

## Title (en)

CLONING AND PRODUCTION OF POLYPEPTIDE ANALOGS OF HUMAN FIBRONECTIN AND METHOD OF USING SUCH POLYPEPTIDE ANALOGS

## Publication

**EP 0451211 A4 19920909 (EN)**

## Application

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## Priority

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## Abstract (en)

[origin: WO9007577A1] This invention provides plasmids for bacterial expression of polypeptides which comprise a substantial portion of the amino acid sequence of, and which have the biological activity of, one of the domains of naturally-occurring human fibronectin, such as the cell binding domain or fibrin binding domain, comprising DNA encoding the polypeptide and DNA encoding suitable regulatory elements positioned relative to the DNA encoding the polypeptide so as to effect expression of the polypeptide in a suitable host cell. In the presently preferred embodiments of the invention, the polypeptide is a 75 kD, 40 kD or 33 kD polypeptide of the cell binding domain, or a 31 kD or 20 kD polypeptide of the fibrin binding domain. The invention also provides methods for producing the polypeptides and pharmaceutical compositions comprising the polypeptides and pharmaceutically acceptable carriers. The polypeptides of this invention may be used to inhibit platelet aggregation, to inhibit thromboxane release from platelets, or to treat a subject with a cerebrovascular disorder, a cardiovascular disorder, a wound, a bacterial infection, a cancer, or to detect a fibrin thrombi. The invention further provides the polypeptides conjugated to thrombolytic agents, growth factors, serum albumin, blood factors, or polyethyleneglycol.

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## IPC 8 full level

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- [X] THE JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 258, no. 5, 10th March 1983, pages 3332-3340, Baltimore, US; M. HAYASHI et al.: "Domain structure of the carboxyl-terminal half of human plasma"
- [X] PROC. NATL. ACAD. SCI. USA, vol. 80, January 1983, pages 137-141; T.E. PETERSEN et al.: "Partial primary structure of bovine plasma fibronectin: Three types of internal homology"
- See references of WO 9007577A1

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