

<b>REQUEST FOR CONTINUED EXAMINATION(RCE)TRANSMITTAL (Submitted Only via EFS-Web)</b>							
Application Number	16/746,028	Filing Date	2020-01-17	Docket Number (if applicable)	066859/542422	Art Unit	1612
First Named Inventor	John Maloney			Examiner Name	Benjamin J. Packard		
<p><b>This is a Request for Continued Examination (RCE) under 37 CFR 1.114 of the above-identified application.</b>            Request for Continued Examination (RCE) practice under 37 CFR 1.114 does not apply to any utility or plant application filed prior to June 8, 1995, or to any design application. The Instruction Sheet for this form is located at WWW.USPTO.GOV</p>							
<b>SUBMISSION REQUIRED UNDER 37 CFR 1.114</b>							
<p>Note: If the RCE is proper, any previously filed unentered amendments and amendments enclosed with the RCE will be entered in the order in which they were filed unless applicant instructs otherwise. If applicant does not wish to have any previously filed unentered amendment(s) entered, applicant must request non-entry of such amendment(s).</p>							
<p><input type="checkbox"/> Previously submitted. If a final Office action is outstanding, any amendments filed after the final Office action may be considered as a submission even if this box is not checked.</p> <p style="margin-left: 40px;"><input type="checkbox"/> Consider the arguments in the Appeal Brief or Reply Brief previously filed on _____</p> <p style="margin-left: 40px;"><input type="checkbox"/> Other _____</p>							
<p><input checked="" type="checkbox"/> Enclosed</p> <p style="margin-left: 40px;"><input type="checkbox"/> Amendment/Reply</p> <p style="margin-left: 40px;"><input checked="" type="checkbox"/> Information Disclosure Statement (IDS)</p> <p style="margin-left: 40px;"><input type="checkbox"/> Affidavit(s)/ Declaration(s)</p> <p style="margin-left: 40px;"><input type="checkbox"/> Other _____</p>							
<b>MISCELLANEOUS</b>							
<p><input type="checkbox"/> Suspension of action on the above-identified application is requested under 37 CFR 1.103(c) for a period of months _____            (Period of suspension shall not exceed 3 months; Fee under 37 CFR 1.17(i) required)</p> <p><input type="checkbox"/> Other _____</p>							
<b>FEES</b>							
<p><b>The RCE fee under 37 CFR 1.17(e) is required by 37 CFR 1.114 when the RCE is filed.</b></p> <p><input checked="" type="checkbox"/> The Director is hereby authorized to charge any underpayment of fees, or credit any overpayments, to            Deposit Account No <input style="width: 100px;" type="text" value="160605"/></p>							
<b>SIGNATURE OF APPLICANT, ATTORNEY, OR AGENT REQUIRED</b>							
<p><input checked="" type="checkbox"/> Patent Practitioner Signature            Applicant Signature</p>							

Doc code: RCEX  
Doc description: Request for Continued Examination (RCE)

PTO/SB/30EFS (02-18)  
Approved for use through 11/30/2020. OMB 0651-0031  
U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

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Signature of Registered U.S. Patent Practitioner			
Signature	/bryan l. skelton/	Date (YYYY-MM-DD)	2020-05-28
Name	Bryan L. Skelton	Registration Number	50893

This collection of information is required by 37 CFR 1.114. 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 12 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.

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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 inspections 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.

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re: John Maloney et al. Confirmation No.: 4075  
Appl. No.: 16/746,028 Group Art Unit: 1612  
Filed: January 17, 2020 Examiner: Benjamin J. Packard  
For: STABLE, HIGHLY PURE L-CYSTEINE COMPOSITIONS FOR INJECTION  
AND METHODS OF USE

Submitted via EFS-Web  
Commissioner for Patents  
P.O. Box 1450  
Alexandria, VA 22313-1450

**INFORMATION DISCLOSURE STATEMENT  
CITATION UNDER 37 C.F.R. § 1.97**

Attached is a list of documents on form PTO-SB08.

It is requested that the Examiner consider these documents and officially make them of record in accordance with the provisions of 37 C.F.R. § 1.97 and Section 609 of the MPEP. By identifying the listed documents, Applicant in no way makes any admission as to the prior art status of the listed documents, but is instead identifying the listed documents for the sake of full disclosure.

Respectfully submitted,

/bryan l. skelton/

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Substitute for form 1449B/PTO				<b>Complete if Known</b>	
<b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b>  <i>(Use as many sheets as necessary)</i>				Application Number	16/746,028
				Filing Date	January 17, 2020
				First Named Inventor	John Maloney
				Art Unit	1612
				Examiner Name	Benjamin J. Packard
Sheet	1	of	4	Attorney Docket Number	066859/542422

U.S. PATENT DOCUMENTS					
Examiner Initials*	Cite No. <sup>1</sup>	Document Number	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number Kind Code <sup>2</sup> (if known)			
	292	US 6,382,442 B1	05-07-2002	Thibault et al.	
	276	US 8,415,337 B1	04-09-2013	Krishna	

NON PATENT LITERATURE DOCUMENTS					
Examiner Initials *	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.			T <sup>2</sup>
	293	"Aluminum in large and small volume parenterals used in total parenteral nutrition," Food and Drug Administration, 21 C.F.R. § 201.323, 89-90, (2003).			
	272	"American Regent Announces the Launch and Availability of Selenious Acid Injection, USP," Press Release, American Regent, Inc., 6 pages, (2019).			
	251	"Cysteine," DrugBank, 23 pages, Exhibit 1016, Petition for Post Grant Review of U.S. Patent No. 10,478,453, Eton Pharmaceuticals, Inc. v. Exela Pharma Sciences, LLC, PGR2020-00064, (PTAB May 19, 2020).			
	298	"ELCYS (Cysteine Hydrochloride)," NDA 210660, Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations, 3 pages, (2019).			
	274	"Neonatal Parenteral Nutrition," Intensive Care Nursery House Staff Manual, UCSF Children's Hospital, pp. 136-142, (2004-2006).			
	280	"Aluminum in Large and Small Volume Parenterals Used in Total Parenteral Nutrition," Federal Register, 63(2):176-185, (1998).			
	246	"AMINOSYN [label information]", Hospira, Inc., 11 pages, Exhibit 1009, Petition for Post Grant Review of U.S. Patent No. 10,478,453, Eton Pharmaceuticals, Inc. v. Exela Pharma Sciences, LLC, PGR2020-00064, (PTAB May 19, 2020).			
	278	"Aminosyn Sulfite Free [drug information]," RX List, 15 pages, Exhibit 1052, Petition for Post Grant Review of U.S. Patent No. 10,478,453, Eton Pharmaceuticals, Inc. v. Exela Pharma Sciences, LLC, PGR2020-00064, (PTAB May 19, 2020).			
	253	"Guidance for Industry: Q8(R2) Pharmaceutical Development," U.S. Dept. of Health and Human Services, FDA, CDER, CBER, 29 pages, (2009).			
	244	"L-CYSTEINE HYDROCHLORIDE - cysteine hydrochloride injection, solution [label information]", Sandoz Inc., 11 pages, Exhibit 1005, Petition for Post Grant Review of U.S. Patent No. 10,478,453, Eton Pharmaceuticals, Inc. v. Exela Pharma Sciences, LLC, PGR2020-00064, (PTAB May 19, 2020).			
	248	"Q3D Elemental Impurities: Guidance for Industry," U.S. Dept. of Health and Human Services, FDA, CDER, CBER, 85 pages, (2015).			
	271	"Selenious Acid Injection [prescribing information]," American Regent, Inc., 8 pages, (2019).			
	273	"Zinc sulfate injection [prescribing information]," American Regent, Inc., 9 pages, (2019).			

Examiner Signature		Date Considered	
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