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This is a communication			DATEM	MAILED: 02/05/97

COMMISSIONER OF PATENTS AND TRADEMARKS

This application has been examined Responsive to communication filed on 10/3/66 This action is made final.
A shortened statutory period for response to this action is set to expire
Part I THE FOLLOWING ATTACHMENT(S) ARE PART OF THIS ACTION:  1.  Notice of References Cited by Examiner, PTO-852.  Notice of Art Cited by Applicant, PTO-1449  Information on How to Effect Drawing Changes, PTO-1474  Information on How to Effect Drawing Changes, PTO-1474  Page 1.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information on How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information On How to Effect Drawing Changes, PTO-1474  Page 2.  Notice of Information On How to Effect Drawing Changes Changes Changes Changes Changes Changes Changes Changes Chang
Part II SUMMARY OF ACTION
1. × C19-75 1-72
of the above, claims 1-13,16, 3 >- 57 244 59-60 are withdrawn from consideration.
2. Clarms
3. Claims have been cancelled.
4. TX Claims 14,15, 17-36, 58 2nd 61-72 are rejected.
S. Claims are objected to.
are subject to restriction or election requirement.
<ol> <li>This application has been filled with informal drawings which are acceptable for examination purposes until such time as allowable subject matter is indicated.</li> </ol>
8. Allowable subject matter having been indicated, formal drawings are required in response to this Office action,
9. The corrected or substitute drawings have been received or These drawings are acceptable, not acceptable (see explanation).
10. The proposed drawing correction and or the proposed additional or substitute sheets of drawings, fried or
11. The proposed drawing correction, fried
12. Acknowledgment is made of the craim for drightly under 35 U.S.C. 113. The certified cap, has been received not been received
been filed in patent application, serial no.
3. Since this application appears to be in condition for attowance except for formal matters, prosecution as to the Herits is crosed in accordance with the practice under Exiparte Quayre, 1935 C.D. 111 451 O.D. 113.
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PTOL-326 (Rev. 7 - 82)

Serial No. 675298

Art Unit 127 -2-

The disclosure is objected to because of the following informalities:

Tables V, VI, XIV and XXX have not been submitted as new drawing Figures 5-8 as applicant contends.

Appropriate correction of the disclosure is required.

Claims 20, 23, 27 and 30 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention. Ordinarly claims should not refer to drawings particularly as here where the material referred to can be easily described without resort to drawings.

35 U.S.C. 101 reads as follows:

Whoever invents or discovers any new and useful process, machine, manufacture, or composition of matter or any new and useful improvement thereof, may obtain a patent therefor, subject to the conditions and requirements of this title.

Claims 14, 15, 17-36, 5B and 61-72 are provisionally rejected under 1, 0.5.c. 101 as claiming the same invention as that of claims 13-24 and 27 of copending application Serial No. 582,185 and claims 1-48 of copending application Serial No. 655,841.

This is a provisional double patenting rejection since the conflicting claims have not in fact been patented.

Contrary to applicant's contention 35 U.S.C. 101 does in fact bar the allowance of the same invention in more than one application.

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Art Unit 127

Claims 14-19, 21, 22, 24, 25, 28, 31-34, 36, 58, 61-66 and 69-71 are rejected under 35 U.S.C. 112, first paragraph, as the disclosure is enabling only for claims limited to the DNA sequence coding for erythropoletin. The recitation of "fragments thereof" the recitation of and/o. "having at least a part of the primary structural conformation and one or more of the biological activities of naturally-occurring erythropoietin\* permits the claims to read on proteins and peptides completly unrelated to erythropoietin. Note for example that those unrelated proteins could possess the common biological activity of being an antigen. See MPEP 706.03(n) and 706.03(z).

Claim 35 is rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention. The use of brakets for indicating different species and strains of erythropoietin DNA sequence is improper since brakets in claims designates excluding the braketed matter.

Claims 14, 15, 61, 62 and 69 are rejected under 35 U.S.C. 112, first paragraph, as the disclosure is enabling only for claims limited the conditions of hybridization. As is clear from Walker et al and Kennell almost any mismatched pair of nucleotides can be

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hybridized to each other given the right set of conditions. See MPEP 706.03(n) and 706.03(z).

The following is a quotation of 35 U.S.C. 103 which forms the basis for all obviousness rejections set forth in this Office action:

A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

Subject matter developed by another person, which qualifies as prior art only under subsection (f) and (g) of section 102 of this title, shall not preclude patentability under this section where the subject matter and the claimed invention were, at the time the invention was made, owned by the same person or subject to an obligation of assignment to the same person.

Claims 14, 17, 18, 21-24, 26, 27, 31-36, 58 and 61-68 are rejected under 35 U.S.C. 103 as being unpatentable over Sue et al (PNAS 80) taken together with Boeslow et al or Woods et al. Sue disclose what they believed to be the 26 amino acid sequence of human erythropoietin amino terminal peptide and indicate that the sequence was known by Goldwasser et al in 1981. Breslow et al and Woods et al disclose screening a cDNA library derived from human adult liver for blood protein sequences with oligonucleotde probes deduced from known amino acid sequence of blood proteins.

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It would be obvious to isolate the human erythropoietin cDNA sequence by utilizing the Sue et al erythropoietin amino acid sequence data to devise oligonucleotide probes for use in screening a cDNA liver library in the manner taught by Breslow et al or Woods et al. The fact that the erythropoietin 26 amino acid terminal peptide sequence of Sue et al differs from that of erythropoietin by designating an Asn instead of a Cys at the seven position is patentably irrelevant since it

Claims 15 and 69-72 are rejected under 35 U.S.C. 103 as being unpatentable over Sue et al taken together with Breslow et al or Woods et al as applied to claims 14, 17, 18, 21-24, 26, 27, 31-36, 58 and 61-68 above, and further in view of Talmadge et al who disclose expressing mammalian proteins using recombinant expression plasmids. It would be obvious to express the erythropoeitin gene sequence by using that sequence in expression plasmids of Talmadge et al.

would not interfer with the preparation of effective

Claims 19 and 20 are rejected under 35 U.S.C. 103 as being unpatentable over Sue et al taken together with Breslow et al or Woods et al as applied to claims 14,17, 18, 21-24, 26, 27, 31-36, 58 and 61-69 above, and further in view of Farber et al who teach a monkey source for erythropoletin. It would be obvious to use

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a cDNA liver library from monkey in place of that from human or use the isolated erythropoietin cDNA sequence for isolating the monkey m RNA for ultimate conversion to monkey cDNA.

Claims 25 and 28-30 are rejected under 35 U.S.C.

103 as being unpatentable over Sue et al taken together with Breslow et al or Woods et al as applied to claims

14, 17, 18, 21-24, 26, 27, 31-36, 58 and 61-68 above, and further in view of Goury et al or Bennetzen et al who disclose the preperfered codons for E. coli and yeast, respectively. It would be obvious to select those codons where the ultimate expression is in E. coli or yeast.

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless-

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claims 14, 15, 17-19, 21, 22, 24, 25, 28, 31-34, 36, 58 and 61-72 are rejected under 35 U.S.C. 102(b) as being anticipated by Talmadge et al who disclose the expression of a mammalion protein using recombinant DNA transformed microorganisms. In view of the recitation of "fragments thereof" and the recitation of "having at least... activities" (see the 35 U.S.C. 112 rejection

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supra) the claims are deemed to embrace the DNA sequences and protein expression method of Talmadge et al.

Claims 69-72 are rejected under 35 U.S.C. 103 as being unpatentable over Talmadge et al and who disclose the basic process of recombinantly expressing and isolating polypeptides as claimed herein. Even it is considered that one or more of the starting materials is novel, the application of an old process to such materials to produce the expected result would still be obvious within the meaning of 35 U.S.C. 103; In reduction, supra; In relarsen, 141 U.S.P.Q. 730 (1964). Whether or not a product produced by the claimed process possesses any unique or unexpected properties is not material to the question of whether or not the process itself would have been obvious.

A.E. Tanenholtz/pww

A/C 703

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ALVIN E. TANENHOLTZ
PRIMARY EXAMINER
ART UNIT 127

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