

Teprotumumab

Targets (1)



This drug entry is a **stub** and has not been fully annotated. It is scheduled to be annotated soon.

IDENTIFICATION

Name	Teprotumumab		
Accession Number	DB06343		
Type	Biotech		
Groups	Investigational		
Biologic Classification	Protein Based Therapies Monoclonal antibody (mAb)		
Description	A fully human IgG1 type monoclonal antibody directed against the human insulin-like growth factor-I receptor.		
Protein chemical formula	Not Available		
Protein average weight	Not Available		
Sequences	Not Available		
Synonyms	Immunoglobulin G1, anti-(human insulin-like growth factor I receptor) (human monoclonal heavy chain), disulfide with human monoclonal light chain, dimer		
External IDs ⓘ	R-1507 / R1507 / RG-1507 / RG1507 / RO-4858696 / RO-4858696-000 / RO-4858696000 / RO4858696 / RO4858696-000 / RV-001 / RV001		
Categories	Amino Acids, Peptides, and Proteins	Blood Proteins Globulins Immunoglobulins	Immunoproteins Proteins Serum Globulins
UNII	Y64GQ0KCOA		
CAS number	1036734-93-6		

PHARMACOLOGY

Indication	Investigated for use/treatment in solid tumors.
Pharmacodynamics	Not Available
Mechanism of	R1507 (formerly called Roche 1) is a fully human antibody which targets the Insulin-like Growth Factor-1 Receptor (IGF-1R). The IGF-1R molecule has been shown to be important in tumor growth

Drugs



for an antibody therapeutic approach. [Genmab Website]

TARGET	ACTIONS	ORGANISM
Insulin-like growth factor 1 receptor	Not Available	Humans



Absorption Not Available

Volume of distribution Not Available

Protein binding Not Available

Metabolism Not Available

Route of elimination Not Available

Half life Not Available

Clearance Not Available

Toxicity Not Available

Affected organisms Not Available

Pathways Not Available

Pharmacogenomic Effects/ADRs [i](#) Not Available

INTERACTIONS

Drug Interactions

**ALL DRUGS**

APPROVED

VET APPROVED

NUTRACEUTICAL

ILLICIT

WITHDRAWN

INVESTIGATIONAL

EXPERIMENTAL

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DRUG	INTERACTION
Abciximab	The risk or severity of adverse effects can be increased when Abciximab is combined with Teprotumumab.
Abituzumab	The risk or severity of adverse effects can be increased when Teprotumumab is combined with Abituzumab.
Adalimumab	The risk or severity of adverse effects can be increased when Adalimumab is combined with Teprotumumab.
Adecatumumab	The risk or severity of adverse effects can be increased when Adecatumumab is combined with Teprotumumab.
Aducanumab	The risk or severity of adverse effects can be increased when Teprotumumab is combined with Aducanumab.
Afelimomab	The risk or severity of adverse effects can be increased when Afelimomab is combined with Teprotumumab.
Alemtuzumab	The risk or severity of adverse effects can be increased when Alemtuzumab is combined with Teprotumumab.
Alirocumab	The risk or severity of adverse effects can be increased when Teprotumumab is combined with Alirocumab.

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	Amatuximab.
AMG 108	The risk or severity of adverse effects can be increased when AMG 108 is combined with Teprotumumab.



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Food Interactions Not Available

REFERENCES

General References Not Available

External Links Wikipedia [Teprotumumab](#)

CLINICAL TRIALS

Clinical Trials

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PHASE	STATUS	PURPOSE	CONDITIONS	COUNT
1	Completed	Basic Science	Cancer, Breast	1
1	Completed	Treatment	Diabetic Macular Edema (DME)	1
1	Completed	Treatment	Neoplasms	2
1	Terminated	Treatment	Neoplasms	2
2	Completed	Treatment	Cancer, Breast	1
2	Completed	Treatment	Non-Squamous Non-Small Cell Lung Cancer	1
2	Completed	Treatment	Ophthalmopathy, Thyroid-Associated / Thyroid Associated Ophthalmopathy	1
2	Completed	Treatment	Sarcomas	1
2	Terminated	Treatment	Lung Cancer Non-Small Cell Cancer (NSCLC)	1
3	Active Not Recruiting	Treatment	Graves' Orbitopathy / Thyroid Eye Disease	1

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PHARMACOECONOMICS

Manufacturers Not Available

Packagers Not Available

Dosage forms Not Available

Prices Not Available

Patents Not Available

PROPERTIES

State Solid

Experimental Properties Not Available

TAXONOMY

Kingdom	Organic Compounds
Super Class	Organic Acids
Class	Carboxylic Acids and Derivatives
Sub Class	Amino Acids, Peptides, and Analogues
Direct Parent	Peptides
Alternative Parents	Not Available
Substituents	Not Available
Molecular Framework	Not Available
External Descriptors	Not Available



TARGETS

1. Insulin-like growth factor 1 receptor

[Details](#)

Kind	Protein
Organism	Humans
Pharmacological action	Unknown
General Function	Protein tyrosine kinase activity
Specific Function	Receptor tyrosine kinase which mediates actions of insulin-like growth factor 1 (IGF1). Binds IGF1 with high affinity and IGF2 and insulin (INS) with a lower affinity. The activated IGF1R is involv...
Gene Name	IGF1R
Uniprot ID	P08069
Uniprot Name	Insulin-like growth factor 1 receptor
Molecular Weight	154791.73 Da

Drug created on March 19, 2008 10:25 / Updated on November 02, 2018 06:16

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