfunction associated to Alzheimer’s disease, HIV-associated dementia, Creutzfeld-Jakob disease, Parkinson’s disease, Huntington’s disease, vascular inflammation, atherosclerosis, diabetes type 2, cardiovascular disease, glaucoma, osteoporosis, myocardial infarction and alcohol abuse treatment, or in risk of suffering from one of said diseases.

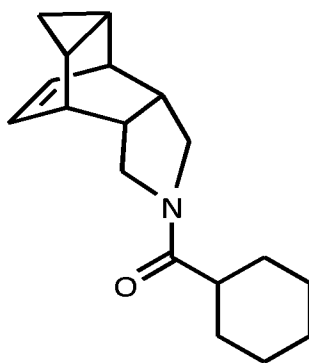
The invention is described below by means of several examples which do not limit, but rather illustrate the invention.

Examples

General

Reagents, solvents and starting products were acquired from commercial sources. The term "concentration" refers to the vacuum evaporation using a Büchi rotavapor. When indicated, the reaction products were purified by "flash" chromatography on silica gel (35-70 μm) with the indicated solvent system. The melting points were measured in a Büchi B-540 or in a MFB 59510M Gallenkamp instruments. IR spectra were run on FTIR Perkin-Elmer Spectrum RX I spectrophotometer using potassium bromide (KBr) pellets or attenuated total reflectance (ATR) technique. Absorption values are expressed as wavenumbers (cm^{-1}); only significant absorption bands are given. The accurate mass analyses were carried out using a LC/MSD-TOF spectrophotometer. The elemental analyses were carried out in a Flash 1112 series Thermofinnigan elemental microanalyzer (A5) to determine C, H, and N.

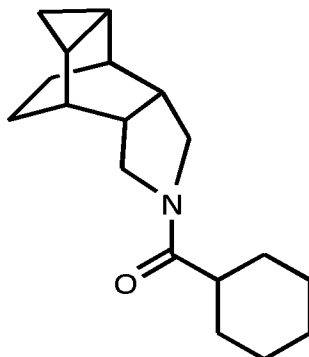
Example 1: cyclohexyl((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-etheno-cyclopropa[f]isoindol-2(1H)-yl)methanone



To a solution of 4-azatetracyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (2.057 g, 10.63 mmol) in EtOAc (150 mL) were added cyclohexanecarboxylic acid (1.239 g, 9.67 mmol), HOBt (1.9585 g, 14.50 mmol), EDC (2.2456 g, 14.50 mmol) and triethylamine (5.9 mL, 42.53 mmol). The reaction mixture was stirred at room temperature overnight. To the resulting suspension was then added water (150 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO_3 solution (150 mL) and brine (150 mL), dried over anhydrous Na_2SO_4 and filtered. Evaporation in *vacuo* of the organics gave the title compound as a yellowish solid (2.4346 g, 94.3% yield). The analytical sample was obtained by crystallization from hot EtOAc (2.0392 g), m.p. 96-97 °C. IR (ATR) ν : 560, 570, 587, 696, 718, 741, 767, 812, 829, 847, 894,

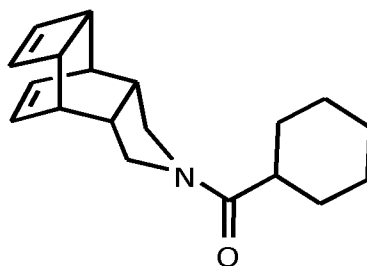
915, 942, 963, 991, 1036, 1089, 1134, 1167, 1209, 1217, 1242, 1272, 1299, 1361, 1380, 1432, 1624, 2849, 2925, 3002, 3040 cm^{-1} . Elemental analysis: Calculated for $\text{C}_{18}\text{H}_{25}\text{NO}$: C 79.66%, H 9.29%, N 5.16%. Found: C 79.64%, H 9.24%, N 5.21%.

Example 2: cyclohexyl((4R,4aR,5aS,6S,6aS)-octahydro-4,6-ethanocyclopropa[f]iso-
5 indol-2(1H)-yl)methanone



A solution of the compound from example 1 (500 mg) and Pd/C (100 mg) in absolute EtOH (85 mL) was hydrogenated at 1 atm for 72 hours. The dark solution was filtered and concentrated in vacuo to give the title compound as a white solid (426 mg, 82.6%
10 yield), m. p. 74-75 °C. IR (ATR) ν : 651, 679, 707, 729, 748, 789, 805, 826, 839, 865, 885, 950, 961, 988, 1016, 1030, 1082, 1113, 1133, 1171, 1205, 1213, 1234, 1264, 1295, 1325, 1346, 1358, 1427, 1471, 1486, 1623, 2846, 2897, 2928, 3009, cm^{-1} . Elemental analysis: Calculated for $\text{C}_{18}\text{H}_{27}\text{NO}$: C 79.07%, H 9.95%, N 5.12%. Found: C 79.15%, H 9.88%, N 5.29%.

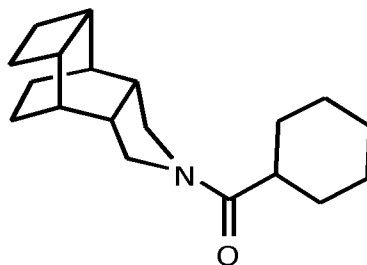
Example 3: cyclohexyl((4R,4aR,6aS,7S,7aS)-1,3,3a,4,4a,6a,7,7a-octahydro-2H-4,7-
15 ethenocyclobuta[f]isoindol-2-yl)methanone



To a solution of 4-azatetracyclo[5.4.2.0^{2,6}.0^{8,11}]trideca-9,12-diene hydrochloride (139 mg, 0.66 mmol) in EtOAc (6 mL) were added cyclohexanecarboxylic acid (77 mg, 0.60
20 mmol), HOBt (122 mg, 0.90 mmol), EDC (139 mg, 0.90 mmol) and triethylamine (0.4 mL, 2.64 mmol). The reaction mixture was stirred at room temperature overnight. To

the resulting suspension was then added water (10 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (10 mL) and brine (10 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave the title compound as a yellowish solid (151 mg, 80.0% yield). The analytical sample was obtained by crystallization from hot EtOAc (97 mg), m. p. 120-121 °C. IR (ATR) ν : 730, 760 788, 808, 967, 994, 1173, 1187, 1214, 1235, 1294, 1361, 1442, 1463, 1617, 2860, 2901, 2922 cm⁻¹. Elemental analysis: Calculated for C₁₉H₂₅NO: C 80.52%, H 8.89%, N 4.94%. Found: C 80.31%, H 8.81%, N 4.97%.

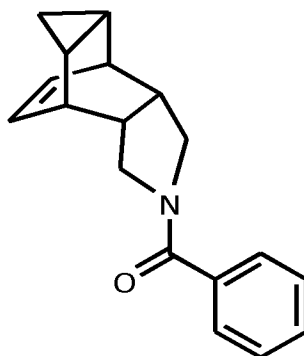
Example 4: cyclohexyl((4R,4aR,6aS,7S,7aS)-decahydro-2H-4,7-ethanocyclobuta[f]isoindol-2-yl)methanone



To a solution of 4-azatetracyclo[5.4.2.0^{2,6}.0^{8,11}]tridecane hydrochloride (235 mg, 1.10 mmol) in EtOAc (10 mL) were added cyclohexanecarboxylic acid (128 mg, 1.0 mmol), HOBt (203 mg, 1.50 mmol), EDC (232 mg, 1.50 mmol) and triethylamine (0.6 mL, 4.40 mmol). The reaction mixture was stirred at room temperature overnight. To the resulting suspension was then added water (10 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (10 mL) and brine (10 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave the title compound as a yellowish solid (253 mg, 80.0% yield). The analytical sample was obtained by crystallization from hot EtOAc (110 mg), m. p. 115-116 °C. IR (ATR) ν : 658, 731, 887, 988, 1133, 1172, 1204, 1236, 1357, 1434, 1440, 1621, 2908, 2925 cm⁻¹. Elemental analysis: Calculated for C₁₉H₂₉NO: C 79.39%, H 10.17%, N 4.87%. Found: C 79.20%, H 10.30%, N 4.72%.

Example 5: ((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)(phenyl)methanone

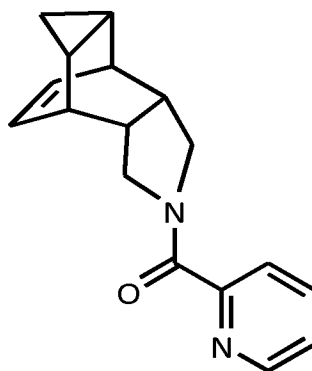
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To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (400 mg, 2.07 mmol) in EtOAc (20 mL) were added benzoic acid (230 mg, 1.88 mmol), HOBt (381 mg, 2.82 mmol), EDC (437 g, 2.82mmol) and triethylamine (1.2 mL, 8.27 mmol).

- 5 The reaction mixture was stirred at room temperature overnight. To the resulting suspension was then added water (20 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (20 mL) and brine (20 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave an orange oil (479 mg, 95.9% yield). Column chromatography (Hexane/Ethyl acetate
- 10 mixture) gave the title compound as a white solid (385 mg), m. p. 65-66 °C. IR (ATR) v: 660, 700, 715, 763, 794, 814, 847, 986, 1029, 1135, 1170, 1231, 1378, 1423, 1572, 1618, 2845, 2865, 2921, 2946 cm⁻¹. Elemental analysis: Calculated for C₁₈H₁₉NO: C 81.47%, H 7.22%, N 5.28%. Found: C 81.52%, H 7.34%, N 5.25%.

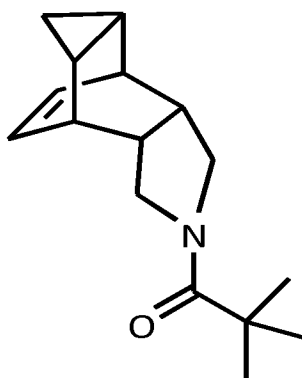
- Example 6:** ((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)(pyridin-2-yl)methanone
- 15



To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (400 mg, 2.07 mmol) in EtOAc (20 mL) were added piridin-2-carboxylic acid (231 mg, 1.88 mmol), HOBt (381 mg, 2.82 mmol), EDC (437 mg, 2.82 mmol) and triethylamine (1.2

mL, 8.27 mmol). The reaction mixture was stirred at room temperature overnight. To the resulting suspension was then added water (20 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (20 mL) and brine (20 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave the title compound as an orange oil (466 mg, 93.1% yield). Column chromatography (Hexane/Ethyl acetate mixture) gave an analytical sample as a white solid (326 mg), m. p. 110-111 °C. IR (ATR) ν : 682, 720, 753, 796, 814, 844, 912, 988, 1041, 1082, 1142, 1165, 1201, 1226, 1269, 1294, 1302, 1340, 1378, 1400, 1441, 1474, 1562, 1585, 1618, 2850, 2870, 2926, 3007, 3048 cm⁻¹. Elemental analysis: Calculated for C₁₇H₁₈N₂O: C 76.66%, H 6.81%, N 10.52%. Found: C 76.47%, H 7.01%, N 10.21%.

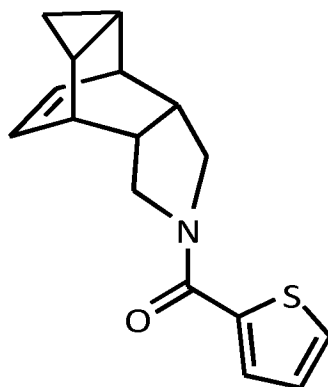
Example 7: 2,2-dimethyl-1-((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)propan-1-one



To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (200 mg, 1.03 mmol) in EtOAc (10 mL) were added pivalic acid (105 mg, 0.94 mmol), HOBt (190mg, 1.41 mmol), EDC (218 mg, 1.41 mmol) and triethylamine (0.6 mL, 4.14 mmol). The reaction mixture was stirred at room temperature overnight. To the resulting suspension was then added water (10 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (10 mL) and brine (10 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave the title compound as a yellowish solid (216 mg, 93.5% yield). The analytical sample was obtained by crystallization from hot EtOAc (69 mg), m. p. 91-92 °C. IR (ATR) ν : 720, 756, 766, 809, 829, 849, 912, 943, 988, 1036, 1069, 1094, 1165, 1193, 1239, 1274, 1340, 1362, 1380, 1405, 1461, 1476, 1507, 1610, 2870, 2896, 2936, 2951, 2992 cm⁻¹.

Elemental analysis: Calculated for C₁₆H₂₃NO: C 78.32%, H 9.45%, N 5.71%. Found: C 78.16%, H 9.52%, N 5.86%.

Example 8: ((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)(thiophen-2-yl)methanone



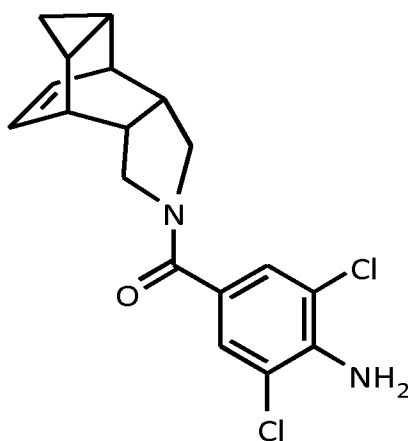
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To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (200 mg, 1.03 mmol) in EtOAc (10 mL) were added 2-thiophenecarboxylic acid (121 mg, 0.94 mmol), HOBt (190mg, 1.41 mmol), EDC (218 mg, 1.41 mmol) and triethylamine (0.6 mL, 4.14 mmol). The reaction mixture was stirred at room temperature overnight. To the resulting suspension was then added water (10 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (10 mL) and brine (10 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave the title compound as a yellowish solid (219 mg, 85.7% yield). The analytical sample was obtained by crystallization from hot EtOAc (87 mg), m. p. 104-105 °C. IR (ATR) ν : 667, 703, 720, 738, 786, 814, 849, 890, 915, 950, 1008, 1031, 1057, 1087, 1132, 1239, 1254, 1279, 1312, 1352, 1380, 1403, 1431, 1519, 1580, 1598, 2921, 2936, 3002, 3037 cm⁻¹. Elemental analysis: Calculated for C₁₆H₁₇NOS: C 70.81%, H 6.31%, N 5.16%. Found: C 70.70%, H 6.28%, N 5.12%.

Example 9: (4-amino-3,5-dichlorophenyl)((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)methanone

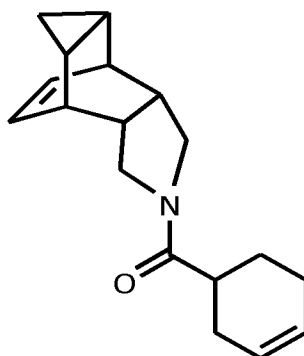
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To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (200 mg, 1.03 mmol) in EtOAc (10 mL) were added 3,5-dichloro-4-aminobenzoic acid (194 mg, 0.94 mmol), HOBt (190mg, 1.41 mmol), EDC (218 mg, 1.41 mmol) and triethylamine (0.6 mL, 4.14 mmol). The reaction mixture was stirred at room temperature overnight. To the resulting suspension was then added water (10 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (10 mL) and brine (10 mL), dried over anhydrous Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave the title compounds as a yellowish solid (322 mg, 89.3% yield). The analytical sample was obtained by crystallization from hot EtOAc (89 mg), m. p. 186-187 °C. IR (ATR) ν : 680, 718, 743, 763, 783, 809, 844, 864, 892, 915, 955, 991, 1034, 1097, 1173, 1223, 1246, 1297, 1347, 1416, 1469, 1501, 1537, 1595, 2875, 2921, 3194, 3240, 3301, 3458 cm⁻¹. Elemental analysis: Calculated for C₁₈H₁₈Cl₂N₂O: C 71.90%, H 5.20%, N 8.02%. Found: C 72.10%, H 5.20%, N 7.92%

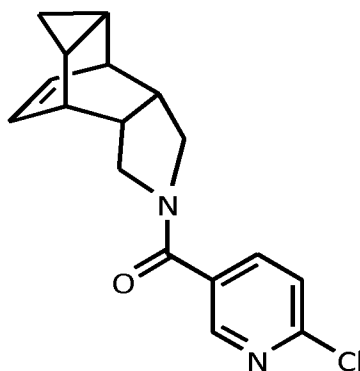
Example 10: cyclohex-3-en-1-yl((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)methanone



To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (200 mg, 1.03 mmol) in EtOAc (10 mL) were added 3-cyclohexene carboxylic acid (119 mg, 0.94 mmol), HOBt (190mg, 1.41 mmol), EDC (218 mg, 1.41 mmol) and triethylamine (0.6 mL, 4.14 mmol). The reaction mixture was stirred at room temperature overnight.

- 5 To the resulting suspension was then added water (10 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (10 mL) and brine (10 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave the title compound as a yellowish solid (259 mg, quantitative yield). Column chromatography (Hexane/Ethyl acetate mixture) gave the compound as a white
10 solid (199 mg), m. p. 78-79 °C. IR (ATR) ν : 682, 710, 763, 816, 839, 854, 887, 915, 940, 981, 1016, 1034, 1087, 1135, 1168, 1203, 1221, 1274, 1292, 1332, 1355, 1380, 1431, 1620, 1651, 2870, 2926, 3022 cm⁻¹. Elemental analysis: Calculated for C₁₈H₂₃NO: C 80.26%, H 8.61%, N 5.20%. Found: C 80.24%, H 8.73%, N 5.19%.

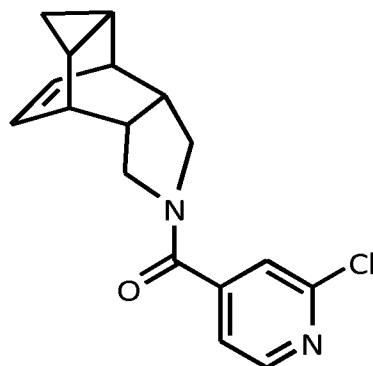
- Example 11:** (6-chloropyridin-3-yl)((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-
15 octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)methanone



- To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (500 mg, 2.58 mmol) in EtOAc (25 mL) were added 6-choloronicotinic acid (370 mg, 2.35 mmol), HOBt (477 mg, 3.53 mmol), EDC (547 mg, 3.53 mmol) and triethylamine (1.4
20 mL, 10.34 mmol). The reaction mixture was stirred at room temperature overnight. To the resulting suspension was then added water (25 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (25 mL) and brine (25 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave the title compound as a yellowish solid (664 mg, 85.6% yield). Column
25 chromatography (Hexane/Ethyl acetate mixture) gave the title compound as a white solid (457 mg), m. p. 101-102 °C. IR (ATR) ν : 712, 736, 759, 793, 814, 835, 924, 940,

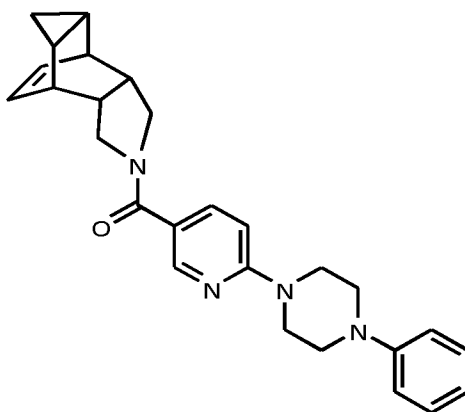
985, 1030, 1097, 1129, 1156, 1174, 1215, 1239, 1251, 1271, 1283, 1350, 1372, 1430, 1455, 1563, 1583, 1612, 2914, 2948, 3002 cm^{-1} . Elemental analysis: Calculated for $\text{C}_{17}\text{H}_{17}\text{ClN}_2\text{O}$: C 67.88%, H 5.70%, N 9.31%. Found: C 68.14%, H 5.84%, N 9.00%.

Example 12: (2-chloropyridin-4-yl)((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)methanone



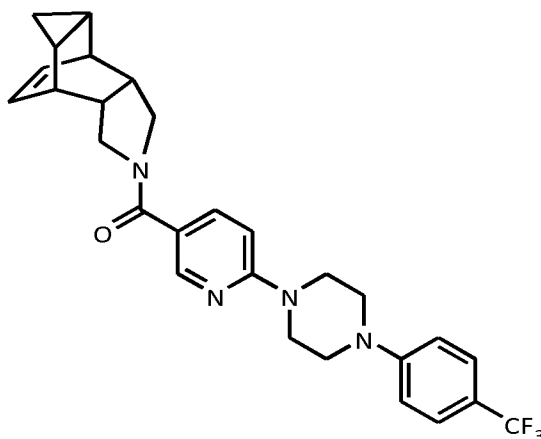
To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (500 mg, 2.58 mmol) in EtOAc (25 mL) were added 6-chloronicotinic acid (370 mg, 2.35 mmol), HOBt (477 mg, 3.53 mmol), EDC (547 mg, 3.53 mmol) and triethylamine (1.4 mL, 10.34 mmol). The reaction mixture was stirred at room temperature overnight. To the resulting suspension was then added water (25 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO_3 solution (25 mL) and brine (25 mL), dried over anhydrous Na_2SO_4 and filtered. Evaporation in *vacuo* of the organics gave the title compound as a yellowish solid (614 mg, 79.1% yield). Column chromatography (Hexane/Ethyl acetate mixture) gave the compound as a white solid (445 mg), m. p. 135-136 °C. IR (ATR) ν : 669, 708, 720, 741, 753, 771, 817, 844, 915, 942, 987, 1041, 1091, 1118, 1163, 1176, 1203, 1232, 1245, 1269, 1287, 1342, 1373, 1437, 1464, 1476, 1530, 1593, 1632, 2868, 2932, 3003, 3057 cm^{-1} . Elemental analysis: Calculated for $\text{C}_{17}\text{H}_{17}\text{ClN}_2\text{O}$: C 67.88%, H 5.70%, N 9.31%. Found: C 67.98%, H 5.77%, N 9.09%.

Example 13: ((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)(6-(4-phenylpiperazin-1-yl)pyridin-3-yl)methanone



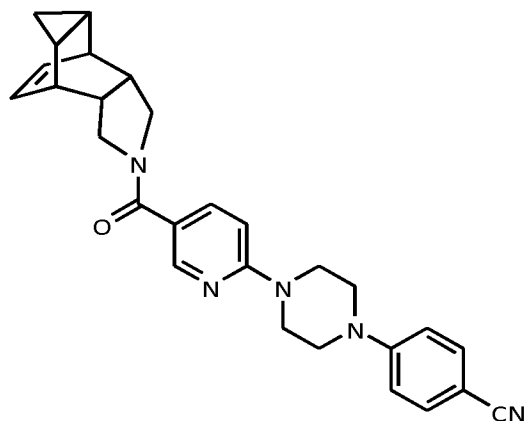
To a solution of the compound from example 11 (100 mg, 0.33 mmol) and 1-phenylpiperazine (60 mg, 0.37 mmol) in DMF (0.5 mL) was added solid K_2CO_3 (82 mg, 0.59 mmol). The resulting suspension was stirred at 90 °C for 48 hours. Water (5 mL) and DCM (5 mL) were added and the phases were separated. The aqueous phase was then extracted with further DCM (2 x 5 mL). The organics were dried over anhydrous Na_2SO_4 , filtered and evaporated in *vacuo* to give a yellowish solid (137 mg). Column chromatography (Hexane/Ethyl acetate mixture) gave the title compound as a white solid (56 mg, 39.4% yield). The analytical sample was obtained by washing with cooled pentane (45 mg), m. p. 90-91 °C. IR (ATR) ν : 661, 695, 739, 754, 814, 822, 845, 948, 987, 1013, 1028, 1041, 1095, 1152, 1227, 1310, 1349, 1349, 1395, 1413, 1491, 1594, 1617, 2847, 2919, 2997 cm^{-1} . Accurate mass: Calculated for $[C_{27}H_{31}N_4O+H]^+$: 427.2494. Found: 427.2492.

Example 14: ((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)(6-(4-(4-(trifluoromethyl)phenyl)piperazin-1-yl)pyridin-3-yl)methanone



To a solution of the compound from example 11 (100 mg, 0.33 mmol) and 1-(4-trifluoromethylphenyl)piperazine (85 mg, 0.37 mmol) in DMF (0.5 mL) was added solid K_2CO_3 (82 mg, 0.59 mmol). The resulting suspension was stirred at 90 °C for 48 hours. Water (5 mL) and DCM (5 mL) were added and the phases were separated. The aqueous phase was then extracted with further DCM (2 x 5 mL). The organics were dried over anh. Na_2SO_4 , filtered and evaporated in *vacuo* to give a yellowish solid (161 mg). Column chromatography (Hexane/Ethyl acetate mixture) gave the title compound as a white solid (52 mg, 31.9% yield). The analytical sample was obtained by washing with cooled pentane (38 mg), m. p. 157-158 °C. IR (ATR) ν : 667, 711, 721, 744, 770, 806, 824, 909, 951, 971, 984, 139, 1070, 1106, 1157, 1199, 1230, 1330, 1354, 1390, 1429, 1493, 1522, 1594, 1615, 2847, 2919 cm^{-1} . Accurate mass: Calculated for $[C_{28}H_{29}F_2N_4O+H]^+$: 475.2294. Found: 475.2366.

Example 15: 4-(4-(5-((4R,4aR,5aS,6S,6aS)-1,2,3,3a,4,4a,5,5a,6,6a-decahydro-4,6-ethenocyclopropa[f]isoindole-2-carbonyl)pyridin-2-yl)piperazin-1-yl)benzonitrile

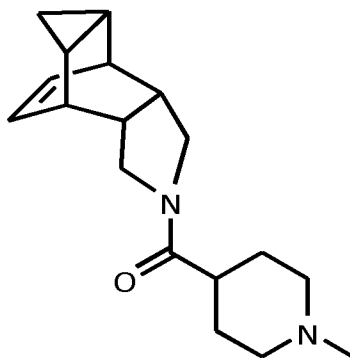


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To a solution of the compound from example 11 (100 mg, 0.33 mmol) and 4-piperazinobenzonitrile (69 mg, 0.37 mmol) in DMF (0.5 mL) was added solid K_2CO_3 (82 mg, 0.59 mmol). The resulting suspension was stirred at 90 °C for 48 hours. Water (5 mL) and DCM (5 mL) were added and the phases were separated. The aqueous phase was then extracted with further DCM (2 x 5 mL). The organics were dried over anh. Na_2SO_4 , filtered and evaporated in *vacuo* to give a yellowish solid (181 mg). Column chromatography (Hexane/Ethyl acetate mixture) gave the title compound as a white solid (72 mg, 48.3% yield), m. p. 160-161 °C. IR (ATR) ν : 656, 692, 713, 742, 773, 811, 912, 951, 1008, 1039, 1176, 1235, 1312, 1392, 1426, 1511, 1537, 1555, 1599,

1648, 1666, 2847, 2925 cm^{-1} . Accurate mass: Calculated for $[\text{C}_{28}\text{H}_{30}\text{N}_5\text{O}+\text{H}]^+$: 452.2444. Found: 452.2445.

Example 16: (1-methylpiperidin-4-yl)((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)methanone



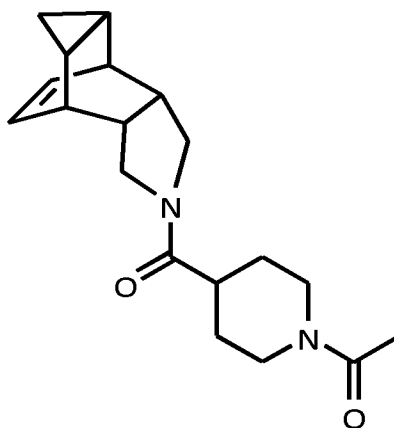
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To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (200 mg, 1.03 mmol) in EtOAc (10 mL) were added 1-methylpiperidine-4-carboxylic acid (135 mg, 0.94 mmol), HOBt (190mg, 1.41 mmol), EDC (218 mg, 1.41 mmol) and triethylamine (0.6 mL, 4.14 mmol). The reaction mixture was stirred at room temperature for 24 h. To the resulting suspension was then added water (10 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO_3 solution (10 mL) and brine (10 mL), dried over anh. Na_2SO_4 and filtered. Evaporation in *vacuo* of the organics gave a mixture of the product and starting materials as a yellowish solid (129 mg). Column chromatography (Dichloromethane/Methanol) gave the title compound as a white solid (86 mg, 32% yield). The analytical sample was obtained by crystallization from hot EtOAc (50 mg), m. p. 100-101 °C. IR (ATR) ν : 718, 767, 819, 835, 850, 876, 915, 987, 1013, 1041, 1067, 1090, 1129, 1150, 1191, 1214, 1250, 1276, 1305, 1359, 1374, 1431, 1447, 1625, 2780, 2857, 2914, 2940, 3328 cm^{-1} . Accurate mass: Calculated for $[\text{C}_{18}\text{H}_{26}\text{N}_2\text{O}+\text{H}]^+$: 287.2118. Found: 287.2113.

Example 17: 1-(4-((4R,4aR,5aS,6S,6aS)-1,2,3,3a,4,4a,5,5a,6,6a-decahydro-4,6-ethenocyclopropa[f]isoindole-2-carbonyl)piperidin-1-yl)ethan-1-one

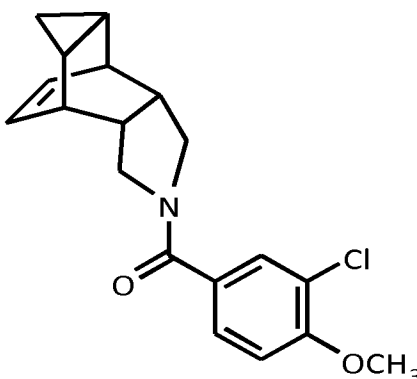
20

44



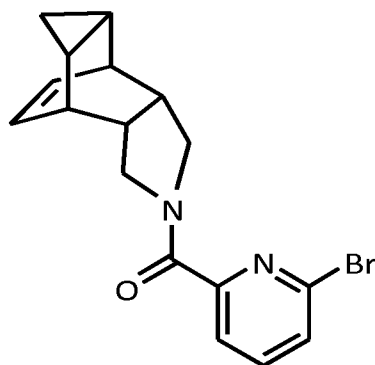
- To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (200 mg, 1.03 mmol) in EtOAc (10 mL) were added 1-acetyl-4-piperidinecarboxylic acid (161 mg, 0.94 mmol), HOBt (190 mg, 1.41 mmol), EDC (218 mg, 1.41 mmol) and triethylamine (0.6 mL, 4.14 mmol). The reaction mixture was stirred at room temperature for 24 h. To the resulting suspension was then added water (10 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (10 mL) and brine (10 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave a mixture of the product and starting materials (182 mg).
- Column chromatography (Hexane/Ethyl acetate) gave the title compound as a white solid (134 mg, 45% yield), m. p. 134-135 °C. IR (ATR) ν : 605, 703, 762, 814, 829, 920, 956, 977, 997, 1041, 1098, 1116, 1168, 1222, 1271, 1307, 1356, 1426, 1620, 1640, 2852, 2925, 2992 cm⁻¹. Elemental analysis: Calculated for C₁₇H₁₇BrN₂O: C 72.58%, H 8.34%, N 8.91%. Found: C 72.75%, H 8.66%, N 8.50%.

- Example 18:** (3-chloro-4-methoxyphenyl)((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)methanone



To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (200 mg, 1.03 mmol) in EtOAc (10 mL) were added 3-chloro-4-methoxybenzoic acid (176 mg, 0.94 mmol), HOBt (190 mg, 1.41 mmol), EDC (218 mg, 1.41 mmol) and triethylamine (0.6 mL, 4.14 mmol). The reaction mixture was stirred at room temperature for 24 h. To the resulting suspension was then added water (10 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (10 mL) and brine (10 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave the title compound as a yellow semisolid (329 mg, quantitative yield). The analytical sample was obtained by column chromatography (Hexane/Ethyl acetate) to obtain the title compound as a waxy solid (164 mg). IR (ATR) ν : 615, 651, 692, 708, 754, 814, 835, 850, 891, 915, 946, 1018, 1059, 1095, 1142, 1183, 1232, 1256, 1294, 1349, 1387, 1418, 1501, 1560, 1599, 1617, 2868, 2925, 3002 cm⁻¹. Accurate mass: Calculated for [C₁₉H₂₀ClNO₂+H]⁺: 300.1255. Found: 300.1251.

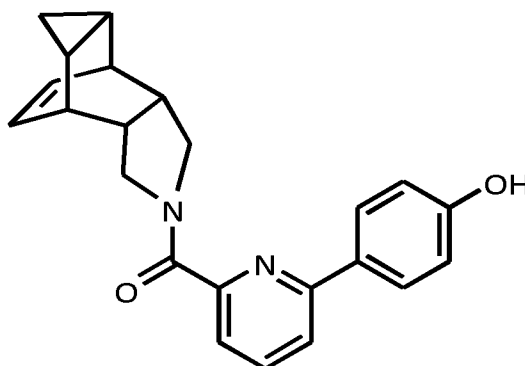
Example 19: (6-bromopyridin-2-yl)((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)methanone



To a solution of 4-azapentacyclo[5.3.2.0^{2,6}.0^{8,10}]dodec-11-ene hydrochloride (300 mg, 1.55 mmol) in EtOAc (15 mL) were added 6-bromopyridine-2-carboxylic acid (285 mg, 1.41 mmol), HOBt (286 mg, 1.41 mmol), EDC (328 mg, 1.41 mmol) and triethylamine (0.9 mL, 6.20 mmol). The round bottomed flask was wrapped with aluminium foil and the reaction mixture was stirred at room temperature for 24 h. To the resulting suspension was then added water (15 mL) and the phases were separated. The organic phase was washed with saturated aqueous NaHCO₃ solution (15 mL) and brine (15 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave the title compound as a yellowish solid (323 mg, 66.3 % yield). The analytical sample was obtained by crystallization from hot EtOAc (209 mg), m. p. 185-186 °C. IR (ATR) ν :

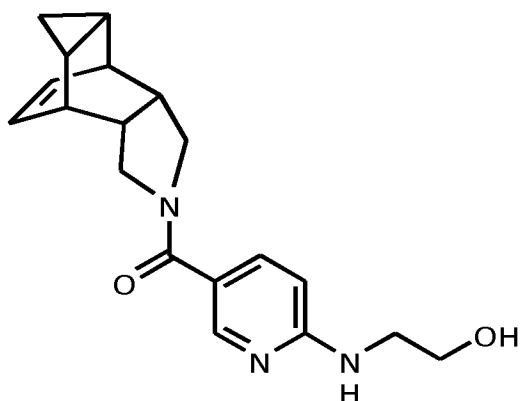
643, 659, 705, 734, 762, 814, 827, 850, 912, 935, 984, 1044, 1080, 1119, 1165, 1199, 1225, 1245, 1274, 1305, 1341, 1392, 1405, 1454, 1545, 1576, 1617, 2857, 2925, 3002, 3043, 3059 cm^{-1} . Elemental analysis: Calculated for $\text{C}_{17}\text{H}_{17}\text{BrN}_2\text{O}$: C 59.14%, H 4.96%, N 8.11%. Found: C 59.31%, H 4.92%, N 7.87%.

- 5 **Example 20:** (6-(4-hydroxyphenyl)pyridin-2-yl)((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)methanone



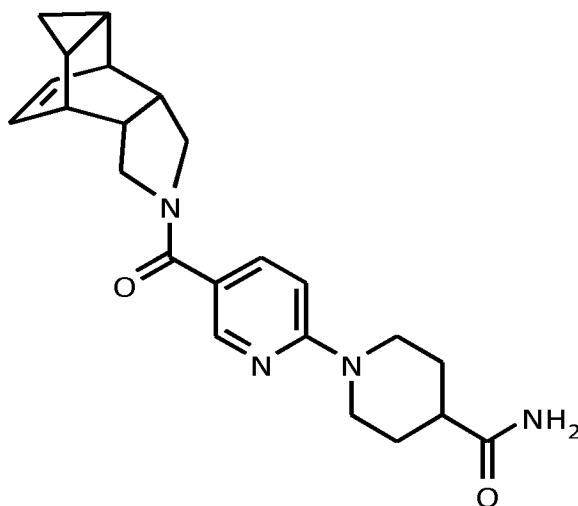
- A mixture of the compound obtained in Example 19 (300 mg, 0.87 mmol), 4-hydroxyphenylboronic acid (132 mg, 0.96 mmol), $\text{Pd}(\text{Ph}_3)_4$ (10 mg, 0.009 mmol) and
- 10 K_2CO_3 (240 mg, 1.74 mmol) in dioxane (3 mL) and water (1.5 mL) was heated at 100°C for 2 h. Then EtOAc (10 mL) was added and the organic phase was washed with water (10 mL), dried over anhydrous Na_2SO_4 and filtered. Evaporation in *vacuo* of the organics gave a mixture of the product and starting materials (461 mg). Column chromatography (Hexane/Ethyl acetate) gave the title compound as a white solid (184
- 15 mg, 59.0 % yield), m. p. $221\text{--}222^\circ\text{C}$. IR (ATR) ν : 630, 654, 711, 754, 822, 840, 915, 946, 992, 1039, 1085, 1103, 1173, 1230, 1276, 1297, 1346, 1380, 1400, 1429, 1460, 1516, 1558, 1586, 1604, 2930, 3002, 3126 cm^{-1} . Accurate mass: Calculated for $[\text{C}_{23}\text{H}_{22}\text{N}_2\text{O}_2+\text{H}]^+$: 359.1754. Found: 359.1750.

- Example 21:** (6-((2-hydroxyethyl)amino)pyridin-3-yl)((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)methanone
- 20



A solution of the compound obtained in example 11 (100 mg, 0.33 mmol) and ethanolamine (0.36 mL, 6 mmol) was stirred at 120 °C for 24 hours. Water (10 mL) and EtOAc (10 mL) were added and the phases were separated. The organic phase was dried
 5 over anhydrous Na_2SO_4 , filtered and evaporated in *vacuo* to give the title compound as a yellowish solid (65 mg, 60.7% yield). The analytical sample was obtained by crystallization from hot EtOAc (59 mg), m. p. 87-88 °C. IR (ATR) ν : 636, 716, 731, 767, 811, 832, 909, 943, 984, 1010, 1044, 1070, 1150, 1232, 1274, 1302, 1343, 1423, 1460, 1488, 1527, 1591, 1612, 2868, 2914, 3002, 3064, 3136, 3271 cm^{-1} . Accurate
 10 mass: Calculated for $[\text{C}_{19}\text{H}_{23}\text{N}_3\text{O}_2+\text{H}]^+$: 326.1863. Found: 326.1867.

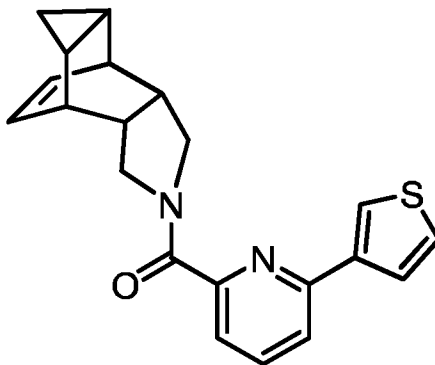
Example 22: 1-(5-((4R,4aR,5aS,6S,6aS)-1,2,3,3a,4,4a,5,5a,6,6a-decahydro-4,6-ethenocyclopropa[f]isoindole-2-carbonyl)pyridin-2-yl)piperidine-4-carboxamide



To a solution of the compound obtained in example 11 (100 mg, 0.33 mmol) and
 15 isonipecotamide (85 mg, 0.66 mmol) in DMF (0.5 mL) was added solid K_2CO_3 (82 mg, 0.59 mmol). The resulting suspension was stirred at 90 °C for 48 hours. Water (5 mL)

and DCM (5 mL) were added and the phases were separated. The aqueous phase was then extracted with further DCM (2 x 5 mL). The organics were dried over anh. Na₂SO₄, filtered and evaporated in *vacuo* to give the title compound as a yellowish solid (110 mg, 84.6% yield), m. p. 162-163 °C. IR (ATR) ν : 630, 692, 723, 773, 811, 845, 943, 982, 1010, 1028, 1041, 1095, 1129, 1178, 1219, 1238, 1312, 1351, 1367, 1408, 1431, 1501, 1540, 1584, 1599, 1687, 1736, 2919, 3152, 3307 cm⁻¹. Accurate mass: Calculated for [C₂₃H₂₈N₄O₂+H]⁺: 393.2285. Found: 393.2285.

Example 23: ((4R,4aR,5aS,6S,6aS)-3,3a,4,4a,5,5a,6,6a-octahydro-4,6-ethenocyclopropa[f]isoindol-2(1H)-yl)(6-(thiophen-3-yl)pyridin-2-yl)methanone

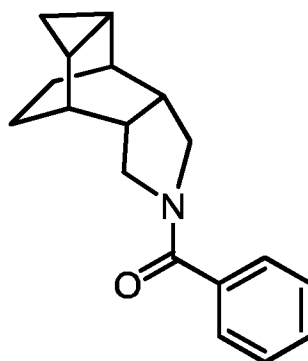


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A mixture of the compound obtained in Example 19 (230 mg, 0.67 mmol), 3-thiopheneboronic acid (93 mg, 0.73 mmol), Pd(Ph₃)₄ (8 mg, 0.007 mmol) and K₂CO₃ (185 mg, 1.34 mmol) in dioxane (2.3 mL) and water (1.3 mL) was heated at 100° C for 2 h. Then EtOAc (10 mL) was added and the organic phase was washed with water (10 mL), dried over anh. Na₂SO₄ and filtered. Evaporation in *vacuo* of the organics gave a mixture of the product and starting materials (238 mg). Column chromatography (Hexane/Ethyl acetate) gave the title compound as a yellowish foamy solid (184 mg, 79.0 % yield), m. p. 47-48 °C. IR (ATR) ν : 615, 646, 703, 716, 752, 791, 829, 845, 863, 915, 948, 987, 1036, 1095, 1178, 1235, 1274, 1343, 1418, 1431, 1460, 1566, 1581, 1617, 1971, 2000, 2051, 2175, 2325, 2868, 2919, 2997 cm⁻¹. Accurate mass: Calculated for [C₂₁H₂₀N₂OS+H]⁺: 349.1369. Found: 349.1374.

20

Example 24: ((4R,4aR,5aS,6S,6aS)-octahydro-4,6-ethanocyclopropa[f]isoindol-2(1H)-yl)(phenyl)methanone



A solution of the compound from example 5 (80 mg) and Pd/C (16 mg) in absolute EtOH (12 mL) was hydrogenated at 1 atm for 24 hours. The suspension was filtered and concentrated in vacuo to give the title compound as a white solid (79 mg, 98.0% yield),
5 m. p. 79-80 °C. IR (ATR) ν : 644, 685, 711, 729, 785, 845, 871, 935, 951, 1010, 1026, 1046, 1075, 1147, 1160, 1248, 1292, 1341, 1418, 1444, 1480, 1576, 1617, 1888, 1961, 2005, 2129, 2201, 2351, 2857, 2925, 2992 cm^{-1} . Accurate mass: Calculated for $[\text{C}_{18}\text{H}_{21}\text{NO}+\text{H}]^+$: 268.1696. Found: 268.1703.

Example 25: Pharmacological data of selected compounds

- 10 The inhibitory activity of the compounds of the invention for the enzymes 11 β -HSD1 and 11 β -HSD2 may be tested with the following protocols:

Determination of the 11 β -HSD1 inhibition in Human Liver Microsomes

- 11 β -HSD1 activity was determined in mixed sex, Human Liver Microsomes (Celsis In-vitro Technologies) by measuring the conversion of ^3H -cortisone to ^3H -cortisol.
- 15 Percentage inhibition was determined relative to a no inhibitor control. 5 μg of Human Liver microsomes was pre-incubated at 37°C for 10 minutes with inhibitor (assay concentrations of 1, 0.5, 0.1, 0.05, 0.01, 0.005, 0.001 and 0.0005 μM) and 1 mM NADPH in a final volume of 90 μL Krebs buffer. 10 μL of 200 nM ^3H -cortisone was then added followed by incubation for at 37°C for a further 30 minutes. The assay was
- 20 terminated by rapid freezing on dry ice. ^3H -cortisone to ^3H -cortisol conversion was determined in 50 μL of the defrosted reaction by capturing liberated ^3H -cortisol on 100 μL 1:8000 dilution anti-cortisol (HyTest Ltd)-coated scintillation proximity assay beads (protein A-coated YSi, GE Healthcare). The percentages of inhibition at every tested

concentration were analyzed using a non-linear regression curve fitting (variable slope) by using the software GraphPad Prism 5.0.

Determination of 11 β -HSD1 inhibition in HEK293 cells stably transfected with the 11 β -HSD1 gene

- 5 11 β -HSD1 activity was determined in cells that are incubated with substrate (cortisone) and product (cortisol) is determined by LC/MS. Cells are plated at 2×10^4 cells/well in a 96-well tissue culture microplate and incubated overnight at @37⁰ C in 5% CO₂ 95% O₂. Compounds to be tested are solubilized in 100% DMSO at 10 mM and serially diluted in water and 10% DMSO to final concentration 100 μ M in 10% DMSO. 10 μ l of
- 10 each test dilution is dispensed into the well of a 96-well microplate (assay concentration of 10 μ M). Low and high control wells contain 10 μ l of 10% DMSO. A solution of DMEM is prepared containing 1% penicillin, 1% streptomycin and 300 nM cortisone. Medium is removed from cell assay plate and DMEM solution (100 μ l) added to each well. Cells are incubated for 2 hours @37⁰ C in 5% CO₂ 95% O₂. Following incubation
- 15 medium is removed from each well into 500 μ l ethyl acetate mixed by vortex and incubated at room temperature for 5 minutes. A calibration curve of known concentrations of cortisol in assay medium is also set up and added to 500 μ l ethyl acetate vortexed and incubated as above. The supernatant is removed to a 96 deep well plate and dried down under liquid nitrogen at 65°C. Each well is solubilised in 100 μ l
- 20 70:30 H₂O:ACN and removed to a 96 well v bottomed plate for LC/MS analysis.

Determination of 11 β -HSD2 inhibition in HEK293 cells stably transfected with the 11 β -HSD2 gene

- 11 β -HSD2 activity was determined in cells that are incubated with substrate (cortisol) and product (cortisone) is determined by LC/MS. Cells are plated at 2×10^4 cells/well
- 25 in a 96-well tissue culture microplate and incubated overnight at @37⁰ C in 5% CO₂ 95% O₂. Compounds to be tested are solubilized in 100% DMSO at 10 mM and serially diluted in water and 10% DMSO to final concentration in 10% DMSO. 10 μ l of each test dilution is dispensed into the well of a 96-well microplate (assay concentrations of 10, 1, 0.1 μ M). Low and high control wells contain 10 μ l of 10% DMSO. A solution of
- 30 DMEM is prepared containing 1% penicillin, 1% streptomycin and 300 nM cortisone. Medium is removed from cell assay plate and DMEM solution (100 μ l) added to each well. Cells are incubated for 2 hours @37⁰ C in 5% CO₂ 95% O₂. Following incubation

medium is removed from each well into 500 μ l ethyl acetate mixed by vortex and incubated at room temperature for 5 minutes. A calibration curve of known concentrations of cortisone in assay medium is also set up and added to 500 μ l ethyl acetate vortexed and incubated as above. The supernatant is removed to a 96 deep well plate and dried down under liquid nitrogen at 65°C. Each well is solubilised in 100 μ l 70:30 H₂O:ACN and removed to a 96 well v bottomed plate for LC/MS analysis.

Following the above described protocols the inhibitory activity of compounds of the present invention has been reported and is shown in the following table 1:

10

TABLE 1

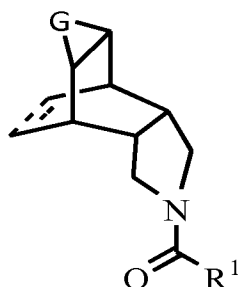
Example	HSD1		HSD2
	IC ₅₀ in HLM ^a	% inh in cells (10 μ M)	IC ₅₀ in cells ^b
1	A	100	C
2	A	100	A
7	C	85	C
9	A	77	B
10	A	41	B
15	B	100	C

^a In the column "HSD1 IC₅₀ in HLM" A means that IC₅₀ is lower than 50 nM, B means that IC₅₀ is at least 50 nM but less than 500 nM and C means that IC₅₀ is at least 500 nM but less than < 1000 nM.

15 ^b In the column "HSD2 IC₅₀ in cells" A means that IC₅₀ is lower than 100 nM, B means that IC₅₀ is at least 100 nM but less than 1000 nM and C means that IC₅₀ is at least 1000 nM.

CLAIMS

1. A compound of formula (I):



5 wherein

R¹ is selected from the group consisting of:

- a) C₃₋₈ cycloalkyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl;
- b) C₃₋₈ cycloalkenyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl;
- 10 c) C₆₋₁₀ aromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy;
- d) C₅₋₁₀ heteroaromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NHR³, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₅₋₁₀ heteroaryl (optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy), phenyl (optionally substituted with one or two groups selected from halogen atoms, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy) and C₅₋₇ heterocyclyl (optionally substituted with a group selected from -COR⁴, -COOR⁴, -CN, -CONHR⁴, -CONH₂ and phenyl which is optionally substituted with one or two groups selected from halogen atoms, CONH₂, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy), wherein the C₅₋₇ heterocyclyl group comprises a nitrogen atom forming part of the heterocycle and is attached to the C₅₋₁₀ heteroaromatic group through said nitrogen atom;
- 15 e) C₃₋₈ heterocyclyl optionally substituted with a group selected from C₁₋₄ alkyl, C₃₋₈ cycloalkyl and -COR⁵; and
- 20 f) -C(CH₃)₂-R²;
- 25

G is a rest selected from the group consisting of $-\text{CH}_2-$, $-\text{CH}_2\text{CH}_2-$ and $-\text{CH}=\text{CH}-$;

R^2 is selected from the group consisting of hydrogen atom and C_{1-3} alkyl

R^3 is selected from the group consisting of C_{1-3} alkyl and C_{1-3} hydroxyalkyl

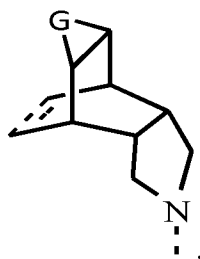
R^4 is selected from the group consisting of C_{1-6} alkyl and phenyl

5 R^5 is a group C_{1-6} alkyl

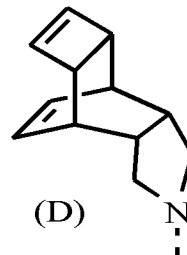
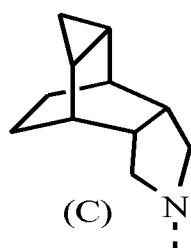
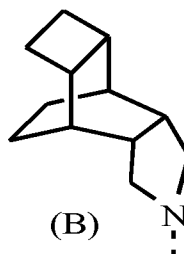
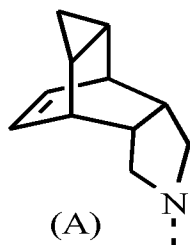
the bond represented by --- is either a single bond or a double bond;

and pharmaceutically stable salts thereof.

2. A compound according to claim 1, wherein the rest

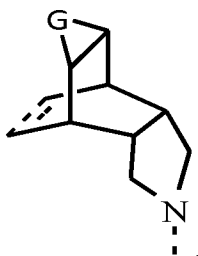


10 is selected from the group consisting of rests (A) to (D):

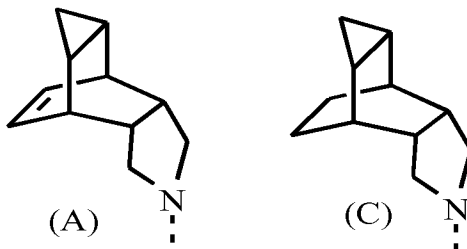


3. A compound according to claim 2, wherein the rest

54



is selected from the group consisting of rests (A) and (C):



- 5 4. A compound according to anyone of claims 1 to 3, wherein G is a rest -CH₂-.
5. A compound according to anyone of claims 1 to 4, wherein the bond represented by  is a double bond.
- 10 6. A compound according to anyone of claims 1 to 4, wherein R¹ is a C₃₋₈ cycloalkyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl.
7. A compound according to anyone of claims 1 to 4, wherein R¹ is a C₃₋₈ cycloalkenyl, optionally substituted with 1 to 3 groups selected from halogen atoms and C₁₋₄ alkyl.
- 15 8. A compound according to anyone of claims 1 to 4, wherein R¹ is a C₆₋₁₀ aromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy.
- 20 9. A compound according to anyone of claims 1 to 4, wherein R¹ is a C₅₋₁₀ heteroaromatic group optionally substituted with 1 to 3 groups selected from halogen atoms, CN, CF₃, NHR³, C₁₋₄ alkyl, C₁₋₄ alkoxy, C₅₋₁₀ heteroaryl (optionally substituted

- with 1 to 3 groups selected from halogen atoms, CN, CF₃, NH₂, OH, C₁₋₄ alkyl and C₁₋₄ alkoxy), phenyl (optionally substituted with one or two groups selected from halogen atoms, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy) and C₅₋₇ heterocyclyl (optionally substituted with a group selected from -COR⁴, -COOR⁴, -CN, -CONHR⁴, -CONH₂ and phenyl which is optionally substituted with one or two groups selected from halogen atoms, CONH₂, CN, OH, CF₃, C₁₋₄ alkyl and C₁₋₄ alkoxy), wherein the C₅₋₇ heterocyclyl group comprises a nitrogen atom forming part of the cycle and is attached to the C₅₋₁₀ heteroaromatic group through said nitrogen atom.
- 10 10. A compound according to anyone of claims 1 to 4, wherein R¹ is a C₃₋₈ heterocyclyl optionally substituted with a group selected from C₁₋₄ alkyl, C₃₋₈ cycloalkyl and COR⁵ wherein R⁵ is a group C₁₋₆ alkyl.
- 15 11. A compound according to anyone of claims 1 to 4, wherein R¹ is a group -C(CH₃)₂-R², wherein R² is selected from the group consisting of hydrogen atom and C₁₋₃ alkyl.
12. A compound according to anyone of claims 1 to 11 for use as a medicament.
- 20 13. A compound according to anyone of claims 1 to 11 for use in the treatment or prevention of a disease or conditions susceptible of improvement by inhibition of 11β-hydroxysteroid dehydrogenase type 1 (11β-HD1).
- 25 14. A compound for use according to claim 13 wherein the disease or condition susceptible of improvement by inhibition of 11β-hydroxysteroid dehydrogenase type 1 (11β-HD1) is selected from the group consisting of obesity, insulin resistance, metabolic syndrome, cognitive dysfunction associated to Alzheimer's disease, HIV-associated dementia, Creutzfeld-Jakob disease, Parkinson's disease, Huntington's disease, vascular inflammation, atherosclerosis, diabetes type 2, cardiovascular disease, 30 glaucoma, osteoporosis, myocardial infarction and alcohol abuse treatment.

15. Use of a compound according to anyone of claims 1 to 11 for the manufacture of a medicament for the treatment or prevention of a disease or conditions susceptible of improvement by inhibition of 11 β -hydroxysteroid dehydrogenase type 1 (11 β -HD1).

5 16. Use according to claim 15 wherein the disease or condition susceptible of improvement by inhibition of 11 β -hydroxysteroid dehydrogenase type 1 (11 β -HD1) is selected from the group consisting of obesity, insulin resistance, metabolic syndrome, cognitive disfunction associated to Alzheimer's disease, HIV-associated dementia, Creutzfeld-Jakob disease, Parkinson's disease, Huntington's disease, vascular
10 inflammation, atherosclerosis, diabetes type 2, cardiovascular disease, glaucoma, osteoporosis, myocardial infarction and alcohol abuse treatment.

17. A method for the treatment or prevention of a disease or conditions susceptible of improvement by inhibition of 11 β -hydroxysteroid dehydrogenase type 1 (11 β -HD1) in a
15 subject in need thereof comprising the administration to said subject of a therapeutically effective amount of a compound as defined in anyone of claims 1 to 11.

18. Method according to claim 17 wherein the disease or condition susceptible of improvement by inhibition of 11 β -hydroxysteroid dehydrogenase type 1 (11 β -HD1) is
20 selected from the group consisting of obesity, insulin resistance, metabolic syndrome, cognitive disfunction associated to Alzheimer's disease, HIV-associated dementia, Creutzfeld-Jakob disease, Parkinson's disease, Huntington's disease, vascular inflammation, atherosclerosis, diabetes type 2, cardiovascular disease, glaucoma, osteoporosis, myocardial infarction and alcohol abuse treatment.

25

19. A pharmaceutical composition comprising a compound as defined in anyone of claims 1 to 11 and a pharmaceutically acceptable excipient.

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2017/059178

A. CLASSIFICATION OF SUBJECT MATTER

INV. C07D401/14 A61K31/403 A61K31/4439 A61K31/454 C07D401/06
C07D409/06 C07D409/14 C07D209/70 C07D209/72 A61P3/00
A61P9/00 A61P25/00 A61P29/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

C07D A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, BIOSIS, CHEM ABS Data, EMBASE, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 3 475 450 A (KABBE HANS-JOACHIM ET AL) 28 October 1969 (1969-10-28) Markush formula; claims; examples -----	1-19
A	WO 2006/129178 A1 (GLENMARK PHARMACEUTICALS SA [CH]; MUTHUPPALANIAPPAN MEYYAPPAN [IN]; BA) 7 December 2006 (2006-12-07) Formulae (1), (g); claims; examples ----- -/--	1-19



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

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"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

19 May 2017

Date of mailing of the international search report

03/07/2017

Name and mailing address of the ISA/

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040,
Fax: (+31-70) 340-3016

Authorized officer

Kirsch, Cécile

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2017/059178

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	MATIAS REY-CARRIZO ET AL: "Easily Accessible Polycyclic Amines that Inhibit the Wild-Type and Amantadine-Resistant Mutants of the M2 Channel of Influenza A Virus", JOURNAL OF MEDICINAL CHEMISTRY, vol. 57, no. 13, 10 July 2014 (2014-07-10) , pages 5738-5747, XP055279820, US ISSN: 0022-2623, DOI: 10.1021/jm5005804 Scheme 1; page 5744, left column, 2nd paragraph; compounds 8-9 -----	1-19

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/EP2017/059178

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 3475450	A	28-10-1969	AT 277275 B 29-12-1969
			BE 706422 A 13-05-1968
			CH 492701 A 30-06-1970
			DE 1670770 A1 23-12-1970
			DE 1670844 A1 11-03-1971
			DK 128901 B 22-07-1974
			ES 347225 A1 01-05-1969
			GB 1151450 A 07-05-1969
			NL 6715534 A 17-05-1968
			SE 314677 B 15-09-1969
			US 3475450 A 28-10-1969

WO 2006129178	A1	07-12-2006	AR 055199 A1 08-08-2007
			AU 2006253842 A1 07-12-2006
			CA 2613678 A1 07-12-2006
			EP 1902034 A1 26-03-2008
			HK 1118806 A1 09-09-2011
			JP 2008545739 A 18-12-2008
			MY 142409 A 30-11-2010
			PE 04312007 A1 27-06-2007
			TW 200716560 A 01-05-2007
			US 2012142748 A1 07-06-2012
			WO 2006129178 A1 07-12-2006
