Author's Accepted Manuscript

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PII: S0014-2999(18)30356-X

https://doi.org/10.1016/j.ejphar.2018.06.025 DOI:

Reference: EJP71855

To appear in: European Journal of Pharmacology

Received date: 14 May 2018 Revised date: 8 June 2018 Accepted date: 19 June 2018

Cite this article as: János G. Filep, Meriem Shekeri and Driss El Kebir, Targeting formyl peptide receptors to facilitate the resolution of inflammation, European Journal of Pharmacology, https://doi.org/10.1016/j.ejphar.2018.06.025

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Targeting formyl peptide receptors to facilitate the resolution of inflammation

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Abstract

The formyl peptide receptors (FPRs) are G protein coupled receptors that recognize a broad range of structurally distinct pathogen and danger-associated molecular patterns and mediate host defense to infection and tissue injury. It became evident that the cellular distribution and biological functions of FPRs extend beyond myeloid cells and governing their activation and trafficking. In recent years, significant progress has been made to position FPRs at check points that control the resolution of inflammation, tissue repair and return to homeostasis.

Accumulating data indicate a role for FPRs in an ever-increasing range of human diseases, including atherosclerosis, chronic obstructive pulmonary disease, asthma, autoimmune diseases and cancer, in which dysregulated or defective resolution are increasingly recognized as critical component of the pathogenesis. This review summarizes recent advances on how FPRs

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