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(Also referred to as FORM PTO-XXXX)

REQUEST FOR SUPPLEMENTAL EXAMINATION TRANSMITTAL FORM

| | Address to: Commissioner for Patents | Attorney Docket No.: | |
|-----|---|--|--------|
| | P.O. Box 1450 Alexandria, VA 22313-1450 | Date: | |
| 1. | | oursuant to 37 CFR 1.610 of patent numberthe first-named inventor is | · |
| 2. | Supplemental examination of claim(s) | is reques | ted. |
| 3. | | ne request (e.g., certificate of mailing (if on a separate shee ubmitted as part of the request) is included. 37 CFR 1.610 | |
| | b. A table of contents for the request is included | d. 37 CFR 1.610(b)(2). | |
| 4. | request for supplemental examination, the fee | s enclosed to cover the fee for processing and treating a for reexamination ordered under 35 USC 257, and the fee t ment over 20 sheets in length (37 CFR 1.20(k)(1 - 3)); | for |
| | b. The Director is hereby authorized to charg to Deposit Account No | e all applicable fees as set forth in 37 CFR 1.20(k)(1 - 3) | |
| | c. Payment by credit card. Form PTO-2038 | is attached. 37 CFR 1.610(a). | |
| 5. | | card, refund must be to the credit card account. | |
| 6. | A copy of the patent for which supplemental ex | camination is requested is included. 37 CFR 1.610(b)(9). | |
| 7. | CD-ROM or CD-R in duplicate, Computer Prog | ram (Appendix) or large table | |
| 8. | Nucleotide and/or Amino Acid Sequence Subn If applicable, items a. – c. are required. | nission | |
| | a. Computer Readable Form (CRF)b. Specification Sequence Listing on: | | |
| | i. CD-ROM (2 copies) or CD-R (2 dii. paper | copies); or | |
| | c. Statements verifying the identity of abo | ve copies | |
| 9. | | on, certificate of extension, supplemental examination certificate, or post grant review certificate that has been issue | |
| 10. | this form. Also included is a statement inclu | is submitted herewith including a listing thereof in Part B of ding an identification and explanation of each submitte I, was inadequately considered, or was incorrect pursu | d |
| 11. | An English language translation of all neces information is included. 37 CFR 1.610(b)(10). | ssary and pertinent parts of each non-English language in | tem of |

[Page 1 of 4]

This collection of information is required by 37 CFR 1.610. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.11 and 1.14. This collection is estimated to take 18 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS.

PTO/SB/59 (01–12)

Approved for use through xxxxxx. OMB xxxxxxxxxxx

U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

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| 12. | | The attached detailed request includes the following items: | | | | | | |
|---------------|--|--|--|--------------------------------------|--|--|--|--|
| | a. An identification of each aspect of the patent for which supplemental examination is sought, including an identification of the structure, material, or acts in the specification that correspond to each means-plus-function or step-plus-function element as set forth in 35 U.S.C. 112(f), in any claim for which supplemental reexamination is requested. 37 CFR 1.610(b)(6). | | | | | | | |
| | 7). | | | | | | | |
| | c. A separate, detailed explanation for each identified issue, discussing how each item of information is relevant to each aspect of the patent identified for examination, and how each item of information raises each issue identified for examination, including an explanation compliant with each of the provisions of 37 CFR 1.610(b)(8)(i)-(ii). | | | | | | | |
| 13. | A summary of the relevant portions of each submitted document that is over 50 pages in length (other than the request) is included. the summary includes the required citations to the particular pages containing the relevant portions. 37 CFR 1.610(b)(11). | | | | | | | |
| 14. | | | mpliance with 37 CFR 3.73(b), which establishe oplemental examination is requested as set forth | | | | | |
| 15. | | supplemental examination, post grant re- | rrent post-patent Office proceedings (ex parte oview, or inter partes review) involving the patent Any prior or concurrent post-patent Office process. | t for which supplemental examination | | | | |
| | | Type of Proceeding | Identifying Number | Filing Date | | | | |
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| | Se | e attached sheet for a listing of additional examination is requested. | prior or concurrent post-patent Office proceeding | ngs for which supplemental | | | | |
| [Page 2 of 4] | | | | | | | | |

| 16. Correspondence Address: Please recognize, or change, the coand for the supplemental examination proceeding to be: | rrespondence address for the file | of the above-identified patent |
|---|---|--------------------------------|
| The address associated with Customer Number: Firm or Individual Name | | OR |
| Address | | |
| City | State | Zip |
| Country | | |
| Telephone | Email | |
| WARNING: Information on this form may become public. C included on this form. Provide credit card information and | redit card information should no authorization on PTO-2038. | ot be |
| Authorized Signature Typed/Printed Name | Date Registration No. | |
| | | |

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REQUEST FOR SUPPLEMENTAL EXAMINATION TRANSMITTAL FORM PART B - LIST OF ITEMS OF INFORMATION

| Pate | ent number for which supplemental examination is requested I | ssue Date | : | |
|------|--|--|-------------------------|---|
| | | • .• | 0.41 | |
| | items of information submitted herewith as part of this request for supplemental exantified patent are included in the following list: | ımınatıon | of the a | bove- |
| luc | Document Description If a U.S. patent or patent application publication. Include patent number, kind code (see MPEP 901.04), and name of patentee or applicant of cited document. If a toreign or international patent or patent application publication, include country code (WPO Standard ST 3), number, kind code (WPO Standard ST 16 if possible), and number of patentee or applicant of cited document. For Japanese patent documents, the indication of the year of the reign of the Emperor must precede the serial number of the patent document. If non-patent literature, include, where available, name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the literature. Include, where available, symposium, catalog, etc.) date, page(s), volume-lease number(s), publisher, city and/or country where published. If a court document, identify the specific court, the designation or numeric designation), the title of the document, and the date submitted in court. For all other materials, include title (if available), author (if applicable), any date shown on document (if not a publication date), and any other descriptive information that would identify the document. | Publication Date MM-DD- YYYY (If applicable) | Translation Attached | Summary under §1.910(b)(11) Attached |
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Privacy Act Statement

The **Privacy Act of 1974 (P.L. 93-579)** requires that you be given certain information in connection with your submission of the attached form related to a patent application or patent. Accordingly, pursuant to the requirements of the Act, please be advised that: (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) furnishing of the information solicited is voluntary; and (3) the principal purpose for which the information is used by the U.S. Patent and Trademark Office is to process and/or examine your submission related to a patent application or patent. If you do not furnish the requested information, the U.S. Patent and Trademark Office may not be able to process and/or examine your submission, which may result in termination of proceedings or abandonment of the application or expiration of the patent.

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether disclosure of these records is required by the Freedom of Information Act.
- 2. A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiations.
- 3. A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
- 4. A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552a(m).
- 5. A record related to an International Application filed under the Patent Cooperation Treaty in this system of records may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant to the Patent Cooperation Treaty.
- A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or his/her designee, during an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2904 and 2906. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- 8. A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S.C. 122(b) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filed in an application which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspection or an issued patent.
- 9. A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.