

DG Inspection/Authorisations Division/Specially Regulated Substances Entity/RD 12.04.1974 Team

Send BY POST to:

Federal Agency for Medicines and Health Products DG Inspection – Authorisations Division – Substances RD 12.04.1974 Team Avenue Galilée 5/03 1210 BRUSSELS

Or

Send BY EMAIL to: <u>substances HAA@fagg-afmps.be</u>
Only when provided with the correct <u>qualified</u> electronic signatures (NO SCANS).

APPLICATION FOR A LICENCE FOR SUBSTANCES R.D. 12.04.1974 FOR SOLELY SCIENTIFIC AND/OR ANALYTICAL PURPOSES

According to art. 1 of the royal decree dated 12 April 1974 regarding some operations in connection with substances with hormonal, anti-hormonal, anabolic, beta-adrenergic, anti-infectious, anti-parasitic or anti-inflammatory effects, last revised by the R.D. dated 4 February 2002 (B.S. dated 28 February 2002).

	RENEWAL OF LICENCE NUMBER: indicate as appropriate: NL/FR							
	NEW LICENCE REQUEST							
	1. IDENTIFICATION OF THE APPLICANT							
Re	Registered office							
	Trade name*:							
	Legal form*:							
	Company number*:							
	Address*:							
	Telephone:							
Accounting information (contact person and address for mailing the invoice).								
Name*:				,				
Address:								
	E mail*:							
E-mail*: VAT-number*:								
Extra information that								
needs to be mentioned on the invoice*								
		a annronriate resnonses)						
	 Activities (indicate the appropriate responses) Note: import and export are regarded as in relation to the Belgian territory. 							
\boxtimes	Be in possession of			Acquiring by payment or at no cost				
	Import							
	Export (please indicate the reason for applying for export):							
	Transport (please indicate the reason for applying for transport):							



^{*:} Mandatory field

	where the substances will be stored and/or being used. f this space is not sufficient, please add an attachment.
Name / department / building / location:	
Address:	
Name / department / building / location:	
Address:	
Name / department / building / location:	
Address:	
Name / department / building / location:	
Address:	
Name / department / building / location:	
Address:	
Name / department / building / location:	
Address:	
Name / department / building / location:	
Address:	
Name / department / building / location:	
Address:	



4. LABORATORY MANAGER (At least 1 mandatory.)						
The laboratory manager(s) is (are) requested to sign the declaration below:						
Hereby declares that the substances to which this permit relates are exclusively used as (reference) substances for analytical and / or scientific purposes within the competence of the laboratory. In this case, I only need only to mention on p. 4 the substances with hormonal, anti-hormonal or beta-adrenergic action.						
Name		Signature				
5. IN THE CASE OF ANOTHER OF THE PERSON(S) RESPO DETERMINED IN THE ABOV date on the legislation on this	NSIBLE FOR	R THE EXECUTION OF THE C NED DECREE (these persons a	DBLIGATIONS AS			
Name		Signature				
6. CONTACTPERSON (for any or questions) Mandatory fields.	communication	ons from the FAMHP such as c	irculars or additional			
Name	E-mail		Tel./mobile			
7. SUBSTANCES and JUSTIFIC	CATION					
Please indicate on the next page the substances for which you are applying for a licence and provide below a brief description of the reason/project for applying for this licence. This will avoid additional questions from our service and potential delays in processing your licence application. The note "for analytical and / or scientific purposes" does not suffice. If this space is not sufficient, please add an attachment.						
REASONS for applying for this licence:						

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SUBSTANCES:

Substances and their corresponding CAS-number for which the licence is requested (art. 1, §1 and §3 of the RD of April 12th 1974, amended last by the R.D. dated 4 February 2002).

<u>Laboratories with analytical and / or scientific purposes will receive a general licence for all substances with an anti-inflammatory, anti-infectious, antiparasitic action.</u> An extension for substances with such an action will then never need to be requested. **ONLY substances with a hormonal, antihormonal or beta-adrenergic action need to be listed below.**

In case of insufficient space, you may add an appendix to the application, to list the relevant substances and their corresponding CAS-number.

SUBSTANCE	CAS- number (Chemical Abstracts Registry Number)



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8. SIGNATURE of **at least one** statutory or legally responsible person of the legal entity (in case of a health organisation or a governmental or educational institution, this can be the managing director, the rector, the principal or the headmaster) **endorsing the entirety of the contents of this completed form**. If this person cannot be found in the CBE, legal proof that this person is authorized to sign must be enclosed.

NOTE to higher educational institutions: contact your environmental- or preventive services for more information regarding a delegation.

Name:	Name:
Position:	Position:
E-mail:	E-mail:
Date:	Date:
Signature:	Signature:

Practical information

- Return the fully completed and signed application form:
 - \circ By POST, only if the original form has been signed (no copies or scans), to:

Federal Agency for Medicines and Health Products DG Inspection – Authorisations Division - Substances RD 12.04.1974 Team Avenue Galilée 5/03 1120 BRUSSELS

Or

 By E-MAIL, ONLY if the form has been provided with ALL the necessary qualified electronic signatures AFTER it has been completed in full, to:

substances HAA@fagg.be

(signature via ID card or see https://economie.fgov.be/fr/themes/line/commerce-electronique/signature-electronique-et).

Company tokens are usually not qualified electronic signatures and may be considered inadmissible.

The signature can be added by double-clicking on the signature field. Once signatures have been added, it is no longer permitted to edit the form, so please complete the form in full before adding the signatures. If the form does need to be changed again, the signatures must be re-applied. Otherwise, the application will be inadmissible.

- The fee amount is indexed annually and can be viewed at any time on the FAMHP website.
- **Every modification** (responsible persons, address, substances...) needs to be communicated to us (by post or e-mail) **within 15 days** by a responsible person mentioned on the licence. The form to be used for this is available on the FAMHP website.



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