

**Detailed guidance for National Scientific-  
Technical Advice (STA) requests:  
(Version 1.4)**

## 1. Introduction:

The DG PRE-Authorisation of the federal agency for medicines and health products (famhp) in Belgium offers applicants the possibility to request national scientific and/or technical (eg. regulatory) advice (STA) related to the research and development aspects of human or veterinary drug products in view of potential clinical trial applications (CTA's), marketing authorization applications (MAA's), introduction of variations to marketed drug products or line extensions. The famhp's main objective in providing STA to applicants at a national level is to promote and facilitate as much as possible the development of new drug products from a regulatory perspective in order to enhance the availability of innovative drug products to patients, especially in therapeutics areas where an unmet medical need exists.

For this purpose the scientific-technical advice unit of the DG PRE-Authorisation offers a centralized and transparent service within the agency which should ensure the processing of national STA requests in a timely fashion while assuring full confidentiality and potential conflict of interest of the involved experts. The famhp also aims to provide a consistent follow-up of previous national and European advices (eg. through its interface with the SAWP and SAWP-V at the EMA) in order to assure the quality and consistency of the national STA's issued by the famhp.

The national STA may be provided to sponsors of clinical trials, pharmaceutical/biotech companies (eg. small biotech spin offs, global companies), research centres, etc.

As the progress in science and subsequently the scientific/regulatory guidelines are constantly evolving over time, the national STA issued by the famhp is to be considered as NOT legally binding with regard to any future, related application (eg. CTA's), neither towards the famhp, nor to the applicant. The advice given by the famhp is based on the questions and information submitted by the applicant at the time of the STA request and cannot account for any future changes in science and the scientific / regulatory guidelines. After an initial STA request has been addressed by the famhp, applicants are free to submit at any time a follow-up STA request (eg. whenever they feel the need to seek advice on new development data that have been obtained, changes in scientific / regulatory guidelines have taken place, etc).

National STA requests may be submitted to the famhp at any time, independently from other ongoing or planned applications (eg. CTA's, MAA's, SAWP advices, PIP's). Nevertheless, applicants are strongly being recommended to seek the famhp's advice well in advance of a forthcoming application. However, the famhp maintains the right to refuse STA requests in case legal procedures are ongoing which are related to the STA request and in which famhp experts may be involved. In this case, the STA request cannot be treated by the famhp as long as the legal procedure has not been closed.

The famhp also wishes to point out that a national STA related to a forthcoming application (eg. CTA, MAA,...) that the applicant plans to submit in the near future is **NOT** to be considered as a preliminary assessment or approval of the planned application.

The objective of this Guidance is to provide applicants with information regarding the planning and submission of national STA requests in order to ensure an effective and efficient handling of the advice requests throughout the procedure. In particular, guidance is given regarding the conditions, timelines and rules of procedure as well as the scope for requesting national scientific-technical advice to the famhp.

## **2. Legal basis:**

Article 6sexies of the Belgian Medicines law of March 25<sup>th</sup> 1964 forms the legal basis of the conditions and rules under which the famhp can provide scientific or technical advice regarding the research and development of medicines in view of future CTA's, MAA's or introduction of variations to marketed drug products. In addition, Article 4 of the Law of July 20<sup>th</sup> 2006 regarding the instauration and functioning of the famhp clearly states that scientific advice to applicants falls within the competence of the agency.

Moreover, Article 13bis of the Belgian Medicines law of March 25<sup>th</sup> 1964 together with the Royal Decree of March 31<sup>th</sup> 2009 forms the legal basis for the fees charged by the famhp to applicants submitting a request for national STA.

The Royal Decree of March 31<sup>th</sup> 2009, in execution of article 6sexies of the Belgian medicines law of March 25<sup>th</sup> 1964, lays down the legal scope, procedures, timelines and fees that are applicable to requests for national STA that are submitted to thefamhp.

The Royal Decree of July 16<sup>th</sup> 2012, modifying the Royal Decree of March 31<sup>th</sup> 2009, defines the new definitions, procedures and fees that are legally applicable to requests for national STA that are submitted to the famhp as of October 18<sup>th</sup> 2012. Cfr. Paragraph 3 and 6 below.

## **3. Definitions - Scope:**

As defined in the Royal Decree July 16<sup>th</sup> 2012 the following types of advice fall within the legal scope of a national STA submitted to the famhp:

### **3.1. STA request Type I:**

An **STA request Type I** is defined as: Advice requests representing **one specific question** regarding:

- (1) scientific issues related to research and development
- (2) technical-regulatory issues for which no (national or European) legislation or guidelines exist or the current legislation or guidelines are insufficient

and for which **no multidisciplinary expertise** would normally be required to address the raised question.

Due to its general simple nature, this type of advice request would normally require only expertise in one specific field and would therefore be addressed in writing. However, on exceptional basis, an individual question may also represent a complex matter that would require in depth expertise. In such case the advice request might be considered by the famhp as a type II or type III STA request (i.e. depending on the complexity of the STA request) since it represents a heavy workload. In such case, the STA request will follow the procedure for type II, III STA requests (cfr. paragraph 6.3).

### 3.2. STA request Type II and III:

A national STA request type II and III can be defined as: Advice requests representing a set of multiple, specific questions regarding:

- (1) scientific issues related to research and development
- (2) technical-regulatory issues for which no (national or European) legislation or guidelines exists or the current legislation or guidelines are insufficient

and for which multidisciplinary expertise is normally required to address the raised question(s).

In general, type II and type III STA requests would typically cover multiple type of questions (eg. related to quality, non-clinical, clinical issues, regulatory, ...) for which multidisciplinary expertise is required.

A detailed list of the different types of STA requests that can be submitted to the famhp is included in table 1 below.

<b>TYPES OF NATIONAL SCIENTIFIC-TECHNICAL ADVICE (STA)</b>	<b>FEE (euro)</b>
<b>TYPE I – Technical-regulatory advice (max. 1 question possible) – written advice (max. 30 calendar days)</b>	
Technical/regulatory advice concerning eg. chemical-pharmaceutical or (pre)-clinical aspects, the statute of a medicinal product, IMP – NIMP statute, namegiving (if umbrella brands), GMP aspects	2.217,91
<b>Technical advice (eg. concerning GMP aspects)</b>	<b>2.217,91</b>
<b>TYPE II – Scientific advice (multiple questions for one domain of expertise possible) – through a scientific advice meeting or in writing upon request (max. 70 calendar days)</b>	
Expertise domain 1: advice concerning chemical-pharmaceutical aspects of a drug product	13.307,48
Expertise domain 2: advice concerning the non-clinical OR clinical aspects of a drug product (i.e. including protocol design, Risk Management plans (RMP's), Benefit/Risk, CTA methodology)	13.307,48
Expertise domain 3: protocol assistance	13.307,48
<b>TYPE III – Mixed advice (i.e. Scientific and technical-regulatory advice, multiple questions possible) – through a scientific advice meeting (max. 70 calendar days)</b>	
<b>TYPE IIIa:</b>	
Mixed advice concerning both technical /regulatory questions (type I) and scientific questions (type II)	17.743,32
Scientific advice concerning multiple expertise domains mentioned above under STA type II: eg. expertise domain 1 (chemical – pharmaceutical aspects) and expertise domain 2 (clinical, non-clinical aspects)	17.743,32
advice concerning early market access aspects of a drug product	17.743,32
<b>TYPE IIIb – Joint STA's:</b>	
Joint advice with other National Competent Authorities (NCA's) within the EU	17.743,32
Joint advice with other HTA bodies within the EU (i.e. joint STA-HTA): advice concerning early market access and/or early reimbursement aspects in combination with STA aspects	17.743,32

Table 1: Types of national scientific-technical advice and corresponding fees

Further detailed information on the maximum legal delays that are applicable for providing the formal famhp advice to the applicant after validation of the STA request, can be found in paragraph 6 in below.

## **Out of scope:**

Depending on the nature and purpose, several type of advice requests / questions fall outside the legal scope of the national STA procedure as these advice request/questions are being addressed by the famhp in a different way, according to different procedures, timelines, etc... Addressing such advice requests/questions normally falls within the informative tasks of the famhp for which no additional fees are being charged.

The following type of advice request/questions are considered out of scope (i.e. non-exhaustive list):

**1. Advice requests related to current legislation:** Advice requests representing one or multiple specific questions regarding technical-regulatory topics for which divergences exist in the current legislation.

**2. Frequently asked questions (FAQ's):**

Frequently asked questions regarding eg. Submission procedures (eg. CTA's, MAA's, MRP's), general dossier requirements, etc., as described in current national and European guidelines, legislation, etc.

**3. Dossier-related questions:**

Questions from applicants which are sent to the famhp in the context of an application submitted in Belgium (eg. a CTA) fall within the procedure for that particular application and are normally addressed by the involved file manager(s). Subsequently, dossier-related questions fall outside the scope of the national STA procedure.

**4. Legal advice:**

Requests for legal advice should be sent directly to the juridical department of the famhp and fall outside the scope of national STA requests.

**5. Presubmission meeting requests related to marketing authorization (MA):**

At current, requests for a presubmission meeting related to new MA applications, line-extensions or variations are only being accepted in case Belgium acts as reference member state (RMS) in the MRP/DCP or when Belgian acts as rapporteur / co-rapporteur in the centralized procedure (CP). Such meetings fall outside the scope of the national STA procedures as they are being addressed in a separate procedure. However, in case that Belgium acts as concerned member state (CMS) in

the MRP/DCP, the famhp offers applicants the opportunity to use the national STA procedures to raise scientific-technical (eg. regulatory) questions in preparation of the MRP/DCP (i.e. as long as the questions fall within the legal scope of a national STA request).

#### **6. Portfolio meetings:**

Applicants are also allowed to request for a **Portfolio Meeting** with the agency. This type of informal and informative meeting typically provides the opportunity for pharmaceutical companies to present an overview of one or several development programs, therapeutics areas in which the company is actively involved, different products (or classes thereof) which are in the development pipeline, etc... In addition, the portfolio meetings allow clarifying the agency's interest for a role in a forthcoming European procedure for specific drug products. This information forms an important basis for the long-term recourse planning of the agency. This type of meeting is considered to fall outside the scope of a national STA request and therefore no fee will be charged to the applicant for a portfolio meeting.

However, in case that the applicant would seek advice on specific scientific and/or technical-regulatory aspects (eg. related to an individual development program), such advice is considered to fall within the scope of a national STA request as this would rather be discussed in a specific advice meeting. Thus, the need for a subsequent specific STA meeting may be the result of a Portfolio Meeting; this advice needs to be applied for separately.

#### **7. Regulatory planning meetings:**

eg. meeting requests from applicants who plan to submit multiple applications (MAA's, variations, etc.) at once.

#### **8. Questions / advice requests concerning healthcare products:**

eg. Questions / advice requests concerning:

- Medical devices
- blood, cells and tissues
- borderline products (eg. food supplements, cosmetics, biocides, etc.)
- .....

#### **9. Questions regarding other (i.e. non-R&D related) aspects of drug products and medical devices used in combination with a medicinal product and falling under the legal statute of a medicinal product:**

eg. Questions concerning the proper use of medicines, publicity issues, counterfeit, etc.

## 4.Areas of advice:

In general, a national STA request submitted to the famhp may cover multiple questions related to a broad range of areas in the field of research and development products. STA requests can also be submitted for medical devices that are used in combination with a medicinal product and in case such combination product falls under the legal statute of a drug product .

A non-exhaustive list of examples is given below:

- Quality aspects
- Non-clinical aspects
- Clinical aspects
- Unmet medical needs
- Early market access
- Switch from Rx – OTC status
- Technical-Regulatory issues (eg. regulatory statute of borderline products)
- Regulatory strategy (eg. for CTA, SAWP advice, PIP, MAA, CHMP re-examination, WHO pre-qualification)
- GXP-related aspects (eg. GMP, GCP, GLP, ...)  
Design and conduct of clinical trials / clinical development programs (eg. integrated protocol designs)
- Pharmacovigilance aspects (eg. PASS/PAES, referrals)
- Risk-management aspects (eg. RMP's, ERA's)
- Benefit / risk aspects
- GMO and Biosafety related aspects of IMP's
- Combination packages
- Drug-device combination products
- New (platform)technologies & concepts (eg. on-site formulation)
- ...

National STA requests may be (1) related to the research & development of specific drug products (i.e. drug products falling within the legal definition of a medicinal product) or a class of drug products or (2) may cover more general, non-product related aspects (eg. regarding genotoxic impurity testing).

National STA requests which are related to a specific drug product or class of drug products may cover different types of drug products: chemical, radiochemical, bio(techno)logical, GMO's, paediatric medicines, geriatric medicines, ATMP's, nanomedicines, etc.

## 5.Timing of STA requests - Type of meetings:

### 5.1Introduction:

Scientific-technical advice can be requested during any stage of the initial development of the medicinal product (eg. before submission of a CTA, MAA, variation, line-extension).

Meetings can also be held with the famhp to discuss pharmacovigilance issues, proposals for changes to labelling or package leaflets, etc.

In conclusion, the following STA requests can be submitted to the famhp throughout the complete product lifecycle:

≤ Phase 1 ( FIM, exploratory CTA's )

≤ Phase 2

≤ Phase 3

**Phase 4: post marketing authorisation (MA)**

## **5.2 Initial / follow-up STA requests:**

As mentioned before, after an **initial STA request** has been addressed by the famhp, applicants are free to submit at any time a **follow-up STA request** (eg. whenever there would be a need to seek advice on new development data that have been obtained, when changes in scientific / regulatory guidelines have taken place, etc).

Independently, of the procedure that was applied during the initial STA request, any follow-up STA request may be treated by the famhp either as a Type I, II or III national advice depending on the scope of the follow-up STA request. Subsequently, the same fees apply for a follow-up STA request as for an **initial STA request** submitted to the famhp.

## **5.3 Clarification requests:**

On some occasions, it is conceivable that the applicant requests for a clarification of the formal advice issued by the famhp in the context of an initial or follow-up STA request. The requests for further clarifications should be submitted within 14 calendar days after the receipt of the Final Scientific Advice (see paragraph 6.3 Procedures and timelines). In such cases, no additional fee will be charged for the clarification given by the famhp. The clarification of an advice may be addressed by the famhp either in writing (i.e. standard approach) or on exceptional basis through a meeting with the applicant. However, such clarification is strictly limited to the questions and issues that were addressed during the formal STA procedure (i.e. either in a face-to-face or teleconference meeting or in writing). Any new questions that have arisen from the initially issued famhp advice should be dealt with in a follow-up advice request.

## **6. Procedure and timelines:**

### **6.1 Introduction:**

In order to allow the famhp to provide a pointed advice within the foreseen time limits, applicants who intend to seek scientific and / or technical (regulatory) advice from the famhp, are requested to strictly follow the procedures described below.

Any official request for national STA falling within the legal scope of a Type I, II or III STA request as mentioned above should be submitted to the Directorate-General PRE-authorisation of the famhp.

These requests should be sent **electronically** to our central mailbox: **sta@fagg-afmps.be**. Alternatively, (eg. in case large electronic files above 5 MB would be submitted), electronic STA requests can also be sent by using the EudraLink system or on CD-rom or USB key to the following address:

**Federal Agency for Medicines and Health Products**

**Directorate-General PRE Marketing Authorisation (desk 8D222)**

**Victor Hortaplein 40/40**

**B-1060 Brussels**

**BELGIUM**

**To the attention of: Ms. Greet Musch**

### **6.2 Payment information:**

Each submitted STA request must be accompanied by payment of the appropriate fee (cfr. Paragraph 9 below). Each type I STA request requires payment of €2.217,91. For type II STA requests a fee of € 13.307,48 is being charged. Each type III STA request requires a payment of € 17.743,32 The fees are applicable to both initial or follow -up STA requests. Payment must be made on the following **bank account number** of the famhp: BE28 6790 0219 4220

### **Contact details of the bank:**

Poste financière

Chaussée d'Anvers 59

B-1100, Bruxelles (Belgium)

IBAN : BE28 6790 0219 4220

BIC : PCHQBEBB

Applicants should clearly mention on the bank statement the **type of STA request** that is being submitted to the famhp and if the STA request concerns an **initial or follow-up STA request**; followed by the **name of the applicant**. The **topic of the STA request** (eg. product name) should also be clearly mentioned.

For **payments from abroad** the transfer fees should be paid by the payer. Applicants are also requested to include a proof of payment (eg. a bank statement) to the STA request since this is verified during the validation of the STA request.

For each initial or follow-up STA requests a separate payment should be made.

In case that an **inappropriate fee** was paid by the applicant which does not correspond to the fees defined in paragraph 9 below (eg. when submitting the STA

request under the wrong procedure), the applicant will be contacted in order to make the appropriate payment.

In case the applicant should ask for a withdrawal of his STA request after payment of the fee (as part of the formal submission to the famhp), the fee will not be refunded by the famhp.

When a formally submitted STA request is declared invalid by the famhp at the end of the validation phase of the procedure (cfr. Paragraph 6.3 below), the fee will be refunded to the applicant.

### 6.3 Procedures and timelines:

In general, national STA requests submitted under the type I, II or III advice procedure will follow during the respective procedure the three steps outlined below:

#### Step 1: Validation phase

During validation of each STA request, the famhp will verify if the following criteria are met:

- the submitted STA request falls within the legal scope of a national STA request
- the submitted STA request has been correctly submitted by the applicant under the appropriate type of STA request
- the appropriate fee has been paid
- all supportive documentation is included in the submitted STA request (cfr. Paragraph 7 below)
- a clear and solid motivation for seeking national scientific-technical advice is provided by the applicant

The validation phase only begins when the Directorate-General PRE-authorisation of the famhp has received the formal STA request (including all supportive documents) and the proof of payment from the bank.

Based on the received supportive documents and depending on the complexity and nature of the raised questions, the famhp will verify if the submitted STA request fulfils the definition of a type I, II or III STA request and subsequently if a meeting at the agency would be necessary (i.e. for type II and type III) or if a written advice will be given (i.e. type I STA advice). Nevertheless, it is the applicant's responsibility to submit the STA request under the appropriate procedure.

The applicant will be informed about the validation of the STA request. Any refusal to validate the STA request will be communicated to the applicant as soon as possible. Grounds for refusing validation of the STA request may be: eg. insufficient / missing motivation of the STA request, insufficient / missing supportive documentation, inappropriate payment of the fee, the requested STA may fall outside the legal scope of national STA, etc.

Shortcomings which cannot be resolved by the applicant will automatically lead to an INVALID application. In contrast, when all shortcomings can be resolved by the applicant during validation phase, the STA request will be declared VALID (= Day 0). As soon as all validation criteria are being met, the valid STA request will enter the evaluation phase of the STA procedure (cfr. step 2).

At current, no legal timelines are applicable for the validation of STA requests as the validation of each STA dossier depends on the completeness and acceptability of the submitted data. Evidently, the famhp tries to validate all incoming STA requests as quickly as practically possible (i.e. normally the validation of each STA request is initiated within 1 week upon receipt of the formal STA request).

### **Step2 : Evaluation phase**

During the evaluation phase of a valid STA request, the appropriate internal and/or external experts of the famhp are being selected and designated by the STA coordinator. Subsequently, the content of the STA request is being evaluated by the designated experts in view of addressing the questions raised by the applicant. The questions contained in the STA request will be formally addressed by the famhp as follows:

#### **Type I STA requests:**

In principle, a valid STA request falling within the definition of type I STA request will be addressed by the famhp in writing, within maximum 30 calendar days<sup>(\*)</sup>. Along with the validated Final Scientific Advice, a qualitative feedback questionnaire will be sent to the applicant (cfr. Paragraph 8 below) which can be completed by the applicant on a voluntary basis.

As mentioned before (cfr. Paragraph 3), this type of advice request will normally be addressed in writing due to its simple nature and the minimal workload associated with such STA request. However, on exceptional basis, a national STA request submitted by the applicant as an type I STA request might represent a very complex question that would require in depth expertise from the famhp. Taken into account the complexity and heavy workload associated with such advice request, such dossier could be considered by the famhp as a type II or type III STA request rather than an type I STA request. In such particular case, the STA request will follow the procedure of a type II or III STA request based on the famhp's decision during the validation phase.

#### **Type II and III STA requests:**

In principle, a valid STA request falling within the definition of a type II or type III STA request will be addressed by the famhp in a face-to-face meeting or through a teleconference meeting with the applicant. Depending on the nature and complexity of the STA request, the applicant is free to propose which type of meeting he prefers with the famhp. However, the famhp generally prefers a face-to-face meeting for discussing these types of STA requests.

Type II and type III STA requests will be addressed by the famhp in a face-to-face meeting or teleconference meeting within maximum 70 calendar days<sup>(\*)</sup>.

As a general rule, the famhp foresees 1.5 hours (max. 2 hours) for advice meetings with applicants. The applicant is strongly encouraged to provide a brief presentation during the meeting with the famhp, however this should be limited (eg. 15 min). Such presentation may cover eg. an overview of the issues to be discussed, background information on the development of the drug product which is relevant to the meeting, etc. The presentation should be electronically sent to the STA unit 14 calendar days prior to the meeting together with the attendant's list.

The meetings are chaired by the famhp and the discussions are normally being held in English. The applicant is free to bring in experts for attending the meeting as long as this is properly being communicated well in advance (i.e. through the attendants list). Nevertheless, it is the responsibility of the famhp to select and designate the internal and/or external experts that will provide the formal advice on behalf of the famhp.

After the STA meeting has taken place, the applicant should send an electronic copy of the company meeting minutes to the famhp within maximum 5 working days following the STA meeting. The meeting minutes should reflect the topics that were orally discussed / clarified during the STA meeting and should complement the supportive documentation provided by the applicant in the briefing package of the STA request prior to the meeting. The company meeting minutes will not be formally reviewed or endorsed by the famhp but will be used as additional supportive information by the involved famhp experts in order to draft the Final Scientific Advice. The validated Final Scientific Advice will be sent as a pdf file to the applicant within maximum 21 calendar days following the formal STA meeting. Along with the validated Final Scientific Advice, a qualitative feedback questionnaire will be sent to the applicant (cfr. paragraph 8 below) which can be completed by the applicant on a voluntary basis.

(\*) Date counting from the day at which the STA request was formally declared VALID by the famhp (= Day 0). These timelines represent the maximum period during which the famhp commits itself to provide a pointed advice to the applicant. Evidently, the famhp aims to address all advice requests from applicants as fast as practically possible after validation.

### **Step 3: Administrative phase**

If no further clarification request from the applicant is received by the famhp within 14 calendar days after sending out ingthe validated Final Scientific Advice , the procedure is officially being closed.

## **7. Content & format of STA requests:**

### **7.1 Introduction:**

Detailed information on the content and format of national STA requests submitted to the famhp can be found on the [website](#).

### **7.2 Application form:**

As part of the required content of a national STA request, applicants are being requested to download and complete the electronic application form (i.e. MS Access format) which is available on the famhp [website](#) and to include the completed electronic document in the STA request.

The electronic application form is intended to contain all essential information related to the national STA request that would provide the famhp with complete and concise data in order to provide a pointed advice.

## **8. Feedback questionnaire for applicants**

As a federal agency, the famhp pays much importance to the quality of the provided national STA's and to the satisfaction of the applicants. In order to improve its national STA service on a continuous basis, applicants are offered the opportunity to fill in a feedback questionnaire after having received the final scientific advice from the famhp. The qualitative feedback questionnaire is intended to ask the applicant's opinion on the following three aspects related to the received national STA:

- Quality of the provided national STA
- Quality of the service provided during the STA procedure
- Consistency of the provided national STA with previous advice(s)

Applicants are being asked to complete the feedback questionnaire on a voluntary basis and send it back afterwards to the famhp. An electronic copy of the document can be found on the famhp [website](#) along with the instructions for filling in and sending back the document to the famhp.

## **9.Fees:**

At current, the following fees<sup>(\*)</sup> apply to requests for national STA submitted to the famhp:

<b>TYPE I: Technical-regulatory advice</b>	<b>2.217,91 euro</b>
<b>TYPE II: Scientific advice</b>	<b>13.307,48 euro</b>
<b>TYPE III: Mixed advice (i.e. scientific and technical-regulatory advice)</b>	<b>17.743,32 euro</b>

The same fees for national STA request will apply independently of the type of applicant by whom the STA request is submitted to the famhp: i.e. sponsors of clinical trials, pharmaceutical/biotech companies (eg. small biotech spin offs, SME's, global companies), academic research centres, etc.

The same fees for national STA request will apply regardless of the area of advice (eg. paediatrics, oncology, etc..) or timing of seeking advice at national level (eg. initial versus follow-up advice).

Detailed instructions on the payment of fees for national STA requests can be found in paragraph 6.2 above.

(\*) Fees for national STA are subject to indexation on a yearly basis. Indexation of the fees takes place at the beginning of each calendar year.

## **10.Contact for further information:**

Any general or specific questions that applicants may have related to the national scientific and/or technical (regulatory) advice service of the famhp can be sent to the following mailbox: [sta@fagg-afmps.be](mailto:sta@fagg-afmps.be)

## **11.Legal texts:**

### **11.1 Laws:**

- Medicines law of March 25<sup>th</sup> 1964:
- Law of July 20<sup>th</sup> 2006 regarding the instauration and functioning of the famhp:

### **11.2 Royal Decrees:**

- Royal decree of March 31<sup>th</sup> 2009 in execution of article 6sexies of the Medicines law of March 25<sup>th</sup> 1964
- Royal Decree of July 16<sup>th</sup> 2012, modifying the Royal Decree of March 31<sup>th</sup> 2009.

## **12.Frequently asked questions (FAQ's):**

A list of frequently asked questions can be found on the [website](#) of the famhp under the section “Scientific-technical Advice”.

## **13.Abbreviations:**

<b>ATMP</b>	<b>advanced therapy medicinal product</b>
<b>CMS</b>	<b>Concerned member state</b>
<b>CP</b>	<b>Centralized Procedure</b>
<b>CTA</b>	<b>Clinical trial application</b>
<b>DCP</b>	<b>Decentralized procedure</b>
<b>EMA</b>	<b>European Medicines Agency</b>
<b>ERA</b>	<b>Environmental Risk Assessment</b>
<b>famhp</b>	<b>federal agency for medicines and health products</b>
<b>FIM</b>	<b>first in man</b>
<b>GMO</b>	<b>Genetically Modified Organism</b>
<b>GMP</b>	<b>Good Manufacturing Practices</b>
<b>GCP</b>	<b>Good Clinical Practices</b>
<b>GLP</b>	<b>Good Laboratory Practices</b>
<b>HTA</b>	<b>Health Technology Assessment</b>
<b>IMP</b>	<b>Investigational Medicinal product</b>
<b>MAA</b>	<b>Marketing authorization application</b>
<b>MRP</b>	<b>Mutual recognition procedure</b>
<b>NCA</b>	<b>National Competent Authority</b>
<b>OTC</b>	<b>Over-the-counter</b>
<b>PAES</b>	<b>Post-Authorisation Efficacy Studies</b>
<b>PASS</b>	<b>Post-Authorisation Safety Studies</b>
<b>PIP</b>	<b>Paediatric investigation plan</b>
<b>RMP</b>	<b>Risk Management Plan</b>
<b>RMS</b>	<b>Reference member state</b>
<b>SAWP</b>	<b>Scientific advice working party</b>
<b>SAWP-V</b>	<b>Scientific advice working party – veterinary products</b>
<b>STA</b>	<b>Scientific-Technical Advice</b>