

99005019261000

Show information officers for pharmaceutical companies

Heruntergeladen am 18.07.2025

<https://fimportal.de/xzufi-services/6018991-99005019261000/L100022>

Modul	Sachverhalt
Leistungsschlüssel	99005019261000
Leistungsbezeichnung I	Show information officers for pharmaceutical companies
Leistungsbezeichnung II	Show information officers for pharmaceutical companies
Typisierung	3a - Bundesaufsichtsverwaltung: Regelung, Land: Vollzug, 2a - Bundesauftragsverwaltung: Regelung, Land: Vollzug
Quellredaktion	Baden-Württemberg
Freigabestatus Katalog	unbestimmter Freigabestatus
Freigabestatus Bibliothek	unbestimmter Freigabestatus
Begriffe im Kontext	
Leistungstyp	
Leistungsgruppierung	
Verrichtungskennung	
SDG-Informationsbereich	

Modul	Sachverhalt
Lagen Portalverbund	
Einheitlicher Ansprechpartner	
Fachlich freigegeben am	
Fachlich freigegeben durch	
Handlungsgrundlage	<p>Arzneimittelgesetz (AMG)</p> <ul style="list-style-type: none"> • § 74a <p>Arzneimittel- und Wirkstoffherstellungsverordnung (AMWHV)</p> <ul style="list-style-type: none"> • § 12
Teaser	<p>If you operate a pharmaceutical company that places finished medicinal products on the market, you must notify the competent authority of an information officer. Any changes must also be reported immediately.</p>
Volltext	<p>If you operate a pharmaceutical company that places finished medicinal products on the market, you must notify the competent authority of an information officer. Any changes must also be reported immediately.</p> <p>The information officer must have the appropriate expertise and reliability .</p> <p>They are responsible, among other things, for ensuring that medicinal products are correctly registered, properly labelled and not misleadingly advertised.</p>
Erforderliche Unterlagen	<ul style="list-style-type: none"> • Employment references (copy) • Proof of training (certified copy) • Curriculum vitae • Certificate of good conduct (document type O for direct transmission from authority to authority) • Form "Declaration of nomination" • Declaration of commitment
Voraussetzungen	<ul style="list-style-type: none"> • Information officers must have the necessary

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	expertise and reliability (see Section 74a of the German Medicinal Products Act)
Kosten	There are no costs for the advert.
Verfahrensablauf	<p>You can notify an information officer in writing or online:</p> <ul style="list-style-type: none"> • You notify the information officer by means of a written application or using the online service. Proof of the person's expertise must be attached to the notification. • Upon receipt, the authority checks the notification formally and for completeness. • If the check reveals that documents are missing, the person making the notification will be contacted and asked to provide the missing documents. • Once the missing documents have been submitted or the formal check has been passed, the competent authority will make a decision. • The notification can be confirmed or rejected. • You will be informed of the decision. • A statement of fees will then be drawn up and sent to you with a request for payment.
Bearbeitungsdauer	
Frist	Any change must be notified in advance. In the event of an unforeseen change of information officer, notification must be given immediately.
weiterführende Informationen	
Hinweise	A fine may be imposed if the obligation to notify is violated.
Rechtsbehelf	<ul style="list-style-type: none"> • Objection • Information on how to lodge an objection will be sent with the decision on your advert.
Kurztext	
Ansprechpunkt	
Zuständige Stelle	

Modul

Sachverhalt

Formulare

Ursprungsportal
