

**PAYMENT FORM FOR THE
TARIFF FOR MARKETING
AUTHORISATION ACCORDING TO
ARTICLE 893 OF LAW NO. 95/2006 AND THE
TARIFF FOR THE ASSESSMENT OF
DOCUMENTATION IN VIEW OF
MARKETING AUTHORISATION
ACCORDING TO MINISTER OF HEALTH
ORDER NO. 888/2014
FOR MEDICINAL PRODUCTS PROPOSED FOR
AUTHORISATION THROUGH MUTUAL
RECOGNITION OR DECENTRALISED
PROCEDURE WITH ROMANIA AS REFERENCE
MEMBER STATE**

Name of the medicinal product:

--

Pharmaceutical form, strength, administration route

Pharmaceutical form:	
Strength:	
Administration route:	

Marketing Authorisation Holder

Name:	
Address:	
City:	
Country:	
Telephone no.:	
Fax no.:	

E-mail address:	
-----------------	--

Status of the medicinal product
--

Authorisation	<input type="checkbox"/>
---------------	--------------------------

Type of authorisation procedure
--

Mutual recognition procedure	<input type="checkbox"/>
Decentralised procedure	<input type="checkbox"/>

Paying company

Name:	
Address:	
City:	
Country:	
Telephone no.:	
Fax no.:	
E-mail address:	
Fiscal code:	
Trade Registry no.:	
IBAN Account no. :	
Bank:	

Proposed form of payment

Lei :	<input type="checkbox"/>
Euro :	<input type="checkbox"/>

Tariff for marketing authorisation according to Article 893 of Law no. 95/2006 on healthcare reform, as republished, with the further amendments

For all types of medicinal products mentioned by Law no. 95/2006 on healthcare reform = 5000 €

Tariff for the assessment of documentation in view of marketing authorisation through European procedures

Activity		The fee in euro currency according to the MHO no. 888/2014 ^{*)}
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State – generic medicinal products [Article 10(1) of Directive 2001/83/EC or Article 708 (1) and (2) of Law 95/2006, as republished, with the further amendments]	<input type="checkbox"/>	
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State – generic medicinal products – different pharmaceutical form, submitted at the same time as the initial application [Article 10(1) of Directive 2001/83/EC or Article 708 (1) and (2) of Law 95/2006, as republished, with the further amendments]		
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State – generic medicinal products – the second and following strengths, submitted at the same time as the initial application [Article 10(1) of Directive 2001/83/EC or Article 708 (1) and (2) of Law 95/2006, as republished, with the further amendments]		

<p>Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State - "hybrid" (mixed) application [Article 10(3) of Directive 2001/83/EC or Article 708 (3) of Law 95/2006, as republished, with the further amendmends]</p>		
<p>Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State - "hybrid" (mixed) application - different pharmaceutical form, submitted at the same time as the initial application [Article 10(3) of Directive 2001/83/EC Article 708 (3) of Law 95/2006, as republished, with the further amendmends]</p>		
<p>Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State - "hybrid" (mixed) application - the second and following strengths, submitted at the same time as the initial application [Article 10(3) of Directive 2001/83/EC or Article 708 (3) of Law 95/2006, as republished, with the further amendmends]</p>		
<p>Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State - "biosimilar medicinal product" [Article 10(4) of Directive 2001/83/EC or Article 708(4) of Law 95/2006, as republished, with the further amendmends]</p>		
<p>Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State - "biosimilar medicinal product" - different pharmaceutical form,</p>		

submitted at the same time as the initial application [Article 10(4) of Directive 2001/83/EC or Article 708 (4) of Law 95/2006, as republished, with the further amendmends]		
Marketing authorisation of medicinal products through procedure or decentralised procedure with Romania as Reference Member State - "biosimilar medicinal product" – the second and following strengths, submitted at the same time as the initial application [Article 10(4) of Directive 2001/83/EC or Article 708(4) of Law 95/2006, as republished, with the further amendmends]		
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State - "bibliographic" application [Article 10(a) of Directive 2001/83/EC or Art. 709 of Law 95/2006, as republished, with the further amendmends]	□	
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State - "bibliographic" application - different pharmaceutical form, submitted at the same time as the initial application [Article 10(a) of Directive 2001/83/EC or Article 709 of Law 95/2006, as republished, with the further amendmends]		
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State - "bibliographic" application - the second and following strengths, submitted at the same time as the initial application [Article 10(a) of Directive 2001/83/EC or Article 709		

of Law 95/2006, as republished, with the further amendmends]		
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State – fixed combination [Art. 10(b) of Directive 2001/83/EC or Article 710 of Law 95/2006, as republished, with the further amendmends]	<input type="checkbox"/>	
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State – fixed combination - different pharmaceutical form, submitted at the same time as the initial application [Article 10(b) of Directive 2001/83/EC or Article 710 of Law 95/2006, as republished, with the further amendmends]		
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State – fixed combination - the second and following strengths, submitted at the same time as the initial application [Article 10(b) of Directive 2001/83/EC or Article 710 of Law 95/2006, as republished, with the further amendmends]		
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State - "informed consent" [Article 10(c) of Directive 2001/83/EC or Article 711 of Law 95/2006, as republished, with the further amendmends]	<input type="checkbox"/>	
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State - "informed consent" - different		

pharmaceutical form, submitted at the same time as the initial application [Article 10(c) of Directive 2001/83/EC or Article 711 of Law 95/2006, as republished, with the further amendmends]		
Marketing authorisation of medicinal products through mutual recognition procedure or decentralised procedure with Romania as Reference Member State - "informed consent" – the second and following strengths, submitted at the same time as the initial application [Article 10(c) of Directive 2001/83/EC or Article 711 of Law 95/2006, as republished, with the further amendmends]		

*) the applicant will fill in the fee in euro currency

Date of application submission (Applicant, NAMMDR)

Representative to Romania/Contact person

Name:	
Address:	
City:	
Country:	
Telephone no.:	
Fax no.:	
E-mail address:	

Signatories assume responsibility for accuracy of data in the present form.

Date.....

Marketing Authorisation Holder/Representative to Romania
Name, signature, stamp