

Chemokines in disease - biology and clinical research

Humana Press - Disease activity, cytokines, chemokines and the risk of incident diabetes in rheumatoid arthritis — Research Nebraska

Table 1 Representative Clinical Trials of Anti-Inflammatory Treatments in Type 1 Diabetes

| Mechanism of action | Drug | Main findings | References |
|--|----------------------------------|--|------------|
| Monoclonal anti-CD3 antibody | Teplizumab | Reduction of C-peptide decline, lower insulin requirements, HbA _{1c} ↓ | 48, 50 |
| Engineered DNA plasmid encoding granulysin (GAP-100) | GAP-100 | ↑ CD4 ⁺ T-cell frequency, reduction in C-peptide presentation, no change in HbA _{1c} or insulin requirements | 51 |
| Proteinase inhibitor | Human kallikrein inhibitor (HAI) | ↑ C-peptide, ↓ granulysin-stimulated IL-1 production, ↓ autoantibody (GAD65) titers | 189 |
| TGF-β antagonist | SB-415286 | HbA _{1c} ↓, insulin requirements ↓ | 52 |
| Anti-inflammatory protein | Alpha-1 antitrypsin (AAT) | ↓ T-cell response to mycoplasma and dendritic cells, ↓ autoantibody titers | 53 |
| Glucocorticoid | Hydrocortisone | Reduction in C-peptide decline, ↓ insulin requirements | 54 |
| Glucocorticoid | Hydrocortisone | Reduction in C-peptide decline, ↓ insulin requirements | 55 |
| Glucocorticoid | Hydrocortisone | Reduction in C-peptide decline, ↓ insulin requirements | 56 |
| Glucocorticoid | Hydrocortisone | Reduction in C-peptide decline, ↓ insulin requirements | 57 |
| Glucocorticoid | Hydrocortisone | Reduction in C-peptide decline, ↓ insulin requirements | 58 |
| Glucocorticoid | Hydrocortisone | Reduction in C-peptide decline, ↓ insulin requirements | 59 |
| Glucocorticoid | Hydrocortisone | Reduction in C-peptide decline, ↓ insulin requirements | 60 |
| Glucocorticoid | Hydrocortisone | Reduction in C-peptide decline, ↓ insulin requirements | 61 |
| Glucocorticoid | Hydrocortisone | Reduction in C-peptide decline, ↓ insulin requirements | 62 |

Description: -

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Receptors, Chemokine

Chemokines

Chemokines -- Physiological effect

Chemokines -- Pathophysiology

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Contemporary immunology

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Notes: Includes bibliographical references and index.

This edition was published in 1999



Filesize: 11.49 MB

Tags: #Corrigendum #to #and #Heart #Disease: #A #Network #Connecting #Cardiovascular #Biology #to #Immune #and #Autonomic #Nervous

Chemokines and Heart Disease: A Network Connecting Cardiovascular Biology to Immune and Autonomic Nervous Systems

The further testing of these changes is likely to facilitate our understanding of the underlying mechanism responsible for ocular damage in Sjögren's syndrome. Today, it is clear that chemokines affect all aspects of immunology and even many unrelated fields, such as tissue development and tumor cell metastasis. The actions of chemokines are controlled by their respective receptors, which, when activated, can uncouple from associated G proteins, internalize, and degrade the ligand.

Chemokines in disease : biology and clinical research (Book, 1999) [perssongroup.materialsproject.org]

Because of its induction or upregulation during CNS pathologies, members of the chemokine system can be used as biological markers. Relation to onset types, disease activity, and synovial fluid leukocytes.

Chemokine Biology Basic Research And Clinical Application Vol 2 Pathophysiology Of Chemokines Progress In Inflammation Research PDF Book

A detailed review of the emerging role of chemokines in viral biology is also presented, with emphasis on HIV biology and novel therapeutic possibilities.

Chemokines and disease

The effectiveness of mobilizing hNSCs with de novo designed agonists may lead to new translational therapeutics for the clinical repair of CNS injuries and other neurodegenerative conditions.

Chemokines In Disease Biology And Clinical Research Contemporary Immunology PDF Book

Mutation Analysis of Receptors and Relationship of Receptor Usage to Disease, Emma J. CCR5 + and CXCR3 + T cells are increased in multiple sclerosis and their ligands MIP- α and IP-10 are expressed in demyelinating brain lesions. The third, 5P14-RANTES, induces significant levels of

CCR5 internalization without detectable signaling activity.

Chemokine Biology Basic Research And Clinical Application Vol 2 Pathophysiology Of Chemokines Progress In Inflammation Research PDF Book

IL-8 in Animal Models of Disease, Akihisa Harada and Kouji Matsushima. Incident DM was defined based on validated algorithms using diagnostic codes and medications. The Role of ELR+-CXC Chemokines in Wound Healing and Melanoma Biology, Ann Richmond, Jing Luan, Jianguo Du, and Hamid Haghnegahdar.

CHEMOKINES IN DISEASE BIOLOGY AND CLINICAL RESEARCH 1999 Edition, 9780896037038, HÉBERT CAROLINE A., SPRINGER

Thus, a consistent increase in MIF levels during TRT therapy suggests its possible association with increased inflammatory activity. Disease activity and clinical assessments occur longitudinally as part of clinical care. Zhou N, Luo Z, Luo J, Fan X, Cayabyab M, Hiraoka M, Liu D, Han X, Pesavento J, Dong CZ, Wang Y, An J, Kaji H, Sodroski JG, Huang Z.

Chemokines in Disease

Two representative SMM-chemokines, RCP168 and RCP188, selective for CXCR4 and CCR5, respectively, showed similar or significantly enhanced binding affinities for their corresponding target receptors. ROS, adenosine, and complement activate mast cells to produce TNF and histamine, leading to leukocyte recruitment from the vessels. Hébert Series Title Copyright 1999 Publisher Humana Press Copyright Holder Springer Science+Business Media New York eBook ISBN 978-1-59259-706-2 DOI 10.

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