

# **EXHIBIT 3**



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**United States Patent** [19]

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**Lin**

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- [54] **PRODUCTION OF ERYTHROPOIETIN**
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- 86/03520 6/1986 WIPO .

**Related U.S. Application Data**

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- [51] **Int. Cl.<sup>5</sup>** ..... A61K 38/18; C12P 21/02
- [52] **U.S. Cl.** ..... 514/8; 435/686; 530/388.7; 530/397; 530/835
- [58] **Field of Search** ..... 435/69.1, 69.4, 435/69.6, 240.2, 320.1, 172.3, 13; 530/350, 380, 397, 834, 835, 23.5, 23.51, 388.7; 514/8

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[57] **ABSTRACT**

Disclosed are novel polypeptides possessing part or all of the primary structural conformation and one or more of the biological properties of mammalian erythropoietin ("EPO") which are characterized in preferred forms by being the product of procaryotic or eucaryotic host expression of an exogenous DNA sequence. Illustratively, genomic DNA, cDNA and manufactured DNA sequences coding for part or all of the sequence of amino acid residues of EPO or for analogs thereof are incorporated into autonomously replicating plasmid or viral vectors employed to transform or transfect suitable procaryotic or eucaryotic host cells such as bacteria, yeast or vertebrate cells in culture. Upon isolation from culture media or cellular lysates or fragments, products of expression of the DNA sequences display, e.g., the immunological properties and in vitro and in vivo biological activities of EPO of human or monkey species origins. Disclosed also are chemically synthesized polypeptides sharing the biochemical and immunological properties of EPO. Also disclosed are improved methods for the detection of specific single stranded polynucleotides in a heterologous cellular or viral sample prepared from, e.g., DNA present in a plasmid or viral-borne cDNA or genomic DNA "library".

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FIG. 1

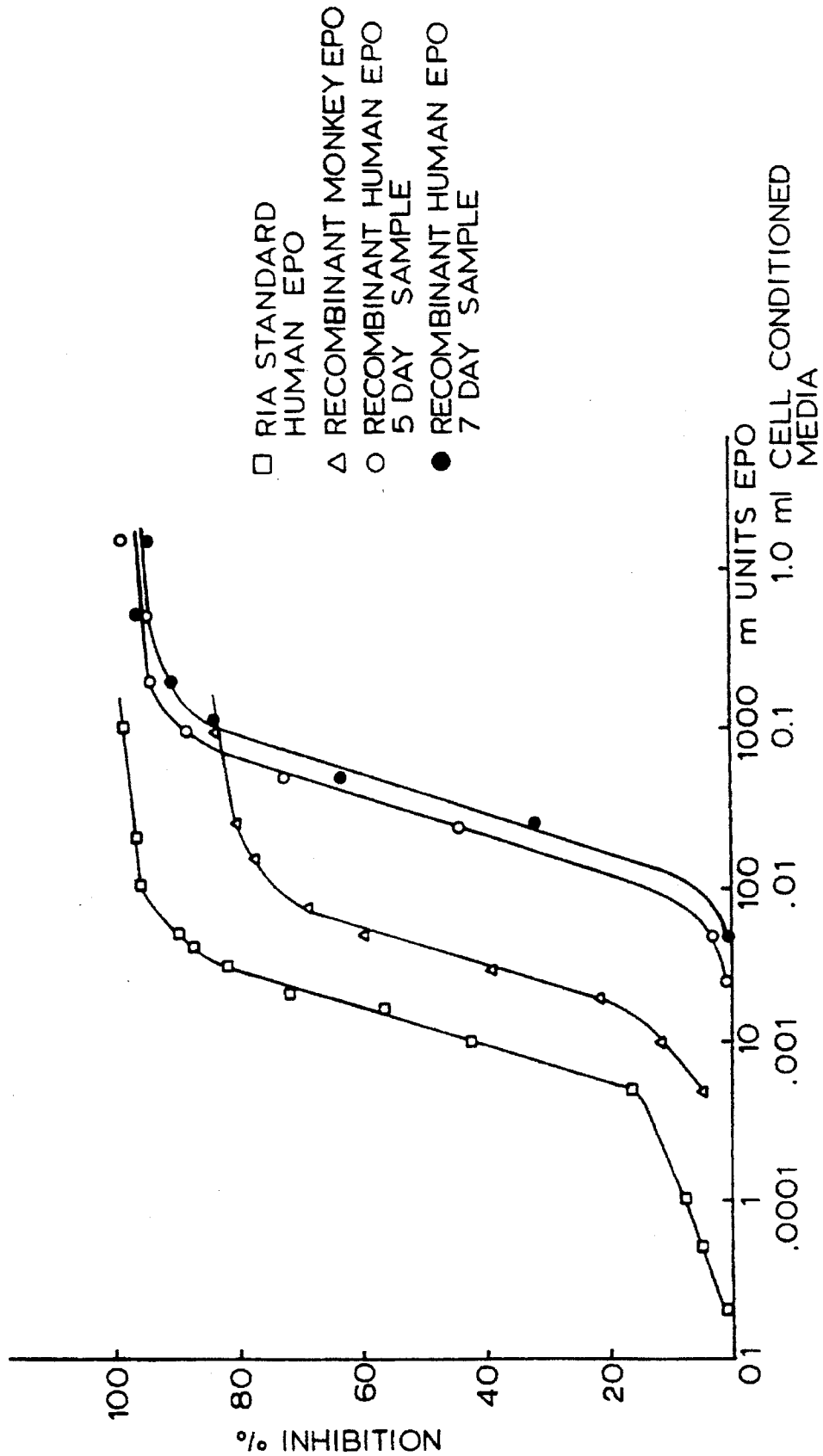
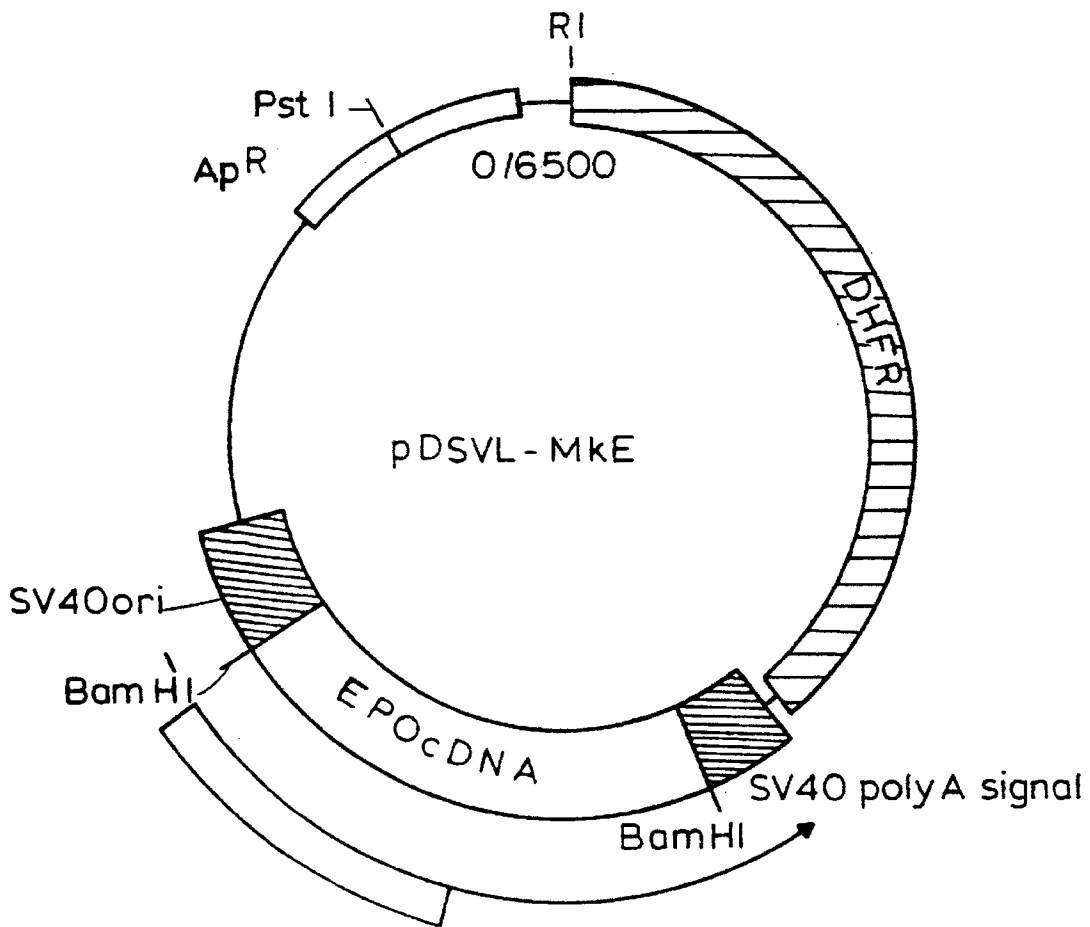


FIG. 2



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FIG. 3

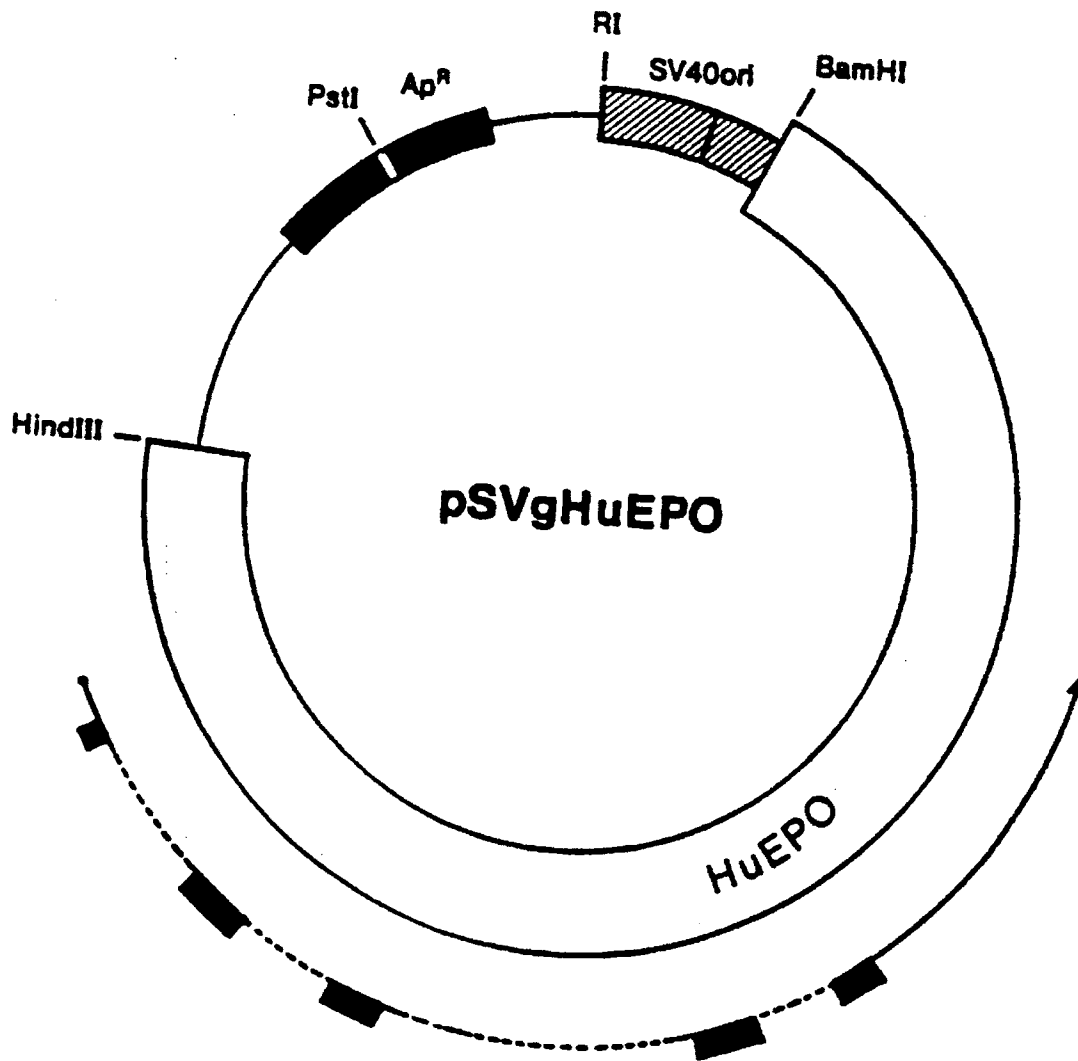


FIG. 4

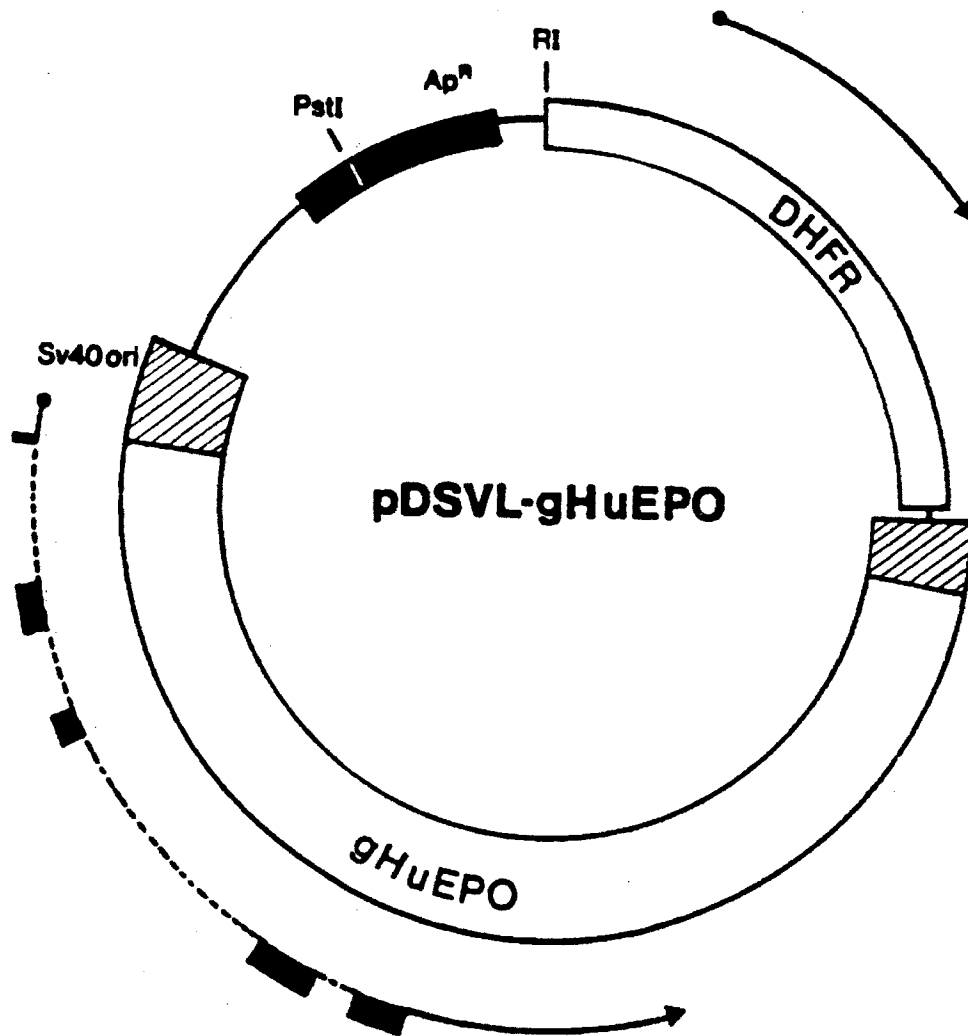


FIG. 5A

**2813A**  
**GATCCCGCCCCCTGGACAGCCGCCCTCTCCTCCAGGCCCGTGGGGCTGGCCCCCTGCC**  
**CGCTGAACCTCCCGGATGAGGACTCCCGGTGTGGTCAACCCCGCCCTAGGTCCCTGAG**

-27  
 Met Gly Val His Glu Cys Pro Ala Trp  
**GGACCCCGCCAGGCCGGAGATG GGG GTG CAC GAA TGT CCT GCC TGG**

-20  
 Leu Trp Leu Leu Ser Leu Val Ser Leu Pro Leu Gly Leu Pro  
**CTG TGG CTT CTC CTG TCT CTC GTG TCG CTC CCT CTG GGC CTC CCA**

-10  
 Val Pro Gly Ala Pro Pro Arg Leu Ile Cys Asp Ser Arg Val Leu  
**GTC CCG GGC GCC CCA CCA CCG CTC ATC TGT GAC AGC CGA GTC CTG**

+1  
 Glu Arg Tyr Leu Leu Glu Ala Lys Glu Ala Glu Asn Val Thr Met  
**GAG AGG TAC CTC TTG GAG GCC AAG GAG GCC GAG AAT GTC ACG ATG**

20  
 Glu Cys Ser Glu Ser Cys Ser Leu Asn Glu Asn Ile Thr Val Pro  
**GCC TGT TCC GAA ACC TGC AGC TTG AAT GAG AAT ATC ACC GTC CCA**

30  
 40



FIG. 5B

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50
ASP THR LYS VAL ASN PHE TYR ALA TIP LYS ARG MET GLU VAL GLY
GAC ACC AAA GTT AAC TTC TAT GCC TGG AAG AGG ATG GAG GTC GGG

60
GLN GLN ALA VAL GLU VAL TIP GLN GLY LEU ALA LEU LEU SER GLU
CAG CAG GCT GTA GAA GTC TGG CAG GCC CTG GCC CTC TCA GAA

70
80
ALA VAL LEU ARG GLY GLN ALA VAL LEU ALA ASN SER SER GLN PRO
GCT GTC CTG CTG CGG GGC CAG GCC GTC TTG GCC AAC TCT TCC CAG CCT

90
PHE GLU PRO LEU GLN LEU HIS MET ASP LYS ALA ILE SER GLY LEU
TTC GAG CCC CTG CAG CTG CAC ATG GAT AAA GCC ATC AGT GGC CTT

100
ARG SER ILE THR THR LEU LEU LEU ARG ALA LEU GLY ALA GLN GLU ALA
CGC AGC ATC ACC ACT ACT CTG CTG CTT CGG GCG GCG CTG GGA GCC CAG GAA GCC

110
ILE SER LEU PRO ASP ALA ALA SER ALA ALA PRO LEU ARG THR ILE
ATC TCC CTC CCA GAT GAT GCG GCC TCG GCT GCT CCA CTC CGA ACC ATC

120
THR ALA ASP THR PHE CYS LYS LEU PHE ARG VAL TYR SER ASN PHE
ACT GCT GAC ACT TTC TGC AAA CTC TTC CGA GTC TAC TCC AAT TTC

130
140

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## FIG. 5C

150      Leu Arg Gly Lys Leu Lys Leu Tyr Thr Gly Glu Ala Cys Arg Arg  
           CUC CGG GGA AAG AAG CTG AAG CTG TAC ACC GGG GAG GCC TGC AGG AGA  
           160

165      Gly Asp Arg OP  
           GGG GAC AGA TGA CCAGGTGCGTCCAGCTGGGCACATCCACCACCTCCCTCACCCAAACA  
           CTGCCGTGCCCACACCCTCCCTCACCTCCCGAACCCTCATCGAGGGGCTCTCAGCTAAG

          CCCCAGCCTGTCCCATGGACACTCCAGTCCAGCCAAATGACATCTCAGGGGCCAGAGGAAAC  
           TGTCCAGAGCACACTGTGAGATCTAAGGATGTCCAGGGCCAACTTGAAGGGCCCCAGAGC  
           AGGAAAGCATTCAGAGAGCAGCTTTAAACTCAGGAGCAGAGACAAATGCAGGGGAAACACCTT  
           GAGCTCACTCGGCCACCTGCAAAATTGATGCCAGGACACGGCTTTGGAGGCCNATTTACCTG  
           TTTTTGCACCTACCATCAGGGACAGGATGACTGGAGAACTTAGGTGGCAGCTGTGACTT  
           CTCAAGGCCTCACGGGCACTCCCTTGGTGGCAAAGACCCCTTGACACTGAGAGAAATATT  
           TTGCCAATCTGCCAGCAGGAAAAAATTAAGGACAGGTTTTTGGAGGTTGGAGGGTACTTGCACG  
           GTGfGTGGGGAAAGCAGGGCCGGJAGGGGTGGAGCTGGGATGCCAGfGAGAACCGTGAAGAC  
           AGGATGGGGGCTGGCCTCTGGTTCTCGTGGGGTCCAAAGCTT

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FIG. 6A

AAGCTTCTGGGCTTCCAGACCAGCTATTTCGGGAACTCAGCAACCCAGGCATCTCTGAGTCTCCGCCCA  
 AGACCGGATGCCCCCAAGGGGAGGGTGTCCGGGAGCCCAAGCCTTTCCCAAGATAGCACGGCTCCGCCCAAGTCCC  
 AAGGGTCCGCAACCGGCTGCCTCCCTCCCGGACCCAGGGCCCCGGGAGCACGCCCCCAATGACCCACACCGC  
 ACGTCTCAGCAGCCCCGCTCACGGCCCCGGGAGCCCTCAACCCAGGCGTCCCTGCCCTGCTGACCCCGG  
 GTGGCCCCCTAGCGACCCCTCACGCAACAGCCTCTCCCGCACCCCGCACCCCGCACACATG  
 CAGATAACAGCCCCGACCCCGGCAAGCCGXAAGATCCCTGGGCCACCCCGGCGCTGCCCTGCCGCTG  
 CCGCGCACCGGCTGTCTCCGAGCCGGACCGGGCCACCGCGCCXGCTCTGCTCCGACACCCCGCCC  
 CTTGGACAGCCCTCTCTCTAGGCCCGTGGGCTGGCCCTGCACCCCGGAGCTTCCCGGATGAGGIX  
 CCCGGTACCGGCGCCCAAGTCCCTAGGGACCCCGGCAAGCCGGGAG ATG GGG GTG CAC G  
 GTGAGTACTCGCGGCTGGGCGCTCCCGGCGCGGTTCCCTGTTTGAAGCGGGGATTTAGCGCCCGCGGCT

-27                    -24  
 Met Gly Val His