

Entry into the European phase (EPO as designated or elected Office)

To the European Patent Office

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Applicant's or representative's reference	42.133062/07
International Filing Date	09.11.2018
International Searching Authority (ISA)	EP
International Preliminary Examining Authority (IPEA)	not applicable
1. Applicant	
Indications concerning the applicant(s) are contained in the international publication or were recorded by the International Bureau after the international publication.	
Changes which have not yet been recorded by the International Bureau are set out here:	
2. Representative Representative 1	
Representative or association of representatives to be listed in the Register of European Patents and to whom communications are to be notified	
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3. Authorisation Representative 1 An individual authorisation is attached. A general authorisation has been registered under No:	mail@dehns.com
3. Authorisation Representative 1 An individual authorisation is attached. A general authorisation has been registered under No: A general authorisation has been filed, but not yet registered. The authorisation filed with the EPO as PCT receiving Office expressly includes	mail@dehns.com
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the following documents	
the application documents published by the International Bureau; where the publication includes a set of claims amended under Article 19 PCT, the latter replaces the originally filed claims	
unless replaced by the amendments attached.	
Comments on the international preliminary examination report established by the EPO as the International Preliminary Examining Authority and/or observations are attached.	
Where necessary, further details should be attached as "Other documents"	
6.2 Proceedings before the EPO as elected Office (PCT II) are to be based on the following documents:	
the documents on which the international preliminary examination report is based, including any annexes	
unless replaced by the amendments attached.	
comments on the international preliminary examination report established by the EPO as the International Preliminary Examining Authority and/or observations are enclosed.	
Where necessary, further details should be attached as "Other documents"	
If the EPO as International Preliminary Examining Authority has been supplied with test reports, these may be used as the basis of proceedings before the EPO.	
6.3 A copy of the results of the search carried out by the authority with which the previous application(s) whose priority is claimed was (were) filed is attached (R. 141(1) EPC).	
7. Translations	
Translations in one of the official languages of the EPO (English, French, German) are attached as crossed below:	
* In proceedings before the EPO as designated or elected Office (PCT I + II):	
7.1 Translation of the international application (description, claims, any text in the drawings) as originally filed, of the abstract as published and of any indication under Rule 13bis.3 and 13bis.4 PCT regarding biological material	
7.2 Translation of the priority application(s) (to be filed only at the EPO's request, Rule 53(3) EPC)	
7.3 It is hereby declared that the international application as originally filed is a complete translation of the previous application (Rule 53(3) EPC)	
* In addition, in proceedings before the EPO as designated Office (PCT I):	
7.4 Translation of amended claims and any statement under Art. 19 PCT, if the claims as amended are to form the basis for the proceedings before the EPO (see Section 6).	
* In addition, in proceedings before the EPO as elected Office (PCT II):	
7.5 Translation of any annexes to the international preliminary examination report	
8. Biological material	
The invention uses and/or relates to biological material deposited under Rule 31 EPC.	
The particulars referred to in Rule 31(1)(c) EPC (if not yet known, the depositary institution and the identification reference(s)) [number, symbols, etc.] of the depositor) are given in the international publication or in the translation submitted in Section 7 on:	
page(s) / line(s)	
The receipt(s) of deposit issued by the depositary institution	
is (are) enclosed.	
will be filed later.	
Waiver of the right to an undertaking from the requester pursuant to Rule 33(2) EPC attached.	
9. Nucleotide and amino acid sequences	
The international application discloses nucleotide and/or amino acid sequences. 9.1 The sequence listing was filed under Rule 5.2(a) PCT, or furnished to the	
EPO as ISA under Rule 13ter.1(a) PCT, or is otherwise available to the EPO, in computer-readable format in accordance with WIPO ST.25.	_

The sequence listing is attached in computer-readable format in accordance with WIPO Standard ST.25 (Rule 163(3) EPC)					
The sequence listing does not include matter which goes beyond the content of the application as filed.					
10. Des	ignation of contracting states				
All the contracting states party to the EPC at the time of filing of the international patent application and designated in the international application are deemed to be designated (see Article 79(1) EPC).					
11. Exte	ension/Validation				
patent non-co which applica	pplication is deemed to be a reques application and the European pater ontracting states to the EPC designates extension or validation agreements ation was filed. However, the request the validation fee, whichever is apposition.	nt granted in respect of it to ated in the international ap were in force on the date of is deemed withdrawn if t	o all plication with on which the he extension		
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12.2 V	Vaivers				
The applicant waives his right to the communication under Rules 161(1) or (2) and 162 EPC.					
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16-7	520 Additional filing fee for the 36th and each subsequent page - entry into EP phase	15	16.00	240.00
	Total:		EUR	2 305.00
17. Ann	otations			

18. Signature(s) of applicant(s) or representative

Place: London

Date: 09 June 2020
Signed by: /Elizabeth Jones/

Association: Dehns

Representative name: Elizabeth Jones
Capacity: (Representative)

Table for section 6 of Form 1200.3

In accordance with the Notice from the European Patent Office dated 26 January 2009 concerning the 2009 fee structure (OJ EPO 2009, 118, and Guidelines for Examination in the EPO, April 2009, A-III, 13.2), the amount of the additional fee (Art. 2, item 1a, Rules relating to Fees) for the pages of this European patent application is calculated as follows:

Documents intended for proceedings before the EPO (R. 159 (1) (b) EPC) and for calculating the additional fee (Art. 2, item 1a, RFees):

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Description:	International application as published	1-38	38
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Drawings:	International application as published	1-7-7/7	7
Abstract:	Default count: one page		1
Total number of page	es		50
Fee-exempt pages (Art. 2, item 1a, RFees) Number of pages to be paid for			-35
			15
			(x 16 EUR per page

Total amount payable EUR 240