SOCIETÀ ITALIANA DI FARMACOLOGIA

Italian Society of Pharmacology
Società Italiana di Farmacologia (SIF)

SIF-EuCP Program

Guidelines for the Certification as European Certified Pharmacologists (EuCP)

current as of: 22.06.2020
revision acknowledged by the EuCP Committee: 14.05.2021

EuCP

Sections

Section 1. Targets

The Italian Society of Pharmacology (SIF) European Certified Pharmacologist (EuCP) – "SIF-EuCP Program" is part of the European Registration Program initiated by the Federation of European Pharmacological Societies (EPHAR) and the European Association for Clinical Pharmacology and Therapeutics (EACPT) for individual pharmacologists or scientists working in the field of pharmacology. It was established to carry out the following tasks:

- To Ensure that "Certified Pharmacologist" designation is restricted to those who satisfy the criteria of EuCP program based on their professional expertise and competence, standards that are equal for all the national societies adhering to EPHAR and EACPT
- To identify, recognize and bring together the expert scientists actively engaged in the multidisciplinary field of pharmacology
- To stimulate professionals in the field of Pharmacology to expand and consolidate personal skills and to increase their qualification for high level positions in an increasingly competitive employment environment, be it academic, industrial, regulatory or self-employed
- To recognize, in addition to scientific expertise, professional competence (portfolio) in the whole discipline of pharmacology
- To issue the "EPHAR EuCP Certificate" by the Italian certifying body SIF and its ratification by the EPHAR EuCP Committee which will certify these individuals as "EuCP" without further evaluation.

Section 2. Association

2.1 Candidates who wish to apply for inclusion in SIF-EuCP Program have to fill out the Application Form which is made available by SIF Secretariat and on-line (https://www.sifweb.org/) in the version approved by "SIF-EuCP Program Certificate Commission" (CC) and ratified by SIF Executive Council (EC).

2.2 A non-refundable fee of € 30.00 is due by each candidate for first application examination (*una tantum*) and for each renewal request (Maintenance of certification) to SIF-EuCP Program. The Register administrative maintenance is self-financed therefore a registration fee of € 300.00 (€ 150.00 for applicants less than 40 years-old) is due after certification approval.

2.3. To be eligible for assessment by the CC the candidate is required to have a minimum but high standard of academic and professional skills in accordance with the provisions and ethical conduct of EPHAR (see http://www.ephar.org/docs/EPHAR_Ethical_Code_of_Conduct.pdf).

The application is possible through two different profiles: "Academic" (essentially Professors and Researchers with a university contract – see also note 2) and "Health Professional" (essentially all public and private professionals, including self-employed professionals, involved in Health and Clinical Research – see also note 3).

The two annexes (annex I and II) identify, through a checklist, which and how many academic and professional competences the candidate has and assign a total qualification score.

Annex III is used for the application aimed at certificate renewal (see Maintenance of Certification) and is also common to both profiles.

All annexes are available at SIF website (https://www.sifweb.org/).

2.4 The fundamental requirements for certification (without these criteria the application will not be taken into consideration by the CC) are:

1) University second cycle qualification[1] (MD, PhD, or MSc) in a relevant subject such as Medicine and Surgery[2], pharmaceutical sciences[2,3], second cycle qualification in biomedical sciences, biology or chemistry achieved at any university in the European Union or in other country and recognized by Italy according to Lisbon Convention (Law n. 148 of 11 July, 2002), and other Italian laws for Nostrification procedure ("ex equipollenza").

The Ministerial Decree 1 September 2016, n. 662 defines the table of correspondence between the Italian and foreign academic positions. (see http://attiministeriali.miur.it/media/303091/allegato_dm_662.pdf).

2) Considering the applicant employment profile:

- Academic [4]: at least three relevant peer review publications in the field of pharmacology research and other activities/credits in accordance with the ANNEX I (assessment check list, see point 2.3.1). Permanent academic personnel with the position of professor or researcher is evaluated exclusively by submission of a Curriculum vitae (see https://europass.cedefop.europa.eu/), with no further application of assessment criteria provided in Annex I [5]
- Health professionals [6]: at least three relevant reports/works/company paper/workshop and other activities/credits in accordance with ANNEX II (assessment check list, see point 2.3.1)

3) current professional engagement in the practice of Pharmacology (experimental, clinical or regulatory) with a proven professional experience and scientific qualifications.

2.4.1) The assessment [7] check lists (ANNEX I or II) screen at least five years of continuous and relevant professional development in pharmacology (experimental, clinical or regulatory) that may also include:

- "Third cycle" Specialisation course in Pharmacology and clinical Toxicology achieved at any university in the European Union or in other country and recognized by Italy according to Lisbon Convention (Law n. 148 of 11 July, 2002), and other Italian laws for Nostrification procedure ("ex equipollenza")

- "Third cycle" Doctoral course in Pharmacological Sciences, Medical and biomedical sciences achieved at any university in the European Union or in other country and recognized by Italy according to Lisbon Convention (Law n. 148 of 11 July, 2002), and other Italian laws for Nostrification procedure ("ex equipollenza")

- Other activities in institutional, pharmaceutical and therapeutic/advisory boards and commissions, events, meetings.

2.4.2) Candidates who meet requirements and who are not members of the SIF may be also accepted if they are registered to a society member of EPHAR or EACPT.

For the inclusion of members from other pharmacological societies (guest societies) a formal agreement shall be in place between SIF as hosting society and the guest society. The guest society should be member of EPHAR and/or EACPT. The agreement has to be approved by the EPHAR EuCP Committee before coming into effect.

2.4.3) The suitability of the candidate for the SIF-EuCP Program must be evaluated by the CC before November, 30th of the year in which the request was submitted and shall be notified in writing, within the same year, both in case of positive or negative result.

i.e. Application in january 2021:

- Answer due by November 30, 2021
- Application in December 2021
- Answer due by November 30, 2022.

2.4.4) In the event that the CC does not approve admission, the Candidate may submit a new application for admission to the Registry until the applicant has satisfied the requirements.

2.4.5) Members of SIF-EuCP Program are "Certified Pharmacologists" and may use the abbreviation "EuCP" (European Certified Pharmacologist).

Section 3. Maintenance of Certification

3.1 On a 5-year basis, the European Certified Pharmacologists (SIF-EuCP Program members) shall re-affirm their certification credentials submitting an application of "Maintenance of certification" using the designated form (see "Maintenance of certification Application Form"). To remain registered, a EuCP must be employed or be active or seek employment in the field of Pharmacology (continuous practice of research in Pharmacology and a continuous professional development i.e: CPD, ECM or ECTS, publications, reports and other academic training such as PhDs, specializations, presentations, teaching activities, publications,

activities in expert committees and/or similar) as required by the EuCP Guidelines (ANNEX III assessment check list).

3.2 Failure to produce evidence to support the maintenance of certification shall result in the revocation or termination of the certification.

Section 4. Membership Fees and costs and SIF-EuCP Program financing

4.1 A non-refundable *una tantum* fee of € 30.00 is due by each candidate for application examination (both admission or renewal application).

4.2 A five-year fee of € 300.00 (€ 150.00 if applicant age is below 40) must be paid by each member to SIF-EuCP Program to access the Register, within 30 days from the receipt of the payment request.

Section 5. Suspension of Members

5.1 Members who do not renew their certification, will be removed from the Register.

5.2 Members may be removed from the Register for ethical reasons. In this case, an appropriate report containing the reasons shall be drawn up by the CC. Suspension decisions could be appealed (see 11.1).

Section 6. The Lead Body - Certification Commission (CC)

6.1 Appointment of the CC

The members of the CC shall be appointed by the SIF EC.

6.2 The Functions of CC are the following:

a) The evaluation of candidate application for inclusion and renewal in the Register

b) the exclusion or removal of a Member from the Register

6.3 Composition of the CC

- The CC is composed of five members, appointed by SIF EC, and holds office for three years. A member of the SIF Secretariat will participate in meetings with only administrative functions.
- The members of the CC shall elect a Chairman and a Secretary
- CC Members may be renewed but they may not hold office consecutively for more than six years (two mandates)
- In the event of resignation or retirement of a member of the CC, he/she will be replaced in accordance with paragraph 6.3.

Section 7. Secretariat Office

The Organizing Secretariat Office of SIF-EuCP Program is entrusted to the secretariat of SIF.

Section 8. Operation of the CC

8.1 **Meetings**

8.1.a) The CC shall meet at least once a year and may be hold also with telematic contact. Travel grants could be ensured to support participation in person.

8.1.b) The meeting is valid if a simple majority of members are present.

8.1.c) Observers and external experts who may assist the CC in the process of evaluating applications for admission~~,~~ without the right to vote, can be invited to the meetings at the discretion of the Chair.

8.2.3 The Chair formulates the agenda and presides over the CC. In his/her absence, the members present at the meeting indicate a deputy chair that shall act as President only for the specific meeting. In each case, a meeting it is considered valid in the presence of a simple majority of members.

8.4 The Secretary, at the request of the Chair or at the request of a simple majority members, convenes the CC (via e-mail) with a notice of at least 15 days, indicating the agenda.

8.5 The CC may elaborate additional operational rules as stated in section 6, 7 and 8 of this Regulation.

8.6 In case a regulation was not prepared, the President's and Secretary's rights and duties must be agreed and/or determined by the Committee from time to time.

Section 9. Functions of the CC

9.1 All issues concerning the SIF-EuCP Program fall within the competence of the CC
9.2 The CC is responsible for:

- preparing an annual report and define fees, in consultation with SIF EC
- managing the SIF-EuCP Program registry and budgets (see section 4)
- including, excluding or removing members from the Register (see sections 2; 3 and 5)
- informing and contributing to the SIF-EuCP Program development in Italy
- choose experts to be convened at meetings, if necessary (see section 8.1.c).

Section 10. Functions of the Secretariat

10.1 The Secretariat is responsible for:

a) keeping all the administrative documents (SIF-EuCP Program register and SIF-EuCP Program members), prepare detailed minutes of all meetings and archive CC documents

b) informing interested applicants on the requirements for the applications

c) notifying and informing applicants of the CC decisions, as quickly as possible

d) preparing certificates when fees have been paid

e) communicating periodically EuCP about the new members of SIF-EuCP Program, including dates of first certification and certification renewal

f) issuing an annual report with the CC analysis and results for submission to SIF Assembly, specifying at least:

- the number of those who applied to be included in the Register, the outcome of requests and the percentage of approved requests
- the names of those who have been removed from the Register for failure to pay the fee or for any other reason
- an evaluation of the quality of the candidates on the basis of what is required for inclusion in the register
- the needs for the continuous professional development of members
- a detailed report on the financial status of the Register
- the composition of the CC and any changes that occurred in the period of the Commission appointment.

g) manage events day by day, including statutory obligations of SIF-EuCP Program and implement decisions of SIF in relation to standards of SIF-EuCP Program and every need that arises from new regulations for members of EuCP.

Section 11. Appeals Against Decisions of the CC Appeal Certification Committee (ACC)

11.1 If a candidate disagrees with the CC's decision, the application must be reconsidered and evaluated by CC within 6 months. In the event that the candidate does not agree after the revaluation made by the CC, he will be able to ask for an appeal by the independent Appeal Certification Committee (ACC) or Probiviris.

11.2 The ACC consists of three members, a Past President of SIF and two members of the SIF Executive Council.

11.3 The decision of the ACC will be binding on all parties.

Section 12. SIF-EuCP Program General Assembly

The SIF-EuCP Program General Assembly must be convened at the SIF Congress and can be called at any time and place, depending on the decisions of the CC. The SIF secretariat will send notice of the assembly – via e-mail – at least 30 days before, to all members and to all those who are entitled to receive notice. All members of SIF-EuCP Program are entitled to participate.

Section 13. Expulsion

13.1 The CC may expel from SIF-EuCP Program every member whose misconduct (which will include but will not be limited to non-payment) is, in the opinion of the CC, adversely affecting SIF-EuCP Program. (see 5.1) Neither SIF-EuCP Program, its officers, employees nor representatives, nor the CC, ACC nor any Member shall have any obligation to the person who has been expelled.

Section 14. Variation of the Regulation

14.1 These Regulations may be changed only when the CC, unanimously, and the SIF EC, with three-fourths majority of the members present, approve the amendment.

14.2 The notification of the request to amend the existing regulation, must be sent by the CC to the Secretariat in writing.

Section 15. Disputes and Controversies

15.1 Except as specified in this document, any dispute or controversy that may arise from the interpretation of the regulations or the powers or the validity of the minutes of a meeting, will be determined by the CC, in agreement with the SIF EC whose decision shall be final and binding upon all members.

Section 16. Legend and Notes

NOTES:

[1] second cycle qualification of relevant foreign higher system, awarded by official institution of foreign higher education system.

[2] Medicina e Chirurgia, Farmacia e Chimica e Tecnologie Farmaceutiche sono Lauree a ciclo unico rispettivamente di 6 e 5 anni.

[3] In Italy there are two equivalent five years degree in pharmaceutical area: pharmacy and chemistry-pharmaceutical technology (CTF).

[4] Sono ipotizzabili due profili per i candidati. Il **profilo accademico** contempla il ruolo di Ricercatore a tempo determinato di cui all'articolo 24, comma 3, lettera a) Legge 240 del 2010 (RTDa). Assegnista di ricerca Contratto ai sensi dell'art. Legge 240/2010– Fonte MIUR (Rev 2019). Borsista di ricerca con almeno un anno di contratto.

[5] **Personale universitario a tempo indeterminato** in Italia: Professore di I fascia; Professore di II fascia; Ricercatore a tempo indeterminato (ruolo ad esaurimento); Assistenti universitari (ruolo ad esaurimento). Ricercatore a tempo determinato di cui all'articolo 24, comma 3, lettera b) Legge 240 del 2010 (RTDb).

[6] Il **profilo non accademico**: professionisti nel campo delle scienze farmaceutiche, chimiche, mediche, biomediche e in attività di libera professione quali titolari di borse di studio o impiegati a tempo indeterminato presso strutture sanitarie pubbliche e private, aziende farmaceutiche, istituti di ricerca pubblici e privati, enti o agenzie regolatorie e di valutazione di tecnologie sanitarie, agenzie o enti di valutazione dei medicinali.

[7] I candidati italiani allegano autocertificazione dei titoli e competenze. (Art. 46 - lettera l, m, n - D.P.R. 28 dicembre 2000, n. 445). Annex IV

ALLEGATI

ANNEX I Check list valutativa delle competenze e delle esperienze (CON PUNTEGGIO) per il profilo "ACCADEMICO"

ANNEX II Check list valutativa delle competenze e delle esperienze (CON PUNTEGGIO) per il profilo "PROFESSIONISTI DELLA SALUTE"

ANNEX III Check list valutativa delle competenze e delle esperienze comuni ai due profili e da utilizzare per la procedura di rinnovo del certificato

ANNEX IV Dichiarazione sostitutiva della CERTIFICAZIONE DEL TITOLO DI STUDIO, delle competenze e delle pubblicazioni

LEGENDA

SIF-EuCP Program: Registro Italiano dei Farmacologi Certificati – Italian Certified Pharmacologists Register.

CC: Certification Commission: The Commission for candidates' evaluation and approval for inclusion in SIF-EuCP Program.

SIF Società Italiana di Farmacologia: the Italian Society of Pharmacology.

EPHAR: The Federation of European Pharmacological Societies.

EACPT: The European Association for Clinical Pharmacology and Therapeutics.

ACC: Appeal Committee (Probiviri): Eminent Pharmacologists' Committee (3 members).

LAST REVISION: 18/04/2020

ANNEX I (Academic PROFILE)

	POINTS
Advanced academic background	
• Specialization (Clinical Pharmacology and Toxicology)[1]	10
• PhD	6
• Other medical specialization[2]	5
• Master	3
Post-graduate 5 years background	
• Pre-clinical Research	7
• Clinical Research	7
Continuous Education (lifelong) (max 5 points)	
• ECTS/Academic advanced courses[3]	4
• Certifications in Toxicology	3
Professional position (applicable at Universities) (The highest)	
• Researcher/lecturer/Assistant Professor[4]	10
• Associate Researcher[5]	8
• Research fellow[6]	6
Professional background in Pharmacology/toxicology field	
• Awards (in the last 15 years)	max 6
• Grants[7] (in the last 15 years)	max 6
• Clinical trial research team membership[8] (in the last 15 years)	max 6
• Scientific societies membership (European Circuit EPHAR) (depending year of registration)	max 10
• Other Scientific societies membership[9]	max 6
• Teaching[10]	max 8
• Editorial boards	max 7
• Peer review Publications (min. 3 in the last three years)	max 10
Total (min 60 POINTS)	

NOTES

La colonna "Points" costituisce un riferimento per il candidato per poter controllare in anticipo quale potrebbe essere il punteggio raggiungibile con i titoli dichiarati nell'ANNEX IV e potrebbe differire dal punteggio finale assegnato dalla Commissione Certificatrice in fase di valutazione.

[1]Specialization: Clinical Pharmacology and Toxicology or Medical Pharmacology /Toxicology; Specializzazione in Farmacologia Medica o Tossicologia
[2]Other medical specialization: in any other medical field; Altre specialità nel campo delle discipline mediche o Specialità in Farmacia Ospedaliera
[3]Corsi di perfezionamento universitari, formazione a distanza certificata (devono essere identificabili nel CV e nell'ANNEX IV ad esempio riportando gli estremi di un attestato di partecipazione).
[4]In Italia: Personale di Ricerca a tempo determinato (RTDa)

[5]Associate Researcher: in Italia Assegnista di ricerca ai sensi dell'art. Legge 240/2010
[6]Research Fellow: Borsista con almeno un anno di contratto; min. one year contract

[7]Grants: Finanziamento di bando competitivo a carattere nazionale e/o internazionale

[8]Clinical trial research team membership-partecipazione ad uno o più trial clinici con i seguenti possibili incarichi:

- Principal Investigator: principal investigator is the professional responsible for leading the clinical trial at the site.
- Investigator**:** An investigator for a clinical trial can hold a number of responsibilities, including but not limited to a serving as a physician, pharmacist, biostatistician, consenting and/or treating participants, conducting data analysis and determining results, or conducting basic research on collected specimens in a laboratory.

[9]Iscrizione ad altre società scientifiche di area medica
[10]Teaching: incarichi di insegnamento e di attività didattica integrativa; partecipazione a commissioni d'esame di profitto, in qualità di cultore della materia

ANNEX II (Health professionals PROFILE)

	POINTS
Advanced academic background	
• Specialization (Clinical Pharmacology and Toxicology)[1]	10
• PhD	6
• Other medical specialization[2]	5
• Master	3
Post-graduate 5 years background	
• Pre-clinical Research	10
• Clinical Research	10
Continuous Education (lifelong)	
• CME courses	max 6
• ECTS/Academic advanced courses[3]	max 8
• Certifications (i.e. Toxicology, quality, etc)	max 6
Professional position (applicable at Pharmaceutical Industry/Hospitals) (the highest)	
• Chairman/Owner[4]	12
• High level Direction (CEO)	11
• Chief[5]	10
• Manager[6]	9
• Hospital general director[7]	10
• Hospital department director[8]	9
• Hospital medic[9]	8
• Hospital medic[10]	7
• Hospital medic[11]	6
• General Practitioner[12]	8
• Hospital Pharmacist Director	9
• Hospital Pharmacist/Biologist[13]	8
• Hospital Pharmacist/Biologist[14]	7
• Fellow Hospital Pharmacist/Biologist[15]	6
• Public Pharmacist[16]	8
• Biomedical laboratory technician[17]	6
Professional background in Pharmacology/toxicology field	
• Awards (in the last 15 years)	max 6
• Grants (in the last 15 years)[18]	max 6
• Clinical trial research team membership [19] (in the last 15 years)	max 8
• Drug repurposing	max 8
• Scientific societies membership (European Circuit EPHAR) (depending year of registration)	max 10
• Other Scientific societies membership[20]	max 6
• Advisory boards (in the last 15 years)	max 6
• Teaching[21]	max 6
• Editorial boards	max 5
• Publications (Peer review, company reports, guidelines, etc.)	max 10
Total (min 60 POINTS)	

NOTES

La colonna "Points" costituisce un riferimento per il candidato per poter controllare in anticipo quale potrebbe essere il punteggio raggiungibile con i titoli dichiarati nell'ANNEX IV e potrebbe differire dal punteggio finale assegnato dalla Commissione Certificatrice in fase di valutazione.

[1]Specialization: Clinical Pharmacology and Toxicology or Medical Pharmacology /Toxicology; Specializzazione in Farmacologia Medica o Tossicologia
[2]Other medical specialization: in any other medical field; Altre specialità nel campo delle discipline mediche o Specialità in Farmacia Ospedaliera
[3]Corsi di perfezionamento universitari, formazione a distanza certificata (devono essere identificabili nel CV e nell'ANNEX IV ad esempio riportando gli estremi di un attestato di partecipazione).

Pharmaceutical
[4]Owner, Titolare Dell'Azienda farmaceutica, Biotecnologica, Farmacista Titolare, Farmacista Amministratore

Pharmaceutical
[5]Chief
CMO, Chief Marketing Officer (Direttore Marketing)
CIO, Chief Information Officer (Direttore informazione)
CKO, Chief Knowledge Officer (Direttore Organizzazione)
CCO, Chief Compliance Officer (Direttore della conformità)

[6]Manager
HR Manager, Human Resource Manager (Direttore del personale)
R&D Manager, Research & Development Manager (Direttore del reparto ricerca & sviluppo)
PM Manager, Product Manager (Responsabile prodotto)
PM Manager, Project Manager (Responsabile di progetto)
PM Manager, Plant Manager (Direttore di stabilimento)

Hospital
[7]Direttore Sanitario, Medico, Medico e Scientifico (Istituti di ricerca, IRCCS, etc)
[8]Primario, Vice Primario, Capo Dipartimento (Aziende Ospedaliere Universitarie)

Hospital
[9] a tempo indeterminato (strutturato), Dirigente Medico, Caposervizio, Resident physician
[10]Medico a tempo determinato, Capo clinica, consulente o in libera professione
[11]Medico specializzando, Medico assistente, Assistant physician
[12]Medico di Medicina Generale
[13]Dirigente Farmacista Ospedaliero, Dirigente Biologo
[14]Farmacista o Biologo specializzato in Farmacologia a tempo determinato, consulente o in libera professione
[15]Farmacista o Biologo Specializzando in Farmacologia con contratto di formazione, consulente o in libera professione
[16]Farmacista dipendente in Farmacia Aperta al Pubblico (Privata o Comunale)
[17]Tecnici di laboratorio universitari od ospedalieri laureati cat. D o EP

[18]Grants: Finanziamento di bando competitivo a carattere nazionale e/o internazionale

[19]Clinical trial research team membership - partecipazione ad uno o più trial clinici con i seguenti possibili incarichi:

- Principal investigator is the professional responsible for leading the clinical trial at the site
- Investigator: An investigator for a clinical trial can hold a number of responsibilities, including but not limited to a serving as a physician, pharmacist, biostatistician, consenting and/or treating participants, conducting data analysis and determining results, or conducting basic research on collected specimens in a laboratory.

- Clinical Research Coordinator/Associate: Clinical Research Coordinators or Associates, often called CRCs or CRAs are a critical research support role that handles the daily conduct of a study. They interact heavily with both participants and investigators, doing things like screening participants, obtaining informed consent, checking eligibility, collecting and entering data, and more.

[20]Appartenenza ad altre Società scientifiche

[21]Teaching: incarichi di insegnamento e di attività didattica integrativa; partecipazione a commissioni d'esame di profitto, in qualità di cultore della materia

ANNEX III Maintenance of certification (All profiles)

POINTS

Advanced academic background

- Specialization (Clinical Pharmacology and Toxicology)
- PhD
- Other medical specialization
- Master

Post-graduate 5 years background

- Pre-clinical Research
- Clinical Research

Continuous Education (lifelong)

- CME courses
- ECTS/Academic advanced courses
- Certifications in Toxicology
- Certifications in Quality, etc

Profession position (applicable at Universities) (The Highest)

- Researcher/Lecturer/Assistant Professor
- Associate Researcher
- Research Fellow

Professional position (applicable at Pharmaceutical Industry/Hospitals) (the highest)

- Chairman/Owner
- High level Direction (CEO)
- Chief
- Manager
- Hospital general director
- Hospital department director
- Hospital medic
- General Practitioner
- Hospital Pharmacist Director
- Hospital Pharmacist/Biologist
- Fellow Hospital Pharmacist/Biologist
- Public Pharmacist
- Biomedical laboratory technician

Professional background in Pharmacology/toxicology field

- Awards (in the last 15 years)
- Grants (in the last 15 years)
- Clinical trial research team membership (in the last 15 years)
- Drug repurposing
- Scientific societies membership (European Circuit EPHAR) (depending year of registration)
- Other Scientific societies membership
- Advisory boards (in the last 15 years)

- Teaching
- Editorial boards
- Peer review Publications (min 3 in the last three years)
- Publications (Peer review, company reports, guidelines, etc.)

Total (min 60 POINTS)

NOTES

La colonna "Points" costituisce un riferimento per il candidato per poter controllare in anticipo quale potrebbe essere il punteggio raggiungibile con i titoli dichiarati nell'ANNEX IV e potrebbe differire dal punteggio finale assegnato dalla Commissione Certificatrice in fase di valutazione.

Please, see specific ANNEX I and ANNEX II notes

APPLICATION FORM

This form is used to send a new request for a comparative evaluation. Fill out all sections, referring to program guidelines and ANNEX documents, if necessary. If you need more space, use a separate sheet of paper and write your name, date of birth and section number. Any omission may result in the return or closure of your request without processing.

1. Preliminary information

First request ☐ Second or more request ☐

maintenance of certification request. Please, consider ANNEX III ☐

2. Identity Information

Write your last and first name(s) in block letters. ▶

Last name (as indicated on the accepted identity document)	First name(s) (as indicated on the accepted identity document)
Last name and first name at birth (if different from the name)	Other names on your educational documents
Date of birth (year/month/day)	Country of birth

State your full mailing address, with the elements of your country's addressing system, to receive our correspondence.

Building number	Street	Apartment	Post Office Box
City	Province, territory or state	Postal code	Country
Email address	Telephone number	Other telephone	
Future mailing address (if applicable)		Date of move (if applicable)	

3. Current professional position

4. Profile

Academic. Please, consider ANNEX I ☐ Health Professionals. Please, consider ANNEX II ☐

5. Evaluation Fee

30 Euro Evaluation Fee is due. Please attach the invoice of payment

6. Declaration, Privacy statement, contacts and payment information

Please, consider ANNEX IV FOR declaration of ANNEX I/ANNEX II titles

Hereby, I declare:
that I have provided all the necessary information and documents, in the required format, for the comparative evaluation and that I have provided only legible, accurate and truthful information and documents (Please see ANNEX IV).

The Italian Society of Pharmacology declares that the personal data communicated by the user are processed in accordance with the provisions of Legislative Decree no. 196/2003, as amended by Legislative Decree no. 101/2018, and EU legislation (EU Regulation 2016/679) as specifically indicated in the privacy policy available on the website of the Company at: https://sif-website.s3.amazonaws.com/uploads/attachment/file/240/Informativa_Privacy_SIF_Generica.pdf that the user, by signing this Agreement, declares to have fully reviewed, understood and accepted.

La Società Italiana di Farmacologia dichiara che i dati personali comunicati dall'utente sono trattati in conformità alle disposizioni del D. Lgs. 196/2003, così come modificato dal D. Lgs. 101/2018, ed alla normativa comunitaria (Regolamento UE 2016/679) secondo quanto indicato specificamente nell'informativa privacy reperibile sul sito internet della Società all'indirizzo: https://sif-website.s3.amazonaws.com/uploads/attachment/file/240/Informativa_Privacy_SIF_Generica.pdf che l'utente, con la sottoscrizione del presente Contratto, dichiara di aver compiutamente visionato, compreso e accettato.

Date_____ Signature _____

Please, send all documents to: **SIF-EuCP Program** c/o Segreteria Organizzativa SIF – Via Giovanni Pascoli, 3 - 20129 Milano tel 02-29520311 - fax 02-700590939 - e-mail: sif.farmacologia@segr.it - object: **application SIF-EuCP Program**

Payments: **Società Italiana di Farmacologia**- Bank: Credito Valtellinese, Agency n.16 di Milano
IBAN: **IT 86 E 05216 01621 000 000 000 530** BIC/SWIFT: **BPCVIT2S**

Further information: sif.farmacologia@segr.it - object: **information SIF-EuCP Program**

Documents Check LIST

☐ Application FORM

☐ CV

☐ ANNEX I

☐ ANNEX II

☐ ANNEX III

☐ ANNEX IV

☐ Invoice evaluation fee