

Small Molecules

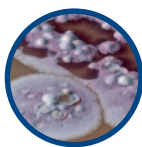
HIGHLIGHTS



Synthetic Small Molecules



Unique Natural Products



Rare Antibiotics



MANUFACTURED IN SWITZERLAND

Full Panel of >1500 Small Molecules
Over 25 Years of Experience
High-end Customer & Technical Support
BULK Quantities

Collaborating with:



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Selected NLRP3 Inflammasome Inhibitors

NEW

MCC950 . Na

AG-CR1-3615

1 mg | 5 mg | 10 mg

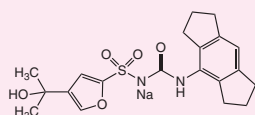
Formula: $C_{20}H_{23}N_2NaO_5S$

MW: 426.5

CAS: 256373-96-3

Potent and selective NLRP3 inflammasome inhibitor.

LIT: A small-molecule inhibitor of the NLRP3 inflammasome for the treatment of inflammatory diseases: R.C. Coll, et al.; Nat. Med. **21**, 248 (2015)



NEW

Isoliquiritigenin

AG-CN2-0459

10 mg | 50 mg

Formula: $C_{15}H_{12}O_4$

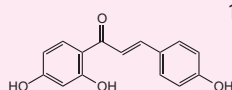
MW: 256.3

CAS: 961-29-5

Source: *Synthetic*

NEW INSIGHTS: Inhibits NLRP3-activated ASC oligomerization. Blocks priming and activation step.

LIT: Isoliquiritigenin is a potent inhibitor of NLRP3 inflammasome activation and diet-induced adipose tissue inflammation: H. Honda, et al.; J. Leukoc. Biol. **96**, 1087 (2014)



Arglabin

AG-CN2-0458

1 mg | 5 mg

BAY 11-7082

AG-CR1-0013

10 mg | 50 mg

Colchicine

AG-CN2-0048

500 mg | 1 g

Glyburide (USP)

AG-CR1-3613

1 g | 5 g | 10 g

(R)-3-Hydroxybutyric acid

AG-CR1-3616

25 mg | 100 mg

(S)-3-Hydroxybutyric acid

AG-CR1-3617

25 mg | 100 mg

LATEST INSIGHT: The ketone metabolite β -hydroxybutyrate blocks NLRP3 inflammasome-mediated inflammatory disease: Y.H. Youm, et al.; Nat. Med. **21**, 263 (2015)

Parthenolide

AG-CN2-0455

10 mg | 50 mg | 250 mg

Resveratrol

AG-CN2-0033

50 mg | 100 mg | 500 mg

LATEST INSIGHT: Resveratrol inhibits NLRP3 inflammasome activation by preserving mitochondrial integrity and augmenting autophagy: Y.P. Chang, et al.; J. Cell Physiol. **230**, 1567 (2015)

Vinpocetine

AG-CN2-0454

20 mg | 100 mg

Selected NLRP3 Inflammasome Activators

The Original Source of MSU!

Monosodium urate [MSU]

AG-CR1-3950 (crystals)

2 mg | 2 x 2 mg

AG-CR1-3951 (ready-to-use solution)

10 mg

Formula: $C_5H_3N_4O_3 \cdot Na$

MW: 167.1 . 23.0

CAS: 1198-77-2

Potent NLRP3 inflammasome activator. **Biological Activity Tested!**

LIT: Gout-associated uric acid crystals activate the NALP3 inflammasome: F. Martinon, et al.; Nature **440**, 237 (2006)

Other Selected Activators

Nigericin . Na

AG-CN2-0020

5 mg | 25 mg

LIT: Cryopyrin activates the inflammasome in response to toxins and ATP: S. Mariathasan, et al.; Nature **440**, 228 (2006)

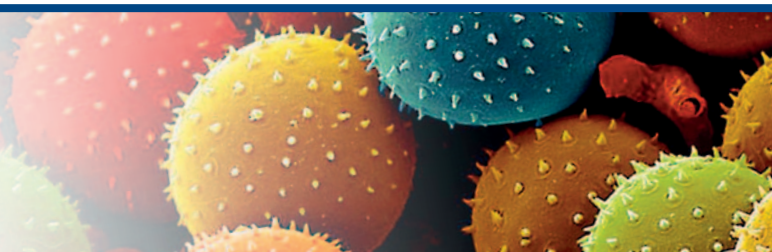
7BIO

MR-C0020

10 mg | 50 mg

LIT: Cutting edge: Necrosis activates the NLRP3 inflammasome: H. Li, et al.; J. Immunol. **183**, 1528 (2009)

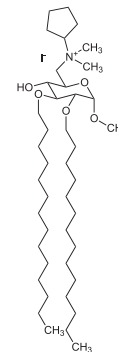
For Compounds from
Fungal Sources
www.adipogen.com



TLR4/CD14 Antagonists – IAXO Compounds

IAXO-101 (CD14/TLR4 Antagonist) (synthetic)
IAXO-102 (CD14/TLR4 Antagonist) (synthetic)
IAXO-103 (CD14/TLR4 Antagonist) (synthetic)
IAXO-201 (Control for IAXO-102)
IAXO-202 (Control for IAXO-101/IAXO-103)

IAX-600-001	1 mg 5 mg
IAX-600-002	1 mg 5 mg
IAX-600-003	1 mg 5 mg
IAX-600-004	1 mg
IAX-600-005	1 mg



TLR Agonists

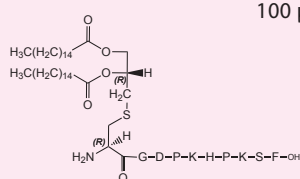
(R)-FSL-1

AG-CP3-0010

Formula: C₈₄H₁₄₀N₁₄O₁₈S

MW: 1666.2

CAS: 322455-70-9 (R/S)



100 µg

Potent stimulator of TLR2/TLR6. The naturally occurring R-stereoisomer is biologically more active than the S-stereoisomer.

Also available: **FSL-1 (AG-CP3-0009)**

Pam₃Cys-Ser-(Lys)₄ (TLR1/2 Complex Agonist)

AG-CP3-0003

2 mg

Poly(I:C) (Endotoxin-free) (TLR3 Agonist)

IAX-200-021

2 mg | 5 mg

MPLA (synthetic) Sterile Solution (TLR4 Agonist)

AG-CU1-0002

100 µg

Gardiquimod (TLR7 Agonist)

AG-CR1-3583

5 mg | 25 mg

Imiquimod (TLR7 Agonist)

AG-CR1-3569

100 mg | 250 mg

Loxoribine (TLR7 Agonist)

AG-CR1-3584

5 mg | 25 mg

R-848 (TLR7/8 Agonist)

AG-CR1-3582

5 mg | 25 mg



TLRpure™ LPS, Lipid A & MPLA

Qualified Purity & TLR4-specific Activity • Ultrapure (no Detectable Protein, RNA & DNA) • Tested in TLR4 KO murine macrophages • Ready-to-use Solutions or Powder • MegaVAX™ BULK Sizes Available • Broadest Range of LPS Chemotype & Serotype Selection



TLRpure™ Endotoxin-free & Sterile ODNs

Qualified Purity & Activity • Endotoxin-free & Sterile • Tested in TLR9 KO murine macrophages • BULK Available for *in vivo* Studies

UNIQUE

Cytosolic DNA Sensor

cGAMP . 2Na

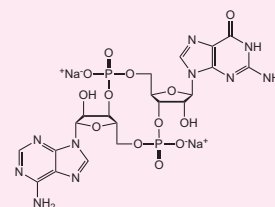
AG-CR1-3588

100 µg | 500 µg

Formula: C₂₀H₂₄N₁₀O₁₃P₂ . 2Na

MW: 674.4 . 46.0

CAS: 849214-04-6 (free acid)



Potent STING ligand. Second messenger in human and mouse cells that triggers interferon production in response to cytosolic DNA.

Antiviral & Anti-inflammatory Compounds

10Z-Hymenialdisine

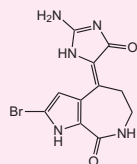
AG-CN2-0067

500 µg | 1 mg

Formula: $C_{11}H_{10}BrN_5O_2$

MW: 324.1

CAS: 82005-12-7

Source: *Axinella carteri*

Potent MEK-1 and NF-κB inhibitor. MARK inhibitor. ATP-competitive kinase inhibitor. Inhibits various pro-inflammatory cytokines and nitric oxide.

BULK

Z-VRPR-FMK . TFA

AG-CP3-0008

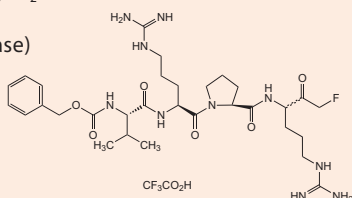
500 µg | 1 mg

Formula: $C_{31}H_{49}FN_{10}O_6 \cdot CF_3CO_2H$

MW: 676.8 . 114.0

CAS: 1381885-28-4 (free base)

Selective MALT1 inhibitor.



BULK

High Purity Betulinic Acid

Betulinic acid

AG-CN2-0417 (>97%)

AG-CN2-0415 (>99%)

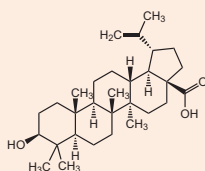
100 mg | 500 mg | 1 g

5 mg | 25 mg | 100 mg

Source: *Platanus acerifolia* bark

Multifunctional natural product with anti-HIV, anticancer, antimalarial and apoptosis inducing properties. Potent DNA topoisomerase II inhibitor and proteasome activator.

BULK available



Actinonin (synth.) (MMP and Meprin A inhibitor)

AG-CN2-0161

5 mg | 25 mg

Aurantimycin A (C5a antagonist)

BVT-0398

1 mg | 5 mg

Boromycin (HIV-1 integrase inhibitor)

AG-CN2-0166

250 µg | 1 mg

Curvularin (Anti-inflammatory compound)

AG-CN2-0147

1 mg | 5 mg

Fuscin (CCR5 receptor antagonist)

AG-CN2-0138

1 mg | 2.5 mg

Helenalin (NF-κB inhibitor)

AG-CN2-0435

500 µg

Malformin A1 (IL-1β inhibitor)

AG-CN2-0169

250 µg | 1 mg

Petasol (HIV-1 Tat transduction inhibitor)

BVT-0439

1 mg | 5 mg

Pyranonigrin A (Potent free radical scavenger)

AG-CN2-0156

250 µg | 1 mg

Rugulosin (HIV-1 integrase inhibitor)

AG-CN2-0124

250 µg | 1 mg

Ruxolitinib (JAK1/JAK2 inhibitor)

AG-CR1-3624

5 mg | 25 mg

Tofacitinib (JAK3 inhibitor, Immunosuppressant)

AG-CR1-3625

5 mg | 25 mg

Triptolide (Potent immunosuppressant)

AG-CN2-0448

1 mg | 5 mg | 10 mg

Wogonin (Anti-inflammatory compound)

AG-CN2-0064

5 mg | 25 mg

NEW

5-Lipoxygenase Inhibitor

Myxochelin A

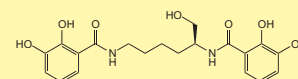
AG-CN2-0470

100 µg | 1 mg

Formula: $C_{20}H_{24}N_2O_7$

MW: 404.4

CAS: 120243-02-9



LIT: Myxochelins target human 5-lipoxygenase: S. Schieferdecker, et al.; J. Nat. Prod. 78, 335 (2015)

THE STANDARDS

Potent Immunosuppressants

Cyclosporin A

AG-CN2-0079

100 mg | 5 x 100 mg | 1 g

Also available: **Cyclosporin C, D & H**

Ascomycin (high purity)

AG-CN 2-0420

1 mg | 5 mg | 25 mg

FTY720 . HCl

AG-CR1-3587

1 mg | 5 mg | 25 mg

Antimicrobial Compounds

UNIQUE

Tropodithietic acid [TDA]

BVT-0152

1 mg | 5 mg

Formula: C₈H₄O₃S₂

MW: 212.3

CAS: 750590-18-2

Source: *Roseobacter gallaeciensis*

Antibiotic. Active against Gram-positive and Gram-negative bacteria. Antifungal and anti-nematodal. Shows antitumor activity. Bacterial signal substance.

Unique Antibiotic Structure



THE STANDARDS

Colistin sulfate (USP Grade)

AG-CN2-0065

100 mg | 1 g | 5 g

Gentamicin sulfate (USP Grade)

AG-CN2-0066

1 g | 5 g | 25 g

Helvolic acid (Potent antibacterial compound)

BVT-0435

1 mg | 5 mg

Meleagrins (Antifouling agent)

AG-CN2-0451

1 mg

White Line Inducing Principle (Diagnostic tool)

AG-CN2-0069

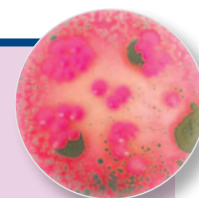
1 mg

PRODUCT NAME	ACTIVITY	PID	SIZE
Aerothionin	Antibiotic & Antimycobacterial	AG-CN2-0453	1 mg
Antibiotic PF 1052	Antibiotic & Antifungal	BVT-0338	500 µg 1 mg
Ambuic acid	Quorum sensing modulator	AG-CN2-0129	250 µg 1 mg
Aranorosin	Antibacterial	AG-CN2-0114	250 µg 1 mg
Aspterric acid	Antibiotic	AG-CN2-0132	1 mg
Aureothin	Potent Antimicrobial	BVT-0303	1 mg 5 mg
Aurodox	Protein biosynthetic (EF-Tu) inhibitor	AG-CN2-0133	1 mg 2.5 mg
Aurofusarin	Mycotoxin	BVT-0392	1 mg 5 mg
Bacimethrin	Antibiotic	BVT-0226	500 µg 1 mg
Chaetoglobosin A	Antibiotic & Anticancer	BVT-0092	1 mg 5 mg
Enterocin	Antibacterial	AG-CN2-0116	250 µg 1 mg
Kirromycin	Protein biosynthesis inhibitor	BVT-0157	1 mg 5 mg
Hormaomycin	Quorum sensing inhibitor	BVT-0107	500 µg 1 mg
Luteoreticulins NEW	Insecticidal & Nematocidal	BVT-0457	500 µg 1 mg
Lysolipin I	Antibiotic & Antifungal	BVT-0037	500 µg 1 mg
Malformin A1	Antibiotic	AG-CN2-0169	250 µg 1 mg
Malformin C (synthetic)	Peptide Antibiotic & Anticancer	AG-CN2-0107	250 µg 1 mg
Nargenicin A1	Antibacterial against Gram-positive	BVT-0204	1 mg 5 mg
Nocardamine	Siderophore	AG-CN2-0150	1 mg 5 mg
Pikromycin	PREP inhibitor	BVT-0400	1 mg 5 mg
Pseurotin A	Antibiotic	BVT-0003	1 mg 5 mg
γ-Rubromycin	Antibiotic & Antiviral	BVT-0007	1 mg 5 mg
Siamycin I	Quorum sensing inhibitor	AG-CN2-0146	1 mg 5 mg
Simocyclinone D8	Bacterial DNA gyrase inhibitor	BVT-0290	500 µg 1 mg
Strobilurin B	Antifungal	BVT-0312	500 µg 1 mg
Thailandolide B	Antibiotic	AG-CN2-0160	1 mg 5 mg
Viomellein	Antibacterial & Insecticidal	BVT-0359	500 µg 1 mg
Zelkovamycin	Antibacterial	AG-CN2-0128	250 µg 1 mg

New Additions! Quorum Sensing Modulators

Comprehensive Panel of over 30 Homoserine-lactones from the Manufacturer!

Visit www.adipogen.com for a Complete Overview!



AMPK Modulators

AMPK (AMP-activated protein kinase) is an enzyme that plays a role in cellular energy homeostasis, regulating several intracellular systems including hepatic fatty acid oxidation and ketogenesis, inhibition of cholesterol synthesis, lipogenesis and triglyceride synthesis, stimulation of skeletal muscle fatty acid oxidation and muscle glucose uptake as well as modulation of insulin secretion by pancreatic β cells.

SELECTED REVIEW: Past strategies and future directions for identifying AMP-activated protein kinase (AMPK) modulators: S.E. Sinnott & J.E. Brenman; Pharmacol. Ther. **143**, 111 (2014)

AICAR

AG-CR1-0061

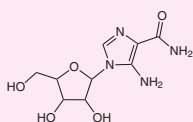
10 mg | 50 mg | 100 mg

Formula: $C_9H_{14}N_4O_5$

MW: 258.2

CAS: 2627-69-2

Cell permeable AMP-activated protein kinase (AMPK) activator. Insulin mimetic.



Other AMPK Modulators

Saquinone (AMPK activator)

AG-CN2-0032

1 mg | 5 mg

(-)-Epigallocatechin gallate [EGCG] (AMPK activator)

AG-CN2-0063

25 mg | 100 mg

Piceatannol

(Promotes AMPK phosphorylation & GLUT4 translocation)

AG-CN2-0086

1 mg | 5 mg | 25 mg

MOTS-c (human) (AMPK Inducer)

AG-CP3-0026

1 mg | 5 mg

NEW

AdipoRon

AG-CR1-0154

10 mg | 50 mg

BULK | Original Source

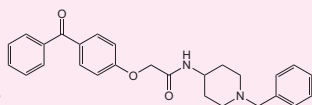
Formula: $C_{27}H_{28}N_2O_3$

MW: 428.5

CAS: 924416-43-3

AdipoR agonist. AMPK & PGC1 α activator. Improves diabetes, glucose and lipid metabolism and insulin sensitivity.

LIT: A small-molecule AdipoR agonist for type 2 diabetes and short life in obesity: M. Okada-Iwabu, et al.; Nature **503**, 493 (2013) • Cell Biology. Ronning after the adiponectin receptors: W.L. Holland & P.E. Scherer; Science **342**, 1460 (2013)



Also available

Compound 112254 (AMPK activator)

AG-CR1-0155

10 mg | 50 mg

Compounds with increased Solubility

AdipoRon . HCl (water soluble)

AG-CR1-0156

Compound 112254 . HCl (water soluble) AG-CR1-0157

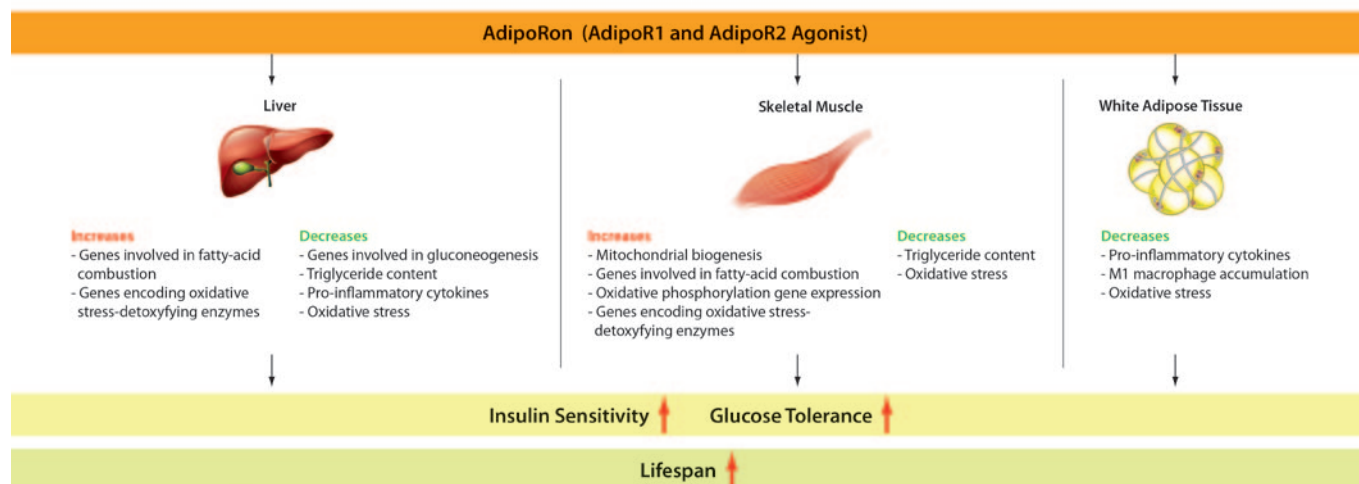


FIGURE: Mechanism of Action of AdipoRon. Adapted from M. Okada-Iwabu, et al.; Nature **503**, 493 (2013)

PPAR Agonists

PPAR γ is a nuclear receptor that is highly expressed in adipose tissue and is a master regulator of adipogenesis. PPAR γ is selectively activated by a class of anti-diabetic agents, the thiazolidinediones (TZD), which ameliorate insulin resistance, increase peripheral glucose utilization and improve serum lipid levels. TZDs were shown to activate PPAR γ , which through PRDM16 and the deacetylase sirtuin1 (SIRT1), promote the browning of white fat, which goes in line with the crosstalk between PPAR γ and the growth factors FGF-1 and FGF-21 in maintaining healthy adipose tissue. Additionally, free fatty acids (FFA) mobilized by adipose triglyceride lipase (ATGL) and hormone sensitive lipase (HSL) activate PPAR α and PPAR δ to promote the expression of brown adipocytes genes.

SELECTED REVIEW: PPAR γ signaling and metabolism: the good, the bad and the future: M. Ahmadian, et al.; Nat. Med. 19, 557 (2013)

NEW

Amorfrutin B

AG-CN2-0464

500 μ g

Formula: C₂₆H₃₂O₄

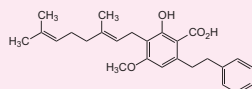
MW: 408.5

CAS: 1174387-94-0

Source: *Amorpha fruticosa*

Natural PPAR γ agonist with potent glucose-lowering properties.

Also available: **Amorfrutin A (AG-CN2-0462)**



Astaxanthin

AG-CN2-0055

5 mg | 25 mg

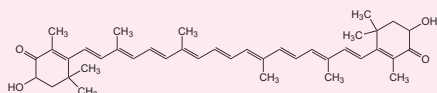
Formula: C₄₀H₅₂O₄

MW: 596.8

CAS: 472-61-7

Source: *Synthetic*

PPAR α agonist and PPAR γ antagonist. Increases insulin sensitivity. Attenuates diabetes associated coagulopathy, oxidative and inflammatory stress. Shows positive effects on obesity and insulin resistance.



Thiazolidinediones (TZDs)

Ciglitazone (Selective PPAR γ agonist)

AG-CR1-0033

1 mg | 5 mg | 25 mg

Pioglitazone (Selective PPAR γ agonist)

AG-CR1-0067

1 mg | 5 mg | 25 mg

Rosiglitazone (Potent PPAR γ agonist)

AG-CR1-3570

25 mg | 100 mg | 1 g

Rosiglitazone . maleate (Potent PPAR γ agonist)

AG-CR1-3571

25 mg | 100 mg | 1 g

Troglitazone (Potent and selective PPAR γ agonist)

AG-CR1-3565

5 mg | 25 mg

Other PPAR Modulators

Asiatic acid (PPAR γ inhibitor)

AG-CN2-0400

5 mg | 25 mg

Curcumin (high purity) (PPAR γ pathway activator)

AG-CN2-0059

10 mg | 50 mg | 250 mg

Genistein (Activates all PPAR isoforms)

AG-CN2-0427

10 mg | 50 mg | 250 mg

GW1929 (Selective PPAR γ agonist)

AG-CR1-0116

1 mg | 5 mg | 25 mg

BULK

Ionomycin (free acid) (PPAR γ ligand with a unique binding mode)

AG-CN2-0416

1 mg | 5 mg

Pseudolaric acid B (PPAR α agonist)

AG-CN2-0083

100 μ g | 1 mg

WY-14643 [Pirinic acid] (Potent PPAR α activator)

AG-CR1-3566

10 mg | 50 mg | 250 mg

Ask for favorable BULK Prices!

BULK

High Purity Streptozotocin

STANDARD Diabetes Inducer

Streptozotocin

AG-CN2-0046-M050

50 mg

AG-CN2-0046-M250

250 mg

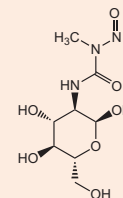
AG-CN2-0046-G001

1 g

Formula: C₈H₁₅N₃O₇

MW: 265.2

CAS: 18883-66-4



α -Glucosidase Inhibitors

Aloeresin A

AG-CN2-0439 500 μ g

Deoxynojirimycin

BVT-0112 1 mg | 5 mg

Luteolin

AG-CN2-0098 5 mg | 25 mg

Pellitorine

AG-CN2-0009 1 mg | 5 mg

Vitexin

AG-CN2-0425 5 mg | 25 mg

N-Methyl-1-deoxynojirimycin

BVT-0130 1 mg

Sirtuin (SIRT) Modulators

AK-7 (Brain-permeable SIRT2 inhibitor)

AG-CR1-3511 5 mg | 25 mg

Hyperforin . DCHA (Potent SIRT1/SIRT2 inhibitor)

AG-CN2-0008 500 μ g | 1 mg

Quercetin . dihydrate (SIRT1 activator)

AG-CN2-0409 1 g | 5 g

Resveratrol (Potent SIRT1 activator)

AG-CN2-0033 50 mg | 100 mg | 500 mg

Sirtinol (Cell permeable SIRT1 inhibitor)

AG-CR1-0055 1 mg | 5 mg | 25 mg

Suramin . 6Na (SIRT1 and SIRT5 inhibitor)

AG-CR1-3575 50 mg | 250 mg | 1 g

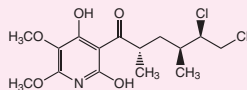
Mitochondrial Modulators

Atpenin A5 (synthetic)

AG-CN2-0100 250 μ g | 1 mgFormula: $C_{15}H_{21}Cl_2NO_5$

MW: 366.2

CAS: 119509-24-9



Potent and specific mitochondrial complex II (succinate-ubiquinone oxidoreductase) inhibitor.

Harzianopyridone (Mitochondrial complex II inhibitor)

AG-CN2-0149 250 μ g | 1 mg

Iromycin A (Mitochondrial electron transport inhibitor)

BVT-0262 500 μ g | 1 mg | 5 mg

PK 11195 (Benzodiazepine receptor antagonist)

AG-CR1-0008 10 mg | 50 mg

Propionyl-L-carnitine (Mitochondrial metabolism stimulator)

AG-CR1-3595 25 mg | 100 mg | 500 mg

Asteltoxin (Mitochondrial respiration inhibitor)

AG-CN2-0441 250 μ g

Citrinin (MPTP activator)

AG-CN2-0101 1 mg | 5 mg | 25 mg

Mitochondrial Membrane Potential Probes

JC-10 (high purity) | JC-1 iodide

see backcover

Potent Nampt/Visfatin Inhibitors from the Manufacturer!

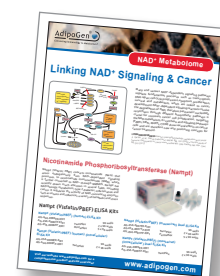
CHS-828

AG-CR1-0064 5 mg | 25 mg

FK-866

AG-CR1-0011 1 mg | 5 mg

Ask for the
**NAD⁺ Metabolome
Product Flyer!**



Reagents for Cardiovascular Research

Agistatin B (Cholesterol biosynthesis inhibitor)

BVT-0223 1 mg | 5 mg

Also available: **Agistatin D & E**

Decarestrictine D (Cholesterol biosynthesis inhibitor)

BVT-0283 1 mg | 5 mg

Beauvericin (ACAT inhibitor)

AG-CN2-0043 1 mg | 5 mg

(3S,6R)-Lateritin (ACAT inhibitor)

AG-CN2-0042 1 mg

(R,R)-Hymeglusin (HMG-CoA synthase inhibitor)

AG-CN2-0103 0.5 mg | 1 mg | 2.5 mg

Lovastatin (HMG-CoA reductase inhibitor)

AG-CN2-0051 5 mg | 25 mg

Penicillide (ACAT inhibitor)

AG-CN2-0122 250 µg | 1 mg

Pyripyropene A (Specific ACAT2 inhibitor)

AG-CN2-0106 250 µg | 1 mg

Sclerotiorin (CETP activity inhibitor)

AG-CN2-0054 0.5 mg | 1 mg | 5 mg

Simvastatin (HMG-CoA reductase inhibitor)

AG-CN2-0052 10 mg | 50 mg

Terpendole C (ACAT1/ACAT2 & CE synth. inhibitor)

AG-CN2-0125 250 µg | 1 mg

Terpendole E (ACAT inhibitor)

AG-CN2-0127 250 µg | 1 mg

Broad Range of Metabolic Research Reagents

Altenusin (MLCK inhibitor)

AG-CN2-0143 1 mg | 5 mg

Amlexanox (Selective TBK1 and IKKε inhibitor)

AG-CR1-3579 10 mg | 50 mg

Amidepsine A (DGAT inhibitor)

AG-CN2-0109 1 mg | 2.5 mg **BULK**

Amidepsine D (DGAT inhibitor)

AG-CN2-0110 1 mg | 2.5 mg

BRS-3 Agonist (Bombesin receptor subtype-3)

AG-CR1-3607 500 µg

Celastrol (Potent leptin sensitizer)

AG-CN2-0460 5 mg | 10 mg | 50 mg

DPVP (Adipogenesis inhibitor)

AG-CR1-3604 1 mg

EM574 (Orexigenic; Motilin receptor agonist)

AG-CN2-0102 250 µg | 1 mg

Emodin (Potent selective 11β-HSD1 inhibitor)

AG-CN2-0457 50 mg | 250 mg

Empagliflozin (SGLT-2 inhibitor)

AG-CR1-3619 10 mg | 50 mg

Geodin (Glucose uptake stimulator)

AG-CN2-0139 1 mg | 5 mg

Itaconic acid (PFKII inhibitor)

AG-CN2-0426 1 g | 5 g

Kaempferitrin (Insulinomimetic/Hypoglycemic)

AG-CN2-0039 1 mg | 5 mg

Linagliptin (DPP4 inhibitor)

AG-CR1-3618 10 mg | 50 mg

Orlistat (DAGLα inhibitor; Antiobesity compound)

AG-CN2-0050 50 mg | 250 mg **BULK**

Salsalate (Anti-inflammatory and antidiabetic)

AG-CR1-3574 1 g | 5 g

Skyrin (Antidiabetic compound)

AG-CN2-0001 1 mg

Stevioside (Antidiabetic compound)

AG-CN2-0077 10 mg | 50 mg **BULK**

BULK

BAIBA

LATEST INSIGHT: The contracting muscle has been shown to act as an endocrine organ, secreting “myokines” that participate in tissue crosstalk. L.D. Roberts, et al. (2014) recently identified the substance BAIBA as a contraction-induced myokine that results in browning of white adipose tissue and increases fat oxidation in the liver.

LIT: β-Aminoisobutyric acid induces browning of white fat and hepatic β-oxidation and is inversely correlated with cardiometabolic risk factors: L.D. Roberts, et al.; Cell Metab. 19, 96 (2014)

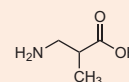
3-Aminoisobutyric acid (BAIBA)

AG-CR1-3596-M250 250 mg | 1 g

Formula: C₄H₉NO₂ · H₂O

MW: 103.1 | 18.0

CAS: 214139-20-5



Also available:

(R)-BAIBA (CDX-A0148) | (S)-BAIBA (CDX-A0147)

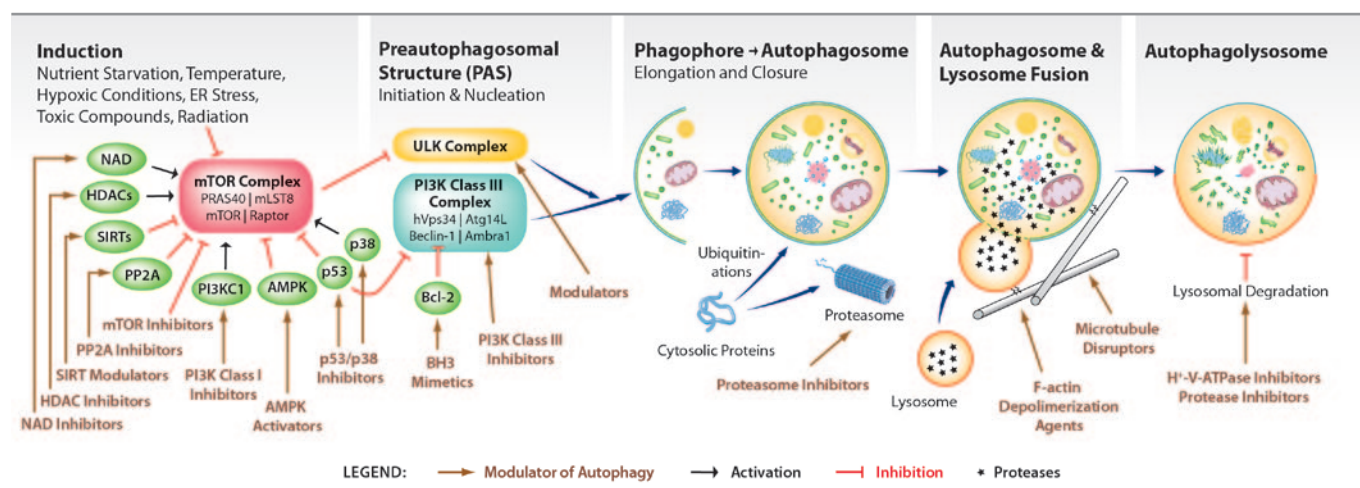
Modulators of Autophagy Signaling

Autophagy represents a homeostatic cellular mechanism for the turnover of organelles and proteins through a lysosome-dependent degradation pathway. Misfolded or aggregated proteins are removed, damaged organelles such as mitochondria, endoplasmic reticulum and peroxisomes are cleared and intracellular pathogens are eliminated. During starvation, autophagy facilitates cell survival through the recycling of metabolic precursors.

Despite the widely accepted role for autophagy in **cellular survival**, it also has been associated with the regulation of

various **cell death pathways**, including regulated (e.g. apoptosis, pyroptosis and necroptosis) and catastrophic (e.g. necrosis) types of cell death. Furthermore it plays a vital role in **innate and adaptive immune mechanisms**. Taken together, autophagy might play a key role in preventing diseases such as cancer, neurodegeneration, cardiomyopathy, diabetes, liver diseases, autoimmune diseases and infections. Strategies aiming to modulate autophagy, such as small molecule activators and inhibitors, might help to elucidate the biological processes behind.

Schematic overview on mammalian macroautophagy, including selected physiological and small molecule modulators



Browse the Catalog Sections for Target-specific Small Molecules!

See Pages 6–8 for AMPK Activators, NAD Inhibitors & SIRT Modulators!

See Pages 12, 19 & 24 for PI3K/mTOR Modulators & HDAC Inhibitors!

See Pages 11 & 14 for Microtubule Disruptors, H⁺-V-ATPase & Proteasome Inhibitors!

PP2A Inhibitors

NEW

NEW Selective PP2A inhibitors

Rubratoxin A

AG-CN2-0092

250 µg | 500 µg

Cytostatin

AG-CN2-0093

250 µg

The Source for Okadaic acids!

Okadaic acid (high purity)

AG-CN2-0056

25 µg | 100 µg | 1 mg

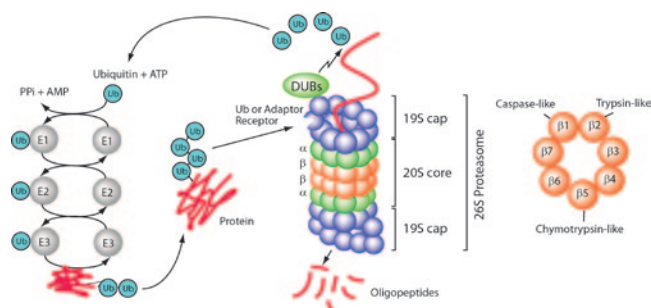
Also available as ammonium, sodium and potassium salts!

Fostriecin

AG-CN2-0057

10 µg

The Ubiquitin-Proteasome System (UPS)



The **ubiquitin-proteasome system (UPS)** and the autophagic-lysosomal pathway are the two major **degradation systems** for both native and misfolded proteins in eukaryotic cells. The regulated proteolysis of bulk and misfolded proteins is strictly controlled by the 26S proteasome complex, which consists of the 19S regulatory cap and the 20S proteasome core. Although eukaryotic 20S proteasomes harbor seven different β -subunits in their two-fold symmetrical $\alpha 7\beta 7\beta 7\alpha 7$ stacked complexes, only three β -subunits per β -ring [**subunits $\beta 1$ (caspase-like), $\beta 2$ (trypsin-like) and $\beta 5$ (chymotrypsin-like)**] are proteolytically active. These three β -subunits are major targets for small molecule proteasome inhibitors. **Proteasome inhibition** has implications in a number of human diseases such as cancer, inflammation and ischemic stroke and is an important therapeutic target.

NEW

Potent 20S Proteasome Inhibitor

Salinosporamide A

AG-CN2-0444-C100

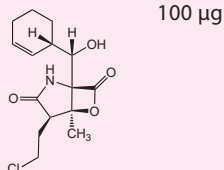
Formula: $C_{15}H_{20}ClNO_4$

MW: 313.8

CAS: 437742-34-2

Source: *Salinispora tropica*

Inhibits all three catalytic activities:
chymotrypsin-like ($EC_{50} = 3.5nM$);
trypsin-like ($EC_{50} = 28nM$); caspase-like ($EC_{50} = 430nM$).

100 μg

Lactacystin

AG-CN2-0104

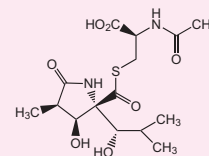
100 μg | 200 μg | 500 μg | 1 mgFormula: $C_{15}H_{24}N_2O_7S$

MW: 376.4

CAS: 133343-34-7

Source: *Streptomyces lactacystiniaeus*

Potent and selective irreversible
proteasome inhibitor. Specifically
inhibits 20S proteasome.



THE STANDARDS

Standard Proteasome Inhibitors – From the Source!

PRODUCT NAME	DESCRIPTION	PID
Bortezomib [PS-341]	Chymotrypsin-like and caspase-like activity inhibitor.	AG-CR1-3602
Epoxomicin	Predominant chymotrypsin-like activity inhibitor.	AG-CN2-0422
clasto-Lactacystin β-lactone	Chymotrypsin-like, trypsin-like and caspase-like activity inhibitor.	AG-CN2-0442
Z-Leu-Leu-Phe-CHO [MG-110]	Chymotrypsin-like activity inhibitor.	AG-CP3-0021
Z-Leu-Leu-Nva-CHO [MG-115]	Chymotrypsin-like activity inhibitor.	AG-CP3-0015
Z-Leu-Leu-Leu-CHO [MG-132]	Chymotrypsin-like and caspase-like activity inhibitor.	AG-CP3-0011
Z-Leu-Leu-Leu-B(OH)₂ [MG-262]	Chymotrypsin-like and caspase-like activity inhibitor.	AG-CP3-0024
b-AP15 [DUB Inhibitor] Solution	Inhibitor of the 19S deubiquitinases (DUBs), ubiquitin-specific-processing protease 14 (USP14) and ubiquitin C-terminal hydrolase isozyme L5 (UCHL5).	AG-CS1-0102

Fluorescent Probes for Proteasome Activity Measurement

Me₄BodipyFL-Ahx₃Leu₃VS	Cell permeable fluorescent proteasome activity probe.	AG-CR1-3601
Ac-Arg-Leu-Arg-AMC	Fluorogenic substrate for the trypsin-like activity of the 20S proteasome.	AG-CP3-0013
Boc-Leu-Arg-Arg-AMC	Fluorogenic substrate for the trypsin-like activity of the 20S proteasome.	AG-CP3-0014
Suc-Leu-Leu-Val-Tyr-AMC	Fluorogenic substrate for measuring the chymotrypsin-like activity of the 20S proteasome.	AG-CP3-0016
Suc-Leu-Tyr-AMC	Fluorogenic substrate for the chymotrypsin-like activity of the 20S proteasome.	AG-CP3-0017
Z-Leu-Leu-Leu-AMC	Fluorogenic substrate for the chymotrypsin-like activity of the 20S proteasome.	AG-CP3-0019
Z-Leu-Leu-Glu-AMC	Fluorogenic substrate for the caspase-like activity of the 20S proteasome.	AG-CP3-0022

PI3K Class I & Class III Inhibitors and mTOR Modulators

BULK

Rapamycin

AG-CN2-0025

100 µg | 1 mg | 5 mg | 25 mg

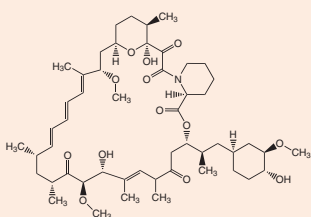
Formula: C₅₁H₇₉NO₁₃

MW: 914.2

CAS: 53123-88-9

Source: *Streptomyces hygroscopicus*

Antitumor compound. Forms a complex with FKBP12 and inhibits the mammalian target of rapamycin (mTOR). Potent immunosuppressant used as an alternative to calcineurin inhibitors.



BULK

LY-294,002

AG-CR1-0108

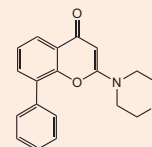
1 mg | 5 mg | 25 mg

Formula: C₁₉H₁₇NO₃

MW: 307.4

CAS: 154447-36-6

Antitumor compound. pan-Class I / II / III PI3K inhibitor. Acts on the ATP-binding site of the enzyme. In solution more stable than wortmannin. Inhibitor of BET bromodomain proteins BRD2, BRD3 and BRD4.



BULK

Wortmannin

AG-CN2-0023

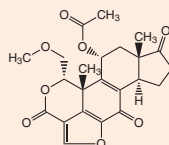
1 mg | 5 mg | 25 mg

Formula: C₂₃H₂₄O₈

MW: 428.4

CAS: 19545-26-7

Antitumor compound. Potent cell permeable and selective inhibitor of phosphatidylinositol 3-kinase (PI3K). Shows similar potency *in vitro* for the class I, II, and III PI3K.



BULK

3-Methyladenine

AG-CR1-3597

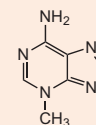
25 mg | 100 mg | 250 mg

Formula: C₆H₇N₅

MW: 149.2

CAS: 5142-23-4

Antitumor compound. Blocks class I, class II and class III PI3Ks, including some downstream targets. Blocks class I PI3K persistently and class III PI3K transiently.



PRODUCT NAME	PID	INHIBITION
AS-252424	SYN-1009	PI3K
AZD-2014	SYN-1167	mTORC1/2
AZD-8055	SYN-1166	PI3K Class I
BEZ235	SYN-1018	PI3K/mTOR
BGT226	SYN-1178	PI3K/Akt
BKM120	SYN-1200	PI3Kα Class I
CH5132799	SYN-1146	PI3Kα Class I
GDC-0032	SYN-1202	PI3K
GDC-0941	SYN-1041	PI3K Class I
GNE-477	SYN-1148	PI3K/mTOR
GNE-490	SYN-1114	pan-PI3K
GNE-493	SYN-1115	pan-PI3K
GSK2126458	SYN-1126	PI3K
IC87114	SYN-1045	PI3K δ Class I
IPI-145	SYN-1175	PI3K Class I
KU-0063794	SYN-1050	mTORC1/2
MLN0128	SYN-1157	mTORC1/2

PRODUCT NAME	PID	INHIBITION
NIBR-17	SYN-1145	pan-PI3K Class I
PF-04979064	SYN-1194	PI3K/mTOR
PI103	SYN-1065	PI3K Class I
PIK-75	SYN-1067	PI3K p110α
PIK-90	SYN-1068	PI3K
PIK-93	SYN-1066	PI3Kα /PI4Kβ
PP242	SYN-1181	mTOR/PI3K
Prodigiosin	AG-CN2-0105	mTORC1/2
RG7422	SYN-1150	PI3K/mTOR
S14161	AG-CR1-3516	PI3K Class I
TASP0415914	SYN-1208	PI3Kγ
TG100115	SYN-1105	PI3Kγ/δ Class I
TGX-221	SYN-1089	PI3K
Viridiol	AG-CN2-0126	PI3K Class III / I
XL388	SYN-1186	mTOR
XL765	SYN-1137	PI3K Class I
ZSTK474	SYN-1099	pan-PI3K

Visit www.adipogen.com for a Broad Panel of Kinase Modulators manufactured by SYNkinase



Broad Panel of Protein Kinase Modulators

BULK

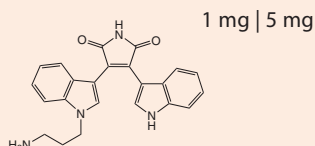
Bisindolylmaleimide III

AG-CR1-0112

Formula: C₂₃H₂₀N₄O₂

MW: 384.4

CAS: 137592-43-9



Also available:

Full Panel of Bisindolylmaleimides I-XI

BULK

Calphostin C

AG-CN2-0430

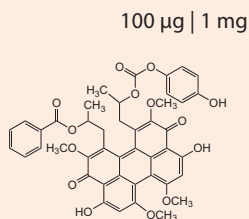
Formula: C₄₄H₃₈O₁₄

MW: 790.8

CAS: 121263-19-2

Source: *Cladosporium cladosporioides* sp.

Potent and highly specific cell permeable PKC inhibitor. Wnt/ β -catenin/lef-1 signaling inhibitor.



BULK

Ibrutinib

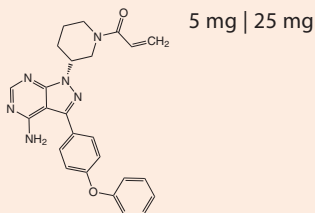
AG-CR1-3620

Formula: C₂₅H₂₄N₆O₂

MW: 440.5

CAS: 936563-96-1

Potent and highly selective Bruton's tyrosine kinase (BTK) inhibitor with antineoplastic activity.



THE SOURCE BULK

Protein Tyrosine Kinase Inhibitors

Herbimycin A | PP1 | PP2 | Radicol

Potent PKC Inhibitors

Staurosporines

see Page 23

Cercosporin (Potent PKC inhibitor)

AG-CN2-0111

1 mg | 5 mg

Chelerythrine chloride (Potent PKC inhibitor)

AG-CR1-0071

1 mg | 5 mg | 25 mg

Anomalin A (Lck inhibitor)

AG-CN2-0006

1 mg

BAY 11-7082 (IkBa kinase inhibitor)

AG-CR1-0013

10 mg | 50 mg

BAY 43-9006 [Sorafenib] (Raf kinase inhibitor)

AG-CR1-0025

1 mg | 5 mg | 25 mg

Debromohymenialdisine (PKC & MEK-1 inhibitor)

AG-CN2-0068

100 μ g

H-8 . 2HCl (PKA and PKG inhibitor)

AG-CR1-0001

10 mg | 50 mg

H-89 . 2HCl (PKA and PKG inhibitor)

AG-CR1-0002

1 mg | 5 mg | 25 mg

(-)-Indolactam V (PKC inducer)

AG-CR1-0041

300 μ g | 1 mg

Ingenol-3-angelate (Specific PKC activator)

AG-CN2-0012

1 mg | 5 mg

MS-1020 (Selective JAK3 inhibitor)

AG-CR1-3501

1 mg

PD 184,352 (MEK-1 inhibitor)

AG-CR1-0029

1 mg | 5 mg

SB202190 (p38 MAPK inhibitor)

AG-CR1-0028

1 mg | 5 mg | 25 mg

SMI-4a (Pim1 & Pim2 inhibitor)

AG-CR1-3503

5 mg

Tripolin A (Aurora A kinase inhibitor)

AG-CR1-3520

1 mg | 5 mg

Tripolin B (Aurora A & B modulator)

AG-CR1-3521

1 mg

CDK, GSK & Dyrk Inhibitors

Alsterpaullone (CDK1/GSK-3 β inhibitor)

AG-CR1-0036

1 mg | 5 mg

6BIO (Potent GSK-3 α/β inhibitor)

AG-CR1-0056

1 mg | 5 mg | 25 mg

(R)-CR8 (CDK1/2/5/7/9, CK1 δ/ϵ & GSK-3 α/β inhibitor)

AG-CR1-0039

1 mg | 5 mg

(S)-CR8 (CDK1/2/5/9, CK1 δ/ϵ & Dyrk1A inhibitor)

AG-CR1-0040

1 mg | 5 mg

Manzamine A (GSK-3 β inhibitor)

AG-CN2-0438

500 μ g

NU6102 (Potent CDK1/CDK2 inhibitor)

AG-CR1-0020

1 mg | 5 mg

Phenylmethylene hydantoin (GSK-3 β inhibitor)

AG-CN2-0041

1 mg

RO-3306 (Selective CDK1 inhibitor)

AG-CR1-3515

1 mg | 5 mg

(R)-Roscovitine (Potent CDK inhibitor)

AG-CR1-0006

1 mg | 5 mg | 50 mg

Also available:

(S)-CR8 | (S)-Roscovitine | (R)-Perharidine 1

Microtubule & F-actin Modulators

THE SOURCE BULK

Latrunculin A

AG-CN2-0027

100 µg | 500 µg

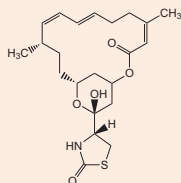
Formula: $C_{22}H_{31}NO_5S$

MW: 421.6

CAS: 76343-93-6

Source: *Latrunculia magnifica*

Actin polymerization inhibitor.
Potent phagocytosis inhibitor.
Anticancer compound.
Inhibits tumor cell invasion.



THE SOURCE BULK

Jasplakinolide

AG-CN2-0037

50 µg | 100 µg

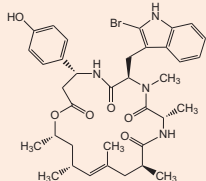
Formula: $C_{36}H_{45}BrN_4O_6$

MW: 709.7

CAS: 102396-24-7

Source: *Jaspis splendens*

Cell permeable, non-fluorescent F-actin probe. Potent inducer of actin polymerization and stabilization. Tool used for autophagy/phagocytosis research.



THE SOURCE BULK

Ilimaquinone

AG-CN2-0038

100 µg

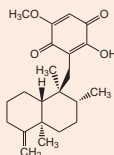
Formula: $C_{22}H_{30}O_4$

MW: 358.5

CAS: 71678-03-0

Source: *Anthelia* sp.

Cytoplasmic microtubule inhibitor. Cell permeable, antimicrobial, anti-inflammatory, antimitotic and cytotoxic compound.



Colcemid (Microtubule assembly inhibitor)

AG-CR1-3567

1 mg | 5 mg

Colchicine (Microtubule assembly inhibitor)

AG-CN2-0048

500 mg | 1 g

Curvulin (Microtubule assembly inhibitor)

BVT-0097

1 mg | 5 mg

Ferulenoil (Stimulator of tubulin polymerization)

AG-CN2-0011

1 mg | 5 mg | 10 mg

Ilimaquinone (Microtubule inhibitor)

AG-CN2-0038

100 µg

Nocodazole (Tubulin depolymerization promoter)

AG-CR1-0019

5 mg | 10 mg | 25 mg | 50 mg

Podophyllotoxin (Microtubule assembly inhibitor)

AG-CN2-0049

100 mg

Paclitaxel (Microtubule assembly stabilizer)

AG-CN2-0045

1 mg | 5 mg | 25 mg | 100 mg

Thiocolchicine (Tubulin polymerization inhibitor)

AG-CN2-0074

5 mg | 25 mg

Vincristine . sulfate (Depolymerizes microtubules)

AG-CN2-0446

5 mg | 25 mg

F-actin Modulators

Cytochalasin H (Actin polymerization inhibitor)

BVT-0447

1 mg | 5 mg

Cytochalasin J (Actin and myosin inhibitor)

BVT-0450

1 mg | 5 mg

Dynasore

AG-CR1-0045

5 mg | 25 mg

Latrunculin B

AG-CN2-0031

1 mg

16-epi-Latrunculin B

AG-CN2-0034

100 µg

Swinholide A

AG-CN2-0035

10 µg | 50 µg

H⁺-V-ATPase Inhibitors

THE SOURCE BULK

Concanamycin A (high purity)

BVT-0237

25 µg | 100 µg | 1 mg

Formula: $C_{46}H_{75}NO_{14}$

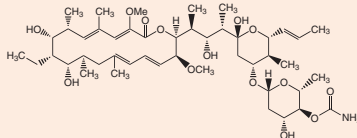
MW: 866.1

CAS: 80890-47-7

Source: *Streptomyces* sp.

Potent and specific
H⁺-ATPase inhibitor.

Also available: **Concanamycin B & C**



THE SOURCE BULK

Bafilomycin A₁ (high purity)

BVT-0252

100 µg | 1 mg

Formula: $C_{35}H_{58}O_9$

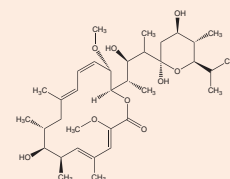
MW: 622.8

CAS: 88899-55-2

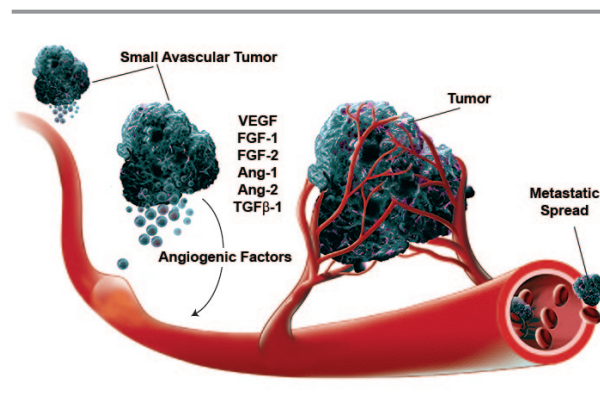
Source: *Streptomyces griseus*

Specific vacuolar-type H⁺-ATPase inhibitor. Distinguishes among different types of ATPases.

Also available: **Bafilomycin B₁ & C₁**



Angiogenesis (VEGF) Modulators



Angiogenesis is a vital and complex biological process, forming new capillaries from pre-existing blood vessels and infusing tissue with supplies of oxygen and nutrients. It plays an important role in physiological conditions such as reproduction, development, wound healing and tissue repair. Aberrant angiogenesis is a crucial mediator in a growing list of diseases such as cancer, chronic inflammatory diseases, atherosclerosis and diabetic retinopathy. The process of angiogenesis is tightly regulated with the involvement of several cell types interacting with each other as well as with the surrounding microenvironment.

Angiogenesis, especially also in tumor, is induced by hypoxia, leading to expression and stabilization of HIF-1 α , a transcription factor that responds to changing oxygen levels, and consequently the transcription of angiogenesis-promoting genes, leading to the upregulation of pro-angiogenic factors, such as VEGF, PDGF, FGF or TGF β . Pro-angiogenic factors activate signaling pathways, such as

PI3K/Akt, Erk1/2, Smad and Notch, which result in endothelial cells (ECs) proliferation and migration of the pre-existing vasculature to sprout and increase vascularization of the tissue. Extensive research continues on anti-angiogenic therapies (biologicals or small molecules) that combat cancer by preventing access to the blood supply that is critical for tumor growth and survival.

SELECTED REVIEWS: Cancer prevention by targeting angiogenesis: A. Albini, et al.; Nat. Rev. Clin. Oncol. 9, 498 (2012) • Angiogenesis in cancer – general pathways and their therapeutic implications; I. Dimova, et al.; JBUON 19, 15 (2014)

PRODUCT NAME	PID	INHIBITION
AG-13958	SYN-1004	VEGF
AMG-Tie2-1	SYN-1008	Tie-2
Axitinib	SYN-1014	VEGFR1-3
BAY 43-9006 [Sorafenib]	AG-CR1-0025	VEGFR2
BMS-2	SYN-1022	VEGFR2
BMS-540215	SYN-1134	VEGFR2
Brivanib alaninate	SYN-1135	VEGFR2
Cabozantinib	SYN-1138	VEGFR2
CYC116	SYN-1034	VEGFR2
E-7080	SYN-1038	VEGFR2/3
Foretinib	SYN-1129	VEGFR2
JNJ-38158471	SYN-1133	VEGFR2
Imatinib mesylate	SYN-1046	PDGFR

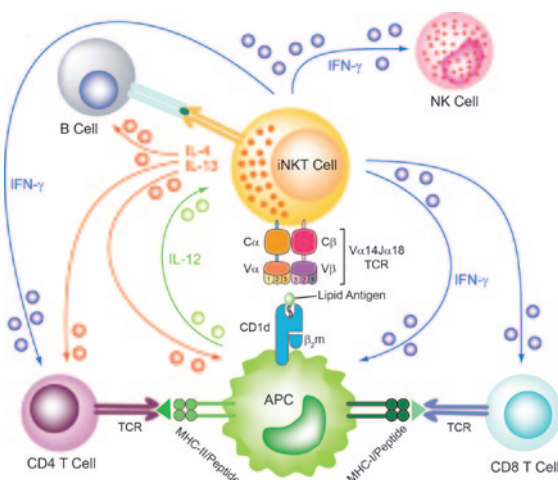
PRODUCT NAME	PID	INHIBITION
Ki20227	SYN-1049	VEGFR2 PDGFR β
Pazopanib . HCl	SYN-1058A	VEGFR1-3 PDGFR
PD-0173074	SYN-1176	FGFR1
Ponatinib	SYN-1116	PDGFR α VEGFR2 FGFR1
Regorafenib	AG-CR1-3623	VEGFR1-3 PDGFR
Salinosporamide A	AG-CN2-0444	VEGF
SAR-131675	SYN-1165	VEGFR3
SU-5402	SYN-1084	VEGFR2 FGFR1
SU-6668	SYN-1085	PDGFR VEGF FGFR
Sunitinib malate	SYN-1086	PDGFR β VEGFR2
Suramin . 6Na	AG-CR1-3575	VEGF
Takeda-6d	SYN-1168	VEGFR2
Tivozanib	SYN-1013	VEGFR1-3
Vandetanib	SYN-1090	VEGFR EGFR

Other Angiogenesis Inhibitors

Ageladine A | Alsterpaullone | BAY 43-9006 | Beauvericin | bpV(phen) | Borrelidin | Ciglitazone | Curcumin | (-)-Epigallocatechin gallate | 17-DMAG | Eupatilin | FK-866 | Fumagillin | Genistein | GW1929 | MS-275 | PLX4720 | Psammaphin A | Pseudolaric acid B | Streptochlorin | Terrein | Thapsigargin | Wortmannin | YC-1

For Compounds from
Marine Sources
www.adipogen.com

CD1d Ligands – Potent iNKT Stimulators



Invariant natural killer T (iNKT) cells are a subset of innate-like lymphocytes that express a characteristic antigen receptor that includes an invariant TCR- α chain and recognize glycolipid antigens bound by the major histocompatibility complex (MHC)-class-I-related protein CD1d. iNKT cells are activated early during a variety of infections and inflammatory diseases and contribute to the subsequent development of adaptive immune responses. Consequently, iNKT cells play a critical role in the development and resolution of inflammatory diseases and represent attractive targets for the development of immunotherapies. In cancer, iNKT cells were attributed a role in immunosurveillance and act as potent activators of antitumor immunity when stimulated with a synthetic agonists.

THE SOURCE

α -Galactosylceramide [KRN7000]

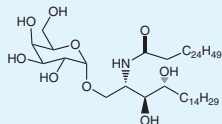
AG-CN2-0013

250 μ g | 1 mgFormula: $C_{50}H_{99}NO_9$

MW: 858.3

CAS: 158021-47-7

Source: Synthetic



α -GalCer Analog I [KBC-007]

AG-CR1-3608

100 μ g

LIT: D.J. Baek, et al.; Chem. Asian J. 5, 1560 (2010)

α -GalCer analog 8

AG-CR1-3622

500 μ g

LIT: T. Lee, et al.; J. Med. Chem. 50, 585 (2007)

OCH (Truncated analog of α -GalCer)

AG-CR1-3593

250 μ g | 1 mg

α -Mannosylceramide

AG-CR1-3594

250 μ g | 1 mg

β -Mannosylceramide

AG-CR1-3621

250 μ g | 1 mg

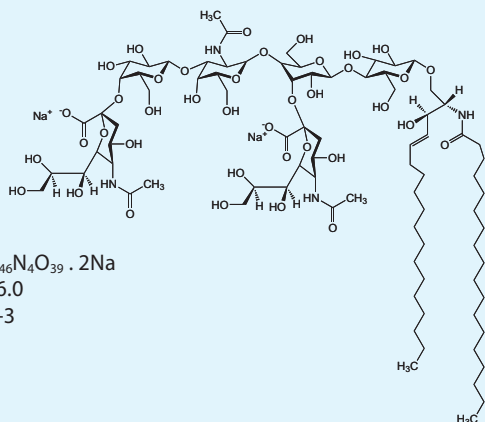
THE SOURCE

Gangliosides

Ganglioside GD1a . disodium salt

AG-CN2-9003

1 mg | 5 mg

Formula: $C_{84}H_{146}N_4O_{39} \cdot 2Na$

MW: 1836.1 . 46.0

CAS: 12707-58-3

Ganglioside GM1 . Na

AG-CN2-9000

1 mg | 5 mg | 10 mg

Ganglioside GM3 . Na

AG-CN2-9002

1 mg

Ganglioside GT1b . 3Na

AG-CN2-9006

1 mg | 5 mg

Ganglioside GQ1b . 4Na

AG-CN2-9007

100 μ g | 500 μ g

Asialo-Ganglioside GM1

AG-CN2-9008

500 μ g | 1 mg

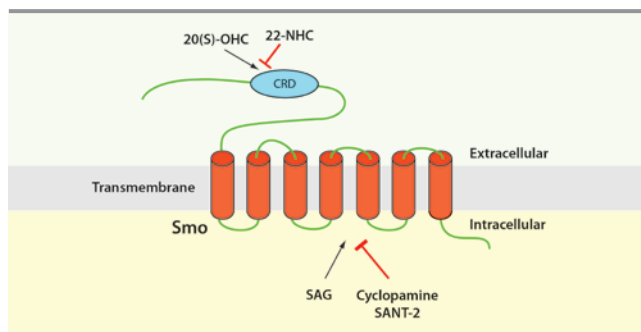
Also available:

Gangliosides GM2, GD3, Asialo-GM2, GD1b

Sonic Hedgehog Modulators

The Hedgehog (Hh) signaling pathway plays key roles in embryonic development, formation and maintenance of cancer stem cells (CSCs) and acquisition of epithelial-to-mesenchymal transition (EMT). Since CSCs and EMT are important biological factors responsible for cancer cell invasion, metastasis, drug resistance and tumor recurrence, inhibition of the Hh signaling pathway is believed to be an important target for cancer therapy. Recently, several small-molecule inhibitors of Hh signaling were developed and synthesized for cancer treatment.

REVIEW: Targeting the Hedgehog signaling pathway for cancer therapy: Y. Li, et al.; Expert Opin. Ther. Targets. 16, 49 (2012)



THE STANDARDS

GANT61

AG-CR1-3561 1 mg | 5 mg

Cyclopamine (Cell permeable Smo inhibitor)

AG-CN2-0028 1 mg | 5 mg

SANT-2 (Cell permeable Smo antagonist)

AG-CR1-3514 1 mg | 5 mg

Curcumin (high purity) (Downregulates Shh & Gli1)

AG-CN2-0059 10 mg | 50 mg | 250 mg

IHR-1 (Smo antagonist)

AG-CR1-3526 1 mg | 5 mg **NEW**

IHR-Nac (Smo antagonist)

AG-CR1-3527 1 mg | 5 mg **NEW**

22-NHC . HCl (Hh inhibitor)

AG-CR1-3523 1 mg | 5 mg

22-NHC inactive (Hh inhibitor control compound)

AG-CR1-3524 1 mg | 5 mg

Resveratrol (Gli1 mRNA expression inhibitor)

AG-CN2-0033 50 mg | 100 mg | 500 mg **BULK**

Robotnikinin (Shh signaling modulator)

AG-CR1-0069 1 mg

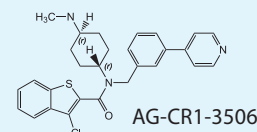
THE SOURCE

SAG – Potent Shh Agonist

LIT: Potent small molecule Hedgehog agonists induce VEGF expression in vitro: K. Seifert, et al.; Bioorg. Med. Chem. 20, 6465 (2012)

SAG AG-CR1-3506
SAG . 2HCl (water soluble) AG-CR1-3585

SAG Analog (LowTox) AG-CR1-3517
SAG Analog (highly active) AG-CR1-3518



Farnesyltransferase Inhibitors

BULK

Manumycin A

BVT-0091 1 mg | 5 mg | 10 mg

Formula: C₃₁H₃₈N₂O₇

MW: 550.6

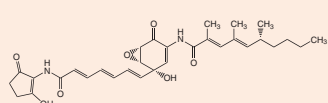
CAS: 52665-74-4

Source: *Streptomyces parvulus*

Potent, selective and competitive cell permeable farnesyltransferase inhibitor.

Also available:

Andrastin A | Manumycin B | Palmarumycin C3



Ubiquitin Related Inhibitors

proTAME Solution (APC/C Ub ligase inhibitor)

AG-CS1-0100 1 mg

MLN4924 Solution (Nedd8 E1 activating enzyme inhibitor)

AG-CS1-0103 1 mg

Apcin (APC/C Ub ligase inhibitor)

AG-CR1-3603 5 mg

Selected Anticancer Small Molecules from the Bench

BULK

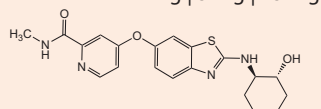
BLZ-945

AG-CR1-3609

Formula: $C_{20}H_{22}N_4O_3S$

MW: 398.5

CAS 953769-46-5



1 mg | 5 mg | 25 mg

Anticancer compound. Potent and selective colony stimulating factor 1 receptor (CSF-1R; c-Fms) inhibitor.

BULK

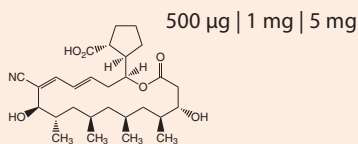
Borrelidin

BVT-0098

Formula: $C_{28}H_{43}NO_6$

MW: 489.6

CAS: 7184-60-3

Source: *Streptomyces* sp.

500 µg | 1 mg | 5 mg

Bacterial, protozoan and mammalian threonyl-tRNA synthetase (THrRS) inhibitor. Antiangiogenic (IC_{50} = 0.8 nM) and anticancer agent.

THE SOURCE

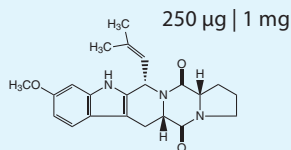
Fumitremorgin C

BVT-0189

Formula: $C_{22}H_{25}N_3O_3$

MW: 379.5

CAS: 118974-02-0

Source: *Aspergillus fumigatus*

250 µg | 1 mg

Mycotoxin. Potent and specific inhibitor of the breast cancer resistance protein (BCRP; ABCG2).

UNIQUE

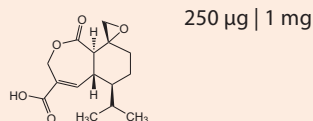
Heptelidic acid

AG-CN2-0118

Formula: $C_{15}H_{20}O_5$

MW: 280.3

CAS: 74310-84-2

Source: *Trichoderma* sp.

250 µg | 1 mg

Potent selective GAPDH inhibitor. Selectively kills high-glycolytic cancer cells through glucose-dependent active ATP deprivation.

UNIQUE

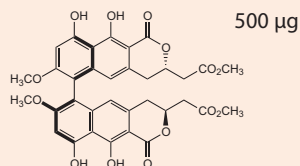
(-)-Viriditoxin

AG-CN2-0471

Formula: $C_{34}H_{30}O_{14}$

MW: 662.6

CAS: 1381782-08-6

Source: *Aspergillus* sp.

500 µg

Bacterial cell division protein FtsZ inhibitor. Apoptosis and autophagy regulator.

Altersolanol A (Anticancer compound)

AG-CN2-0436

1 mg

Avarol (Anticancer compound)

AG-CN2-0044

1 mg

Asperphenamate (Anticancer compound)

AG-CN2-0171

250 µg | 1 mg

Averantin (Antitumor compound)

BVT-0169

1 mg | 5 mg

Becatecarin (Topo II inhibitor)

BVT-0258

250 µg | 1 mg

Bikaverin (Anticancer compound)

AG-CN2-0130

250 µg | 1 mg

BQ-788 . Na (Potent ETBR antagonist)

AG-CP3-0012

500 µg | 1 mg

Decoyinine (GMP synthetase inhibitor)

BVT-0030

1 mg

2,5-Dimethyl-celecoxib (Apoptosis inducer)

AG-CR1-3599

5 mg | 25 mg

Elaiophylin (Autophagy inhibitor and Apoptosis inducer)

BVT-0185

1 mg | 5 mg

Etoposide (DNA topoisomerase II inhibitor)

AG-CR1-3572

25 mg | 100 mg | 5 x 100 mg

Funalenone (MMP-1 inhibitor)

AG-CN2-0137

1 mg

Harzianum A (Anticancer compound)

AG-CN2-0117

250 µg | 1 mg

Neoxaline (Cell cycle inhibitor at G2/M phase)

AG-CN2-0154

1 mg | 5 mg

OM173-αA (DNMT3B inhibitor)

AG-CN2-0158

1 mg | 5 mg

Ophiobolin A (Calmodulin antagonist)

AG-CN2-0431

100 µg | 1 mg

PD 150,606 (Calpain-1/-2 inhibitor)

AG-CR1-0066

5 mg | 25 mg

Piperlongumine (Apoptosis/Necrosis inhibitor)

AG-CN2-0024

10 mg | 50 mg

PJ-34 (Potent PARP inhibitor)

AG-CR1-0100

1 mg | 5 mg | 25 mg

Pyridoxatin (MMP-2 inhibitor)

AG-CN2-0123

250 µg | 1 mg

Trypacidin (Necrosis inhibitor)

BVT-0442

1 mg | 5 mg

Zerumbone (Potent antitumor compound)

AG-CN2-0004

10 mg | 50 mg

HDAC & HAT Inhibitors

For HDAC6 Inhibitors see backcover

Apicidin (HDAC inhibitor)

AG-CN2-0087 1 mg | 5 mg

Butyrolactone 3 (Specific Gcn5 inhibitor)

AG-CR1-3522 2 mg

C646 (p300/CBP HAT inhibitor)

AG-CR1-3508 1 mg | 5 mg

Dihydrochlamydocin (HDAC inhibitor)

AG-CN2-0115 250 µg | 1 mg

5-Hydroxymethylcytosine (Epigenetic marker) **NEW**
BVT-0229 10 mg

MS-275 (HDAC1 inhibitor)

AG-CR1-0032 1 mg | 5 mg | 25 mg

Psammaplin A (Class I HDAC inhibitor)

AG-CN2-0088 100 µg | 500 µg

Splitomicin (Yeast HDAC Sir2p inhibitor)

AG-CR1-0088 1 mg | 5 mg | 25 mg

Trichostatin A (HDAC inhibitor)

AG-CN2-0108 1 mg | 5 mg

Vorinostat [SAHA] (HDAC inhibitor)

SYN-3006 5 mg | 10 mg | 50 mg | 100 mg

pan-Caspase Inhibitors

Q-VD-OPh (pan-Caspase inhibitor)

AG-CP3-0006 1 mg | 3 x 1 mg | 5 mg
For Negative Control see Q-VE-Oph (AG-CP3-0007)

Z-VAD-FMK (cell permeable)

(pan-Caspase inhibitor) **BULK**
AG-CP3-0002 1 mg | 5 mg

Selected Antioxidants, ROS, NO and H₂S Reagents

Eupatilin

AG-CN2-0432

5 mg | 25 mg

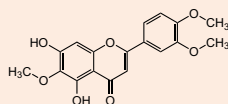
Formula: C₁₈H₁₆O₇

MW: 344.3

CAS: 22368-21-4

Source: *Artemisia* sp.

Selective inhibitor of 5-lipoxygenase. PI3K Class I, MKK3/6 and MKK4 inhibitor. Antioxidant, antidiabetic, anticancer, anti-inflammatory and immunosuppressive compound.



MTSSL

AG-CR1-3573

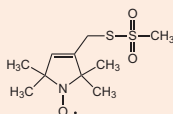
10 mg | 50 mg

Formula: C₁₀H₁₈NO₃S₂

MW: 264.3

CAS: 81213-52-7

Highly reactive, thiol-specific spin-label. Used to label cysteine residues in proteins (site-directed labeling, SDS-labeling).



Auranofin (5-Lipoxygenase inhibitor)

AG-CR1-3611 25 mg | 100 mg

7-Azido-4-methylcoumarin (H₂S probe)

AG-CR1-3525 1 mg | 5 mg

Bradykinin (eNOS/NOSIII activator)

AG-CP3-0025 5 mg | 25 mg

Cinnamtannin B-1 (Potent antioxidant)

AG-CN2-0428 5 mg

COX-2 Inhibitor 8c

AG-CR1-3606 1 mg

DETA NONOate (NO donor)

AG-CR1-3614 5 mg | 25 mg

DMNQ (ROS inducer)

AG-CR1-3598 5 mg | 10 mg | 25 mg

H₂S Donor 5a

AG-CR1-3510 5 mg | 25 mg

Luteolin (Potent free radical scavenger)

AG-CN2-0098 5 mg | 25 mg

iNOS Modulators

1400W (Selective iNOS/NOSII inhibitor)

AG-CR1-0018 5 mg | 25 mg | 100 mg

Xanthorrhizol (Potent COX-2 & iNOS/NOSII inhibitor)

AG-CN2-0090 1 mg

THE STANDARDS

Receptor Agonists and Antagonists

PRODUCT NAME	ACTIVITY	PID	SIZE
Amauromine	CB1 receptor antagonist	AG-CN2-0113	250 µg 1 mg
epi-Aszonalenin A	Substance P inhibitor	AG-CN2-0163	1 mg 5 mg
Bilobalide	GABA(A) receptor antagonist	AG-CN2-0026	10 mg 50 mg
Chaetoviridin A	Monoamine oxidase inhibitor	BVT-0419	1 mg 5 mg
Cyclophenin	AChE inhibitor	AG-CN2-0134	1 mg 5 mg
Debromohymenialdisine	Potential anti-Alzheimer's agent	AG-CN2-0068	100 µg
EM574 [Motilide]	Motilin receptor agonist	AG-CN2-0102	250 µg 1 mg
Finasteride	GABA activity enhancer	AG-CR1-3589	25 mg 100 mg 500 mg
Fulvic acid	Tau and Aβ aggregation inhibitor	AG-CN2-0135	1 mg 5 mg
Hyperforin . DCHA	TRPC6 channel activator	AG-CN2-0008	500 µg 1 mg
20-Hydroxyecdysone	GABA(A) receptor modulator	AG-CN2-0072	5 mg 10 mg 50 mg
Ionomycin (free acid)	Potent Ca ²⁺ ionophore	AG-CN2-0416	1 mg 5 mg
Luteolin	Monoamine transporter activator	AG-CN2-0098	5 mg 25 mg
MTEP	Potent mGluR5 antagonist	AG-CR1-0022	5 mg 25 mg
NG 012	NGF potentiator	AG-CN2-0155	1 mg 5 mg
Papaverine . HCl	PDE10A inhibitor	AG-CN2-0414	1 g 5 g
Pseurotin D	Neuroleptic agent	BVT-0426	1 mg 5 mg
Pimprinine	Monoamine oxidase inhibitor	BVT-0297	1 mg 5 mg
Quinolactacin A	AChE inhibitor	AG-CN2-0164	1 mg 5 mg
Territrem B	AChE inhibitor	AG-CN2-0142	500 µg 1 mg
Thiocolchicoside	GABA(A) receptor antagonist	AG-CN2-0076	1 mg 5 mg 25 mg
Sceptrin . 2HCl	mAChR Inhibitor	AG-CN2-0440	500 µg
SNC80	δ-Opioid receptor agonist	AG-CR1-0017	5 mg 25 mg
SB366791	Potent TRPV1 antagonist	AG-CR1-0034	5 mg 25 mg
Umbellulone	Selective TRPA1 activator	AG-CN2-0085	10 mg
URMC-099	MLK-3 inhibitor	SYN-1211	1 mg 5 mg 10 mg 50 mg



Key Constituents of *Hypericum perforatum*

Hypericin

AG-CN2-0449

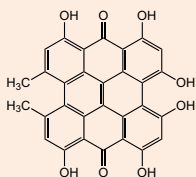
1 mg | 5 mg | 10 mg

Formula: C₃₀H₁₆O₈

MW: 504.5

CAS: 548-04-9

Photosensitive pigment/chromophore with bright red fluorescence emission (λ_{max}: 594nm). Antiviral, anticancer and antidepressant agent.



Hyperforin . DCHA

AG-CN2-0008

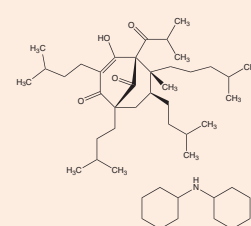
500 µg | 1 mg

Formula: C₃₅H₅₁O₄ . C₁₂H₂₄N

MW: 535.8 . 182.3

CAS: 238074-03-8

Anti-depressant, anti-Alzheimer's, anti-malarial, anti-inflammatory, anticancer and anti-angiogenic compound. Specific activator of TRPC6 channels. Potent SIRT1 and SIRT2 inhibitor.



Neurotoxins

BULK

Tetrodotoxin

AG-CN2-0452 (free base)
AG-CN2-0450 (citrate, water soluble)

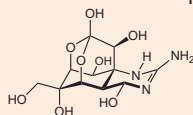
Formula: $C_{11}H_{17}N_3O_8$

MW: 319.3

CAS: 4368-28-9 (citrate free)

Source: *Tetraodontidae*

Potent neurotoxin. Selective, reversible excitable voltage-gated sodium channel (VGSC) blocker.



1 mg
1 mg

Convulxin (GPVI Receptor agonist)

AG-CN2-0465

50 µg

Roquefortine C (Potent neurotoxin)

AG-CN2-0002

1 mg

BVT -0425

500 µg | 1 mg

Verruculogen (Neurotoxin)

BVT -0443

1 mg | 5 mg

Lead Compounds for Neurodegenerative Diseases

Anti-Prion Agents – Protein Aggregation Inhibitors

6-Amino-8-trifluoromethylphenanthridine

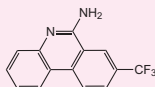
MR-C0031

1 mg | 5 mg | 25 mg

Formula: $C_{14}H_9F_3N_2$

MW: 262.2

CAS: 651055-83-3



6-Aminophenanthridine

MR-C0029

1 mg | 5 mg | 25 mg

Chloroguanabenz acetate

MR-C0036

1 mg | 5 mg

Selective N- & P/Q-type Ca^{2+} Channel Agonist

GV-58

MR-C0035

1 mg | 5 mg

Anti-Alzheimer's Disease (AD) Agent

Leucettine L41

MR-C0023

1 mg | 5 mg | 25 mg

Alzheimer's Disease (AD) Accelerator

Aftin-4

MR-C0014

1 mg | 5 mg | 25 mg

Aftin-5

MR-C0015

1 mg | 5 mg | 25 mg

THE SOURCE BULK

Model Compound for Sensory Studies

Pellitorine

AG-CN2-0009

1 mg | 5 mg

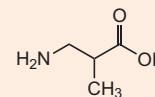
Formula: $C_{14}H_{25}NO$

MW: 223.4

CAS: 18836-52-7

Source: *Synthetic*

Tingling-inducing agent. Excellent stable model compound for sensory studies. Exerts same profile as the unstable compound hydroxy- α -sanshool.



For Compounds from
Plant Sources

www.adipogen.com



ROCK Inhibitors & Stem Cell Modulators

Human embryonic stem cells (hESCs) and human induced pluripotent stem cells (hiPSCs) promise new avenues for medical innovation. These human cells share many similarities with mouse counterparts, including pluripotency and they exhibit several unique properties. A technical problem for various cellular manipulations is the phenomenon of dissociation-induced apoptosis, which is unique to human pluripotent stem cells. This apoptosis is suppressed by ROCK inhibitors and brought a revolutionary change to this troublesome situation.

REVIEW: Lonely death dance of human pluripotent stem cells: ROCKing between metastable cell states: M. Ohgushi & Y. Sasai; Trends Cell Biol. 21, 274 (2011)

Y-27632 . 2HCl

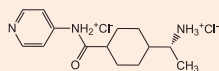
AG-CR1-3564

1 mg | 5 mg | 25 mg

Formula: C₁₄H₂₁N₃O . 2HCl

MW: 247.3 . 73.0

CAS: 129830-38-2



Enhances the survival and cloning efficiency of dissociated human embryonic stem (hES) cells without affecting their pluripotency. Increases survival rate of hES cells undergoing cryopreservation. Prevents apoptosis.

GSK429286A (Potent ROCK-I inhibitor)

SYN-1071

1 mg | 5 mg | 10 mg | 50 mg | 100 mg

IWR-1-endo (Wnt pathway signaling inhibitor)

AG-CR1-3581

5 mg | 25 mg

MS-275 (Promotes differentiation of ES cells)

AG-CR1-0032

1 mg | 5 mg | 25 mg

Pifithrin-α . HBr (p53 inhibitor)

AG-CR1-0004

5 mg | 10 mg | 25 mg

NU6102 (Potent CDK1/CDK2 inhibitor/ROCK-II inhibitor)

AG-CR1-0020

1 mg | 5 mg

Shz-1 (Enhancer of regenerative repair)

AG-CR1-3502

1 mg | 5 mg

SR3677 (Potent ROCK-II and ROCK-I inhibitor)

SYN-1083

1 mg | 5 mg | 10 mg | 50 mg | 100 mg

Trichostatin A (Promotes differentiation of ES cells)

AG-CN2-0108

1 mg | 5 mg

LATEST REVIEW: Rho Kinase (ROCK) inhibitors and their therapeutic potential: Y. Feng, et al.; J. Med. Chem. (Epub ahead of print) (2015)

Other Selected Inhibitors

Apicidin (Promotes differentiation of ES cells)

AG-CN2-0087

1 mg | 5 mg

BIX 01294 (free base) (Stem cell inducer)

AG-CR1-0051

1 mg | 5 mg | 25 mg

BULK

Notch Processing Inhibitors

Compound E

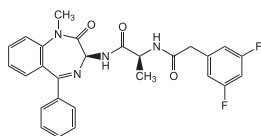
AG-CR1-0081

250 µg | 1 mg | 5 mg

Formula: C₂₇H₂₄F₂N₄O₃

MW: 490.5

CAS: 209986-17-4



Non-competitive γ-secretase inhibitor. Notch processing inhibitor.

DAPT

AG-CR1-0016

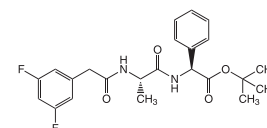
5 mg | 25 mg

Formula: C₂₃H₂₆F₂N₂O₄

MW: 432.5

CAS: 208255-80-5

Cell permeable γ-secretase inhibitor. Notch processing inhibitor.



BULK

High Purity IBMX

Enhances Differentiation of 3T3-L1 Cells

IBMX

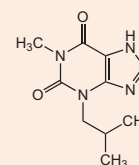
AG-CR1-3512

500 mg | 1 g

Formula: C₁₀H₁₄N₄O₂

MW: 222.3

CAS: 28822-58-4



COMMODITIES out of the Box

BULK from Original Source

High Purity Detergents

MEGA-8 | n-Octyl- β -D-thioglucoopyranoside | n-Dodecyl- α -D-maltoside | n-Dodecyl- β -D-maltoside

Protease Inhibitors

AEBSF | Amastatin | Antipain | Leupeptin | Paptstatin

HSP90 Inhibitors

Geldanamycin | 17-AAG | 17-DMAG

Gene Cloning Tools

IPTG (animal-free, dioxane-free) | X-Gal

Cell Selective Agents

G418 . sulfate | Puromycin

AC & sGC Activators

Forskolin | YC-1

Tumor Promoters

Phorbol 12-myristate 13-acetate [TPA/PMA] | Thapsigargin

Staurosporine & Analogs

Staurosporine | K-252a | K-252c

Ecdysteroid Agonists Gene Expression Inducer

Ponasterone A | Muristerone A | Makisterone A | Ecdysone | 20-Hydroxyecdysone | Halofenozide | Methoxyfenozide

HIF Inhibitors

Chetomin | Echinomycin

Bile Acid Nuclear Receptor Modulators

6-ECDCA | Chenodeoxycholic acid | Ursodeoxycholic acid | Stigmasterol

Potent Platinum-based Antineoplastic Agents

Carboplatin | Cisplatin | Oxaliplatin

Other Products

Thaxtomin A (Plant cell necrosis inducer)

BVT-0206 1 mg | 5 mg

Actinomycin D (RNA synthesis inhibitor)

BVT-0089 5 mg | 25 mg | 100 mg

4-Aminophenylphosphate . Na

AG-CR1-3586 10 mg | 50 mg | 100 mg

(+)-Brefeldin A (Protein transport inhibitor)

AG-CN2-0018 5 mg | 10 mg | 25 mg

(+)-Aphidicolin

BVT-0307

Formula: $C_{20}H_{34}O_4$

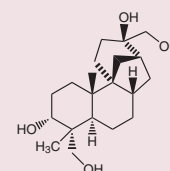
MW: 338.5

CAS: 38966-21-1

Source: *Phoma* sp. BS 7210

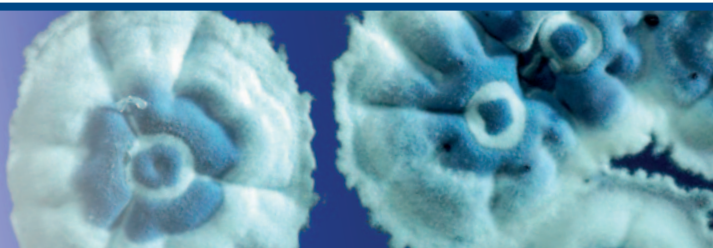
Antiviral and antineoplastic agent. Specific DNA polymerase α and δ inhibitor in eukaryotic cells and in some viruses of animal origin.

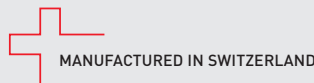
1 mg | 5 mg | 25 mg



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JC-10 (high purity)

AG-CR1-3600

MITOCHONDRIA

1 mg | 5 mg

Shows improved solubility compared to JC-1 in aqueous media. Useful for determining MMP in cells by flow cytometry, fluorescence microscopy and in microplate-based fluorescent assays. Accumulates in mitochondria of cells with a polarized mitochondrial membrane. Generates an orange J-aggregate emission profile ($\lambda_{ex} = 540 \text{ nm}$ / $\lambda_{em} = 590 \text{ nm}$) in healthy cells and a shift to green emission ($\lambda_{ex} = 490 \text{ nm}$ / $\lambda_{em} = 525 \text{ nm}$) in injured or dead cells.

Wavelength Maxima: $\lambda_{ex} = 510 \text{ nm}$ | $\lambda_{em} = 525 \text{ nm}$.

Also available: **JC-1 iodide (AG-CR1-3568)**

LysoGlow84™

AG-CMA-1005

ACIDIC ORGANELLES

1 mg

Cell/membrane permeable, non-toxic, pH-sensitive fluorescent dye (blue-green range) for live imaging of pH alteration in acidic organelles, lysosomes, autophagolysosomes, vesicles, cells, tissue and even small, transparent whole animals without side effects. Useful as intracellular pH indicator with a wide range of fluorescence (pH 3 to pH 13).

Wavelength Maxima:

$\lambda_{ex} = 320 \text{ nm}$ | $\lambda_{em} = 440 \text{ nm}$ (pH 3-6) and 400 nm (pH 8-13)

Also available: **Ageladine A . TFA (AG-CMA-1001)**

NEW

Potent & Selective HDAC6 Inhibitors

HDAC6 deacetylates tubulin, Hsp90 and the core histones (H2A, H2B, H3, H4). Histone deacetylases act via the formation of large multiprotein complexes. HDAC6 plays an important role in microtubule-dependent cell motility, transcriptional regulation, degradation of misfolded proteins and cell cycle and is involved in autophagy, inflammation, cancer and neurodegeneration.

PRODUCT NAME	PID	SIZE	HDAC6 INHIBITION	SELECTIVITY TOWARDS OTHER HDACS
Tubastatin A [TubA]	AG-CR1-3900	1 mg 5 mg	IC ₅₀ =15nM	Other HDACs (IC ₅₀ = >15μM), HDAC8 (IC ₅₀ =0.9μM)
Nexturastat A	AG-CR1-3901	1 mg 5 mg	IC ₅₀ =5.02nM	Other HDACs (IC ₅₀ =3-10μM)
Nexturastat B	AG-CR1-3902	1 mg 5 mg	IC ₅₀ =3nM	HDAC1 (IC ₅₀ =0.9μM)
ACY-775	AG-CR1-3903	1 mg 5 mg	IC ₅₀ =7.5nM	HDAC1-9 (IC ₅₀ =1-10μM)
DMAPB	AG-CR1-3904	1 mg 5 mg	IC ₅₀ =114nM	Other HDACs (IC ₅₀ =1-8μM)
PMPH	AG-CR1-3905	1 mg 5 mg	IC ₅₀ =11nM	HDAC1 (IC ₅₀ =1.5μM)
DABPH	AG-CR1-3906	1 mg 5 mg	IC ₅₀ =12nM	HDAC1 (IC ₅₀ =6.8μM)
MBIMPH	AG-CR1-3907	1 mg 5 mg	IC ₅₀ =9nM	Other HDACs (IC ₅₀ =0.1-12μM)
MBIMPH F-Analog 1 . HCl	AG-CR1-3908	1 mg 5 mg	IC ₅₀ =3nM	Other HDACs (IC ₅₀ =0.03-20μM)
MBIMPH F-Analog 2	AG-CR1-3909	1 mg 5 mg	IC ₅₀ =5nM	Other HDACs (IC ₅₀ =0.2-11μM)