

Title (en)
HUMAN CRABP-I AND CRABP-II.

Title (de)
MENSCHLISCHES CRABP-I UND CRABP-II.

Title (fr)
CRABP-I ET CRABP-II HUMAINES.

Publication
EP 0640093 A4 19960605 (EN)

Application
EP 93910828 A 19930427

Priority
• US 9303936 W 19930427
• US 87484792 A 19920428

Abstract (en)
[origin: WO9322331A1] The sequences encoding two isoforms of human cellular retinoic acid binding proteins, CRABP-I and CRABP-II, have been cloned and sequenced and are set forth with their corresponding amino acid sequences in SEQ ID NOS. 1-4. The identification of human CRABP nucleic and amino acid sequences provides the basis for the construction of recombinant human CRABP vectors and expression constructs. Human CRABP can also be synthesized or produced ex vivo, e.g. in bacterial or other production systems. Ligand binding assays, including recombinant and chimeric receptor reporter assays, and direct and competition hybridization assays employing the human CRABP sequences herein described can be used to test drugs for retinoic induction and tissue specificity for pathologies in which retinoids are implicated. Immunoassays utilizing antibodies or binding fragments produced to human CRABP can also be used to test patient tissues for the presence and levels of CRABP for diagnosis and to monitor treatment. The identification of the nucleic and amino acids sequences for human CRABP-I and CRABP-II also contributes to the elucidation of the function and interaction of the retinoid-binding proteins. The CRABP-II gene, isolated from a human placenta genomic library, spans 6 kilobases and includes 4 exons. One major transcription initiation site was mapped to an A residue 137 nucleotides upstream of the ATG initiation codon. CRABP-II mRNA was rapidly induced within 2-6 hours in culture by retinoic acid, primarily due to an increased rate of transcription which required on-going synthesis. The human CRABP-II gene is thus apparently transcriptionally regulated by a newly synthesized regulator protein. Once the CRABP-II is produced, message stabilization may provide means by which elevated CRABP-II in mRNA is maintained.

IPC 1-7
C07H 21/04; C12Q 1/68; C07K 15/26; C12N 15/12; C12N 15/63; G01N 33/53

IPC 8 full level
A61K 39/395 (2006.01); **C07H 21/04** (2006.01); **C07K 14/705** (2006.01); **C12N 15/09** (2006.01); **C12N 15/12** (2006.01); **C12P 21/02** (2006.01); **C12Q 1/68** (2006.01); **G01N 33/53** (2006.01); **G01N 33/566** (2006.01)

CPC (source: EP)
C07K 14/70567 (2013.01); **C12Q 1/6897** (2013.01); **G01N 2333/70567** (2013.01); **G01N 2500/00** (2013.01)

Citation (search report)
• [A] EP 0325849 A2 19890802 - SALK INST FOR BIOLOGICAL STUDI [US] & US 4981784 A 19910101 - EVANS RONALD M [US], et al
• [X] ELLER M S ET AL: "THE MOLECULAR CLONING AND EXPRESSION OF TWO CRABP CDNAS FROM HUMAN SKIN.", EXP CELL RES 199(2). 1992. 328-336. CODEN: ECREAL ISSN: 0014-4827
• [DX] ASTROM, ANDERS ET AL: "Molecular cloning of two human cellular retinoic acid-binding proteins (CRABP). Retinoic acid-induced expression of CRABP -II but not CRABP -I in adult human skin in vivo and in skin fibroblasts in vitro", J. BIOL. CHEM. (1991), 266(26), 17662-6 CODEN: JBCHA3;ISSN: 0021-9258
• [DA] ANDERS ASTRÖM ET AL.: "Retinoic acid and synthetic analogs differentially activate retinoic acid receptor dependent transcription", BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS, vol. 173, no. 1, 30 November 1990 (1990-11-30), ORLANDO, FL US, pages 339 - 345, XP000570308, DOI: doi:10.1016/S0006-291X(05)81062-9
• [PX] ANDERS ASTRÖM ET AL.: "Structure of the human cellular retinoic acid-binding protein II gene", JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 267, no. 35, 15 December 1992 (1992-12-15), MD US, pages 25251 - 25255
• See references of WO 9322331A1

Cited by
US6092619A

Designated contracting state (EPC)
AT BE CH DE DK ES FR GB GR IE IT LI LU MC NL PT SE

DOCDB simple family (publication)
WO 9322331 A1 19931111; AU 4118393 A 19931129; CA 2134550 A1 19931111; EP 0640093 A1 19950301; EP 0640093 A4 19960605; JP H08500963 A 19960206

DOCDB simple family (application)
US 9303936 W 19930427; AU 4118393 A 19930427; CA 2134550 A 19930427; EP 93910828 A 19930427; JP 51945593 A 19930427