

99005025016000

# Apply for recognition as a pharmaceutical consultant

Heruntergeladen am 14.06.2025

<https://fimportal.de/xzufi-services/6022571/L100022>

Modul	Sachverhalt
Leistungsschlüssel	99005025016000
Leistungsbezeichnung I	Apply for recognition as a pharmaceutical consultant
Leistungsbezeichnung II	Apply for recognition as a pharmaceutical consultant
Typisierung	3 - Bundesaufsichtsverwaltung: Regelung
Quellredaktion	Baden-Württemberg
Freigabestatus Katalog	unbestimmter Freigabestatus
Freigabestatus Bibliothek	unbestimmter Freigabestatus
Begriffe im Kontext	
Leistungstyp	
Leistungsgruppierung	
Verrichtungskennung	
SDG-Informationsbereich	
Lagen Portalverbund	
Einheitlicher Ansprechpartner	

Modul	Sachverhalt
Fachlich freigegeben am	
Fachlich freigegeben durch	
Handlungsgrundlage	
Teaser	<p>according to pharmaceutical legislation, pharmaceutical companies may only appoint persons with the relevant expertise to provide full-time healthcare professionals with specialist information on medicinal products. These experts are pharmaceutical counsellors.</p> <p>The necessary expertise for this position is held by</p>
Volltext	<p>according to pharmaceutical legislation, pharmaceutical companies may only appoint persons with the relevant expertise to provide full-time healthcare professionals with specialist information on medicinal products. These experts are pharmaceutical counsellors.</p> <p>The necessary expertise for this position is held by</p> <ul style="list-style-type: none"> <li>• Pharmacists or persons with a certificate of having passed an examination after completing a university degree in the following areas <ul style="list-style-type: none"> <li>• pharmacy,</li> <li>• chemistry</li> <li>• biology,</li> <li>• human medicine or</li> <li>• veterinary medicine,</li> </ul> </li> <li>• Pharmacy assistants and persons who have completed training as a technical assistant in <ul style="list-style-type: none"> <li>• pharmacy,</li> <li>• chemistry</li> <li>• biology,</li> <li>• human</li> <li>• or veterinary medicine,</li> </ul> </li> <li>• Pharmaceutical sales representatives.</li> </ul> <p>The competent authority may recognise as sufficient an examination or completed training that is at least equivalent to one of the training courses of the aforementioned persons.</p>

## Modul

## Sachverhalt

### Erforderliche Unterlagen

- signed curriculum vitae in tabular form on the professional career
  - all professional qualifications obtained
  - officially certified copy of the final certificate or diploma
  - A document detailing the exact course and content of the training, including details of subjects and hours
  - if applicable, confirmation from the current or future employer with details of the place of work
  - if applicable, certificate of equivalence, e.g. for professional qualifications obtained in third countries
  - proof of name changes, if applicable, if your name has changed since the documents were issued
  - copy of identity document or current registration certificate, if applicable

### Voraussetzungen

- You are
  - A pharmacist,
  - Assistant pharmacist or pharmacist assistant or
  - You have a comparable degree or training in the following areas
    - pharmacy,
    - chemistry
    - biology
    - human
    - or veterinary medicine or
- You are a pharmaceutical sales representative.

The examination regulations of the individual training and study subjects are relevant for the recognition of comparable studies or comparable training.

Exceptions for which no application is required:

- You have
  - a foreign degree in the study programmes under the Medicinal Products Act or
  - a foreign professional qualification in the professions under the Medicinal Products Act.
  - Your degree or professional qualification has been recognised as equivalent by a competent German authority.

Modul	Sachverhalt
Kosten	Fees in accordance with the state fee schedule
Verfahrensablauf	In Baden-Württemberg, the central responsibility for the recognition of pharmaceutical consultants in accordance with the Medicinal Products Act lies with the Baden-Württemberg Drug Monitoring Centre at the Regional Council of Tübingen.
Bearbeitungsdauer	Depending on the individual case, you should expect a processing time of 4 - 6 weeks from receipt of the complete documents.
Frist	There are no deadlines to be observed.
weiterführende Informationen	
Hinweise	<p>Please note the following:</p> <p>All documents that are not written in German must also be available in translated form. Translations are accepted if they have been translated by a</p> <ul style="list-style-type: none"> <li>• Germany,</li> <li>• the other contracting countries of the European Economic Area (EEA) or</li> <li>• Switzerland</li> </ul> <p>sworn interpreter or translator appointed or sworn in Germany, the other contracting states of the European Economic Area (EEA) or Switzerland.</p> <p>Translations produced outside Germany, the EEA or Switzerland are generally not recognised. Exception: Translations already produced in a third country must be submitted to an interpreter or translator appointed or sworn in Germany to check their accuracy and then submitted here.</p>
Rechtsbehelf	• Administrative court action
Kurztext	
Ansprechpunkt	

Modul	Sachverhalt
Zuständige Stelle	
Formulare	
Ursprungsportal	