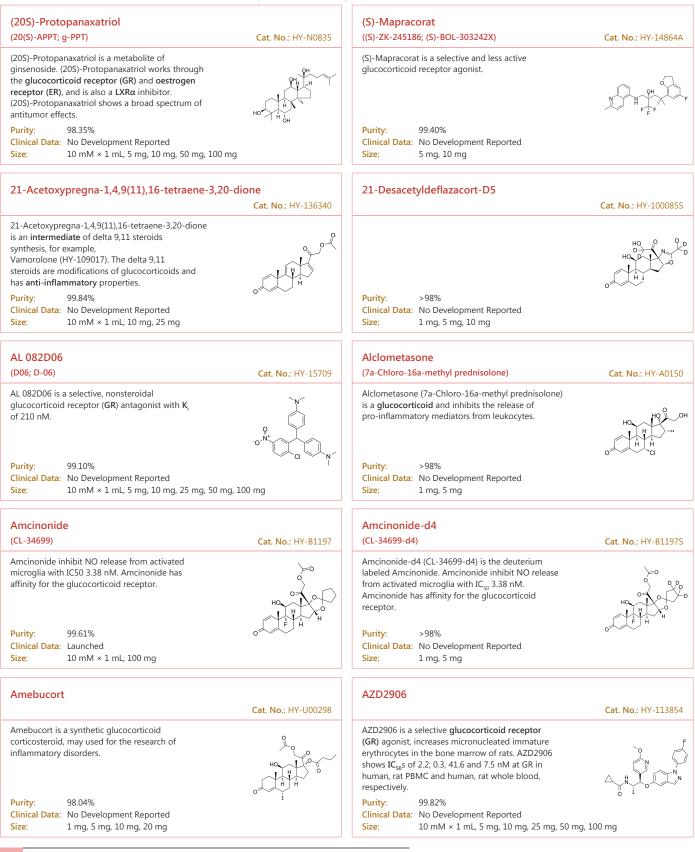


Glucocorticoid Receptor

Glucocorticoid Receptor (GR, or GCR) also known as NR3C1 (nuclear receptor subfamily 3, group C, member 1) is the receptor to which cortisol and other glucocorticoids bind. The GR is expressed in almost every cell in the body and regulates genes controlling the development, metabolism, and immune response. When the glucocorticoid receptor binds to glucocorticoids, its primary mechanism of action is the regulation of gene transcription. The unbound receptor resides in the cytosol of the cell. After the receptor is bound to glucocorticoid, the receptor-glucorticoid complex can take either of two paths. The activated GR complex up-regulates the expression of anti-inflammatory proteins in the nucleus or represses the expression of pro-inflammatory proteins in the cytosol by preventing the translocation of other transcription factors from the cytosol into the nucleus. Dexamethasone is an agonist, and RU486 and cyproterone acetate are antagonists of the GR. Also, progesterone and DHEA have antagonist effects on the GR.

Glucocorticoid Receptor Inhibitors, Agonists, Antagonists, Activators & Modulators



AZD5423 AZD9567 Cat. No.: HY-108243 Cat. No.: HY-120012 AZD5423 is an inhaled, potent, selective, and AZD9567 (compound 15) is a potent, oral active, non-steroidal glucocorticoid receptor (GR) non-steroidal and selective glucocorticoid receptor modulator (SGRM). AZD5423 effectively reduces modulator (SGRM), with an IC₅₀ of 3.8 nM. allergen-induced responses in subjects with mild Exhibits excellent efficacy in the streptococcal cell wall (SCW) reactivation model of joint allergic asthma. inflammation. Purity: 99 85% Purity: 9971% Clinical Data: Phase 2 Clinical Data: No Development Reported 5 mg, 10 mg Size: 10 mM × 1 mL, 5 mg, 10 mg, 50 mg, 100 mg Size: BAY 1003803 **Beclometasone** Cat. No.: HY-145351 (Beclomethasone) Cat. No.: HY-B1540 BAY 1003803 is a glucocorticoid receptor agonist Beclometasone (Beclomethasone) is a prototype for the topical treatment of psoriasis or severe glucocorticoid receptor agonist. atopic dermatitis. Purity: > 98% **Purity:** 95 44% Clinical Data: No Development Reported Clinical Data: Launched 10 mM × 1 mL, 25 mg, 50 mg, 100 mg Size: 1 mg, 5 mg Size: Beclometasone dipropionate Beclometasone dipropionate-d10 Cat. No.: HY-13571A Cat. No.: HY-13571AS1 Betamethasone dipropionate, the prodrug of Beclometasone dipropionate-d10 is the deuterium Betamethasone, is an orally active and potent labeled Beclometasone dipropionate. Betamethasone glucocorticoid with anti-inflammatory and dipropionate, the prodrug of Betamethasone, is an immunosuppressive activity. Betamethasone appears orally active and potent glucocorticoid with to be an effective inhibitor of LPS-induced anti-inflammatory and immunosuppressive activity. inflammation and MMP release. 99.92% Purity: >98% Purity: Clinical Data: Launched Clinical Data: No Development Reported Size: 10 mM × 1 mL, 250 mg Size: 1 mg, 5 mg Beclometasone dipropionate-d6 **Beclomethasone 17-propionate** (Beclomethasone-17-monopropionate; 17-BMP) Cat. No.: HY-13571AS Cat. No.: HY-136239 Beclometasone dipropionate-d6 is deuterium labeled Beclomethasone 17-propionate Beclometasone dipropionate. Betamethasone (Beclomethasone-17-monopropionate), an active dipropionate, the prodrug of Betamethasone, is an metabolite of Beclomethasone dipropionate orally active and potent glucocorticoid with (HY-13571), is a glucocorticoid receptor (GR) anti-inflammatory and immunosuppressive activity. agonist. >98% Purity: >98% Purity: Clinical Data: No Development Reported

Beclomethasone-d5

Size:

Beclomethasone-d5 is the deuterium labeled Beclometasone. Beclometasone (Beclomethasone) is a prototype glucocorticoid receptor agonist.

Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg

1 mg, 5 mg

Clinical Data: No Development Reported 10 mM × 1 mL, 5 mg, 10 mg, 25 mg, 50 mg Size

Betamethasone

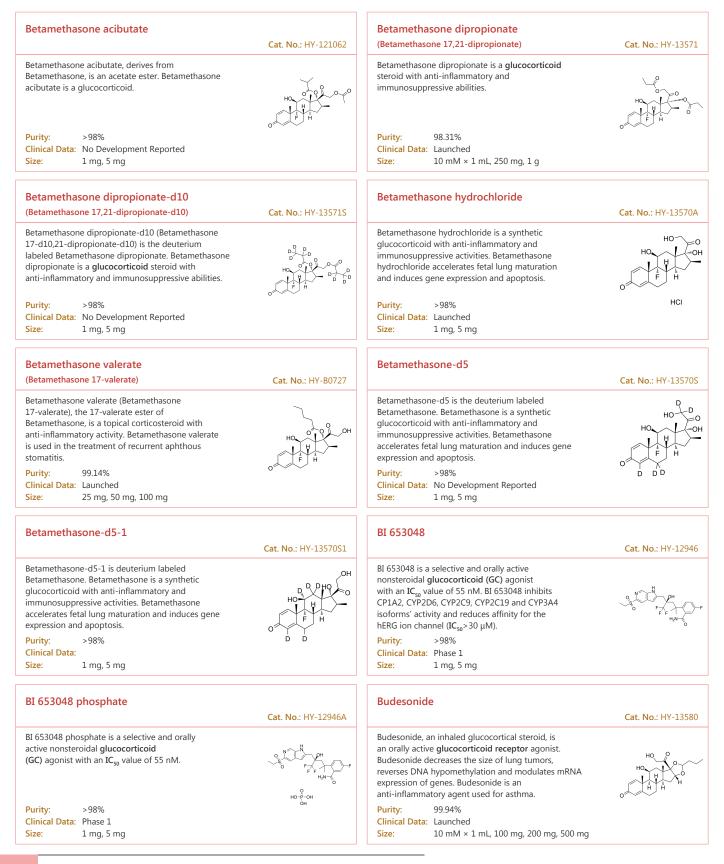
Betamethasone is a synthetic glucocorticoid with anti-inflammatory and immunosuppressive activities. Betamethasone accelerates fetal lung maturation and induces gene expression and apoptosis.

Purity: 99.97% Clinical Data: Launched Size: 10 mM × 1 mL, 100 mg, 500 mg

Cat. No.: HY-B1540S



Cat. No.: HY-13570



Budesonide impurity C **Budesonide-d8** Cat. No.: HY-100087 Cat. No.: HY-13580S Budesonide impurity C is an impurity of Budesonide-d8 is the deuterium labeled Budesonide. Budesonide, Budesonide, an inhaled glucocortical Budesonide, an inhaled glucocortical steroid, is steroid, is an orally active glucocorticoid an orally active glucocorticoid receptor agonist. Budesonide decreases the size of lung tumors, receptor agonist. reverses DNA hypomethylation and modulates mRNA expression of genes. Purity: > 98% Purity: >98% Clinical Data: No Development Reported Clinical Data: No Development Reported 1 mg, 5 mg, 10 mg Size: 1 mg, 5 mg Size: C108297 Ciclesonide Cat. No.: HY-125096 (RPR251526) Cat. No.: HY-B0625 C108297 is a selective glucocorticoid receptor Ciclesonide (RPR251526) is a glucocorticoid with an potent anti-inflammatory activity. Ciclesonide (GR) modulator (GR binding K, 0.7 nM; GR reporter gene functional K, 0.6 nM). C108297 attenuates can be used for asthma research. obesity by reducing caloric intake and increasing lipolysis and fat oxidation, and in addition attenuates inflammation. > 98% Purity: **Purity:** 9945% Clinical Data: No Development Reported Clinical Data: Launched 1 mg, 5 mg 10 mM × 1 mL, 50 mg, 100 mg Size: Size: Ciclesonide-d7 **Clobetasone butyrate** Cat. No.: HY-B0625S Cat. No.: HY-B1616 Clobetasone butyrate is a synthetic glucocorticoid Ciclesonide-d7 is the deuterium labeled Ciclesonide. Ciclesonide (RPR251526) is a and has topical anti-inflammatory activity glucocorticoid with an potent anti-inflammatory especially in skin. Clobetasone butyrate can be activity. Ciclesonide can be used for asthma used to relieve corticosteroid-responsive research. dermatoses, including atopic dermatitis and psoriasis. 99.24% Purity: >98% **Purity:** Clinical Data: No Development Reported Clinical Data: Launched Size: 1 mg, 10 mg Size: 10 mM × 1 mL, 100 mg Clobetasone butyrate-d7 Corticosterone (17-Deoxycortisol; 11β,21-Dihydroxyprogesterone; Kendall's compound B) Cat. No.: HY-B1616S Cat. No.: HY-B1618 Clobetasone butyrate-d7 is the deuterium labeled Corticosterone is an adrenocortical steroid that Clobetasone butyrate. Clobetasone butyrate is a has modest but significant activities as a mineralocorticoid and a glucocorticoid. synthetic glucocorticoid and has topical anti-inflammatory activity especially in skin.

>98% Purity: Clinical Data: No Development Reported Size: 1 mg, 5 mg

Corticosterone-d4

Corticosterone-d4 is the deuterium labeled Corticosterone. Corticosterone is an adrenocortical steroid that has modest but significant activities as a mineralocorticoid and a glucocorticoid.

Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg



Cat. No.: HY-B1618S1

99.70% **Purity:** Clinical Data: Phase 3 10 mM × 1 mL, 50 mg, 100 mg Size:

Corticosterone-d8

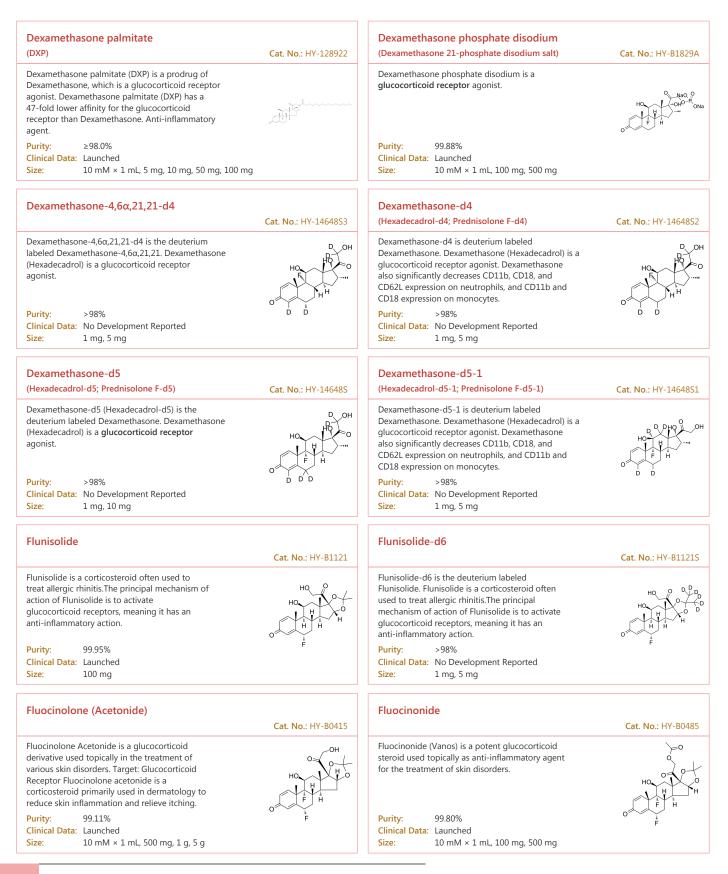
Corticosterone-d8 is the deuterium labeled Corticosterone. Corticosterone is an adrenocortical steroid that has modest but significant activities as a mineralocorticoid and a glucocorticoid.

Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg

Cat. No.: HY-B1618S



Dagrocorat hydrochloride		Dazucorilant	
(PF-00251802 hydrochloride)	Cat. No.: HY-16718A	(CORT113176)	Cat. No.: HY-132811
Dagrocorat (PF-00251802) hydrochloride is an orally active and selective high-affinity partial agonist of the glucocorticoid receptor. Purity: 99.85% Clinical Data: No Development Reported Size: 5 mg, 10 mg, 25 mg, 50 mg, 100 mg		Dazucorilant (CORT113176) is a selective and high affinity non-steroidal glucocorticoid receptor (GR) modulator with a K ₁ value 1 nM in vitro. Dazucorilant can be used for the research of neurological disorders. Purity: >98% Clinical Data: No Development Reported Size: 5 mg, 10 mg, 25 mg, 50 mg, 100 mg	
Deflazacort	Cat. No .: HY-13609	Deflazacort-D5	Cat. No. : HY-13609S
Deflazacort, a glucocorticoid, is an inactive prodrug and is converted rapidly to the active metabolite 21-desacetyldeflazacort. Deflazacort is used as an anti-inflammatory and immunosuppressant. Purity: 99.73% Clinical Data: Launched Size: 10 mM × 1 mL, 50 mg, 100 mg		Deflazacort-D5 is the deuterium labeled Deflazacort. Deflazacort, a glucocorticoid, is an inactive prodrug and is converted rapidly to the active metabolite 21-desacetyldeflazacort. Deflazacort is used as an anti-inflammatory and immunosuppressant. Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg, 10 mg	
Deflazacort-d5-1	Cat. No.: HY-13609S2	Deflazacort-D7	Cat. No .: HY-13609S1
Deflazacort-d5-1 is the deuterium labeled Deflazacort. Deflazacort, a glucocorticoid, is an inactive prodrug and is converted rapidly to the active metabolite 21-desacetyldeflazacort. Deflazacort is used as an anti-inflammatory and immunosuppressant. Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg		Deflazacort-D7 is the deuterium labeledDeflazacort. Deflazacort, a glucocorticoid, is aninactive prodrug and is converted rapidly to theactive metabolite 21-desacetyldeflazacort.Deflazacort is used as an anti-inflammatory andimmunosuppressant.Purity:>98%Clinical Data:No Development ReportedSize:1 mg, 5 mg, 10 mg	
Desisobutyryl-ciclesonide (CIC-AP; Ciclesonide active principle)	Cat. No.: HY-111490	Desonide	Cat. No.: HY-B0248
Desisobutyryl-ciclesonide is the active metabolite of Ciclesonide. Desisobutyryl-ciclesonide has affinity for the glucocorticoid receptor . Purity: 99.53% Clinical Data: No Development Reported		Desonide is a nonfluorinated corticosteroid anti-inflammatory agent used topically for dermatoses. Target: Glucocorticoid Receptor Desonide is a low-potency topical corticosteroid that has been used for decades in the treatment of steroid-responsive dermatoses . Purity: 99.45% Clinical Data: Launched	
Clinical Data: No Development Reported Size: 10 mM × 1 mL, 5 mg, 10 mg, 25 mg		Size: 10 mM × 1 mL, 10 mg, 50 mg, 100 mg, 200 mg	mg, 500 mg
Dexamethasone (Hexadecadrol; Prednisolone F)	Cat. No.: HY-14648	Dexamethasone acetate (Dexamethasone 21-acetate; Hexadecadrol acetate)	Cat. No.: HY-14648A
Dexamethasone (Hexadecadrol) is a glucocorticoid receptor agonist. Dexamethasone also significantly decreases CD11b, CD18, and CD62L expression on neutrophils, and CD11b and CD18 expression on monocytes.		Dexamethasone acetate (Dexamethasone 21-acetate) is a glucocorticoid receptor agonist. Dexamethasone acetate has the potential for ophthalmic infections treatment.	
Purity: 99.86% Clinical Data: Launched Size: 10 mM × 1 mL, 500 mg, 1 g, 5 g		Purity: 98.24% Clinical Data: Launched Size: 10 mM × 1 mL, 500 mg, 1 g, 5 g	



Fluorometholone

Fluorometholone, a synthetic glucocorticoid, is a alucocorticoid receptor agonist with anti-inflammatory and anti-allergic properties. Fluorometholone can be used for the research of dry eye.

Purity: 99 49% Clinical Data: Launched Size: 10 mM × 1 mL, 25 mg, 50 mg, 100 mg

Fluticasone furoate

Cat. No.: HY-15234

Cat. No.: HY-B1893

Fluticasone furoate is a topical, intranasal, enhanced-affinity synthetic trifluorinated corticosteroid with a K_d of 0.3 nM. Fluticasone furoate has potent anti-inflamatory and anti-asthmatic activity, and low systemic exposure. Purity: 99.06%

Clinical Data: Launched 10 mM × 1 mL, 10 mg Size:

Fluticasone propionate-d3

Fluticasone propionate-d3 is the deuterium labeled Fluticasone propionate. Fluticasone propionate, a potent topical anti-inflammatory corticosteroid, is a selective glucocorticoid receptor agonist, with an absolute affinity (K_p) of 0.5 nM.

Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg

Fosdagrocorat (PF-04171327)

Fosdagrocorat (PF-04171327) is a dissociated glucocorticoid receptor agonist.

99.14% Purity: Clinical Data: No Development Reported 10 mM × 1 mL, 5 mg, 10 mg, 25 mg, 50 mg, 100 mg Size:

Glucocorticoid receptor agonist-1

Glucocorticoid receptor agonist-1 is a potent glucocorticoid receptor agonist with an IC₅₀ of 2.8 nM extracted from patent WO2017210471A1, compound 41.

Purity: >98% Clinical Data: No Development Reported Size: 5 mg, 10 mg

Fluticasone (propionate)

Fluticasone propionate, a potent topical anti-inflammatory corticosteroid, is a selective glucocorticoid receptor agonist, with an absolute affinity (K_p) of 0.5 nM. Fluticasone propionate shows little or no activity at other steroid receptors. Anti-viral activity. Purity: 99 97% Clinical Data: Launched

10 mM × 1 mL, 10 mg, 50 mg

Fluticasone furoate-d3

Size:

Fluticasone furoate-d3 is deuterium labeled Fluticasone furoate. Fluticasone furoate is a topical, intranasal, enhanced-affinity synthetic trifluorinated corticosteroid with a Kd of 0.3 nM.

Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg

Fluticasone propionate-d5

Fluticasone propionate-d5 is deuterium labeled Fluticasone (propionate). Fluticasone propionate, a potent topical anti-inflammatory corticosteroid, is a selective glucocorticoid receptor agonist, with an absolute affinity (KD) of 0.5 nM.

>98% Purity: Clinical Data: No Development Reported Size: 1 mg, 5 mg

Glucocorticoid receptor agonist

Glucocorticoid receptor agonist is a potent Glucocorticoid receptor agonist. IC50 value: Target:.

99.56% **Purity:** Clinical Data: No Development Reported 10 mM × 1 mL, 5 mg, 10 mg Size:

Glucocorticoid receptor-IN-1

Glucocorticoid receptor-IN-1 (Compound WX002) is a selective glucocorticoid receptor (GR) modulator with anti-inflammatory effect.

Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg

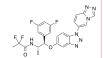
www.MedChemExpress.com

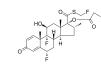


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Cat. No.: HY-14234

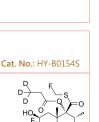






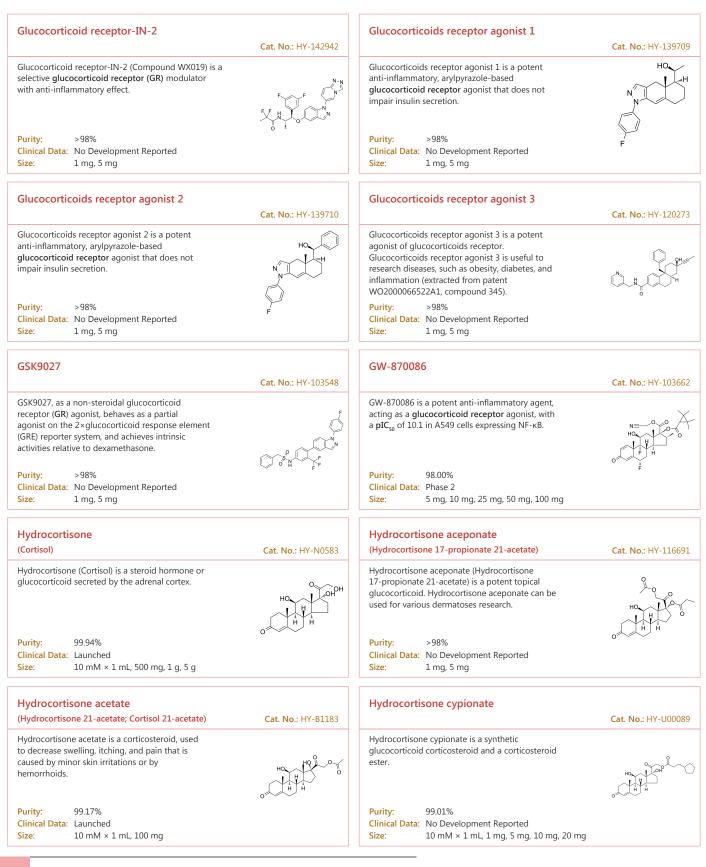
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Cat. No.: HY-B0154



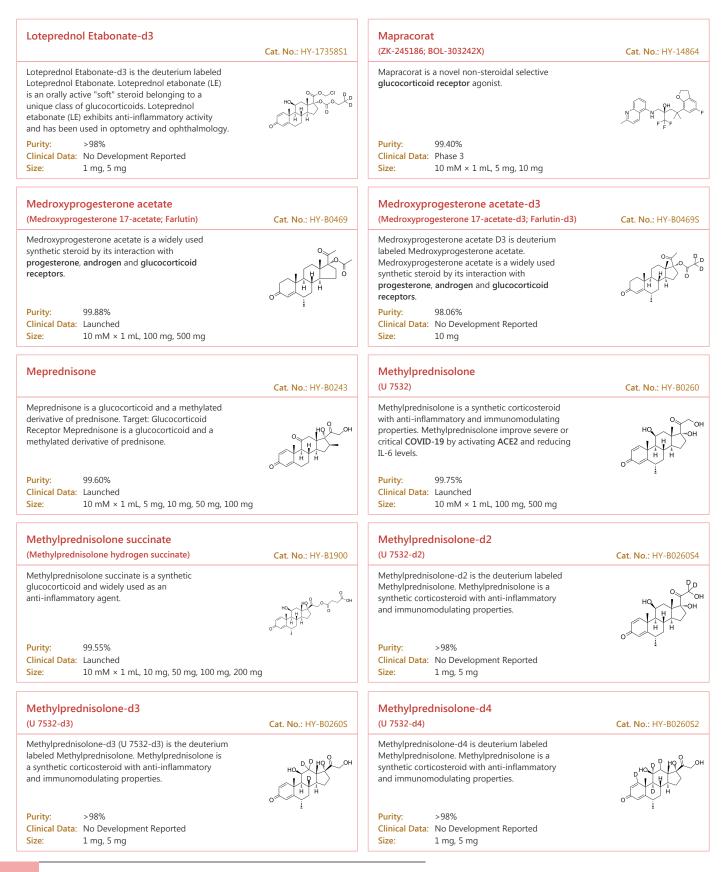
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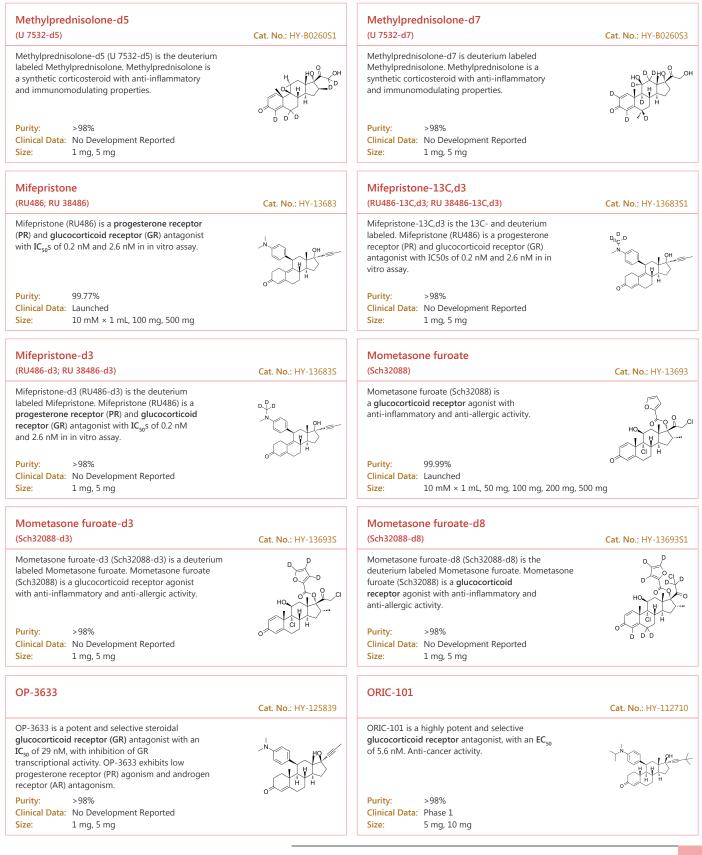
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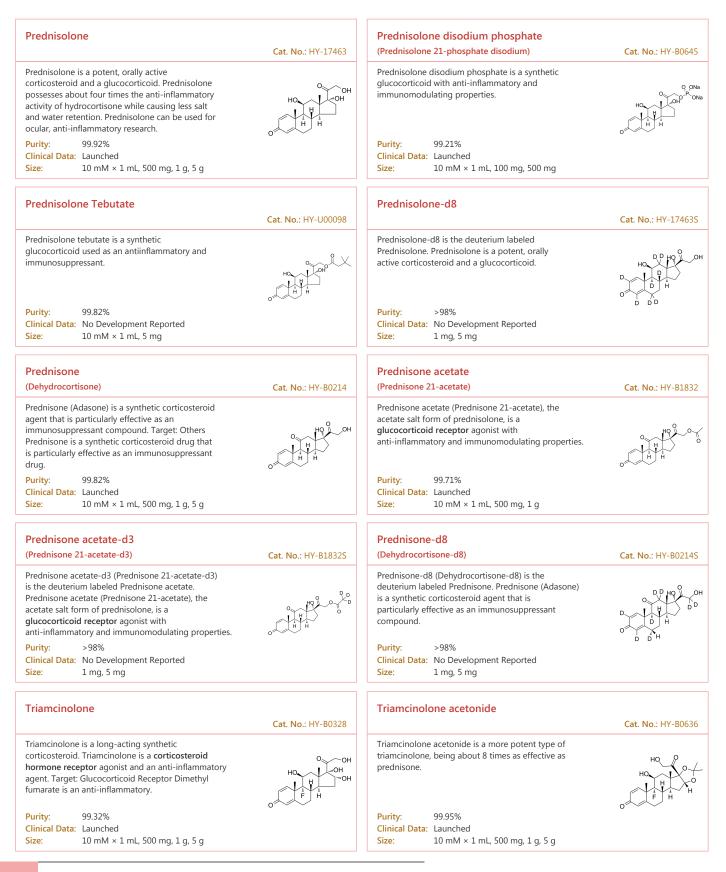


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Hydrocortisone hemisuccinate (Hydrocortisone 21-hemisuccinate)	Cat. No. : HY-B1402	Hydrocortisone phosphate (Hydrocortisone 21-phosphate; Cortisol 21-phosphate)	Cat. No.: HY-B1155
Hydrocortisone hemisuccinate (Hydrocortisone 21-hemisuccinate), a physiological glucocorticoid, is an orally active steroidal anti-inflammatory drug (SAID).		Hydrocortisone phosphate (Hydrocortisone 21-phosphate), a physiological glucocorticoid, and is an orally active steroidal anti-inflammatory drug (SAID).	
Purity: 99.76% Clinical Data: Launched Size: 10 mM × 1 mL, 25 mg, 50 mg, 100 mg		Purity:>98%Clinical Data:LaunchedSize:1 mg, 5 mg	
Hydrocortisone-d2		Hydrocortisone-d3	
(Cortisol-d2) Hydrocortisone-d2 is the deuterium labeled Hydrocortisone. Hydrocortisone (Cortisol) is a steroid hormone or glucocorticoid secreted by the adrenal cortex.	Cat. No.: HY-N058355	(Cortisol-d3) Hydrocortisone-d3 (Cortisol-d3) is the deuterium labeled Hydrocortisone. Hydrocortisone (Cortisol) is a steroid hormone or glucocorticoid secreted by the adrenal cortex.	
Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg	Ĵ	Purity:>98%Clinical Data:No Development ReportedSize:1 mg, 5 mg	
Hydrocortisone-d4 (Cortisol-d4)	Cat. No. : HY-N0583S2	Hydrocortisone-d7 (Cortisol-d7)	Cat. No.: HY-N0583S1
Hydrocortisone-d4 (Cortisol-d4) is the deuterium labeled Hydrocortisone. Hydrocortisone (Cortisol) is a steroid hormone or glucocorticoid secreted by the adrenal cortex.		Hydrocortisone-d7 (Cortisol-d7) is the deuterium labeled Hydrocortisone. Hydrocortisone (Cortisol) is a steroid hormone or glucocorticoid secreted by the adrenal cortex.	
Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg	0, ~ ~	Purity:>98%Clinical Data:No Development ReportedSize:1 mg, 5 mg	ם ם מ
JTP-117968	Cat. No .: HY-125265	LEO 134310	Cat. No .: HY-144397
JTP-117968, a novel selective glucocorticoid receptor modulator (a non-steroidal SGRM, IC _{so} of 6.8 nM), exhibits improved transrepression/transactivation dissociation.	Received a second secon	LEO 134310 is a selective, non-steroidal glucocorticoid receptor (GR) agonist optimized for topical treatment., LEO 134310 showed high affinity (EC_{s0} of 14 nM) in a GR binding assay. LEO 134310 can be used for skin diseases.	citore
Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg	< <u>,</u> ∼ 0	Purity:>98%Clinical Data:No Development ReportedSize:1 mg, 5 mg	
Loteprednol Etabonate	Cat. No .: HY-17358	Loteprednol Etabonate D5	Cat. No.: HY-17358S
Loteprednol etabonate (LE) is an orally active "soft" steroid belonging to a unique class of glucocorticoids. Loteprednol etabonate (LE) exhibits anti-inflammatory activity and has been used in optometry and ophthalmology.		Loteprednol Etabonate D5 is a deuterium labeled Loteprednol etabonate. Loteprednol etabonate (LE) is an orally active "soft" steroid belonging to a unique class of glucocorticoids. Loteprednol etabonate (LE) exhibits anti-inflammatory activity and has been used in optometry and ophthalmology.	
Purity: 99.90% Clinical Data: Launched Size: 10 mM × 1 mL, 10 mg, 50 mg, 100 mg		Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg	







Triamcinolone acetonide-d6		Triamcinolone acetonide-d7	
	Cat. No.: HY-B0636S3		Cat. No.: HY-B0636S
Triamcinolone acetonide-d6 is deuterium labeled Triamcinolone acetonide.		Triamcinolone acetonide-d7 is the deuterium labeled Triamcinolone acetonide. Triamcinolone acetonide is a more potent type of triamcinolone, being about 8 times as effective as prednisone.	
Purity:>98%Clinical Data:No Development ReportedSize:1 mg, 5 mg		Purity:>98%Clinical Data:No Development ReportedSize:1 mg, 5 mg	0, ~ ~
Triamcinolone acetonide-d7-1	Cat. No. : HY-B0636S2	Triamcinolone Benetonide	Cat. No.: HY-U00043
Triamcinolone acetonide-d7-1 is deuterium labeled Triamcinolone acetonide.		Triamcinolone benetonide is a synthetic glucocorticoid corticosteroid with anti-inflammatory activity.	
Purity:>98%Clinical Data:No Development ReportedSize:1 mg, 5 mg	Ď	Purity:>98%Clinical Data:No Development ReportedSize:1 mg, 5 mg	-
Vamorolone		Velsecorat	
(VBP15)	Cat. No.: HY-109017	(AZD7594; AZ13189620)	Cat. No.: HY-111453
Vamorolone (VBP15) is a first-in-class, orally active dissociative steroidal anti-inflammatory drug and membrane-stabilizer. Vamorolone improves muscular dystrophy without side effects. Vamorolone shows potent NF-kB inhibition and substantially reduces hormonal effects.	HQ OH	AZD7594 is a potent selective nonsteroidal glucocorticoid receptor modulator, with an IC _{so} of 0.9 nM.	Át COM
Purity: 99.12% Clinical Data: Phase 2 Size: 10 mM × 1 mL, 5 mg, 10 mg, 50 mg	Ŭ	Purity: 99.60% Clinical Data: Phase 2 Size: 10 mM × 1 mL, 5 mg, 10 mg, 25 mg, 50 mg, 100 mg	
Zavacorilant		ZK 216348	
	Cat. No.: HY-139556	((+)-ZK 216348)	Cat. No.: HY-123352
Zavacorilant is capable of modulating glucocorticoid receptor (GR).		ZK 216348 ((+)-ZK 216348) is a nonsteroidal selective glucocorticoid receptor agonist with an IC_{s0} of 20.3 nM. ZK 216348 also binds to Progesterone and mineralocorticoid receptors with IC_{s0} s of 20.4 nM and 79.9 nM, respectively.	
Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg	/ N ^{=J} - 1 S N≂/	Purity:>98%Clinical Data:No Development ReportedSize:1 mg, 5 mg	<u></u>