



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

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Information Technology

Electronic Application Form Data Exchange Standard 3.0

Supplementary Specification Annex 4 "Initial Veterinary Application Form"
v1.21.0.0



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1. Introduction

1.1. How to read this document

In association with this document the **maa_vet_schema.xsd** file contains the description of the structure of all the concepts used for this annex. This will enable you to construct a data extraction/generation script to populate the relevant information to/from your systems.

The "Chapters" refer to the paragraph number of the paper application form. The "Sections" refer to the paragraph numbering of this document.

Some diagrams are too large to describe the whole hierarchy on only one page. Therefore, the diagrams are split in sub sections that might not be in line with the paper document chapters.

In order to find your way back in this document when starting from paper document refer to the chapters' labels and numbering.

The information provided in this document focuses only on the initial veterinary application form information and how it is mapped with the DES 3.0 standard.

Description and definition of the DES 3.0 Concepts used in this document can be found in the DES 3.0 Technical Guide document.

1.2. Sections Components

Each section is split in three components that show different aspects of the DES 3.0 standard applied to the application form.

1.2.1. The Elements Mapping Table

This table describes the mapping between the paper form fields of a specific chapter and the elements of the DES 3.0 Model.

The table consists in 4 columns:

- **Element Id:** The id of the field used in business rules. <paragraph>-<numeric order>
Ex: 264-1
- **Label:** The label of the field in the application form is sometimes preceded by a chapter numbering.
- **DES 3.0 Mapping:** It is the corresponding mapped-to element in the DES 3.0 model. It contains at least one mapped-to element. The mapping shows the hierarchy from the root element to the leaf element with the parent-child link represented by the "/" sign.
- **RDM Mapping:** It is the corresponding mapped-to attribute in the RDM Model. The mapping shows how to get the information in the RDM relational model through links between the technical concepts represented by the ">" sign.

The minimal notation is always "<technical concept parent>/" in the common context and "<mapped-to element>" in the DES 3.0 column.

The description of the technical concept parent is in the DES 2.0 supplementary specifications sections 7.1 and 7.2

If there is no mapping, the DES 1.0 element remains in the form specific part of the model.

- **Remarks:** Contains any relevant information concerning the element values, format or business rules.

Colours

Text: Tells that the elements are not part of the RDM 3.0 and can be found only in DES 3.0 with no similarity in terms of definition.

Text: Tells that there is no existing mapping between the DES 3.0 and RDM 3.0. The missing mapping can be of two kinds.

- "ignored" based on the decision of the RDM team not to map the element
- "not mapped": The RDM 3.0 may contain more or less elements because the RDM 3.0 draft came after the DES 3.0.

Text: Tells that the RDM element is an additional linked entity comparing to the DES 3.0 hierarchy.

Note: The EUTCT controlled terms used in the RDM 3 are **not** always published yet. That's why some of the DES lists only provide a "short-name" which does not directly corresponds to a CTL term id in RDM model.

1.2.2. The Business Rules Table

All the rules are gathered with their corresponding Element Tree Diagram (ETD) and are defined as follows:

Element: The name of the element mentioned in the (ETD)

Default Cardinality: Cardinality that applies by default. It corresponds to the cardinality of the concerned element in the ETD.

Rule: Description of the condition to be evaluated.

Effect: if the condition is evaluated to true then the effect is applied.

1.2.3. The Element Tree Diagram (ETD)

The data structure constraints are captured in a graphic approach to facilitate the reading and assessment by the business.

The model used refers to the one used by W3C (World Wide Web Consortium XML Specification DTD for its publication standard issued in 1998.

The model is called "Element Tree Diagram" (ETD)

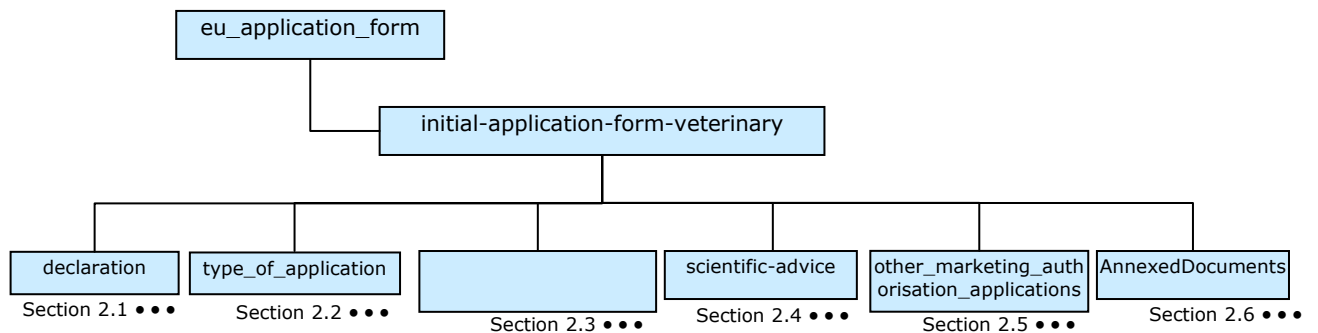
The diagrams of this version reflect the DES 3.0 standard described in the DES 3.0 Supplementary Specifications document. The ETD shows which are the concepts involved in the mapping of all the application form fields in the Element Mapping Table and the hierarchical constraints between them.

2. Initial Application form

The “initial-application-form-veterinary” is the highest level of the form specific model that represents the paper form. All sections are fully mapped to the Reference Data Model core concepts and common application form concepts.

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/ maa:initial-application-form-veterinary/			
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2.1	DECLARATION AND SIGNATURE	maa:declaration	Application	See Section 2.1
E2.2	TYPE OF APPLICATION	maa:type_of_application	MP Procedure	See section 2.2
E2.3	MARKETING AUTHORISATION APPLICATION PARTICULARS	maa:marketing_authorisation_application_particulars		See Section 2.3
E2.4	SCIENTIFIC ADVICE	maa:scientific-advice	App Scientific Advice	See Section 2.4
E2.5	OTHER MARKETING AUTHORISATION APPLICATIONS	maa:other_marketing_authorisation_applications		See Section 2.5
E2.6	5. ANNEXED DOCUMENTS (WHERE APPROPRIATE)	maa:AnnexedDocuments		See Section 2.6

Element Tree Diagram

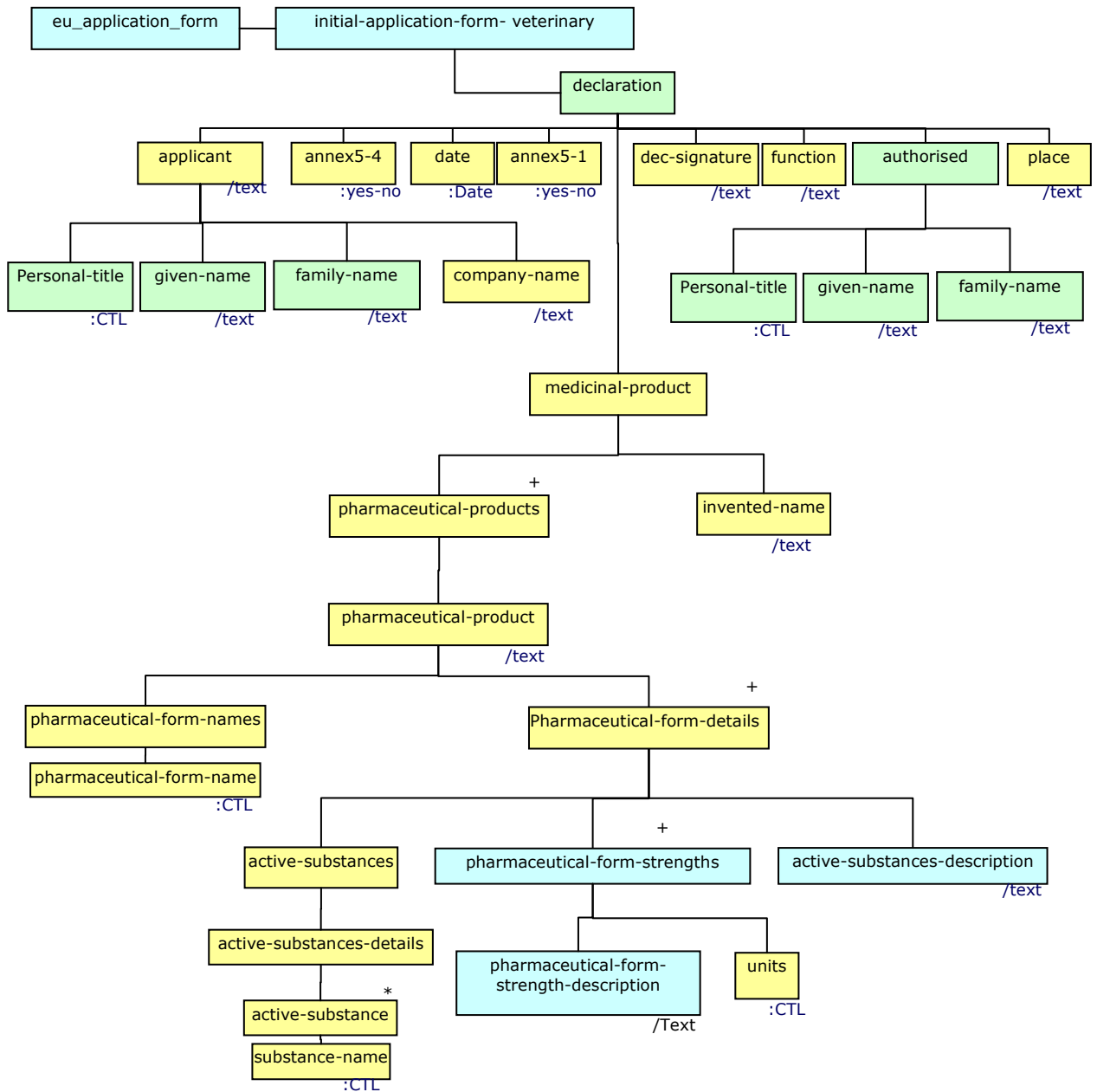


2.1. DECLARATION AND SIGNATURE

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa: initial-application-form-veterinary maa:declaration/	Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E21-1	Product (invented) name	rdm:medicinal-product/rdm:invented-name	Medicinal Product > Medicinal Product Group > invented name	
E21-2	Pharmaceutical form(s)	rdm:medicinal-product/rdm:pharmaceutical-products/ rdm:pharmaceutical-product/rdm:pharmaceutical-form-names/rdm:pharmaceutical-form-name	Pharmaceutical Dose Form CTL > term id	
E21-3	Strength(s)	rdm:medicinal-product/rdm:pharmaceutical-products/rdm:pharmaceutical-product/rdm:pharmaceutical-form-details/rdm pharmaceutical-form-strengths/rdm:pharmaceutical-form-strength-description		
E21-3a	Units	rdm:medicinal-product/rdm:pharmaceutical-products/rdm:pharmaceutical-product/rdm:pharmaceutical-form-details/rdm pharmaceutical-form-strengths/rdm:units		
E21-4	Active Substance(s)	rdm:medicinal-product/rdm:pharmaceutical-products/rdm:pharmaceutical-product/rdm:pharmaceutical-form-details/rdm:active-substances-description		
E21-5	Active Substance	rdm:medicinal-product/rdm:pharmaceutical-products/rdm:pharmaceutical-product/rdm:pharmaceutical-form-details/rdm:active-substances/rdm: active-substances-details/rdm:active-substance/rdm:substance-name	Ingredient > Substance CTL, Ingredient > IngredientRole CTL	
E21-6	Applicant details:			
E21-7	Name	maa:applicant/company-name	Role > Party > Organisation > Name	
E21-8	Person authorized for communication*, on behalf of the Applicant:			
E21-9	Title	maa:authorised/rdm:personal-title	Role > Party > Person > Personal title	
E21-10	First name	maa:authorised/rdm:given-name	Role > Party > Person > given name	
E21-11	Surname	maa:authorised/rdm:family-name	Role > Party > Person > family name	
E21-12	Signatory	maa:dec-signature	Not mapped	
E21-13	Title	maa:signature/rdm:personal-title	Role > Party > Person > Personal title	
E21-14	First name	maa:signature/rdm:given-name	Role > Party > Person > given name	
E21-15	Surname	maa:signature/rdm: family-name	Role > Party > Person > family name	
E21-16	Function	maa:function	Manufacturer MP> functions performed	
E21-17	Address	maa:signature/rdm:address	signature address	
E21-18	Date	maa:date	signature date	
E21-19	<i>Note: please attach letter of authorisation for communication/signing on behalf of the applicant in annex 5.4</i>	maa:annex5-4		

E21-20	Note: if fees have been paid, attach proof of payment in Annex5.1 - see information on fee payments in the Notice to Applicants, Volume2A, chapter 7.	maa:annex5-1		
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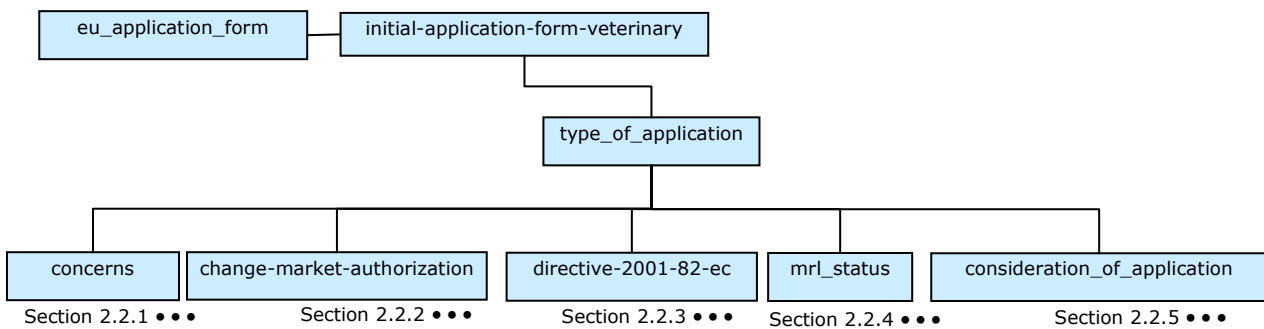
Element Tree Diagram



2.2. TYPE OF APPLICATION

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/	Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E22-1	THIS APPLICATION CONCERNS	maa:concerns		See section 2.2.1
E22-2	Is this an application for a change to your existing marketing authorisation leading to an extension as referred to in Annex II of Regulations (EC) no. 1084/2003 or 1085/2003, or any national legislation, where applicable?	maa:change-market-authorization		See section 2.2.2
E22-3	This application is submitted in accordance with the following article in Directive 001/82/EC or Regulation (EC) No 726/2004	maa:directive-2001-82-ec		See section 2.2.3
E22-4	MRL status (only for food producing species)	maa:mrl_status		See section 2.2.4
E22-5	Consideration of this application is also requested under the following article in Directive 2001/82/EC or Regulation (EC) no. 726/2004	maa:consideration_of_application		See section 2.2.5

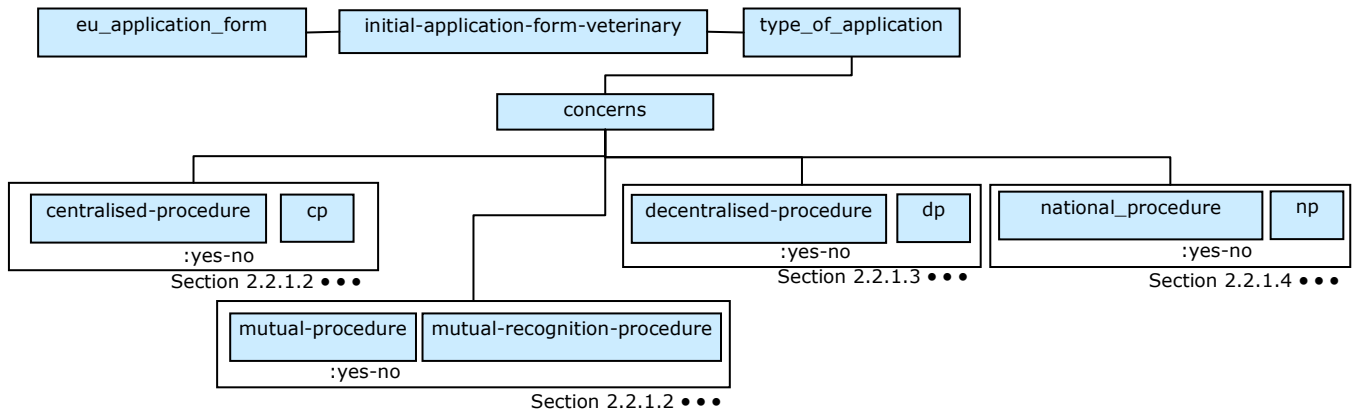
Element Tree Diagram



2.2.1. THIS APPLICATION CONCERNS

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:concerns/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E221-1	A CENTRALISED PROCEDURE (according to Regulation (EC) No 726/2004)	maa:centralised-procedure maa:cp		B221-1
E221-2	A MUTUAL RECOGNITION PROCEDURE (according to Article 32(2) of Directives 2001/82/EC)	maa:mutual-procedure maa:mutual-recognition-procedure		B221-1
E221-3	A DECENTRALISED PROCEDURE (according to Article 32(3) of Directive 2001/82/EC)	maa:decentralised-procedure maa:dp		B221-1
E221-4	1.1.4 A NATIONAL PROCEDURE	maa:national_procedure maa:np		B221-1, B25-1

Element Tree Diagram

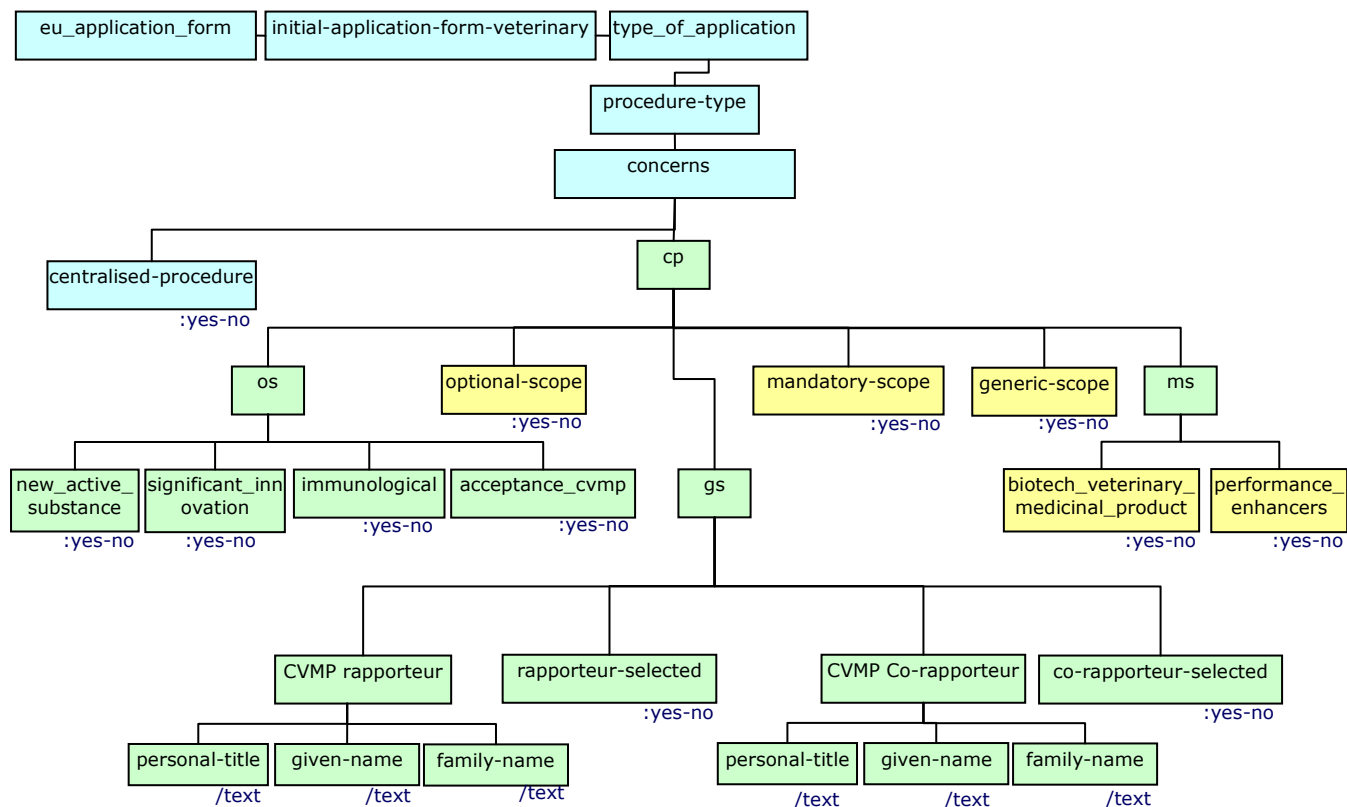


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B221-1	E221-1 to E221-4	Mandatory.	Mutually Exclusive.	

2.2.1.1. A CENTRALISED PROCEDURE (according to Regulation (EC) No 726/2004)

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:concerns/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2211-1	A CENTRALISED PROCEDURE (according to Regulation (EC) No 726/2004)	maa:centralised-procedure	Procedure Type CTL (Value="centralised")	
E2211-2	"Mandatory scope" (Article 3(1))	maa:cp/maa:mandatory-scope	Basis for Eligibility CTL	B2211-1
E2211-3	Annex (1) (Biotech veterinary medicinal product)	maa:cp/maa:ms/maa:biotech_veterinary_medicinal_product	Basis for Eligibility CTL	B2211-2
E2211-4	Annex (2) (performance enhancers)	maa:cp/ maa:ms/ maa:performance_enhancers	Basis for Eligibility CTL	B2211-2
E2211-5	"Optional scope" (Article 3(2))	maa:cp/maa:optional-scope	Basis for Eligibility CTL	B2211-1
E2211-6	Article 3(2)(a) (new active substance)	maa:cp/maa:os/ maa:new_active_substance	Basis for Eligibility CTL	B2211-3
E2211-7	Article 3(2)(b) (significant innovation or interest of animal health at Community level)	maa:cp/maa:os/ maa:significant_innovation	Basis for Eligibility CTL	B2211-3
E2211-8	Article 3 - Immunological veterinary medicinal products for the treatment of animal diseases subject to Community prophylactic measures	maa:cp/maa:os/ maa:immunological	Basis for Eligibility CTL	B2211-3, B2211-6
E2211-9	Date of acceptance/confirmation by CVMP	maa:cp/maa:os/ maa:acceptance_cvmp	chmp acceptance date	B2211-6
E2211-10	"Generic of a centrally authorised veterinary medicinal product" (Article 3(3))	maa:cp/maa:generic-scope		B2211-1
E2211-11	CVMP Rapporteur	maa:cp/maa:gs/ maa:rapporteur-selected	Role > Party> Party Type CTL (Value="Person")	B2211-4
E2211-12	Title	maa:cp/maa:gs/maa:rapporteur/maa:personal-title	Role > Party > Person > Personal Title	B2211-4
E2211-13	First name	maa:cp/maa:gs/maa:rapporteur/maa:given-name	Role > Party > Person > Given Name	B2211-4
E2211-14	Surname	maa:cp/maa:gs/maa:rapporteur/maa:family-name	Role > Party > Person > Family Name	B2211-4
E2211-15	CVMP Co-rapporteur	maa:cp/maa:gs/maa:co-rapporteur-selected	Role > Party> Party Type CTL (Value="Person")	B2211-5
E2211-16	Title	maa:cp/maa:gs/maa:co-rapporteur/maa:personal-title	Role > Party > Person > Personal Title	B2211-5
E2211-17	First name	maa:cp/maa:gs/maa:co-rapporteur/maa:given-name	Role > Party > Person > Given Name	B2211-5
E2211-18	Surname	maa:cp/maa:gs/maa:co-rapporteur/maa:family-name	Role > Party > Person > Family Name	B2211-5

Element Tree Diagram

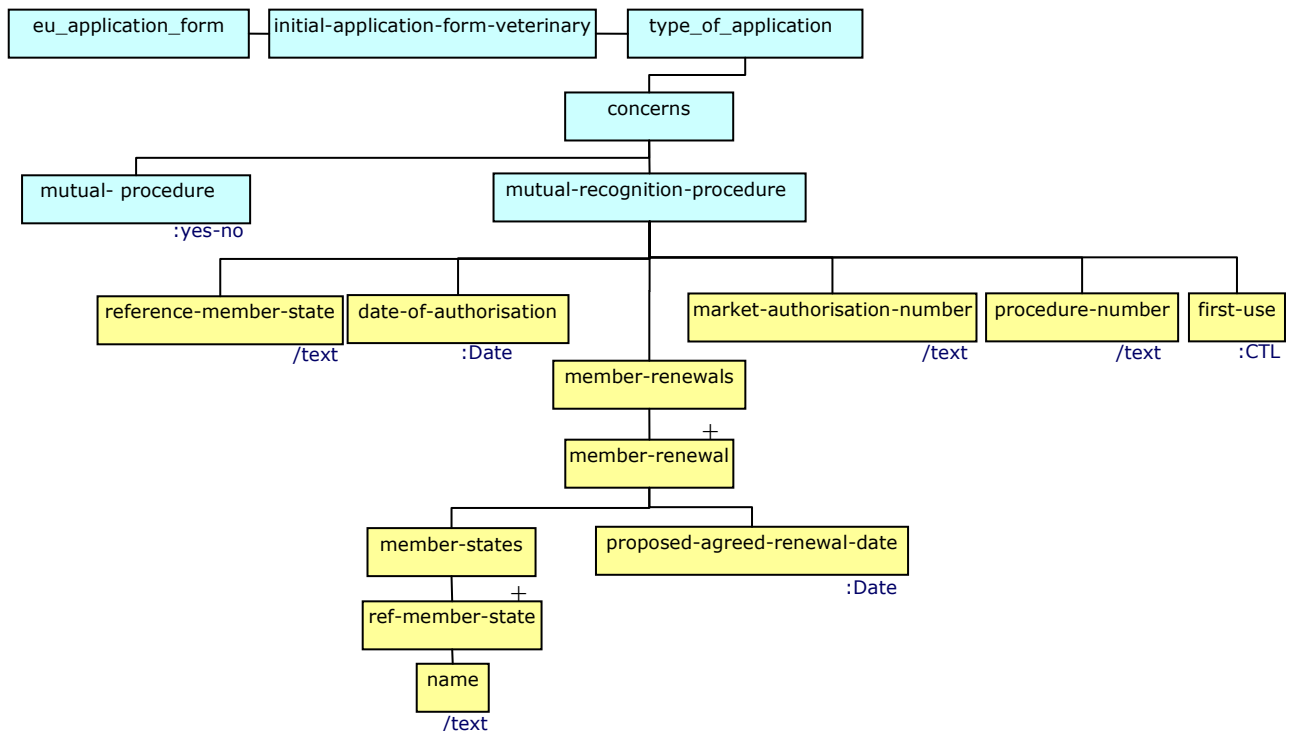


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2211-1	E2211-2, E2211-5, E2211-10	Mandatory.	Fields are mutually exclusive.	Can only have one of them active at a time.
B2211-2	E2211-3 to E2211-4	Mandatory if E2211-2 is selected.	Fields are mutually exclusive.	Can only have one of them active at a time.
B2211-3	E2211-6, E2211-7, E2211-8	Mandatory if E2211-5 is selected.	Fields are mutually exclusive.	Can only have one of them active at a time.
B2211-4	E2211-11 to E2211-14	E2211-11 is mandatory.	If E2211-11, then the rest are mandatory.	
B2211-5	E2211-15 to E2211-18	E2211-15 is mandatory, rest are optional.	If E2211-15, then the rest are mandatory.	
B2211-6	E2211-8, E2211-9	Optional	If E2211-8 is selected, then E2211-9 is available	

2.2.1.2. A MUTUAL RECOGNITION PROCEDURE

Common DES 3.0 Context			Common RDM Entry point	
maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:concerns/			Application > MP Procedure >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2212-1	1.1.2 A MUTUAL RECOGNITION PROCEDURE	maa:mutual-procedure	Procedure Type CTL (Value=" mutual-recognition ")	
E2212-2	Reference Member State	maa:mutual-recognition-procedure/rdm:reference-member-state	Role > Country CTL	
E2212-3	Date of authorisation	maa:mutual-recognition-procedure/rdm:date-of-authorisation	previous auth date	
E2212-4	Marketing authorisation number	maa:mutual-recognition-procedure/rdm:market-authorisation-number	previous auth number	
E2212-5	Procedure number:	maa:mutual-recognition-procedure/rdm:procedure-number	procedure number	
E2212-6	First use	maa:mutual-recognition-procedure/rdm:first-use (Value=1)	Procedure Use CTL	B2212-1
E2212-7	Repeat use (Please also complete section 4.2)	maa:mutual-recognition-procedure/rdm:first-use (Value=2)	Procedure Use CTL	B2212-1, B222-2
E2212-8	Wave	maa:mutual-recognition-procedure/rdm:member-renewals/rdm:member-renewal	Procedure Use > waive id	B2212-2
E2212-9	Concerned Member State (specify)	maa:mutual-recognition-procedure/rdm:member-renewals/rmd:member-renewal/rdm:member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2212-10	Proposed/Agreed common renewal date	maa:mutual-recognition-procedure/rdm:member-renewals/rmd:member-renewal/rdm:proposed-agreed-renewal-date	Procedure Use > Proposed / agreed common renewal date	

Element Tree Diagram

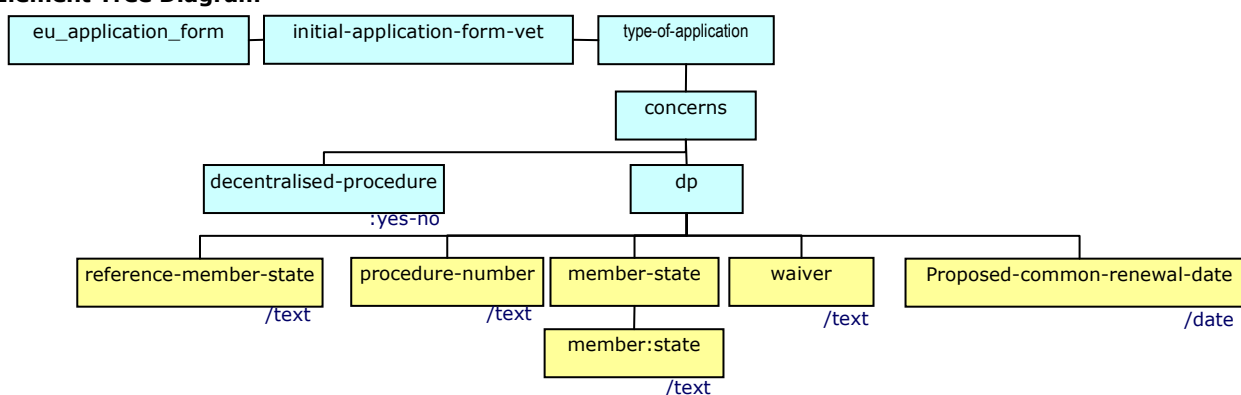


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2212-1	E2212-6 , E2212-7	Mandatory.	Mutually Exclusive.	
B2212-2	E2212-7 to E2212-10	If E2212-7 is selected, then E2212-8 to E2212-10 are mandatory.		

2.2.1.3. A DECENTRALISED PROCEDURE

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:concerns/		Application > MP Procedure >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2213-1	1.1.3 A DECENTRALISED PROCEDURE	maa:decentralised-procedure	Procedure Type CTL (Value="decentralised")	
E2213-2	(according to Article 28(3) of Directives 2001/82/EC)			
E2213-3	Reference Member State	maa:reference-member-state	Procedure Use > Role > Country CTL	
E2213-4	Procedure number:	maa:procedure-number	Procedure number	
E2213-5	Concerned Member State (specify)	maa:member-state/ maa:member-state	Procedure Use > Role > Country CTL	
E2213-6	If a waiver or amendment or PSUR-cycle is applied for, to harmonize with a substance birthdate, please specify:	maa:waiver	Procedure use > psur cycle waiver	
E2213-7	Proposed Common Renewal Date	maa:proposed-common-renewal-date	proposed-common-renewal-date	

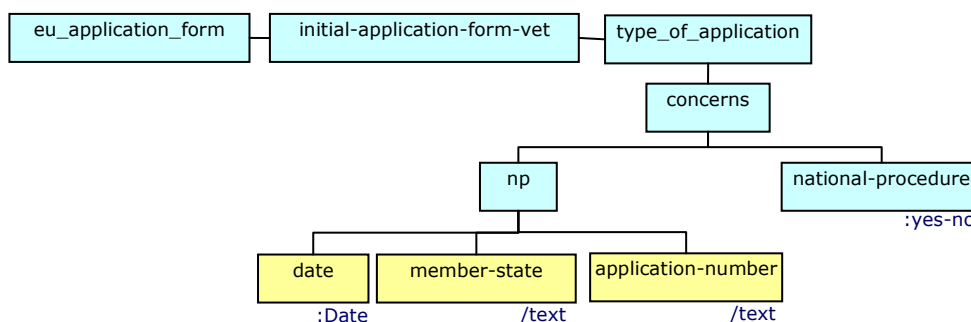
Element Tree Diagram



2.2.1.4. A NATIONAL PROCEDURE

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-human/maa:concerns/		Application > MP Procedure >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2214-1	1.1.4 A NATIONAL PROCEDURE	maa:national_procedure	Procedure Type CTL (Value="national")	
E2214-2	Member State	maa:member-state	Role > Country CTL	
E2214-3	Application number (if available)	maa:application_number	previous app number	
E2214-4	Date	maa:date		

Element Tree Diagram

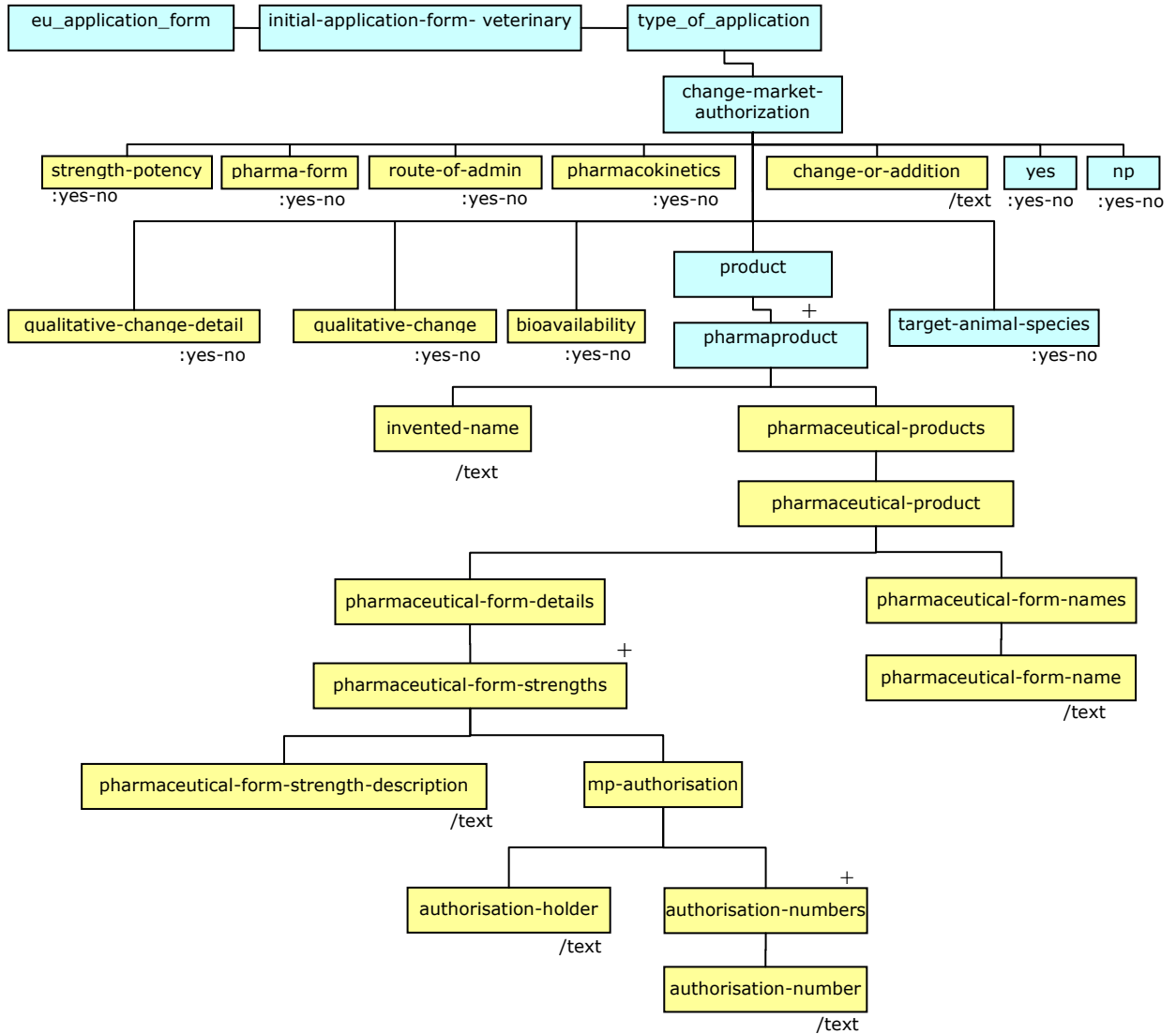


2.2.2. Is this an application for a change to your existing marketing authorisation leading to an extension as referred to in Annex I of Commission Regulation (EC) no. 1234/2008, or any national legislation, where applicable?

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:change-market-authorization		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remark
E222-1	Yes (complete sections below <i>and also complete 1.4</i>)	maa:yes(Value=1)	change in existing ma	B222-1, B222-2
E222-2	No (complete section 1.3 and 1.4.)	maa:no (Value=0)	change in existing ma	B222-1, B222-2
E222-3	Change of bioavailability	maa:bioavailability	Difference CTL	B222-2
E222-4	Change of pharmacokinetics	maa:pharmacokinetics	Difference CTL	B222-2
E222-5	Change or addition of a new strength/potency	maa:strength-potency	Difference CTL	B222-2
E222-6	Change or addition of a new pharmaceutical form	maa:pharma-form	Difference CTL	B222-2
E222-7	Change or addition of a new route of administration	maa:route-of-admin	Difference CTL	B222-2
E222-14	Change or addition of a food-producing target animal species	maa:target-animal-species	Difference CTL	B222-2
E222-8	Qualitative change in declared active substance <u>not defined as a new active substance</u>	maa:qualitative-change	Difference CTL	B222-2, B222-3
E222-9	Replacement by a different salt/ester, complex/derivative (same therapeutic moiety)	maa:qualitative-change- detail	Difference CTL	B222-3, B222-4
E222-10	Replacement by a different isomer, mixture of isomers, of a mixture by an isolated isomer	maa:qualitative-change- detail	Difference CTL	B222-3, B222-4
E222-11	Replacement of a biological substance or product of biotechnology	maa:qualitative-change- detail	Difference CTL	B222-3, B222-4
E222-12	modification of the vector used to produce the antigen or the source material, including a new master cell bank from a different source, where the clinical/safety characteristics are not significantly different;	maa:qualitative-change- detail	Difference CTL	B222-3, B222-4
E222-13	change to the extension solvent or the ratio of herbal drug to herbal drug preparation	maa:qualitative-change- detail	Difference CTL	B222-3, B222-4
E222-14	Product (invented) name	maa:product/ rdm:pharmaproduct/ rdm:invented-name	Reference Medicinal Product > Medicinal Product>Pharmaceutical Product>Ingredient>Subs	B222-2
E222-15	Pharmaceutical form(s)	maa:product/ rdm:pharmaproduct/ rdm:pharmaceutical-products/ rdm:pharmaceutical-product / rdm:pharmaceutical-form- names/rdm:pharmaceutical- form-name	Reference Medicinal Product > Medicinal Product > Pharmaceutical Dose Form CTL	B222-2
E222-16	Strength(s)	maa:product/ rdm:pharmaproduct/ rdm:pharmaceutical-products/ rdm:pharmaceutical-product / rdm:pharmaceutical-form- details/rdm:pharmaceutical-form- strengths/rdm: pharmaceutical- form-strength-description	Reference Medicinal Product > Medicinal Product > Pharmaceutical Product > Ingredient	B222-2
E222-17	Marketing authorisation holder	maa:product/ rdm:pharmaproduct/ rdm:pharmaceutical-products/ rdm:pharmaceutical-product / rdm:pharmaceutical-form- details/rdm:pharmaceutical-form- strengths /rdm:mp- authorisation/rdm:authorisat ion- holder	Role > Party > Organisation > Name	B222-2

E222-18	Marketing authorisation number	maa:product/ rdm:pharmaproduct/ rdm:pharmaceutical-products/ rdm:pharmaceutical-product / rdm:pharmaceutical-form- details/rdm:pharmaceutical- form-strengths /rdm:authorisation-numbers/ rdm:authorisation-number	MP Authorisation > authorization number	B222-2
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Element Tree Diagram

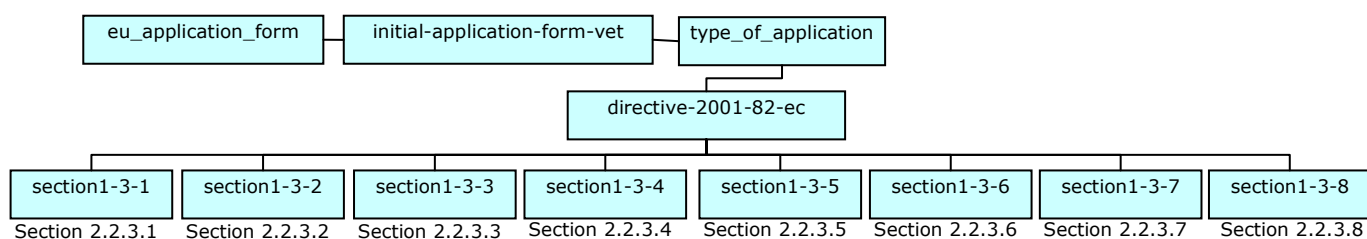


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B222-1	E222-1, E222-2	Mandatory.	Mutually Exclusive.	
B222-2	E222-1, E222-3 to E222-8, E222-14 to E222-19	E223-1 is Mandatory, rest are optional.	If E222-1 is selected, the rest are mandatory.	
B222-3	E222-8 to 13	Optional	If E222-8 is selected, then the rest are required and visible	
B222-4	E222-8 to 32	Optional	Mutually Exclusive	

2.2.3. THIS APPLICATION IS SUBMITTED IN ACCORDANCE WITH THE FOLLOWING ARTICLE IN DIRECTIVE 2001/82/EC

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/		Application > MP Procedure >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E223-1	Article 12(3) application, (i.e. dossier with administrative, quality, safety and efficacy data*)	maa:section1-3-1		B223-1
E223-2	Article 13(1) <u>Generic application</u>	maa:section1-3-2		B223-1
E223-3	Article 13(3) <u>hybrid application</u>	maa:section1-3-3		B223-1
E223-4	Article 13(4) <u>Similar biological application</u>	maa:section1-3-4		B223-1
E223-5	Article 13a - Well established Veterinary use	maa:section1-3-5		B223-1
E223-6	Article 13b - Fixed combination application	maa:section1-3-6		B223-1
E223-7	Article 13c - Informed consent application	maa:section1-3-7		B223-1
E223-8	Article 13d - Immunological Veterinary Medicinal Product for which the results of certain trials are not being submitted	maa:section1-3-8		B223-1

Element Tree Diagram

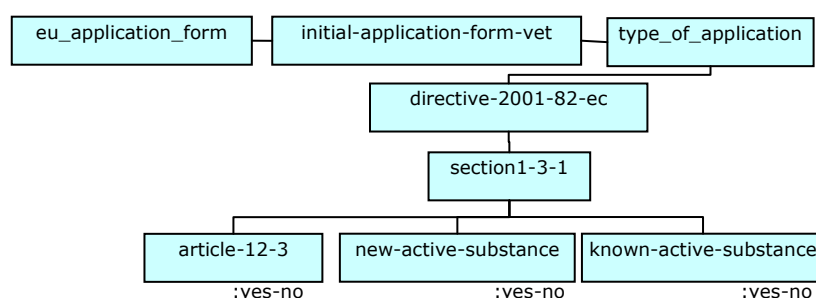


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B223-1	E223-1 to E223-8	Mandatory.	Mutually Exclusive.	

2.2.3.1. Article 12(3) application, (i.e. dossier with administrative, quality, safety and efficacy data*)

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-1/	Application		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2231-1	Article 12(3) application, (i.e. dossier with administrative, quality, safety and efficacy data*)	maa:article-12-3	Basis for Eligibility CTL	B223-1,
E2231-2	New active substance	maa:new-active-substance		B223-1, B223-2
E2231-3	Known active substance	maa:known-active-substance		B223-1, B223-2

Element Tree Diagram

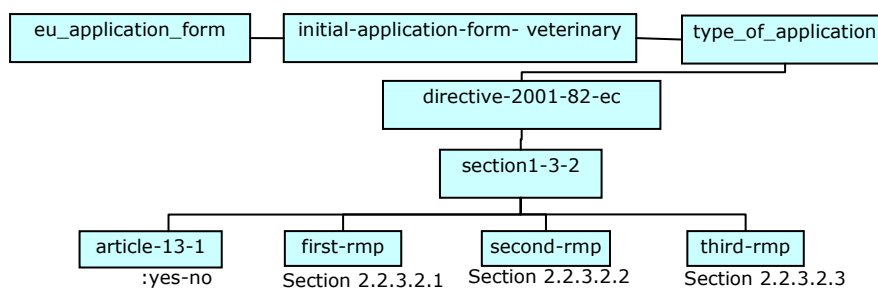


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B223-1	E2231-1 to 3	Optional	If E2231-1 is selected, the rest become required.	
B222-2	E2231-2 to 3	Optional	Mutually Exclusive.	

2.2.3.2. Article 13(1) Generic application

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-2/		Application > MP Procedure >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2232-1	Article 13(1) Generic application	maa:article-13-1	Basis for Eligibility CTL	B2232-1
E2232-2	Veterinary medicinal product which is or has been authorised in accordance with Union provisions in force for not less than 6/10 years in the EEA	maa:first-rmp		B2232-1
E2232-3	Veterinary medicinal product authorised in the Union/Member State where the application is made or European reference medicinal product:	maa:second-rmp		B2232-1
E2232-4	Veterinary medicinal product which is or has been authorised in accordance with Union provision in force and to which bioequivalence has been demonstrated by appropriate bioavailability studies	maa:third-rmp		

Element Tree Diagram

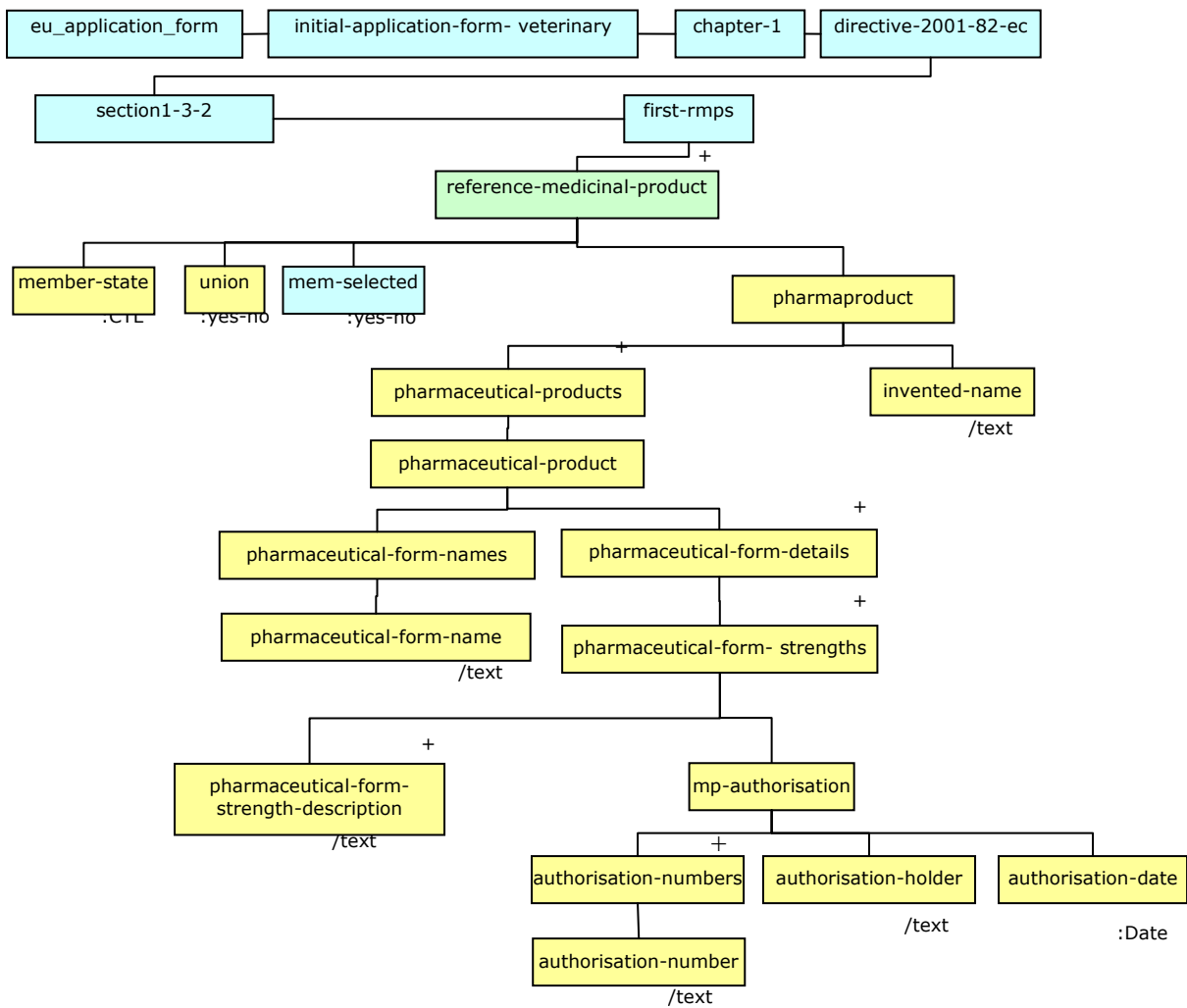


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2232-1	E2232-1 to 3	Optional	If E2232-1 is selected, the rest become required	

2.2.3.2.1. Veterinary medicinal product which is or has been authorised for not less than 6/10 years in the EEA

	Common DES 3.0 Context		Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-2/maa:first-rmp/maa:reference-medicinal-product/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E22321-1	Product (Invented) name	rdm:pharmaproduct/rdm:invented-name	Reference Medicinal Product > Medicinal Product > Medicinal Product Name	
E22321-2	Pharmaceutical form(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product / rdm:pharmaceutical-form-names/rdm:pharmaceutical-form-name	Reference Medicinal Product > Medicinal Product > Pharmaceutical Product > Pharmaceutical Dose Form CTL	
E22321-3	Strength(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:pharmaceutical-form-strength-description	Ingredient	
E22321-4	Marketing authorisation holder	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-holder	Role > Party> Organisation> Name	
E22321-5	Date of Authorisation	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:mp-authorisation/rdm:authorisation-date	MP Authorisation > authorisation date	
E22321-6	Marketing authorisation granted by			
E22321-7	Union	rdm:community	MP Authorisation > Country CTL	B22321-1
E22321-8	Member State(EEA)	rdm:mem-selected		B22321-1, B22321-2
E22321-9	Member State(EEA)	rdm:member-state	MP Authorisation > Country CTL	B22321-1, B22321-2
E22321-10	Marketing authorization number(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-numbers/rdm:authorisation-number	MP Authorisation > authorisation number	

Element Tree Diagram

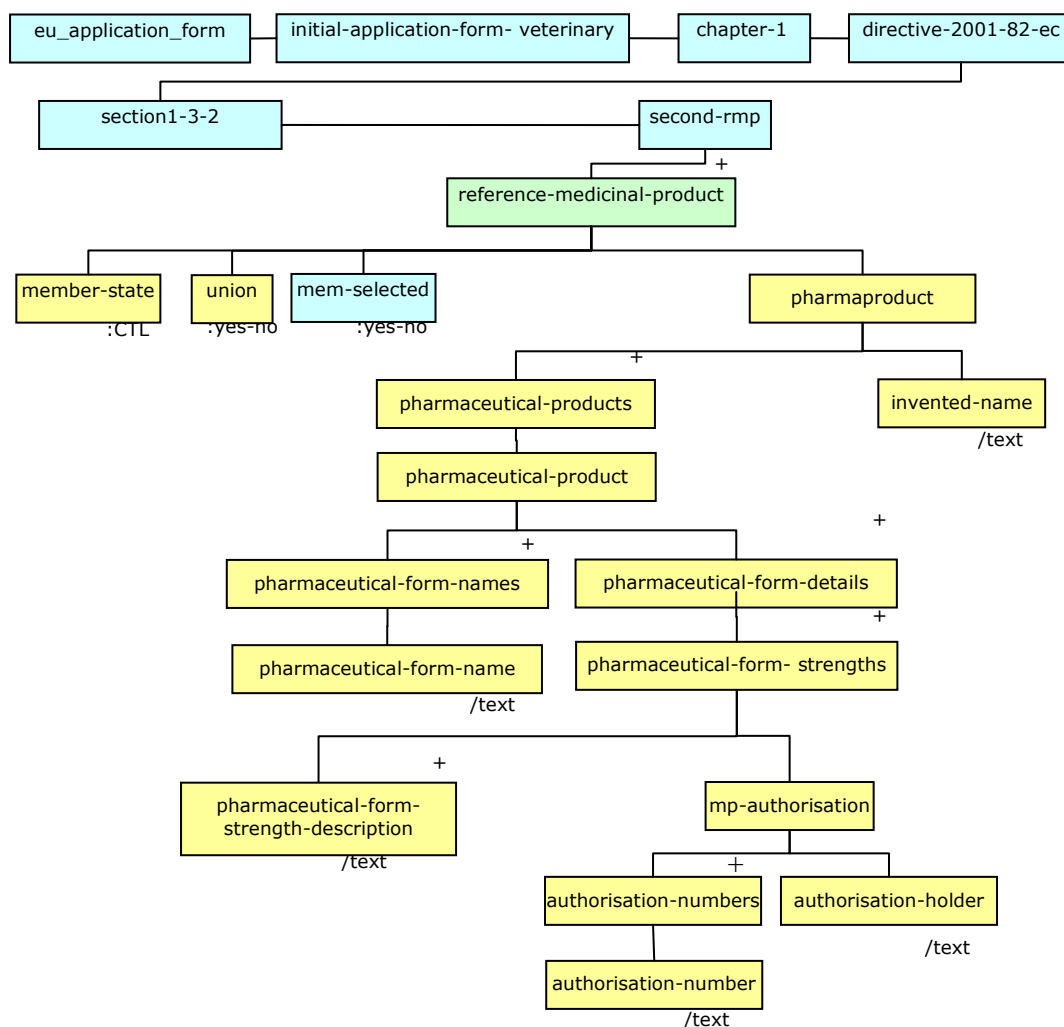


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B22321-1	E22321-7 to 8	Required	Mutually Exclusive	
B22321-2	E22321-8 to 9	Optional	If E22321-7 is selected, E22321-9 is required.	

2.2.3.2.2. Veterinary medicinal product authorised in the Union/Member State where the application is made or European reference medicinal product

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-2/maa:second-rmp/maa:reference-medicinal-product/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E22322-1	Product (Invented) name	rdm:pharmaproduct/rdm:invented-name	Reference Medicinal Product > Medicinal Product > Medicinal Product Name	
E22322-2	Pharmaceutical form(s)	rdm:pharmaproduct/rdm: pharmaceutical-products/rdm: pharmaceutical-product / rdm: pharmaceutical-form-names/rdm: pharmaceutical-form-name	Reference Medicinal Product > Medicinal Product > Pharmaceutical Product > Pharmaceutical Dose Form CTL	
E22322-3	Strength(s)	rdm: pharmaproduct/ rdm: pharmaceutical-products/rdm: pharmaceutical-product /rdm: pharmaceutical-form-details/rdm: pharmaceutical-form-strengths/rdm: pharmaceutical-form-strength-description	Ingredient	
E22322-4	Marketing authorisation holder	rdm: pharmaproduct/ rdm: pharmaceutical-products/rdm: pharmaceutical-product /rdm: pharmaceutical-form-details/rdm: pharmaceutical-form-strengths /rdm:mp-authorisation/ rdm:authorisation-holder	Role > Party> Organisation> Name	
E22322-5	Marketing authorisation number	rdm: pharmaproduct/ rdm: pharmaceutical-products/rdm: pharmaceutical-product /rdm: pharmaceutical-form-details/rdm: pharmaceutical-form-strengths /rdm:mp-authorisation/ rdm:authorisation-numbers/rdm:authorisation-number	MP Authorisation > authorisation number	
E22322-6	Marketing authorisation granted by			
E22322-7	Union	rdm:community	MP Authorisation > Country CTL	B22322-1
E22322-8	Member State(EEA)	rdm:mem-selected		B22322-1, B22322-2
E22322-9	Member State(EEA)	rdm:member-state	MP Authorisation > Country CTL	B22322-1, B22322-2

Element Tree Diagram

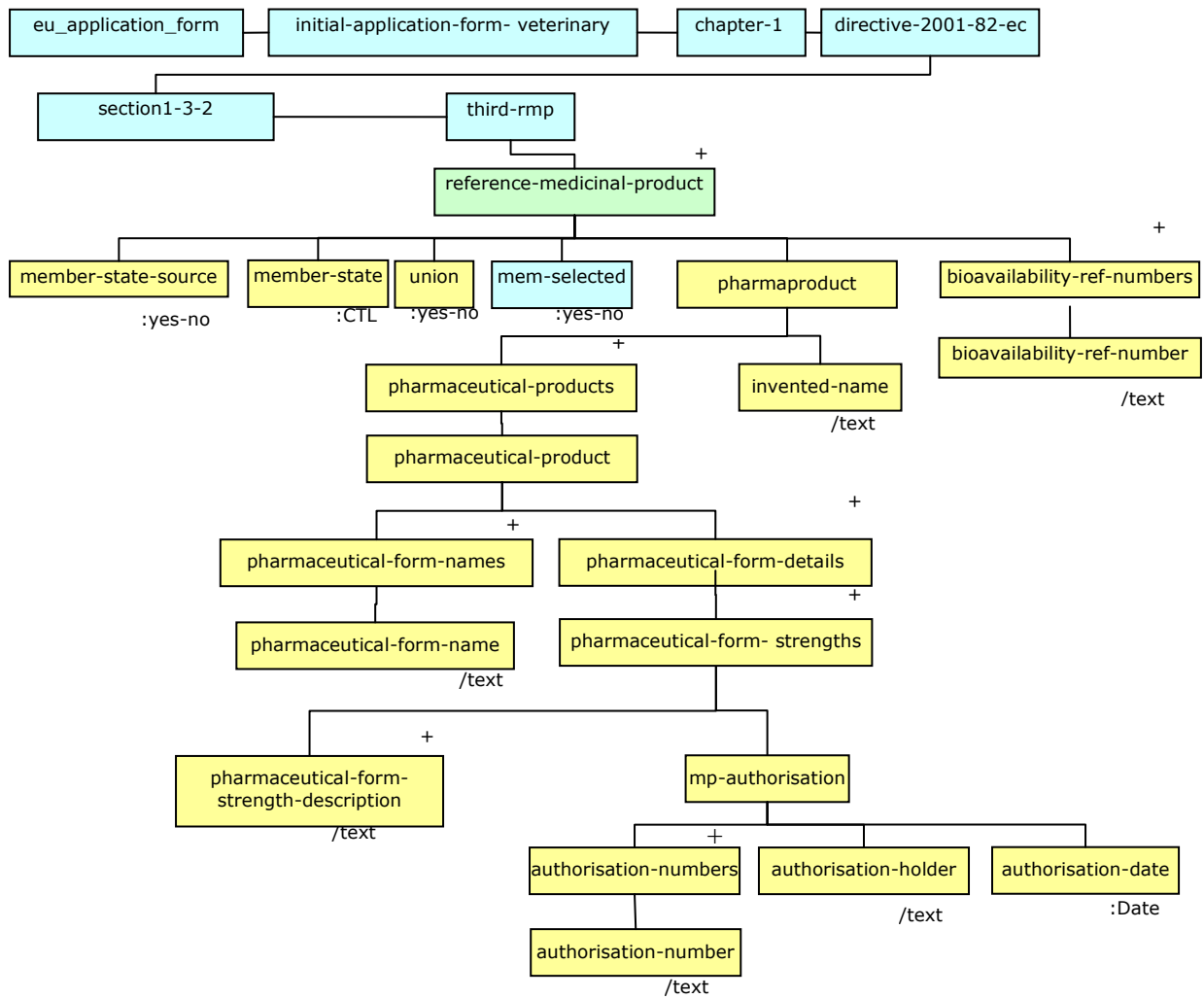


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B22322-1	E22322-7 to 8	Required	Mutually Exclusive	
B22322-2	E22322-8 to 9	Optional	If E22322-7 is selected, E22322-9 is required.	
B22322-2	E22322-10	Not visible	When E221-2 or E221-3 is selected, then E22322-10 is visible and mandatory	

2.2.3.2.3. Veterinary medicinal product which is or has been authorised in accordance with Union provision in force and to which bioequivalence has been demonstrated by appropriate bioavailability studies:

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-2/maa:third-rmp/maa:reference-medicinal-product/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E22323-1	Product (Invented) name	rdm:pharmaproduct/rdm:invented-name	Reference Medicinal Product > Medicinal Product > Medicinal Product Name	
E22323-2	Pharmaceutical form(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product / rdm:pharmaceutical-form-names/rdm:pharmaceutical-form-name	Reference Medicinal Product > Medicinal Product > Pharmaceutical Product > Pharmaceutical Dose Form CTL	
E22323-3	Strength(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:pharmaceutical-form-strength-description	Ingredient	
E22323-4	Marketing authorization holder	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-holder	Role > Party> Organisation > Name	
E22323-5	Member State of source	rdm:member-state-source	Country CTL	
E22323-6	Marketing authorization granted by			
E22323-7	Union	rdm:community	MP Authorisation > Country CTL	B22323-1
E22323-8	Member State(EEA)	rdm:mem-selected		B22323-1, B22323-2
E22323-9	Member State(EEA)	rdm:member-state	MP Authorisation > Country CTL	B22323-1, B22323-2
E22323-10	Bioavailability study(ies) reference number(s)	rdm:bioavailability-ref-numbers/rdm:bioavailability-ref-number		
E22323-11	Marketing authorisation number	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-numbers/rdm:authorisation-number	MP Authorisation > authorisation number	
E22323-12	Date of authorisation	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:mp-authorisation/rdm:authorisation-date	MP Authorisation > authorisation date	

Element Tree Diagram

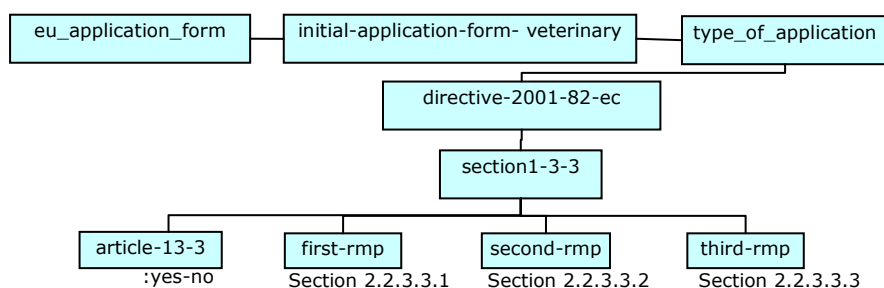


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B22323-1	E22323-7 to 8	Required	Mutually Exclusive	
B22323-2	E22323-8 to 9	Optional	If E22323-7 is selected, E22323-9 is required.	

2.2.3.3. Article 13(3) hybrid application

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-3/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2233-1	Article 13(3) hybrid application	maa:article13-3	Basis for Eligibility CTL	
E2233-2	Reference veterinary medicinal product which is or has been authorised for not less than 6/10 years in the EEA	maa:first-rmp		B2233-1
E2233-3	Reference veterinary medicinal product authorised in the Community/Member State where the application is made	maa:second-rmp		B2233-1
E2233-4	Veterinary medicinal product used in bioequivalence studies, where applicable	maa:third-rmp		B2233-1

Element Tree Diagram

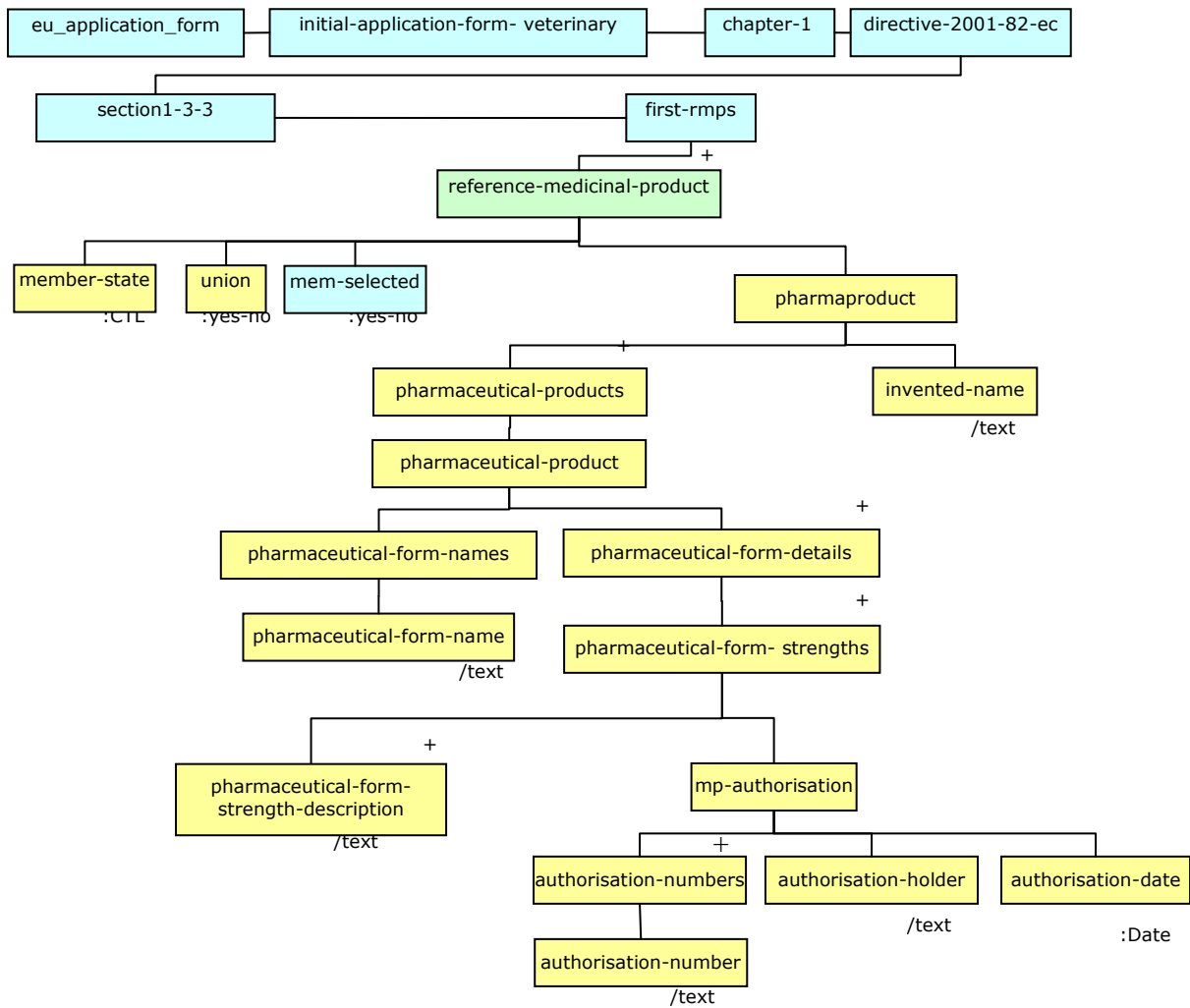


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2233-1	E2233-1 to 4	Optional	If E2233-1 is selected, the rest become required	

2.2.3.3.1. Veterinary medicinal product which is or has been authorised in accordance with Union provision in force for not less than 6/10 years in the EEA

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-3/maa:first-rmp/maa:reference-medicinal-product/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E22331-1	Product (Invented) name	rdm:pharmaproduct/rdm:invented-name	Reference Medicinal Product > Medicinal Product > Medicinal Product Name	
E22331-2	Pharmaceutical form(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product / rdm:pharmaceutical-form-names/rdm:pharmaceutical-form-name	Reference Medicinal Product > Medicinal Product > Pharmaceutical Product > Pharmaceutical Dose Form CTL	
E22331-3	Strength(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:pharmaceutical-form-strength-description	Ingredient	
E22331-4	Marketing authorisation holder	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-holder	Role > Party> Organisation> Name	
E22331-5	Date of Authorisation	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:mp-authorisation/rdm:authorisation-date	MP Authorisation > authorisation date	
E22331-6	Marketing authorisation granted by			
E22331-7	Union	rdm:community	MP Authorisation > Country CTL	B22331-1
E22331-8	Member State(EEA)	rdm:mem-selected		B22331-1, B22331-2
E22331-9	Member State(EEA)	rdm:member-state	MP Authorisation > Country CTL	B22331-2
E22331-9	Marketing authorisation number(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-numbers/rdm:authorisation-number	MP Authorisation > authorisation number	

Element Tree Diagram

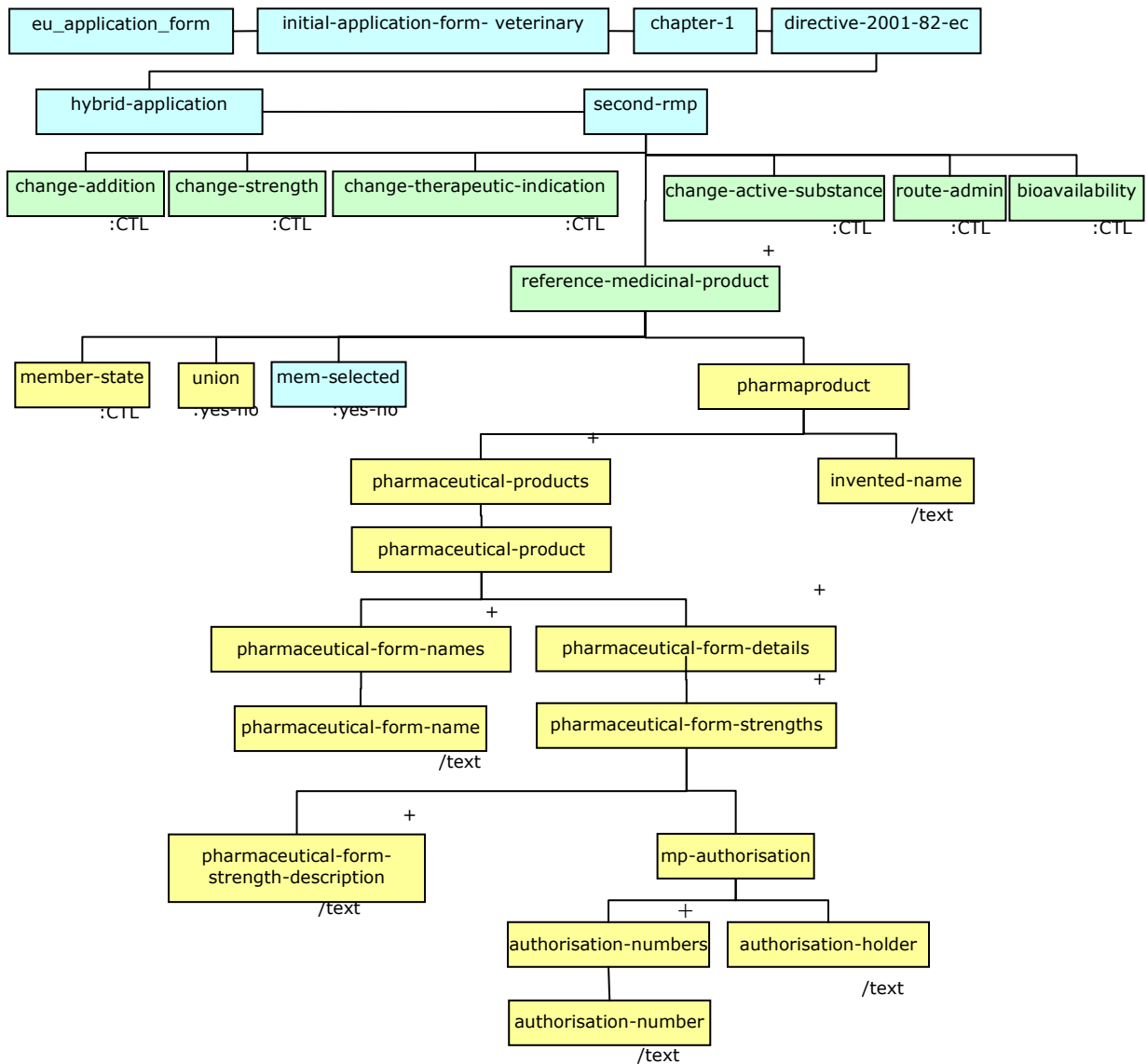


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B22331-1	E22331-7 to 8	Optional	Mutually Exclusive	
B22331-2	E22331-8 to 9	Optional	If E22331-8, then E22331-9 is required	

2.2.3.3.2. Veterinary medicinal product authorised in the Union/Member State where the application is made or European reference medicinal product

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-3/maa:second-rmp		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E22332-1	Product (Invented) name	maa:reference-medicinal-product/rdm:pharmaproduct/rdm:invented-name	Reference Medicinal Product > Medicinal Product > Medicinal Product Name	
E22332-2	Pharmaceutical form(s)	maa:reference-medicinal-product/rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-names/rdm:pharmaceutical-form-name	Reference Medicinal Product > Medicinal Product > Pharmaceutical Product > Pharmaceutical Dose Form CTL	
E22332-3	Strength(s)	maa:reference-medicinal-product/rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:pharmaceutical-form-strength-description	Ingredient	
E22332-4	Marketing authorisation holder	maa:reference-medicinal-product/rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-holder	Role > Party> Organisation> Name	
E22332-5	Marketing authorisation number	maa:reference-medicinal-product/rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-numbers/rdm:authorisation-number	MP Authorisation > authorisation number	
E22332-6	Marketing authorisation granted by			
E22332-7	Union	maa:reference-medicinal-product/rdm:community	MP Authorisation > Country CTL	B22332-1
E22332-8	Member State(EEA)	maa:reference-medicinal-product/rdm:mem-selected		B22332-1, B22332-2
E22332-9	Member State(EEA)	maa:reference-medicinal-product/rdm:member-state	MP Authorisation > Country CTL	B22332-2
E22333-10	Difference(s) compared to this reference veterinary medicinal product			
E22333-11	Changes in the active substance(s)	maa:change-active-substance	Difference CTL	B22423-3
E22333-12	Change in therapeutic indications	maa:change-therapeutic-indication	Difference CTL	B22423-3
E22333-13	Change in strength (quantitative change to the active substance(s))	maa:change-strength	Difference CTL	B22423-3
E22333-14	Change in pharmaceutical form	maa:change-addition	Difference CTL	B22423-3
E22333-15	Change in route of administration	maa:route-admin	Difference CTL	B22423-3
E22333-16	Bioequivalence cannot be demonstrated through bioavailability studies	maa:bioequivalence	Difference CTL	B22423-3
E22333-17	Member States	maa:reference-medicinal-product/rdm:member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	B22423-4

Element tree Diagram

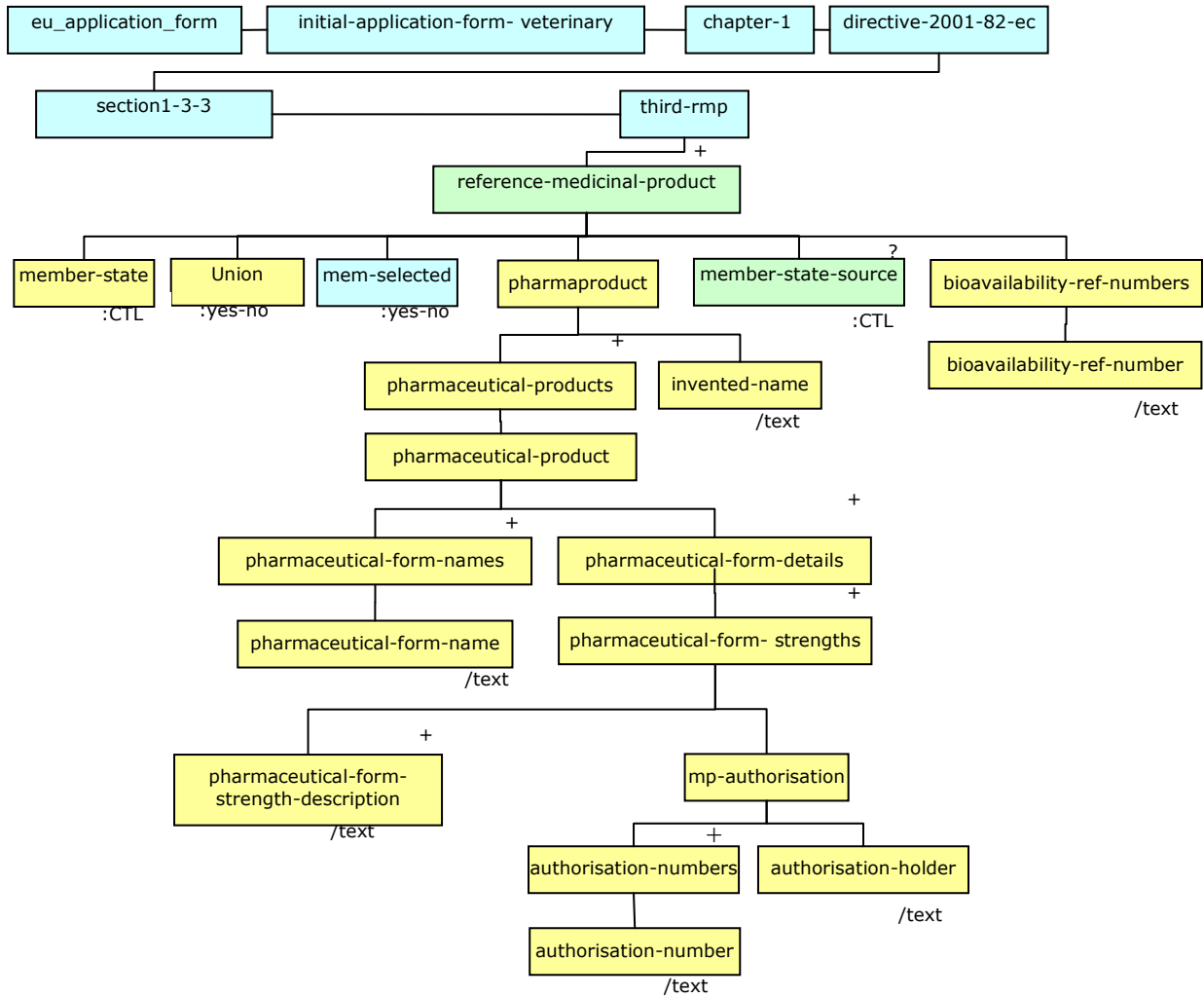


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B22332-1	E22333-7 to 8	Optional	Mutually Exclusive	
B22332-2	E22333-8 to 9	Optional	If E22333-8 is selected, then E22332-9 is required	
B22423-3	E22333-11 to 16	Mandatory	Mutually Exclusive	
B22423-4	E22333-17,	E22423-17 is not visible.	When E221-2 or E221-3 is selected, then E22333-17 is visible and mandatory	

2.2.3.3.3. Veterinary Medicinal Product which is or has been authorised in accordance with Union provisions in force used for the demonstration of bioequivalence (if applicable) and/or in other studies.

	Common DES 3.0 Context		Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-3/maa:third-rmp/maa:reference-medicinal-product		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E22333-1	Product (Invented) name	rdm:pharmaproduct/rdm:invented-name	Reference Medicinal Product > Medicinal Product > Medicinal Product Name	
E22333-2	Pharmaceutical form(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product / rdm:pharmaceutical-form-names/rdm:pharmaceutical-form-name	Reference Medicinal Product > Medicinal Product > Pharmaceutical Product > Pharmaceutical Dose Form CTL	
E22333-3	Strength(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:pharmaceutical-form-strength-description	Ingredient	
E22333-4	Marketing authorization holder	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-holder	Role > Party> Organisation> Name	
E22333-4a	Marketing authorisation number	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-numbers/rdm:authorisation-number	MP Authorisation > authorisation number	
E22333-5	Member State of source	rdm:member-state	Country CTL	
E22333-6	Marketing authorisation granted by			
E22333-7	Union	rdm:community	MP Authorisation > Country CTL	B22333-1
E22333-8	Member State(EEA)	rdm:mem-selected		B22333-1, B22333-2
E22333-9	Member State(EEA)	rdm:member-state	MP Authorisation > Country CTL	B22333-2
E22333-10	Study reference number	rdm:bioavailability-ref-numbers/rdm:bioavailability-ref-number	Reference Medicinal Product > study ref number	

Element Tree Diagram

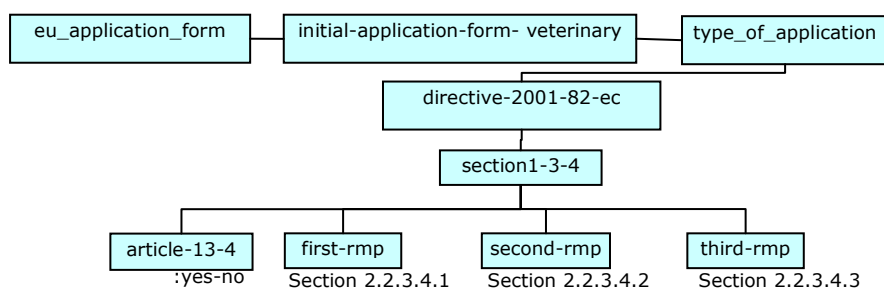


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B22333-1	E22333-7 to 8	Optional	Mutually Exclusive	
B22333-2	E22333-8 to 9	Optional	If E22333-8, then E22333-9 is required	

2.2.3.4. Article 13(4) Similar biological application

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-4/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2234-1	Article 13(4) <u>Similar biological application</u>	maa:article-13-4	Basis for Eligibility CTL	B2234-1
E2234-2	<i>Reference product which is or has been authorised for not less than 6/10 years in the EEA</i>	maa:first-rmp		B2234-1
E2234-3	<i>Reference veterinary medicinal product authorised in the Community/Member State where the application is made</i>	maa:second-rmp		B2234-1
E2234-4	<i>Veterinary medicinal product used in bioequivalence studies where applicable</i>	maa:third-rmp		B2234-1

Element Tree Diagram

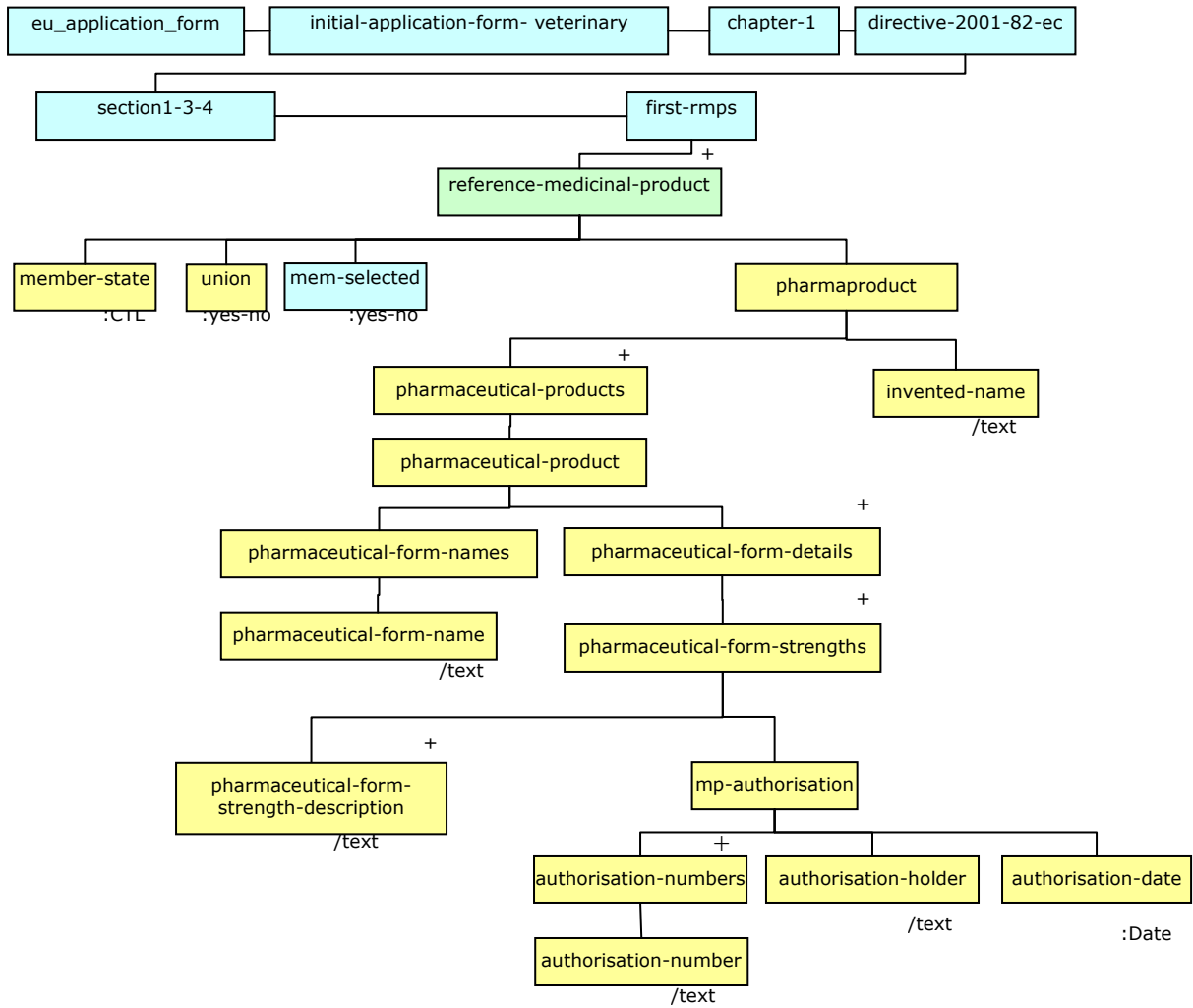


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2234-1	E2234-1 to 4	Optional	If E2234-1 is selected, the rest become required	

2.2.3.4.1. Reference veterinary medicinal product which is or has been authorised in accordance with Union provisions in force for not less than 6/10 years in the EEA

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-4/maa:first-rmp/maa:reference-medicinal-product/	Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E22341-1	Product (Invented) name	rdm:pharmaproduct/rdm:invented-name	Reference Medicinal Product > Medicinal Product > Medicinal Product Name	
E22341-2	Pharmaceutical form(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product / rdm:pharmaceutical-form-names/rdm:pharmaceutical-form-name	Reference Medicinal Product > Medicinal Product > Pharmaceutical Product > Pharmaceutical Dose Form CTL	
E22341-3	Strength(s)	rdm:pharmaproduct/ rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:pharmaceutical-form-strength-description	Ingredient	
E22341-4	Marketing authorisation holder	rdm:pharmaproduct/ rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-holder	Role > Party> Organisation > Name	
E22341-5	Date of Authorisation	rdm:pharmaproduct/ rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:mp-authorisation/rdm:authorisation-date	MP Authorisation > authorisation date	
E22341-6	Marketing authorisation granted by			
E22341-7	Union	rdm:community	MP Authorisation > Country CTL	B22341-1
E22341-8	Member State(EEA)	rdm:mem-selected		B22341-1, B22341-2
E22341-9	Member State(EEA)	rdm:member-state	MP Authorisation > Country CTL	B22341-2
E22341-10	Marketing authorisation number	rdm:pharmaproduct/ rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-numbers/rdm:authorisation-number	MP Authorisation > authorisation number	

Element Tree Diagram

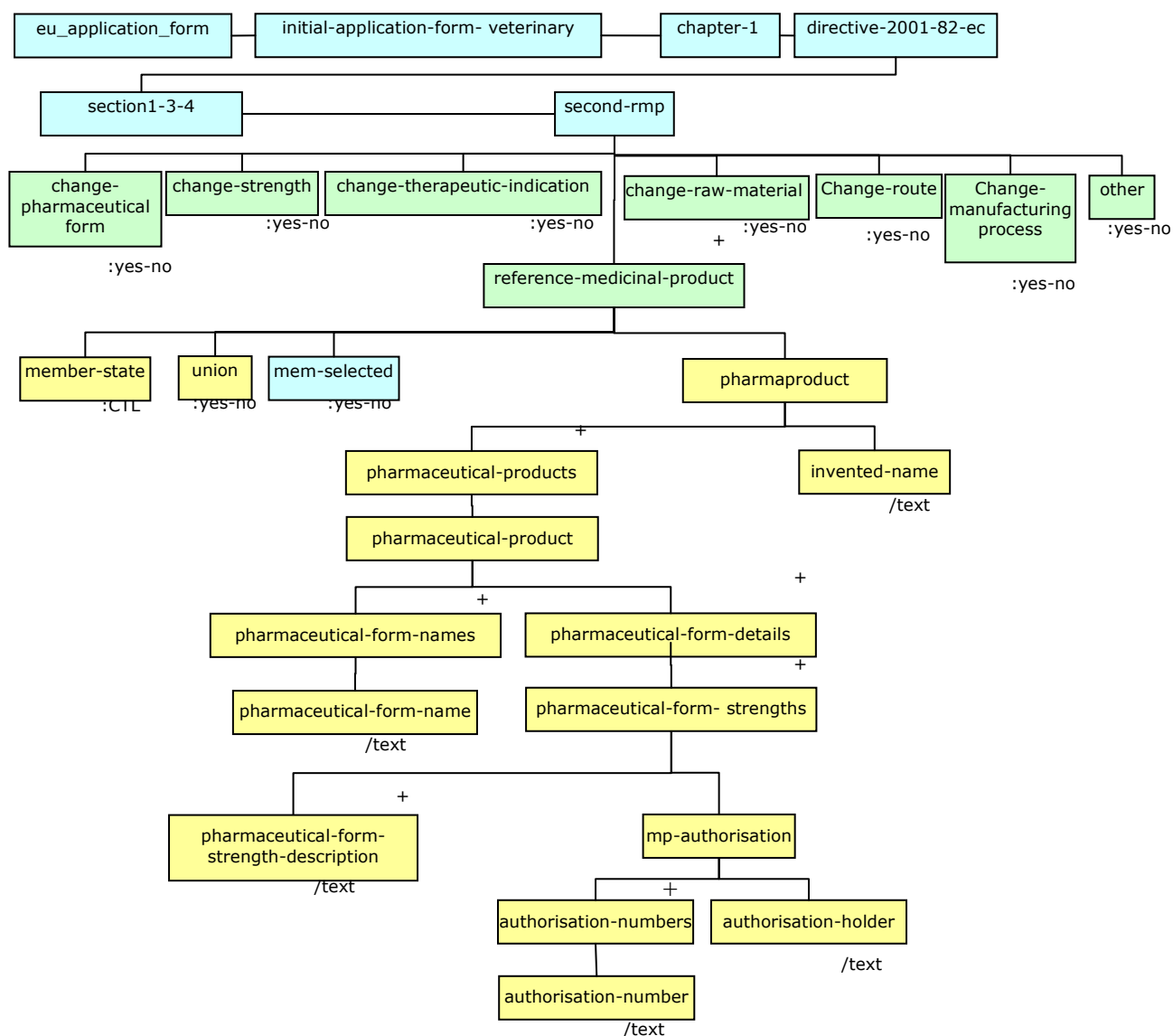


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B22341-1	E22341-7 to 8	Optional	Mutually Exclusive	
B22341-2	E22341-8 to 9	Optional	If E22341-8, then E22341-9 is required	

2.2.3.4.2. Reference veterinary medicinal product authorised in the Union/Member State where the application is made or European reference medicinal product

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-4/maa:second-rmp		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E22342-1	Product (Invented) name	maa:reference-medicinal-product/rdm:pharmaproduct/rdm:invented-name	Reference Medicinal Product > Medicinal Product > Medicinal Product Name	
E22342-2	Pharmaceutical form(s)	maa:reference-medicinal-product/rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-names/rdm:pharmaceutical-form-name	Reference Medicinal Product > Medicinal Product > Pharmaceutical Product > Pharmaceutical Dose Form CTL	
E22342-3	Strength(s)	maa:reference-medicinal-product/rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:pharmaceutical-form-strength-description	Ingredient	
E22342-4	Marketing authorisation holder	maa:reference-medicinal-product/rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-holder	Role > Party> Organisation> Name	
E22342-5	Marketing authorisation number	maa:reference-medicinal-product/rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-numbers/rdm:authorisation-number	MP Authorisation > authorisation number	
E22342-6	Marketing authorisation granted by			
E22342-7	Union	maa:reference-medicinal-product/rdm:community	MP Authorisation > Country CTL	B22342-1
E22342-8	Member State(EEA)	maa:reference-medicinal-product/rdm:mem-selected		B22342-1, B22342-2
E22342-9	Member State(EEA)	maa:reference-medicinal-product /rdm:member-state	MP Authorisation > Country CTL	B22342-2
E22342-10	Difference(s) compared to this reference medicinal product			
E22342-11	change-raw material	rdm:change-raw-substance		B22342-3
E22342-12	change-manufacturing process	rdm:manufacturing		B22342-3
E22342-13	change-therapeutic indication	rdm:change-therapeutic-indication		B22342-3
E22342-14	change-pharmaceutical form	rdm:change-addition		B22342-3
E22342-15	Change-strength	rdm:change-strength		B22342-3
E22342-16	Change - route	rdm:route-admin		B22342-3
E22342-17	other	rdm:other		B22342-3

Element Tree Diagram

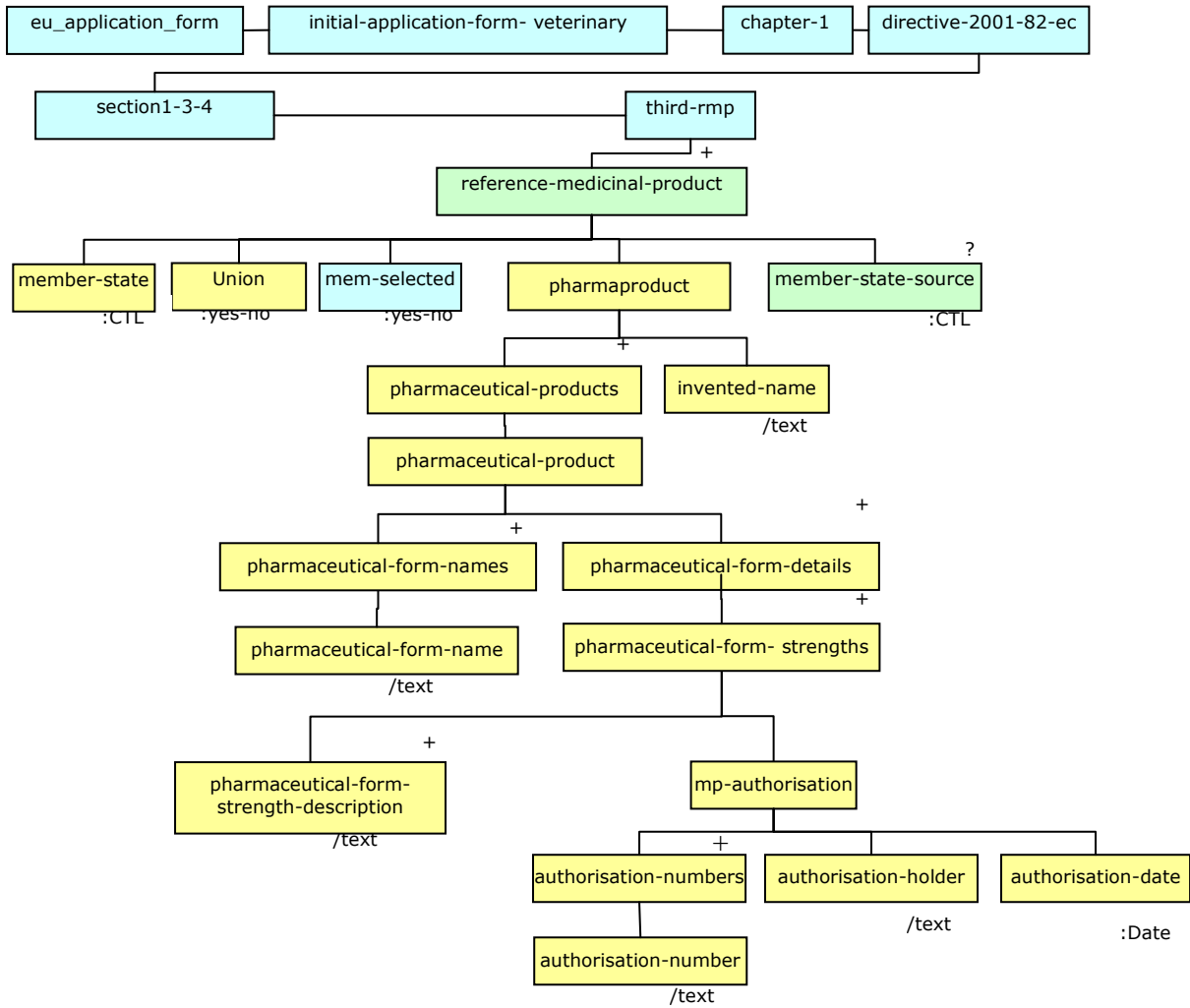


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B22342-1	E22342-7 to 8	Optional	Mutually Exclusive	
B22342-2	E22342-8 to 9	Optional	If E22342-8, then E22342-9 is required	
B22342-3	E22342-11 to 17	Mandatory	Mutually Exclusive	

2.2.3.4.3. Veterinary medicinal product which is or has been authorised in accordance with Union provisions in force and to which comparability tests and studies have been conducted

		Common DES 3.0 Context	Common RDM Entry point	
		maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-4/maa:third-rmp/maa:reference-medicinal-product/	Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E22343-1	Product (Invented) name	rdm:pharmaproduct/rdm:invented-name	Reference Medicinal Product > Medicinal Product > Medicinal Product Name	
E22343-2	Pharmaceutical form(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product / rdm:pharmaceutical-form-names/rdm:pharmaceutical-form-name	Reference Medicinal Product > Medicinal Product > Pharmaceutical Product > Pharmaceutical Dose Form CTL	
E22343-3	Strength(s)	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:pharmaceutical-form-strength-description	Ingredient	
E22343-4	Marketing authorization holder	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-holder	Role > Party> Organisation> Name	
E22343-5	Date of Authorisation	rdm:products/rdm:product/rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:mp-authorisation/rdm:authorisation-date	MP Authorisation > authorisation date	
E22343-6	Marketing authorisation granted by			
E22343-7	Union	rdm:community	MP Authorisation > Country CTL	B22343-1
E22343-8	Member State(EEA)	rdm:mem-selected		B22343-1, B22343-2
E22343-9	Member State(EEA)	rdm:member-state	MP Authorisation > Country CTL	B22343-2
E22343-10	Marketing authorisation number	rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product /rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths /rdm:mp-authorisation/rdm:authorisation-numbers/rdm:authorisation-number	MP Authorisation > authorisation number	

Element Tree Diagram

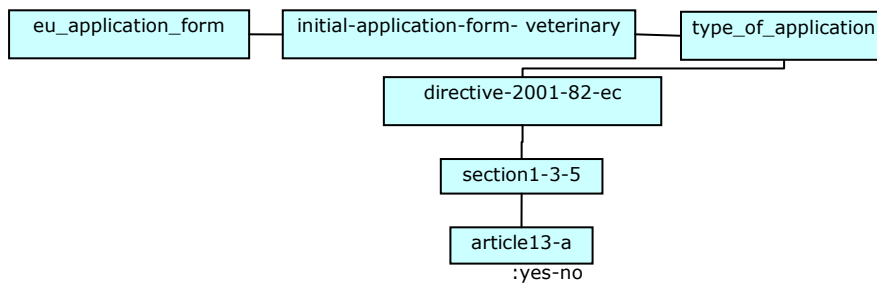


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B22343-1	E22343-7 to 8	Optional	Mutually Exclusive	
B22343-2	E22343-8 to 9	Optional	If E22343-8, then E22343-9 is required	

2.2.3.5. Article 13a - Well established Veterinary use

	Common DES 3.0 Context	Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-5/	Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2235-1	Article 13a - Well established Veterinary use	maa:article13-a	Basis for Eligibility CTL	

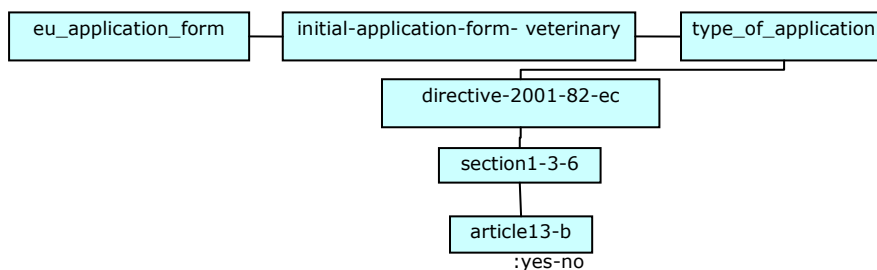
Element tree Diagram



2.2.3.6. Article 13b - Fixed combination application

	Common DES 3.0 Context	Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-6/	Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2236-1	Article 13b - Fixed combination application	maa:article13-b	Basis for Eligibility CTL	

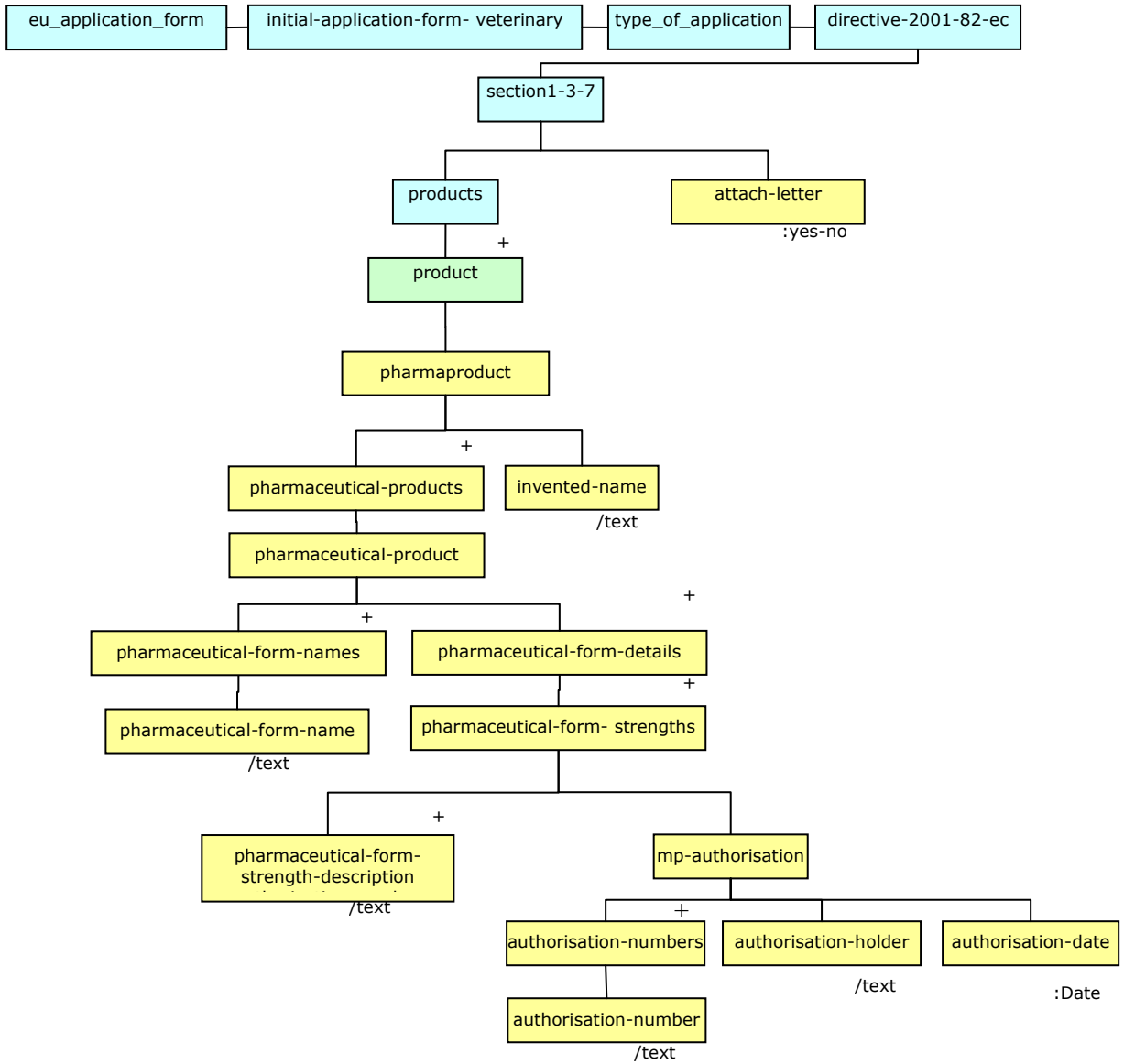
Element Tree Diagram



2.2.3.7. Article 13c - Informed consent application

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-7/	Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2237-1	Article 13c - Informed consent application	maa:article13-c	Basis for Eligibility CTL	
E2237-2	Product (Invented) name	rdm:products/rdm:product/rdm:pharma product/rdm:invented-name	Reference Medicinal Product > medicinal product name	
E2237-3	Pharmaceutical form(s)	rdm:products/rdm:product/rdm:pharmaproduct/rdm: pharmaceutical-products/rdm: pharmaceutical-product / rdm: pharmaceutical-form-names/rdm: pharmaceutical-form-name	Reference Medicinal Product >Pharmaceutical Dose Form CTL	
E2237-4	Strength(s)	rdm:products/rdm:product/rdm: pharmaproduct/ rdm: pharmaceutical-products/rdm: pharmaceutical-product /rdm: pharmaceutical-form-details/rdm: pharmaceutical-form-strengths/rdm: pharmaceutical-form-strength-description	Ingredient	
E2237-5	Marketing authorisation holder	rdm:products/rdm:product/rdm: pharmaproduct/ rdm: pharmaceutical-products/rdm: pharmaceutical-product /rdm: pharmaceutical-form-details/rdm: pharmaceutical-form-strengths /rdm:mp-authorisation/ rdm:authorisation-holder	Role > Party	
E2237-6	Marketing authorisation number	rdm:products/rdm:product/rdm: pharmaproduct/ rdm: pharmaceutical-products/rdm: pharmaceutical-product /rdm: pharmaceutical-form-details/rdm: pharmaceutical-form-strengths /rdm:mp-authorisation/ rdm:authorisation-numbers/rdm:authorisation-number	MP Authorisation > authorisation number	
E2237-7	Attach letter of consent from marketing authorisation holder of the authorised product	maa:attach-letter		

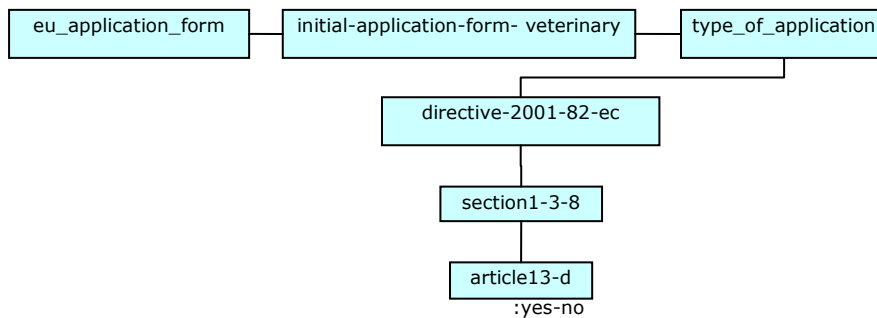
Element Tree Diagram



2.2.3.8. Article 13d - Immunological Veterinary Medicinal Product for which the results of certain trials are not being submitted

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:directive-2001-82-ec/maa:section1-3-8/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2235-1	Article 13d - Immunological Veterinary Medicinal Product for which the results of certain trials are not being submitted	maa:article13-d	Basis for Eligibility CTL	

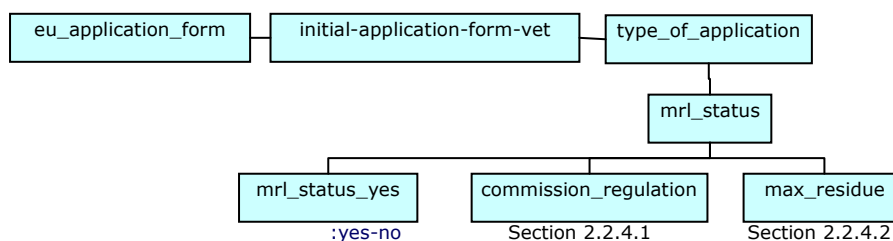
Element Tree Diagram



2.2.4. MRL status (only for food producing species)

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:mrl_status/		Application > MP Procedure >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E223-1	Maximum Residue Limits (MRL) according to Commission Regulation (EU) No 37/2010	maa:commission_regulation		
E223-2	Application for a Maximum Residue Limit has been made to the EMA	maa:max_residue		
E223-3	Check box	maa:mrl_status_yes		B223-1

Element Tree Diagram

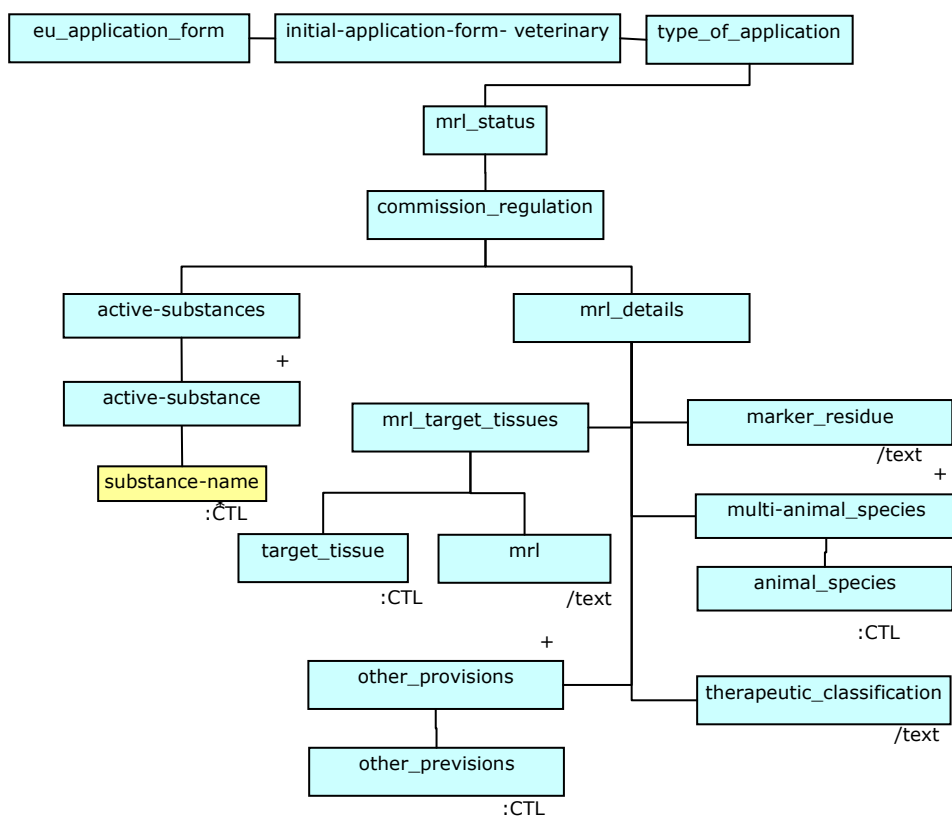


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B223-1	E223-3	Optional	If E223-3 is selected then E2241-1 to E2242-5 is visible	E2241-1 to E2242-5

Maximum Residue Limits (MRL) according to Commission Regulation (EU) No 37/2010

	Common DES 3.0 Context	Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:mrl_status/maa:commission_regulation/	Application > MRL		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2241-1	Active Substance(s)	maa:active-substances/maa:active-substance/rdm:substance-name	Ingredient > Substance CTL, Ingredient > IngredientRole CTL	B223-1
E2241-2	Marker residue	maa:mrl_details/maa:marker_residue		B223-1
E2241-3	Animal species	maa:mrl_details/maa:multi-animal_species/maa:animal_species	Target Population > Species CTL	B223-1
E2241-4	MRL	maa:mrl_details/maa:mrl_target_tissues/maa:mrl	Maximum Residue Limit	B223-1
E2241-5	Target Tissue	maa:mrl_details/maa:mrl_target_tissues/maa:target_tissue	Tissue CTL	B223-1
E2241-6	Other Provisions	maa:mrl_details/maa:other_provisions/maa:other_provisions		B223-1
E2241-7	Therapeutic classification	maa:mrl_details/maa:therapeutic_classification		B223-1

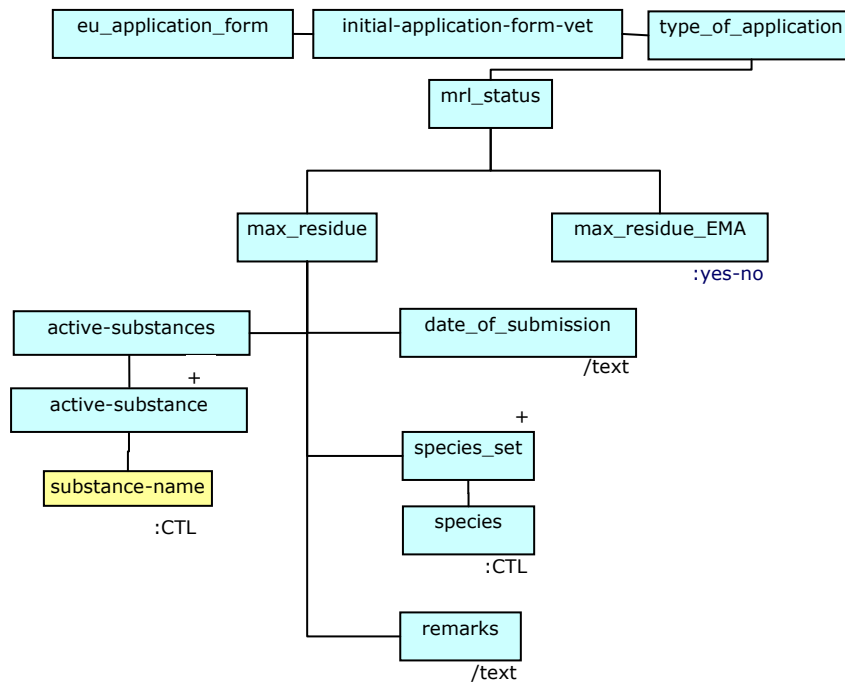
Element Tree Diagram



2.2.4.1. Application for a Maximum Residue Limit has been made to the EMA

Common DES 3.0 Context			Common RDM Entry point	
maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:mrl_status			Application > MRL >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2242-1	Active Substance(s)	maa:max_residue/maa:active-substances/maa:active-substance/rdm:substance-name	Ingredient > Substance CTL, Ingredient > IngredientRole CTL	B223-1
E2242-2	Date of Submission	maa:max_residue/maa:date_of_submission	Emea submission date	B223-1
E2242-3	Species	maa:max_residue/maa:species_set/maa:species	Species CTL	B223-1
E2242-4	Remarks	maa:max_residue/maa:remarks	mrl remarks	B223-1
E2242-5	Application for a Maximum Residue Limit has been made to the EMA: Yes – not applicable	maa:max_residue_EMA		B223-1

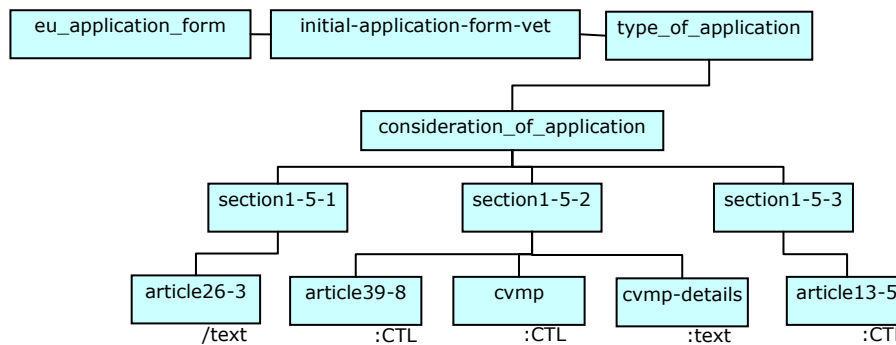
Element Tree Diagram



2.2.5. Consideration of this application is also requested under the following article in Directive 2001/82/EC or Regulation (EC) no. 726/2004

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:type_of_application/maa:consideration_of_application/	Application > MP Procedure >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E225-1	Exceptional Circumstances <i>Note: according to Article 26(3) of Directive 2001/82/EC and Article 39(7) of Regulation (EC) No 726/2004</i>	maa:section1-5-1/ maa:article26-3		B225-1
E225-2	Accelerated Review <i>Note: centralised procedure only according to Regulation (EC) No 726/2004 Article 39(8)</i>	maa:section1-5-2/ maa:article39-8		B225-1, B225-2
E225-3	Date of acceptance by CVMP	maa:section1-5-2/maa:cvmp		B225-2
E225-4	Free text field	maa:section1-5-2/maa:cvmp-details		B225-2
E225-5	Article 13(5) of Directive 2001/82/EC (one year of data exclusivity for an extension to another food-producing species within five years of the initial authorisation)	maa:section1-5-3/ maa:article13-5		B225-1

Element Tree Diagram

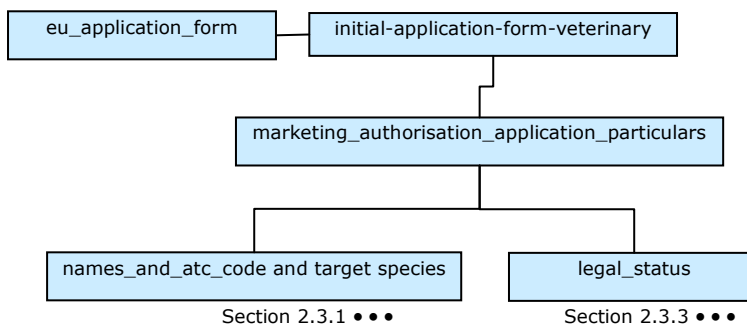


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B225-1	E225-1, 2, 4	Optional	Mutually Exclusive	
B225-2	E225-2, E225-3, E225-4	Optional	If E225-2 is selected, E225-3 and E225-4 are visible	

2.3. MARKETING AUTHORISATION APPLICATION PARTICULARS

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E23-1	NAME(S), ATC VET CODE and TARGET SPECIES	maa:names_and_atc_code		
E23-2	STRENGTH, PHARMACEUTICAL FORM, ROUTE OF ADMINISTRATION, CONTAINER AND PACK SIZES			
E23-3	LEGAL STATUS	maa:legal_status		
E23-4	MARKETING AUTHORISATION HOLDER / CONTACT PERSONS / COMPANY			
E23-5	MANUFACTURERS			
E23-6	QUALITATIVE AND QUANTITATIVE COMPOSITION			

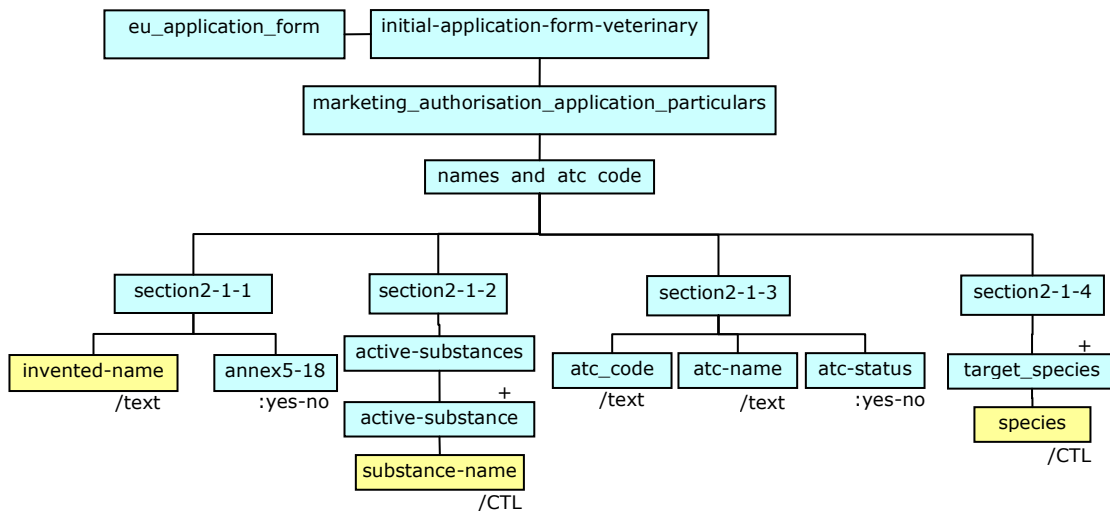
Element Tree Diagram



2.3.1. NAME(S), ATC VET CODE AND TARGET SPECIES

	Common DES 3.0 Context	Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:names_and_atc_code/	Application > MP Procedure >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E231-1	Proposed (invented) name of the veterinary medicinal product in the European Union / Member State / Iceland / Liechtenstein / Norway	maa:section2-1-1/ maa:invented-name	Substance CTL	
E231-2	If different (invented) names in different Member States are proposed in a mutual recognition or decentralised procedure, these should be listed in Annex 5.18	maa:section2-1-1/ maa:annex5-18		
E231-3	Name of the active substance(s)	maa:section2-1-2/ maa:active-substances/ maa:active-substance/ rdm:substance-name	Substance CTL	
E231-4	Pharmacotherapeutic group (Please use current ATC vet code)			
E231-5	Group	maa:section2-1-3/ maa:atc/maa:atc_name	ATC Group CTL	
E231-6	ATC Vet code	maa:section2-1-3/ maa:atc/maa:atc_code	ATC Group CTL	
E231-7	Please indicate if the application for the ATC Vet Code is still pending	maa:section2-1-3/ maa:atc/maa:atc-status		
E231-8	Target Species	maa:section2-1-4/ maa:target_species/ maa:species	Species CTL	

Element Tree Diagram

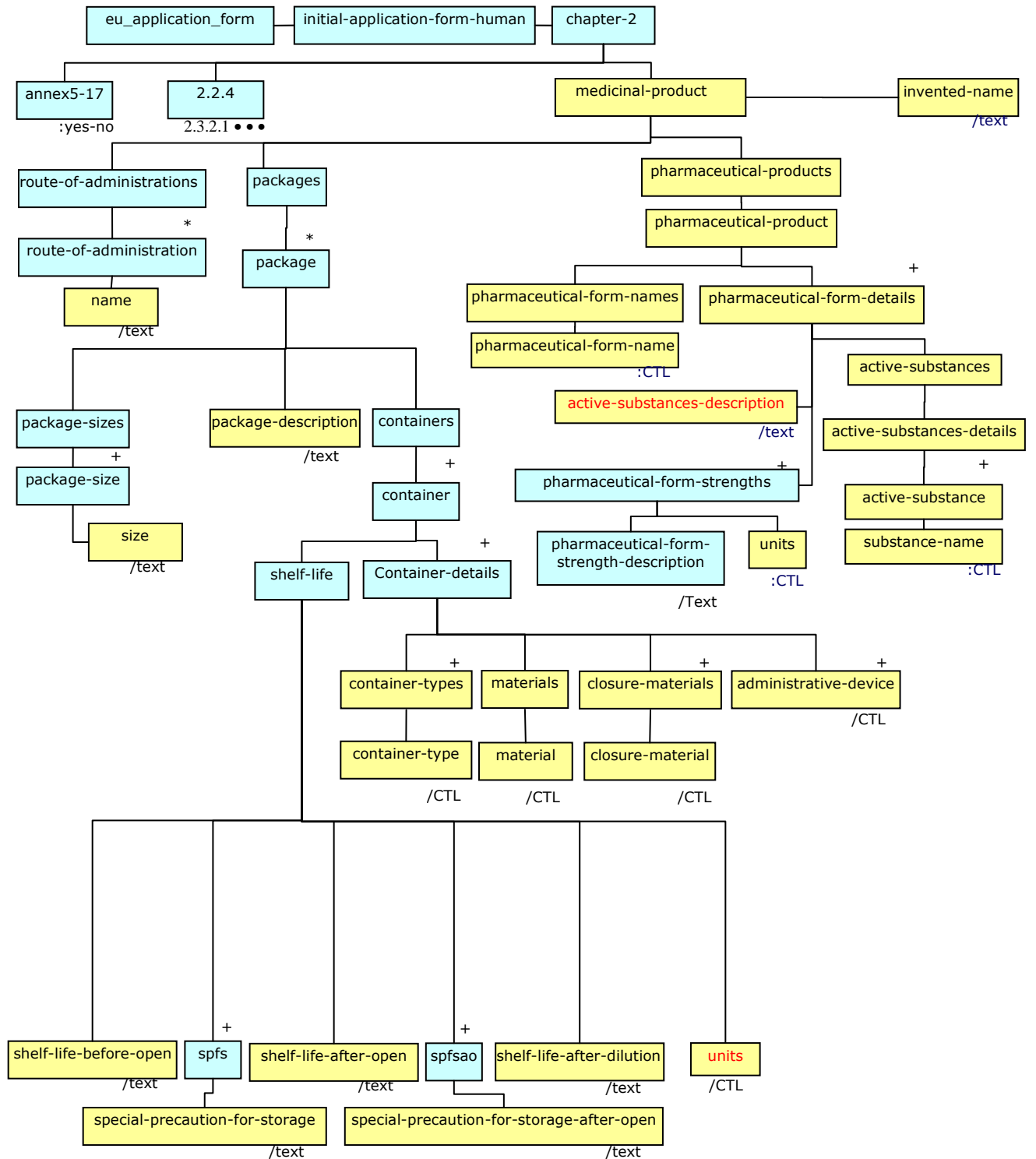


2.3.2. STRENGTH, PHARMACEUTICAL FORM, ROUTE OF ADMINISTRATION, CONTAINER AND PACK SIZES

	Common DES 3.0 Context		Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/		Application > Medicinal Product >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E232-1	Strength and pharmaceutical form (use current list of standard terms - European Pharmacopoeia)			
E232-2	Pharmaceutical form(s)	rdm:medicinal-product/ rdm:pharmaceutical-products/ rdm:pharmaceutical-product/ rdm:pharmaceutical-form-names/ rdm:pharmaceutical-form-name	Pharmaceutical Product > Pharmaceuticla Dose Form CTL	
E232-3	Strength(s)	rdm:medicinal-product/ rdm:pharmaceutical-products/ rdm:pharmaceutical-product/rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/ rdm:pharmaceutical-form-strength-description		
E232-3a	Units	rdm:medicinal-product/ rdm:pharmaceutical-products/ rdm:pharmaceutical-product/rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:units		
E232-4	Active Substance(s)	rdm:medicinal-product/ rdm:pharmaceutical-products/ rdm:pharmaceutical-product/rdm:pharmaceutical-form-details/rdm:active-substances-description		
E232-5	Active Substance	rdm:medicinal-product/ rdm:pharmaceutical-products/ rdm:pharmaceutical-product/rdm:pharmaceutical-form-details/rdm:active-substances/rdm:active-substances-details/rdm:active-substance/rdm:substance-name	Ingredient > Substance CTL, Ingredient > IngredientRole CTL	
E232-6	Route(s) of administration (use current list of standard terms - European Pharmacopoeia)			
E232-7	Route of Administration	rdm:medicinal-product/maa:route-of-administrations/rdm:route-of-administration/rdm:name	Pharmaceutical Product > Route of Administration CTL	
E232-7a	Target species	rdm:medicinal-product/maa:route-of-administrations/rdm:route-of-administration/rdm:target_species/rdm:species	Species CTL	
E232-8	Container, closure and administration device(s), including description of material from which it is constructed. (use current list of standard terms - European Pharmacopoeia)	rdm:medicinal-product/maa:packages/rdm:package/rdm:package-description	Package > Package Description	
E232-9	For each type of pack give:			
E232-10	Package size	rdm:medicinal-product/ maa:packages/rdm:package/ rdm:package-size/rdm:size	Package > package Size	
E232-11	<i>Note: For mutual recognition and decentralised procedures, all package sizes authorised in the Reference Member State should be listed</i>			
E232-12	Description	rdm:medicinal-product/ maa:packages/rdm:package/rdm:package-description		
E232-13	Container	rdm:medicinal-product/maa:packages/rdm:package/rdm:containers/rdm:container/rdm:container-details/rdm:container-type	Package > Outer Container > Container CTL	
E232-14	Material	rdm:medicinal-product/maa:packages/rdm:package/rdm:containers/rdm:container/rdm:container-details/ rdm:material	Package > Immediate Container > Container CTL	
E232-15	Closure	rdm:medicinal-product/maa:packages/rdm:package	Package > Immediate Container > Container	B232-1

		/rdm:containers/rdm:container/ rdm:container-details/rdm: closure- material	CTL	
E232-16	Administration Device	rdm:medicinal- product/maa:packages/rdm:package /rdm:containers/rdm:container/ rdm:container-details/ rdm:administrative-device	Package > Outer Container > Administration Device > Administration Device CTL	B232-1
E232-17	Proposed shelf life (before first opening container)	rdm:medicinal-product/ maa:packages/rdm:package/rdm:co ntainers/rdm:container/rdm:shelf- life/rdm:shelf-life-before-open	Shelf Life > Shelf Life Type CTL	
E232-18	Proposed shelf life (after first opening container)	rdm:medicinal-product/ maa:packages/rdm:package/rdm:co ntainers/rdm:container/rdm:shelf- life/rdm:shelf-life-after-open	Shelf Life > Shelf Life Type CTL	B232-1
E232-19	Proposed shelf life (after reconstitution or dilution)	rdm:medicinal-product/ maa:packages/rdm:package/rdm:co ntainers/rdm:container/rdm:shelf- life/rdm:shelf-life-after-dilution	Shelf Life > Shelf Life Type CTL	B232-1
E232-20	Proposed storage conditions	rdm:medicinal-product/ maa:packages/rdm:package/rdm:co ntainers/rdm:container/rdm:shelf- life/rdm:spf/rdm:special- precaution-for-storage	Precaution for Storage > Special Precaution for Storage CTL	B232-1
E232-21	Proposed storage conditions after first opening	rdm:medicinal-product/ maa:packages/rdm:package/rdm:co ntainers/rdm:container/rdm:shelf- life/rdm:spf/rdm:special- precaution-for-storage-after-open	Precaution for Storage > Special Precaution for Storage CTL	B232-1
E232-22	Attach a list of Mock-ups or Samples/specimens sent with the application, as appropriate (see Annex 5.17)	maa:annex5-17		B232-1
E232-23	units	rdm:medicinal-product/ maa:packages/rdm:package/rdm:co ntainers/rdm:container/rdm:shelf- life/rdm:shelf-life-before-open-units		
E232-24	units	rdm:medicinal-product/ maa:packages/rdm:package/rdm:co ntainers/rdm:container/rdm:shelf- life/rdm:shelf-life-after-open-units		
E232-25	units	rdm:medicinal-product/ maa:packages/rdm:package/rdm:co ntainers/rdm:container/rdm:shelf- life/rdm:shelf-life-after-dilution-units		

Element Tree Diagram

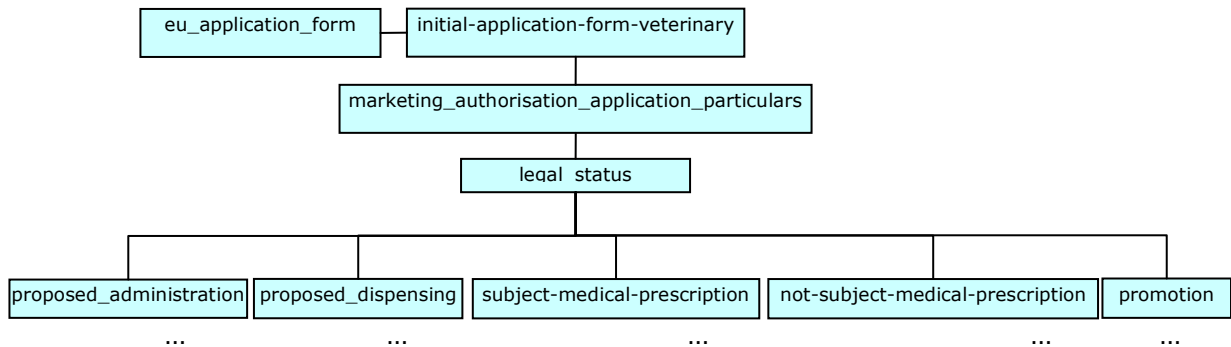


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B232-1	E232-11, E232-12, E232-17, E232-18, E232-20- E232-24	Optional.	These fields are optional.	

2.3.3. LEGAL STATUS

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:legal_status/		Application > Medicinal Product >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E233-1	Proposed administration	maa:proposed_administration		
E233-2	Proposed dispensing/classification	maa:proposed_dispensing		
E233-3	For veterinary products subject to medical prescriptions	maa:subject-medical-prescription		
E233-4	Supply for products <u>not</u> subject to medical prescription	maa:not-subject-medical-prescription		
E233-5	Promotion for product <u>not</u> subject to medical prescription	maa:promotion		

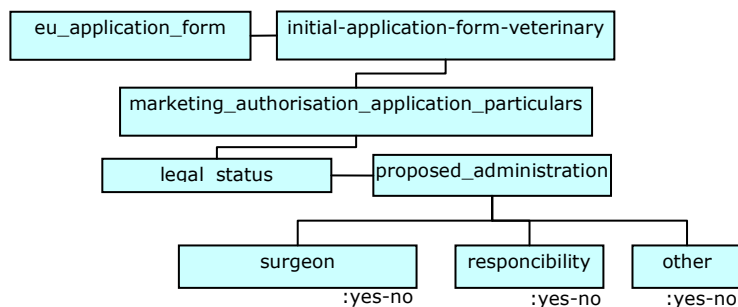
Element Tree Diagram



2.3.3.1. Proposed administration

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:legal_status/maa:proposed_administration/		Application > Medicinal Product	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2332-1	Only by a veterinary surgeon	maa:surgeon	Route of Administration CTL	
E2332-2	By a veterinary surgeon or under their direct responsibility	maa:responsibility	Route of Administration CTL	
E2332-3	Other	maa:other	Route of Administration CTL	

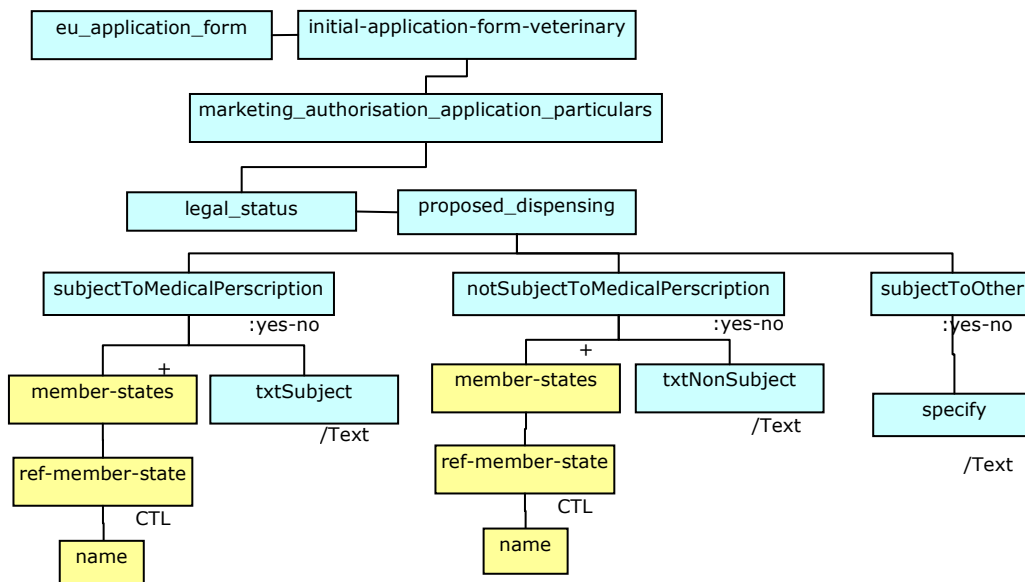
Element Tree Diagram



2.3.3.2. Proposed dispensing/classification

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:legal_status/maa:proposed_dispensing/		Application >Package Legal Status >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2332-1	Subject to medical prescription	maa:subjectToMedicalPerscription	Legal Status for the Supply CTL	
E2332-2	txtSubject	rdm:subject-package-size		
E2332-2	Member states	maa:subjectToMedicalPerscription-details/rdm:subject-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2332-3	Not subject to medical prescription	maa:notSubjectToMedicalPerscription	Legal Status for the Supply CTL	
E2332-2	txtNonSubject	rdm:non-subject-package-size		
E2332-4	Member states	maa:notSubjectToMedicalPerscription-details/maa:non-subject-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2332-5	Subject to other controls	maa:subjectToOther	Legal Status for the Supply CTL	
E2332-6	Specify	maa:specify	Legal Status for the Supply CTL	

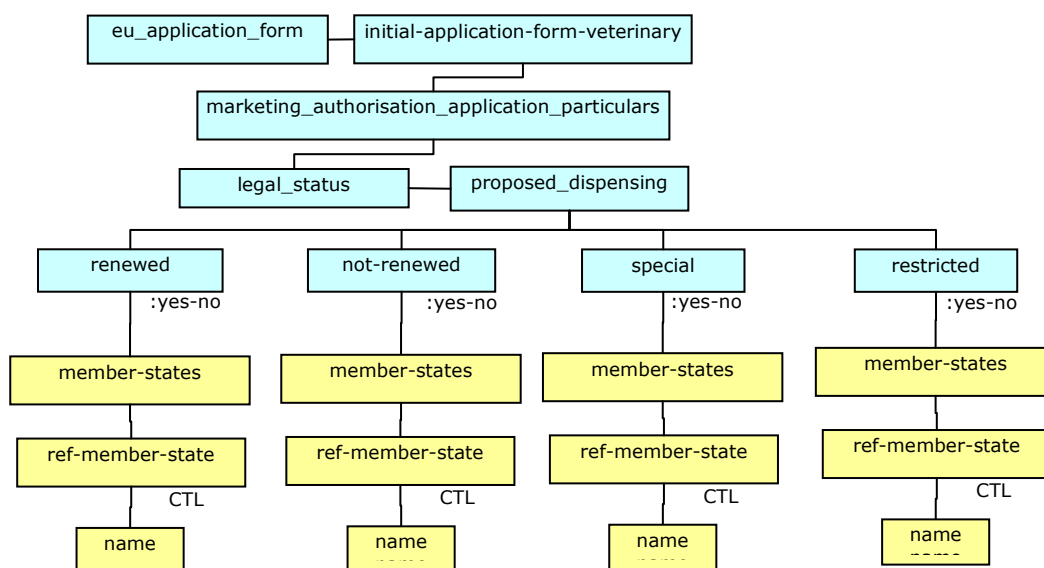
Element Tree Diagram



2.3.3.3. For veterinary products subject to medical prescriptions

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:legal_status/maa:subject-medical-prescription/		Application >Package Legal Status >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2333-1	Veterinary product on prescription which may be renewed (if applicable)	maa:renewed	Legal Status for the Supply CTL	
E2333-2	Member states	maa:renewed-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2333-3	Veterinary product on prescription which may not be renewed (if applicable)	maa:not-renewed	Legal Status for the Supply CTL	
E2333-4	Member states	maa:not-renewed-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2333-5	Veterinary product on special prescription	maa:special	Legal Status for the Supply CTL	
E2333-6	Member states	maa:special-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2333-7	Veterinary product on restricted prescription	maa:restricted	Legal Status for the Supply CTL	
E2333-8	Member states	maa:restricted-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	

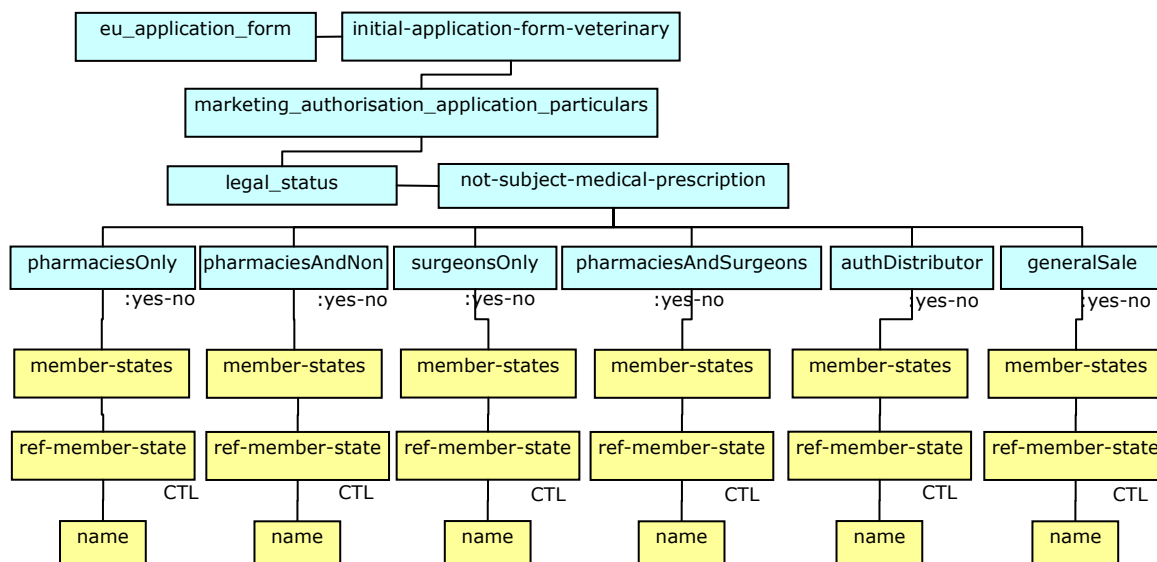
Element Tree Diagram



2.3.3.4. Supply for products not subject to medical prescription

		Common DES 3.0 Context	Common RDM Entry point	
		maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:legal_status/maa:not-subject-medical-prescription/	Application >Package Legal Status>	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2334-1	Supply through pharmacies only	maa:pharmaciesOnly	Legal Status for the Supply CTL	
E2334-2	Member states	maa:pharma-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2334-3	Supply through non-pharmacy outlets and pharmacies (if applicable)	maa:pharmaciesAndNon	Legal Status for the Supply CTL	
E2334-4	Member states	maa:non-harma-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2334-5	Supply/administration by veterinary surgeons only	maa:surgeonsOnly	Legal Status for the Supply CTL	
E2334-6	Member states	maa:admin-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2334-7	Supply by pharmacies and/or veterinary surgeons for animals under their care	maa:pharmaciesAndSurgeons	Legal Status for the Supply CTL	
E2334-8	Member states	maa:vet-care-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2334-9	Supply through authorised distributor	maa:authDistributor	Legal Status for the Supply CTL	
E2334-10	Member states	maa:distributor-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2334-11	General sale	maa:generalSale	Legal Status for the Supply CTL	
E2334-12	Member states	maa:generalSale-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	

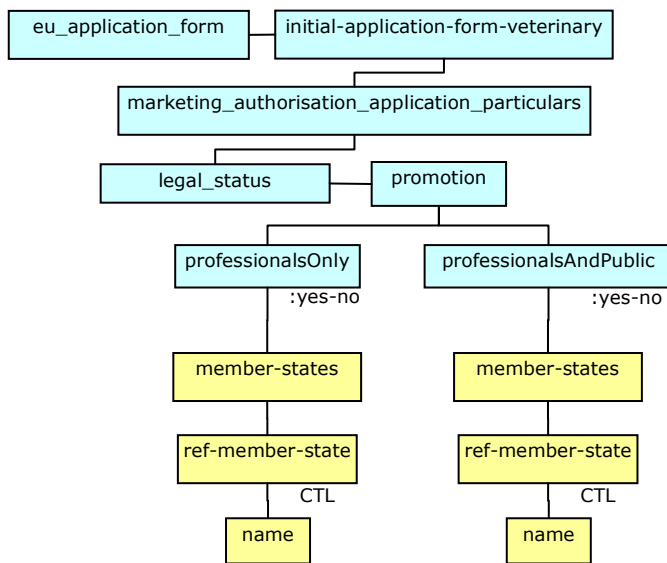
Element Tree Diagram



2.3.3.5. Promotion for product not subject to medical prescription

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:legal_status/maa:promotion/		Application > Package Legal Status>	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2335-1	Promotion to veterinary care professionals only	maa:professionalsOnly	Legal Status for the Supply CTL	
E2335-2	Member states	maa:promotion-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	
E2335-3	Promotion to the general public and veterinary professionals	maa:professionalsAndPublic	Legal Status for the Supply CTL	
E2335-4	Member states	maa:public-member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	

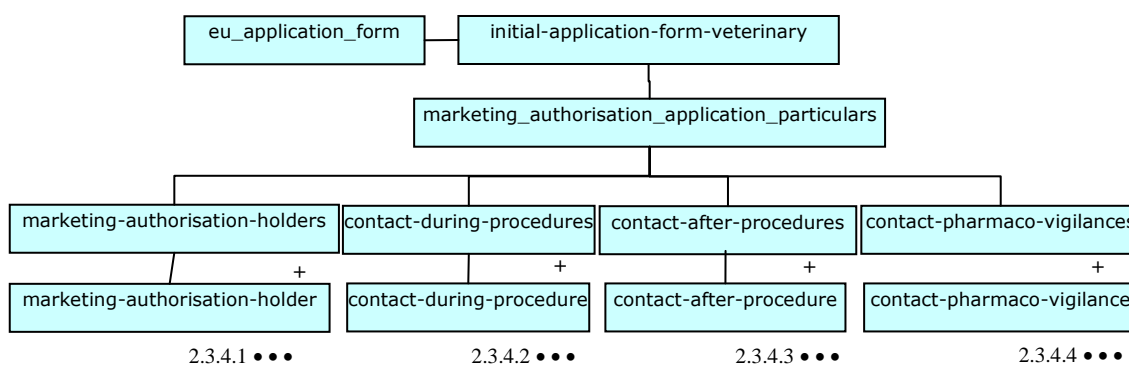
Element Tree Diagram



2.3.4. MARKETING AUTHORISATION HOLDER/ CONTACT PERSONS/ COMPANY

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:c:marketing_authorisation_application_particulars/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E234-1	Proposed marketing authorisation holder/person legally responsible for placing the product on the market in the European Union/each MS	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder		See Section 2.3.4.1
E234-2	Person/company authorised for communication on behalf of the applicant during the procedure in the European Union /each MS	maa:contact-during-procedures/maa:contact-during-procedure/		See Section 2.3.4.2
E234-3	Person/company authorised for communication between the marketing authorisation holder and the competent authorities after authorisation if different from 2.4.2 in European Union /each MS	maa:contact-after-procedures/maa:contact-after-procedure/		See Section 2.3.4.3
E234-4	Qualified person in the EEA for Pharmacovigilance	maa:contact-pharmaco-vigilances/maa:contact-pharmaco-vigilance/		See Section 2.3.4.4

Element Tree Diagram

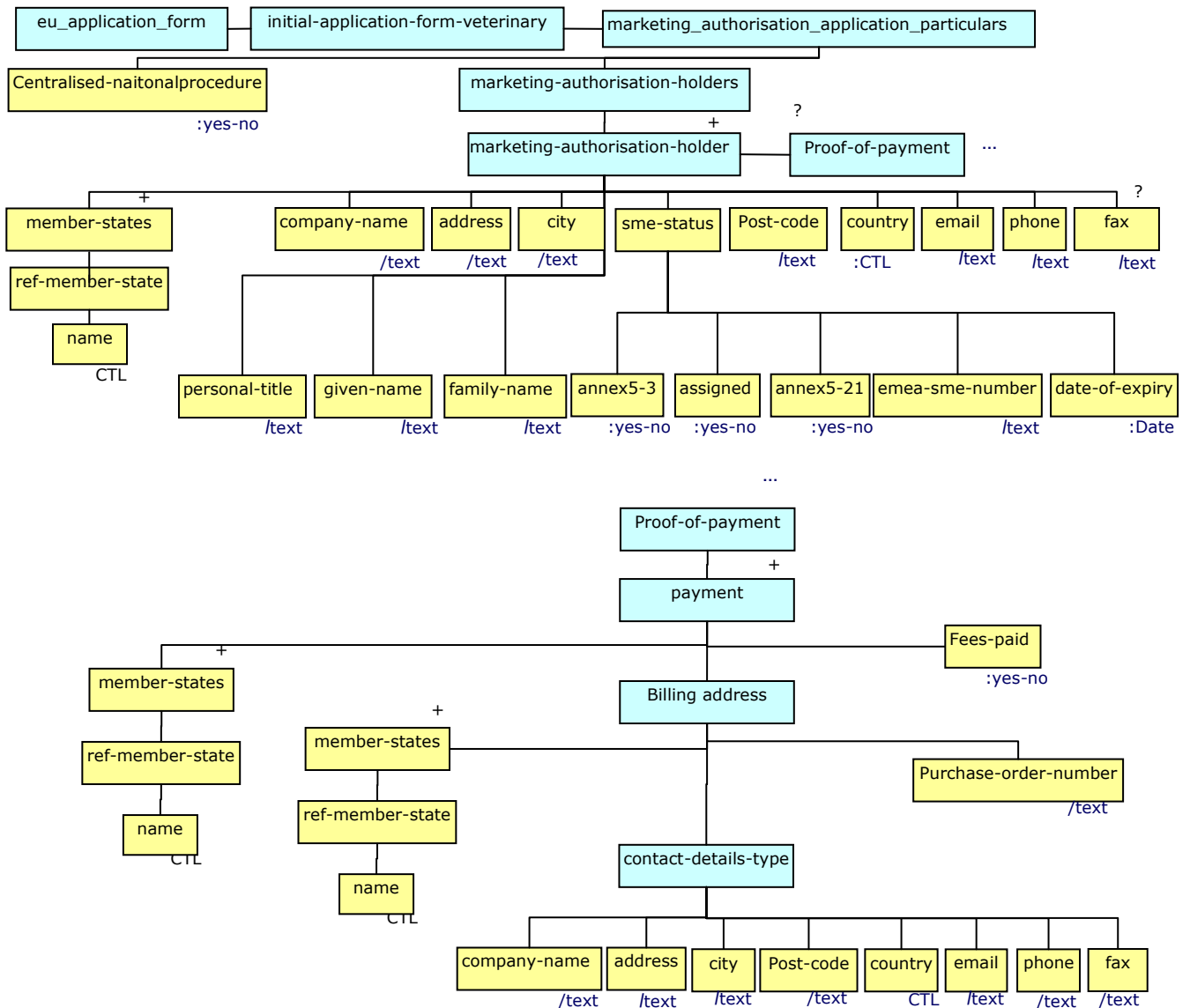


2.3.4.1. Proposed marketing authorisation holder/person legally responsible for placing the product on the market in the Community/each MS

		Common DES 3.0 Context	Common RDM Entry point	
		maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/	Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2341-1	National procedure	maa:centralised-nationalprocedure(value=1)	Procedure Type CTL (Value="national")	B2341-3, B2342-1, B2343-2, B2344-1
E2341-2	Centralized procedure	maa:centralised-nationalprocedure(value=2)	Procedure Type CTL (Value="centralised")	B2341-3, B2342-1, B2343-2, B2344-1
E2341-3	Member states	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	B2341-3
E2341-4	Company name	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:company-name	MAH for Placing Product> Role>Party>Organisation>Name	
E2341-5	Address1	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:address	MAH for Placing Product>Role>Party>Contact Details > Address	
E2341-6	Address2	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:city	MAH for Placing Product>Role>Party>Contact Details > city	
E2341-7	Postcode	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:post-code	MAH for Placing Product>Role>Party>Contact Details > Address> post code	
E2341-8	Country	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:country	MAH for Placing Product>Role>Party>Contact Details > Address > Country CTL	
E2341-9	Telephone	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:phone	MAH for Placing Product>Role>Party>Contact Details > Electronic Contact > electronic contact	
E2341-10	Telefax	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:fax	MAH for Placing Product>Role>Party>Contact Details > Electronic Contact > electronic contact	
E2341-11	E-mail	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:email	MAH for Placing Product>Role>Party>Contact Details > Electronic Contact > electronic contact	
E2341-12	Contact person at this address			B2341-4
E2341-13	Title	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:personal-title	MAH for Placing Product>Role>Party>Person> Personal Title	B2341-4
E2341-14	First name	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:given-name	MAH for Placing Product>Role>Party>Person> given name	B2341-4
E2341-15	Surname	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:family-name	MAH for Placing Product>Role>Party>Person> family name	B2341-4
E2341-16	Attach proof of establishment of the applicant in the EEA (Annex 5.3)	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:sme-status/rdm:annex5-3		
E2341-17	Has SME status been assigned by the EMEA?			
E2341-18	Yes	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:sme-status/rdm:assigned	MAH for Placing Product> has sme assigned to emea	B2341-1, B2341-2
E2341-19	No	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:sme-status/rdm:assigned	MAH for Placing Product> has sme assigned to emea	B2341-1
E2341-20	EMEA-SME Number	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:sme-status/rdm:emea-sme-number	Party Identification>identification number	B2341-2
E2341-21	Date of expiry	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:sme-status/rdm:date-of-expiry	Party Identification>expiry date	B2341-2
E2341-22	Attach copy of the "Qualification of SME Status" (Annex 5.21)	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/rdm:sme-status/rdm:annex5-21		

E2341-23	Proof of payment (when relevant)			
E2341-24	Have all relevant fees been prepaid to competent authorities?			
E2341-25	Yes	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/maa:payment /maa:fees-paid (1)		B2341-5
E2341-26	No	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment / maa:fees-paid (0)		B2341-5
E2341-27	For Member States	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment / member-states/ref-member-state/name		B2341-5
E2341-28	Billing address (when relevant)			
E2341-28a	For Member States	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment /maa: billing-details / member-states/ref-member-state/name		B2341-5
E2341-29	Company Name	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment /maa: billing-details/rdm:contact-details-type/company-name		B2341-5
E2341-30	VAT Number	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment / maa:billing-details/maa:VAT-number		B2341-5
E2341-31	Address1	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment / maa:billing-details/rdm:contact-details-type/rdm:address	Role>Party>Contact Details > Address	B2341-5
E2341-32	Address2	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment / maa:billing-details/rdm:contact-details-type/rdm:city	Role>Party>Contact Details > city	B2341-5
E2341-33	Postcode	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment / maa:billing-details/rdm:contact-details-type/rdm:postcode	Role>Party>Contact Details > postcode	B2341-5
E2341-34	Country	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment / maa:billing-details/rdm:contact-details-type/rdm:country	Role>Party>Contact Details > Address > Country CTL	B2341-5
E2341-35	Telephone	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment / maa:billing-details/rdm:contact-details-type/rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	B2341-5
E2341-36	Telefax	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment / maa:billing-details/rdm:contact-details-type/rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	B2341-5
E2341-37	E-mail	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment / maa:billing-details/rdm:contact-details-type/rdm:email	Role>Party>Contact Details> Electronic Contact > electronic contact	B2341-5
E2341-38	Purchase Order Number	maa:marketing-authorisation-holders/maa:marketing-authorisation-holder/maa:Proof-of-payment/ maa:payment / maa: billing-details/Purchase-order-number		B2341-5

Element Tree Diagram

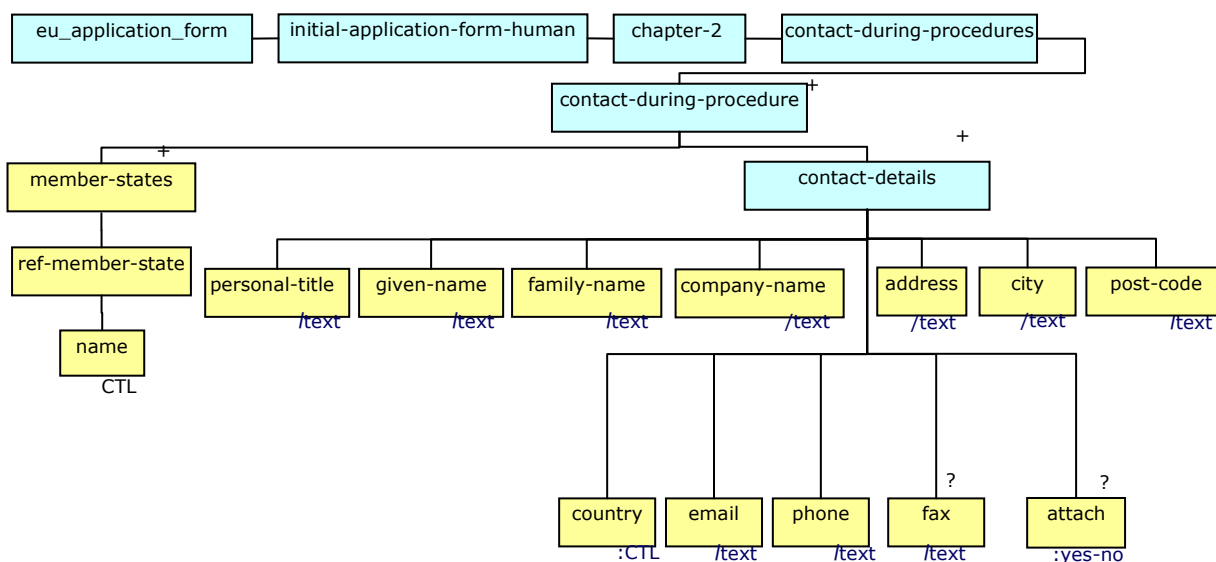


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2341-1	E2341-18, E2341-19	Mandatory.	Mutually Exclusive.	
B2341-2	E2341-18, E2341-20 to E2341-22	Mandatory.	If E2341-18 is selected, then the rest are mandatory, else they are hidden.	
B2341-3	E2341-1, E2341-2, E2341-3	hidden	If E2341-2 is selected then E2341-3 is visible If E2341-1 is selected then E2341-3 is invisible	
B2341-4	E2341-1, E2341-2, E2341-12, E2341-15,	hidden	If E2341-2 is selected then E2341-12, ..., E2341-15 is visible If E2341-1 is selected then E2341-12, E2341-15 is invisible	
B2341-5	E2341-27 to E2341-38	hidden	If E2341-25 is selected, then E2341-27 to E2341-38 are hidden If E2341-26 is selected, then E2341-27 to E2341-38 are invisible	

2.3.4.2. Person/company authorised for communication on behalf of the applicant during the procedure in the European Union/each MS

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:contact-during-procedures/maa:contact-during-procedure/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2342-1	Member states	rdm:member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	B2342-1
E2342-2	Title	rdm:contact-details/rdm:personal-title	Role>Party>Person> Personal Title	
E2342-3	First name	rdm:contact-details/rdm:given-name	Role>Party>Person> given name	
E2342-4	Surname	rdm:contact-details/rdm:family-name	Role>Party>Person> family name	
E2342-5	Company name	rdm:contact-details/rdm:company-name	Role>Party>Organisation>Name	
E2342-6	Address1	rdm:contact-details/rdm:address	Role>Party>Contact Details > Address	
E2342-7	Address2	rdm:contact-details/rdm:city	Role>Party>Contact Details > city	
E2342-8	Postcode	rdm:contact-details/rdm:post-code	Role>Party>Contact Details > postcode	
E2342-9	Country	rdm:contact-details/rdm:country	Role>Party>Contact Details > Address > Country CTL	
E2342-10	Telephone	rdm:contact-details/rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	
E2342-11	Telefax	rdm:contact-details/rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	
E2342-12	E-mail	rdm:contact-details/rdm:email	Role>Party>Contact Details> Electronic Contact > electronic contact	
E2342-13	If different to 2.4.1 above, Attach letter of authorisation (Annex 5.4)	rdm:contact-details/rdm:attach		

Element Tree Diagram

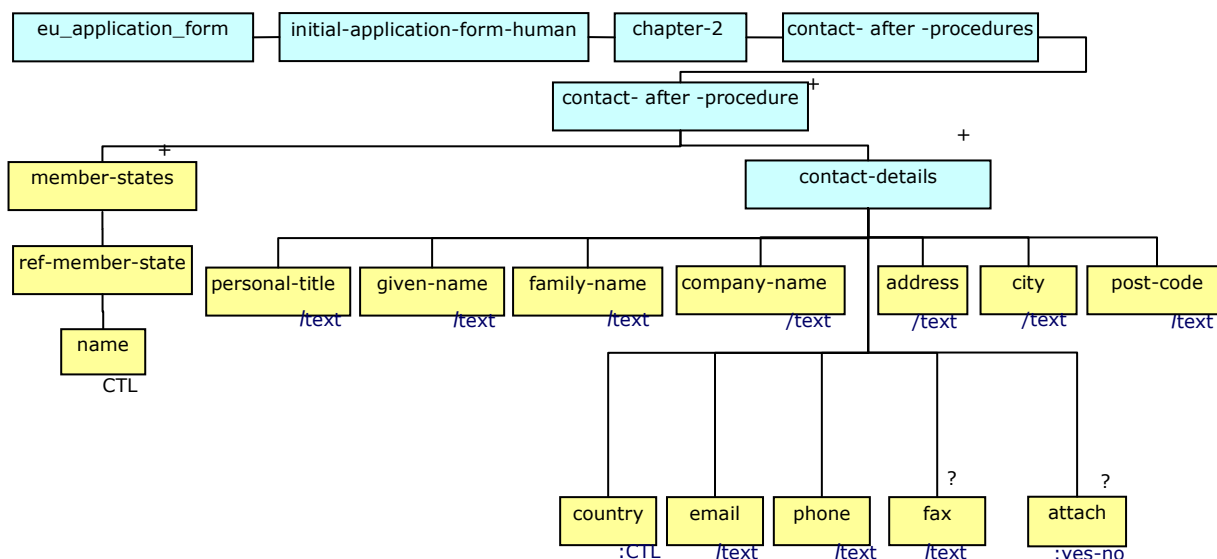


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2342-1	E2341-1	hidden	If E2341-2 is selected then E2342-1 is visible If E2341-1 is selected then E2342-1 is invisible	

2.3.4.3. Person/company authorised for communication between the marketing authorisation holder and the competent authorities after authorisation if different from 2.4.2 in European Union/each MS

	Common DES 3.0 Context		Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/ maa: marketing_authorisation_application_particulars/maa:contact-after-procedures/maa:contact-after-procedure/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2343-1	Member states	rdm:member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	B2343-2
E2343-2	Title	rdm:contact-details/rdm:personal-title	Role>Party>Person> Personal Title	B2343-1
E2343-3	First name	rdm:contact-details/rdm:given-name	Role>Party>Person> given name	B2343-1
E2343-4	Surname	rdm:contact-details/rdm:family-name	Role>Party>Person> family name	B2343-1
E2343-5	Company name	rdm:contact-details/rdm:company-name	Role>Party>Organisation>Name	B2343-1
E2343-6	Address1	rdm:contact-details/rdm:address	Role>Party>Contact Details > Address	B2343-1
E2343-7	Address2	rdm:contact-details/rdm:city	Role>Party>Contact Details > city	B2343-1
E2343-8	Postcode	rdm:contact-details/rdm:post-code	Role>Party>Contact Details > Address> post code	B2343-1
E2343-9	Country	rdm:contact-details/rdm:country	Role>Party>Contact Details > Address > Country CTL	B2343-1
E2343-10	Telephone	rdm:contact-details/rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	B2343-1
E2343-11	Telefax	rdm:contact-details/rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	B2343-1
E2343-12	E-mail	rdm:contact-details/rdm:email	Role>Party>Contact Details> Electronic Contact > electronic contact	B2343-1
E2343-13	If different to 2.4.1 above, attach a letter of authorisation (Annex 5.4)	rdm:contact-details/rdm:attach		B2343-1

Element Tree Diagram

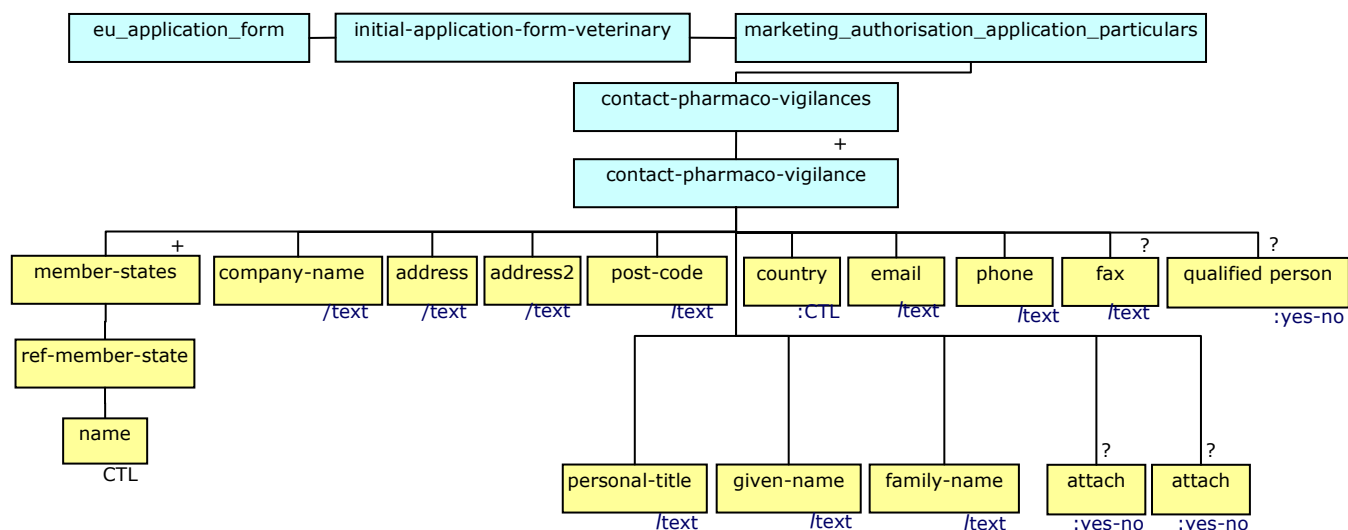


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2343-1	E2343-1 to E2343-13	Optional.	Elements should be optional.	
B2343-2	E2343-1	hidden	If E2341-2 is selected then E2343-1 is visible If E2341-1 is selected then E2343-1 is invisible	

2.3.4.4. Qualified person in the EEA for Pharmacovigilance

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa: initial-application-form-veterinary /maa:marketing_authorisation_application_particulars/maa:contact-pharmaco-vigilances/maa:contact-pharmaco-vigilance/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2344-1	Member states	rdm:member-states/rdm:ref-member-state/rdm:name	Procedure Use > Role > Country CTL	B2344-1
E2344-2	Title	rdm:personal-title	Role>Party>Person> Personal Title	
E2344-3	First name	rdm:given-name	Role>Party>Person> given name	
E2344-4	Surname	rdm:family-name	Role>Party>Person> family name	
E2344-5	Company name	rdm:company-name	Role>Party>Organisation>Name	
E2344-6	Address1	rdm:address	Role>Party>Contact Details > Address	
E2344-7	Address2	rdm:city	Role>Party>Contact Details > city	
E2344-8	Postcode	rdm:post-code	Role>Party>Contact Details > Address> post code	
E2344-9	Country	rdm:country	Role>Party>Contact Details > Address > Country CTL	
E2344-10	Telephone	rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	
E2344-11	Telefax	rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	
E2344-12	E-mail	rdm:email	Role>Party>Contact Details> Electronic Contact > electronic contact	
E2344-12	The above-mentioned qualified person resides ⁶ in the EEA	rdm:residesInEEA		
E2344-13	The qualified person is registered with Eudravigilance	rdm:qualifiedPerson		
E2344-14	Attach detailed description of the pharmacovigilance system (Annex 5.20)	rdm:attach		

Element Tree Diagram

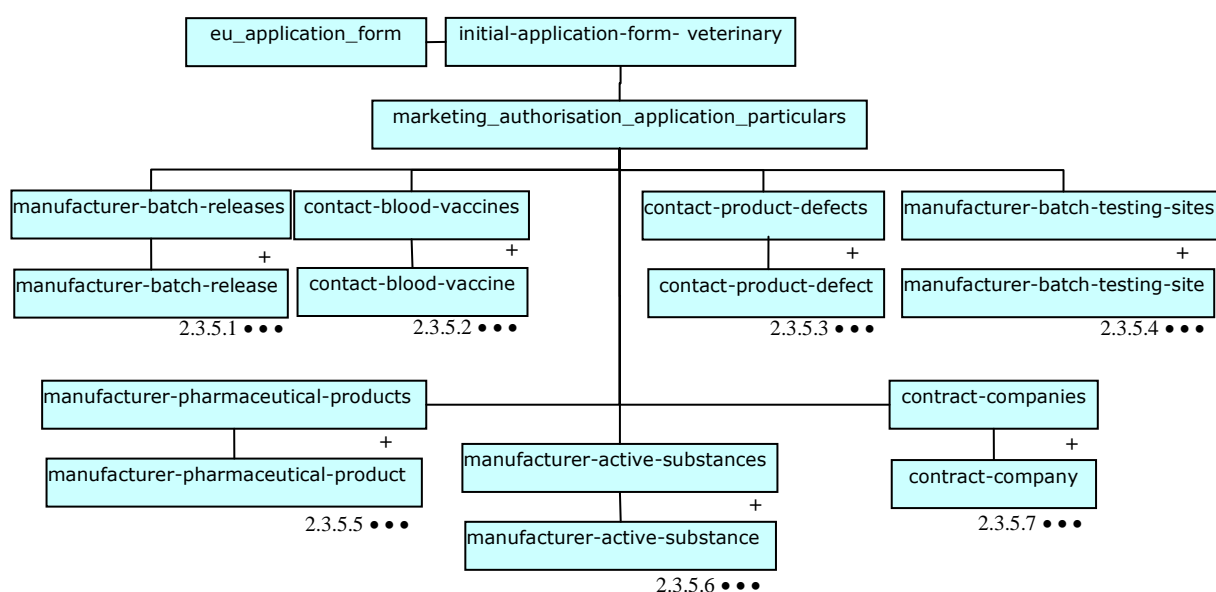


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2344-1	E2343-1	hidden	If E2341-2 is selected then E2344-1 is visible If E2341-1 is selected then E2344-1 is hidden	

2.3.5. MANUFACTURERS

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary /maa:marketing_authorisation_application_particulars/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E235-1	Authorised manufacturer(s) (or importer) responsible for batch release in the EEA in accordance with Article 55 and Article 53 of Directive 2001/82/EC (as shown in the package leaflet and where applicable in the labelling or Annex II of the Commission Decision)	maa:manufacturer-batch-releases/maa:manufacturer-batch-release/		See Section 2.3.5.1
E235-2	<u>For Vaccines:</u> Details of the state laboratory or laboratory designated for that purpose (OMCL) where the official batch protocol review (Article 81 of Directive 2001/82/EC) or the official control authority batch release Article 82 of Directive 2001/82/EC) takes place.	maa:contact-blood-vaccines/maa:contact-blood-vaccine/		See Section 2.3.5.2
E235-3	Contact person in the EEA for product defects and recalls	maa:contact-product-defects/maa:contact-product-defect		See Section 2.3.5.3
E235-4	Batch control/Testing arrangements. Sites in EEA or in countries where an MRA or other Community arrangements apply where batch control/testing takes place (if different from 2.5.1) as required by Article 55 of Directive 2001/82/EC	maa:manufacturer-batch-testing-sites/maa:manufacturer-batch-testing-site/		See Section 2.3.5.4
E235-5	Manufacturer(s) of the veterinary medicinal product and site(s) of manufacture: <i>(Note: including manufacturing sites of any diluent/solvent presented in a separate container but forming part of the veterinary medicinal product)</i>	maa:manufacturer-pharmaceutical-products/maa:manufacturer-pharmaceutical-product/		See Section 2.3.5.5
E235-6	Manufacturer(s) of the active substance(s) and site(s) of manufacture <i>Note: All manufacturing sites involved the manufacturing process of each source of active substance, including quality control/ in-process testing sites, should be listed. Broker or supplier details alone are not acceptable. For biotech products include all sites of storage of master and working cell bank and preparation of working cell banks.</i>	maa:manufacturer-active-substances/maa:manufacturer-active-substance		See Section 2.3.5.6
E235-7	Contract companies used for clinical trial(s) on bioavailability or bioequivalence or used for the validation of blood product manufacturing processes. For each contract company, state where analytical tests are performed and where clinical data are collected and give:	maa:contract-companies/maa:contract-study		See Section 2.3.5.7

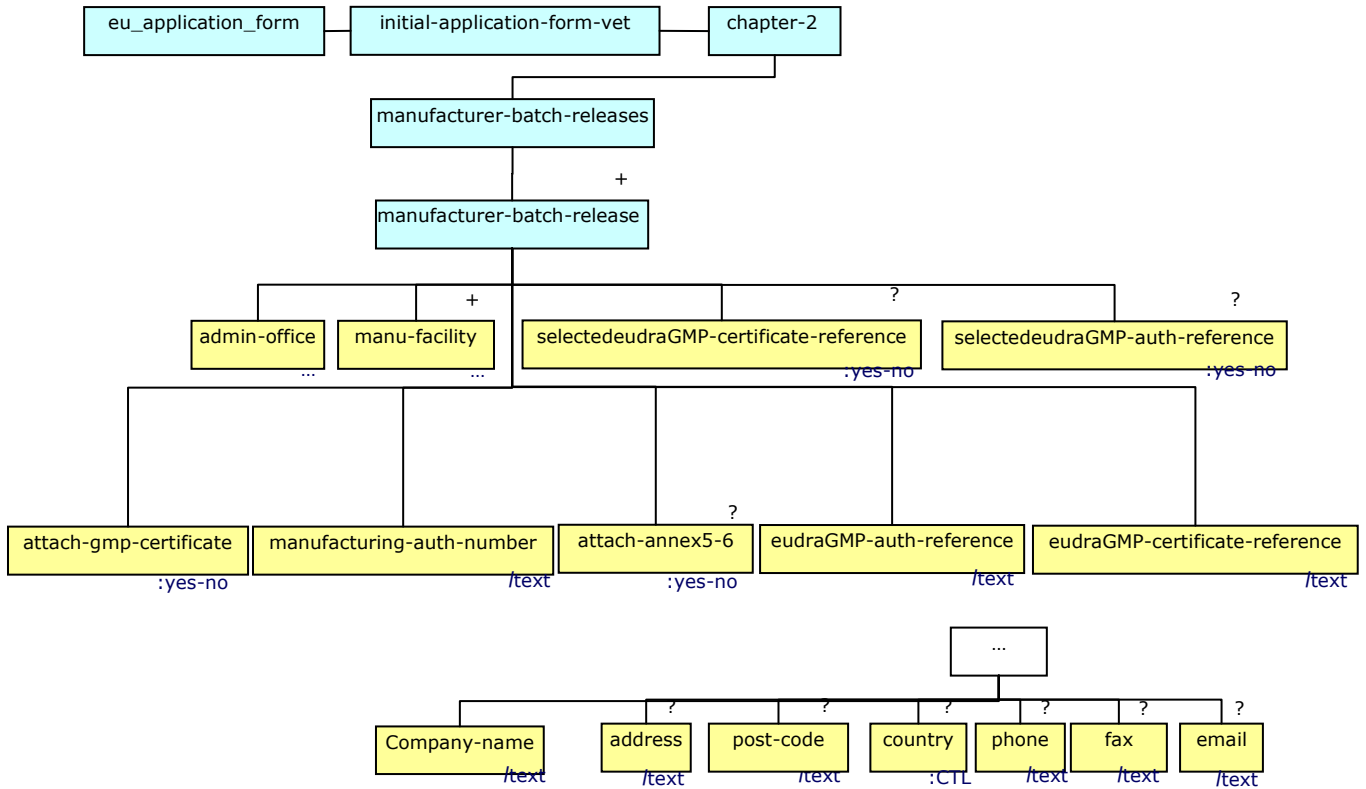
Element Tree Diagram



2.3.5.1. Authorised manufacturer(s) (or importer) responsible for batch release in the EEA in accordance with Article 55 and Article 53 of Directive 2001/82/EC (as shown in the package leaflet and where applicable in the labelling or Annex II of the Commission Decision)

	Common DES 3.0 Context		Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:manufacturer-batch-releases/maa:manufacturer-batch-release/		Application > Manufacturer Batch Release	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2351-0	Company description	rdm:package-responsible-manufacturer		
E2351-1	Do you have admin address and manufacturer address	maa:contact-details/rdm:admin-manu-address		B2351-5
E2351-1a	Company name	rdm:/contact-details/ rdm:admin-office/ rdm:company-name	Role>Party>Organisation>Name	B2351-5
E2351-2	Admin Office Address 1	rdm:/contact-details/ rdm:admin-office/rdm:address	Role>Party>Contact Details > Address	B2351-5
E2351-2a	Admin Office Address 2	rdm:/contact-details/ rdm:admin-office/rdm:city	Role>Party>Contact Details > City	B2351-5
E2351-3	Postcode	rdm:/contact-details/ rdm:post-code	Role>Party>Contact Details > Address> post code	B2351-5
E2351-4	Admin Office Country	rdm:/contact-details/ rdm:admin-office/rdm:country	Role>Party>Contact Details > Address > Country CTL	B2351-5
E2351-5	Admin Office Telephone	rdm:/contact-details/ rdm:admin-office/rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	B2351-5
E2351-6	Admin Office Telefax	rdm:/contact-details/ rdm:admin-office/rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	B2351-5
E2351-7	Admin Office E-mail	rdm:/contact-details/ rdm:admin-office/rdm:email	Role>Party>Contact Details > Electronic Contact > electronic contact	B2351-5
E2351-7a	Company name	rdm:/contact-details/ rdm:manu-facility/ rdm:company-name	Role>Party>Organisation>Name	B2351-5
E2351-8	Manufacturing Facility Address 1	rdm:/contact-details/ rdm:manu-facility/rdm:address	Role>Party>Contact Details > address	B2351-5
E2351-8a	Manufacturing Facility Address 2	rdm:/contact-details/ rdm:manu-facility/rdm:city	Role>Party>Contact Details > city	B2351-5
E2351-8b	Post code	rdm:/contact-details/ rdm:manu-facility/rdm:postcode	Role>Party>Contact Details > Address> post code	B2351-5
E2351-9	Manufacturing Facility Country	rdm:/contact-details/ rdm:manu-facility/rdm:country	Role>Party>Contact Details > Address > Country CTL	B2351-5
E2351-10	Manufacturing Facility Telephone	rdm:/contact-details/ rdm:manu-facility/rdm:phone	Role>Party>Contact Details > Address > Country CTL	B2351-5
E2351-11	Manufacturing Facility Telefax	rdm:/contact-details/ rdm:manu-facility/rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	B2351-5
E2351-12	Manufacturing Facility E-mail	rdm:/contact-details/ rdm:manu-facility/rdm:email	Role>Party>Contact Details > Electronic Contact > electronic contact	B2351-5
E2351-13	Manufacturing Authorisation number	rdm:manufacturing-auth-number	Manufacturing auth number	
E2351-14	Attach copy of manufacturing authorisation(s) (Annex 5.6)	rdm:attach-annex5-6		B2351-1
E2351-15	Enter EudraGMP manufacturing authorisation reference	rdm:selectedeudraGMP-auth-reference		B2351-1, B2351-3
E2351-16	Enter EudraGMP manufacturing authorisation reference	rdm:eudraGMP-auth-reference	Eudragmp auth ref	B2351-3
E2351-17	Attach latest GMP certificate (Annex 5.9)	rdm:attach-gmp-certificate		B2351-2
E2351-18	Enter EudraGMP certificate reference number	rdm:selectedeudraGMP-certificate-reference		B2351-2, B2351-4
E2351-19	Enter EudraGMP certificate reference number	rdm:eudraGMP-certificate-reference	Eudragmp certificate ref no	B2351-4

Element Tree Diagram

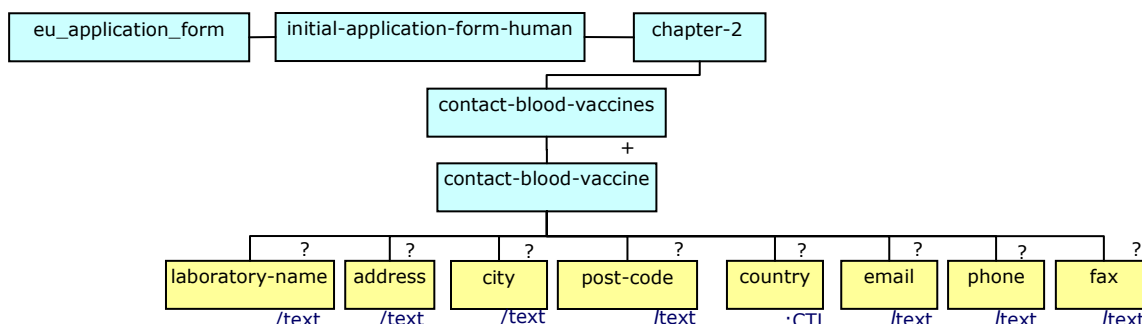


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2351-1	E2351-14, E2343-15	Mandatory.	Mutually Exclusive.	
B2351-2	E2351-17, E2351-18	Mandatory.	Mutually Exclusive.	
B2351-3	E2351-15, E2351-16	Mandatory, Optional.	If E2351-15 is selected, E2351-16 is mandatory.	
B2351-4	E2351-18, E2351-19	Mandatory, Optional.	If E2351-18 is selected, E2351-19 is mandatory.	
B2351-5	E2351-1 to E2351-12	mandatory and visible	E2351-1 is yes then E2351-1a to E2351-12 are visible else E2351-1a to E2351-7 are hidden	

2.3.5.1.1 Official batch release For Vaccines: Details of Official Medicines Control Laboratory (OMCL) or laboratory designated the purpose of official batch release (in accordance with Article 81 and 82 of Directive 2001/82/EC as amended)

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:contact-blood-vaccines/maa:contact-blood-vaccine/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E23511-1	Laboratory name	rdm:laboratory-name	Role>Party>Organisation>Name	B23511-1
E23511-2	Address1	rdm:address	Role>Party>Contact Details > Address	B23511-1
E23511-3	Address2	rdm:city	Role>Party>Contact Details > city	
E23511-4	Postcode	rdm:post-code	Role>Party>Contact Details > Address > post code	
E23511-5	Country	rdm:country	Role>Party>Contact Details > Address > Country CTL	B23511-1
E23511-6	Telephone	rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	B23511-1
E23511-7	Telefax	rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	B23511-1
E23511-8	E-mail	rdm:email	Role>Party>Contact Details> Electronic Contact > electronic contact	B23511-1

Element Tree Diagram

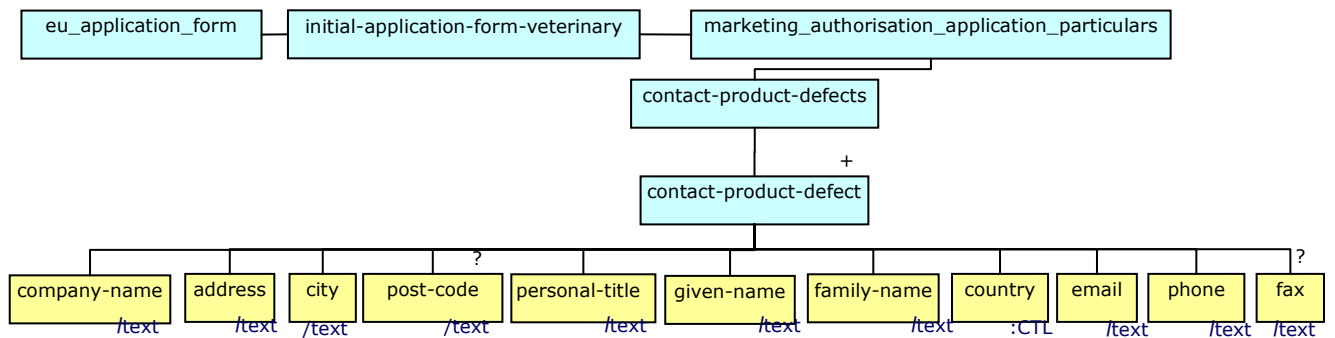


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B23511-1	E23511-1 to E23511-8	Optional	Elements should be optional	

2.3.5.2. Contact person in the EEA for product defects and recalls

Common DES 3.0 Context			Common RDM Entry point	
maa:eu_application_form/ maa:initial-application-form- veterinary/maa:marketing_authorisation_applicati on_particulars/maa:contact-product-defects/ maa:contact-product-defect/			Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2353-1	Company name	rdm:company-name	Role>Party>Organisation>Name	
E2353-2	Title	rdm:personal-title	Role>Party>Person> Personal Title	
E2353-3	First name	rdm:given-name	Role>Party>Person> given name	
E2353-4	Surname	rdm:family-name	Role>Party>Person> family name	
E2353-5	Address1	rdm:address	Role>Party>Contact Details > Address	
E2353-6	Address2	rdm:city	Role>Party>Contact Details > city	
E2353-7	Postcode	rdm:post-code	Role>Party>Contact Details > Address> post code	
E2353-8	Country	rdm:country	Role>Party>Contact Details > Address > Country CTL	
E2353-9	24H Telephone	rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	
E2353-10	Telefax	rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	
E2353-11	E-mail	rdm:email	Role>Party>Contact Details> Electronic Contact > electronic contact	

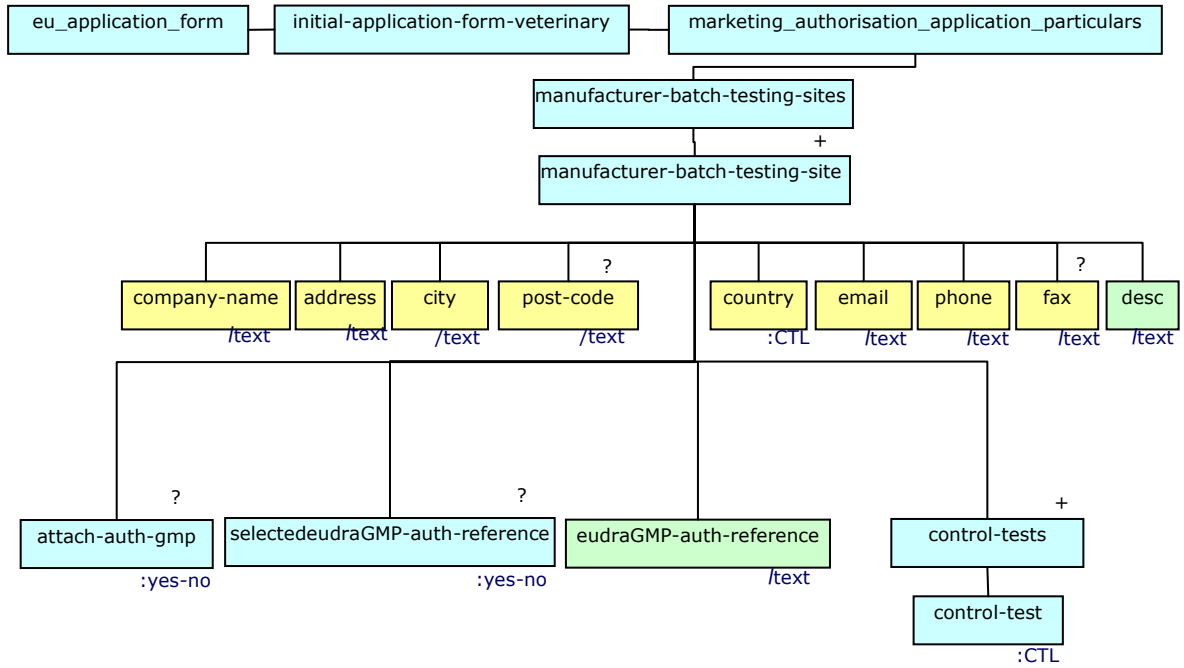
Element Tree Diagram



2.3.5.2.1 Batch control/Testing arrangements Sites in EEA or in countries where an MRA or other European Union arrangements apply where batch control/testing takes place as required by Article 55 of Directive 2001/82/EC.

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/ maa:marketing_authorisation_application_particulars/ maa:manufacturer-batch-testing-sites/ maa:manufacturer-batch-testing-site/	Application > Batch Control Arrangement >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2353-1	Company name	rdm:company-name	Role>Party>Organisation>Name	
E2353-2	Address1	rdm:address	Role>Party>Contact Details > Address	
E2353-3	Address2	rdm:city	Role>Party>Contact Details > city	
E2353-4				
E2353-5	Country	rdm:country	Role>Party>Contact Details > Address > Country CTL	
E2353-6	Telephone	rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	
E2353-7	Telefax	rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	
E2353-8	E-mail	rdm:email	Role>Party>Contact Details> Electronic Contact > electronic contact	
E2353-9	control-test	rdm:control-tests/ rdm:control-test		
E2353-10	Brief description of control tests carried out by the laboratory(ies) concerned. (note: please see the 'Compilation of Union Procedures on Inspections and Exchange of Information' document, (see pages - Interpretation of the Union Format for Manufacturer/Importer Authorisation): http://www.ema.europa.eu/docs/en_GB/document_library/Regulatory_and_procedural_guideline/2009/10/WC500004706.pdf Site(s)	rdm:desc	control test carried	
E2353-11	Attach copy of manufacturing authorisation(s) or other proof of GMP compliance (Annex 5.6)	rdm:attach-auth-gmp		B2353-1
E2353-12	Enter EudraGMP manufacturing authorisation reference	rdm:selectedeudraGMP-auth-reference		B2353-1, B2353 B2353-2
E2353-13	Enter EudraGMP manufacturing authorisation reference	rdm:eudraGMP-auth-reference	Batch Control Arrangement > eudragmp auth ref	B2354-2

Element Tree Diagram



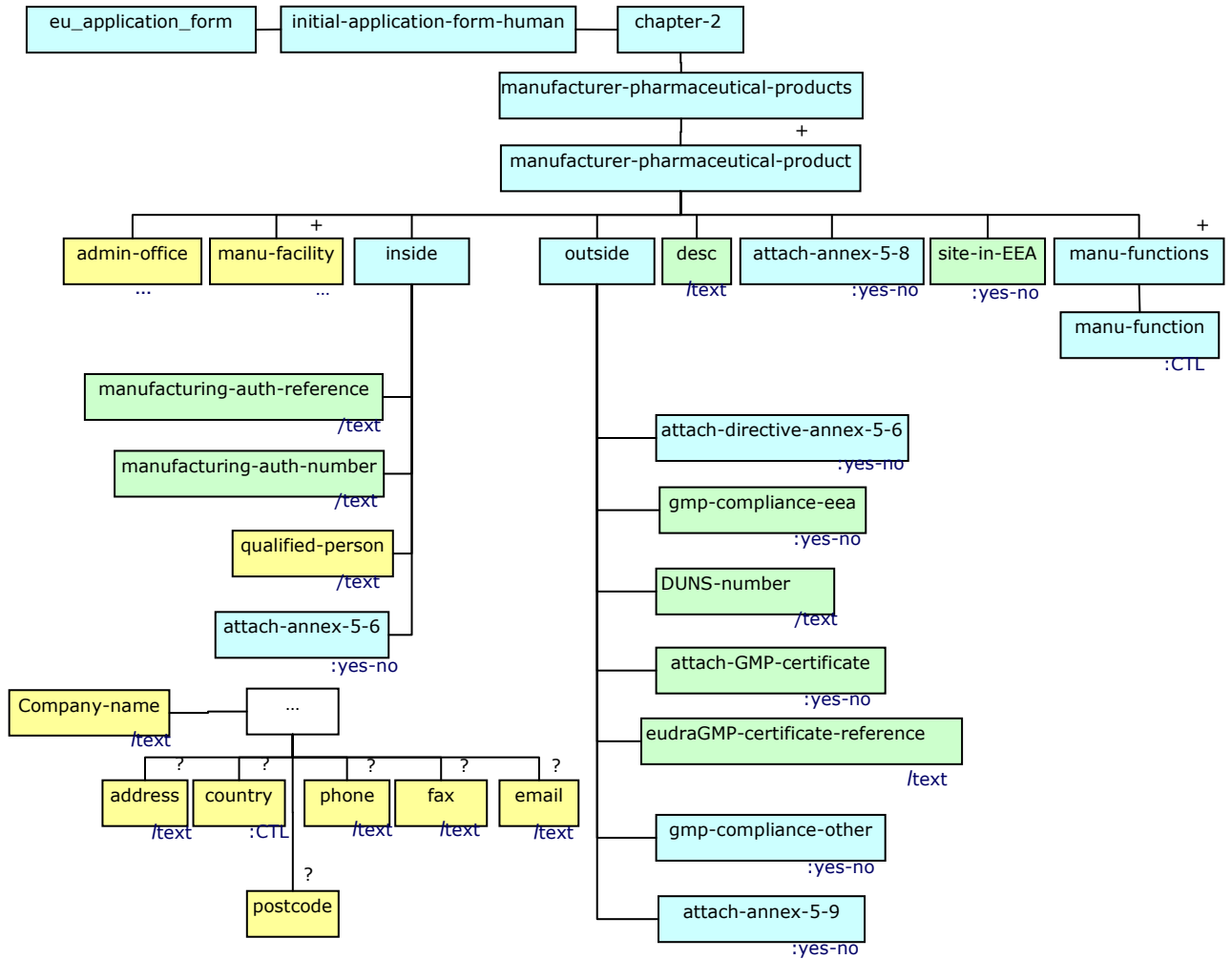
Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2354-1	E2352-10, E2352-11	Mandatory.	Elements are mutually exclusive.	
B2354-2	E2352-11, E2352-12	E2352-9 is Mandatory, E2352-10 is Optional.	If E2352-11 is selected then E2352-12 is mandatory.	

2.3.5.3. Manufacturer(s) of the veterinary medicinal product and site(s) of manufacture:
(Note: including manufacturing sites of any diluent/solvent presented in a separate container but forming part of the veterinary medicinal product)

	Common DES 3.0 Context		Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/ maa:marketing_authorisation_application_particulars/maa:manufacturer-pharmaceutical-products/maa:manufacturer-pharmaceutical-product/		Application > Manufacturer MP >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2355-1	Do you have admin address and manufacturer address	maa:contact-details/rdm:admin-manu-address		B2355-12
E2355-1a	Company name	rdm:/contact-details/ rdm:admin-office/ rdm:company-name	Role>Party>Organisation>Name	B2355-12
E2355-2	Admin Office Address	rdm:/contact-details/ rdm:admin-office/rdm:address	Role>Party>Contact Details > Address	B2355-12
E2355-2a	Admin Office Address 2	rdm:/contact-details/ rdm:admin-office/rdm:city	Role>Party>Contact Details > City	B2355-12
E2355-3	Postcode	rdm:/contact-details/ rdm:post-code	Role>Party>Contact Details > Address> post code	B2355-12
E2356-4	Admin Office Country	rdm:/contact-details/ rdm:admin-office/rdm:country	Role>Party>Contact Details > Address > Country CTL	B2355-12
E2356-5	Admin Office Telephone	rdm:/contact-details/ rdm:admin-office/rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	B2355-12
E2356-6	Admin Office Telefax	rdm:/contact-details/ rdm:admin-office/rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	B2355-12
E2356-7	Admin Office E-mail	rdm:/contact-details/ rdm:admin-office/rdm:email	Role>Party>Contact Details > Electronic Contact > electronic contact	B2355-12
E2355-7a	Company name	rdm:/contact-details/ rdm:manu-facility/ rdm:company-name	Role>Party>Organisation>Name	B2355-12
E2356-8	Manufacturing Facility Address	rdm:/contact-details/ rdm:manu-facility/ rdm:address	Role>Party>Contact Details > Address	B2355-12
E2356-8a	Manufacturing Facility Address 2	rdm:/contact-details/ rdm:manu-facility/rdm:city	Role>Party>Contact Details > city	B2355-12
E2356-8b	Postcode	rdm:/contact-details/ maa:role/maa:contact-details/rdm:manu-facility/rdm:postcode	Role> Party > Contact Details > postcode	B2355-12
E2356-9	Manufacturing Facility Country	rdm:/contact-details/ rdm:manu-facility/ rdm:country	Role>Party>Contact Details > Address > Country CTL	B2355-12
E2356-10	Manufacturing Facility Telephone	rdm:/contact-details/ rdm:manu-facility/rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	B2355-12
E2356-11	Manufacturing Facility Telefax	rdm:/contact-details/ rdm:manu-facility/rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	B2355-12
E2356-12	Manufacturing Facility E-mail	rdm:/contact-details/ rdm:manu-facility/rdm:email	Role>Party>Contact Details > Electronic Contact > electronic contact	B2355-12
E2355-13	Brief description of functions performed. (note: please see the 'Compilation of Union Procedures on Inspections and Exchange of Information' document, (see pages - Interpretation of the Union Format for Manufacturer/Importer Authorisation): http://www.ema.europa.eu/docs/en_GB/document_library/Regulatory_and_procedural_guideline/2009/10/WC500004706.pdf Site(s)	rdm:desc		
E2355-13a	Manufacturer functions	rdm:manu-functions/ rdm:manu-function		
E2355-14	Site is in the EEA:	rdm:site-in-EEA	is site in eea	B2355-1, B2355-2
E2355-15	Site is outside the EEA:	rdm:site-in-EEA	is site in eea	B2355-1, B2355-4, B2355-5
E2355-16	Manufacturing authorisation	rdm:inside/	manufacturing auth number	B2355-2,

	number	rdm:manufacturing-auth-number		B2355-3
E2355-17	Attach copy of manufacturing authorisation(s) (Annex 5.6)	rdm:inside/rdm:attach-annex-5-6		
E2355-18	Enter EudraGMP Manufacturing Authorisation reference	rdm:inside/rdm:manufacturing-auth-reference	eudragmp auth ref	B2355-2, B2355-3
E2355-19	Name of qualified person	rdm:inside/rdm:qualified-person	Role>Party>Person> given name	B2355-4
E2355-20	Attach document equivalent of manufacturing authorisation in accordance with Article 8(k) of Directive 2001/83/EC (Annex 5.6)	rdm:outside/rdm:attach-directive-annex-5-6		B2355-4
E2355-21	Has the site been inspected for GMP compliance by an EEA authority or by an authority of countries where MRA or other Community arrangements apply within the terms of the agreement?			B2355-4
E2355-22	D-U-N-S number	rdm:outside/DUNS-number		
E2355-23	Yes	rdm:outside/ rdm:gmp-compliance-eea	inspected by eea authority	B2355-4, B2355-5, B2355-6, B2355-7
E2355-24	No	rdm:outside/ rdm:gmp-compliance-eea	inspected by eea authority	B2355-4, B2355-5, B2355-6 B2355-9
E2355-25	Attach latest GMP certificate	rdm:outside/rdm:attach-GMP-certificate		B2355-4, B2355-7, B2355-8 B2355-9
E2355-26	Enter EudraGMP certificate reference number:	rdm:outside/ rdm:eudraGMP-certificate-reference	eudragmp certificate ref no	B2355-4, B2355-7, B2355-8, B2355-9
E2355-27	Has the site been inspected for GMP compliance by any other authority (including those of countries where MRA or other European Union arrangements apply but not within their respective territory)?			B2355-4,
E2355-28	Yes	rdm:outside/ rdm:gmp-compliance-other	inspected by other authority	B2355-4, B2355-5, B2355-9, B2355-10
E2355-29	No	rdm:outside/rdm:gmp-compliance-other	inspected by other authority	B2355-4, B2355-5, B2355-9
E2355-30	Please provide summary information in Annex 5.9 (and, if available a GMP certificate or a statement from the competent authority which carried out the inspection)	rdm:outside/rdm:attach-annex-5-9		B2355-4, B2355-10
E2355-31	Attach flow chart indicating the sequence and activities of the different sites involved in the manufacturing process, including testing sites (Annex 5.8)	rdm:attach-annex-5-8		

Element Tree Diagram



Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2355-1	E2355-14, E2355-15	Mandatory.	Mutually Exclusive.	
B2355-2	E2355-14, E2355-16 to E2355-18	Mandatory.	If E2355-14 is selected, then the other fields are required.	
B2355-3	E2355-16, E2355-18	Optional.	Mutually Exclusive.	
B2355-4	E2355-15, E2355-20 to E2355-30	invisible	If E2355-15 is selected, then the other fields are visible.	
B2355-5	E2355-15, E2355-23 to E2355-24, E2355-28 to E2355-29	Optional.	If E2355-15 is selected, then the other fields are mandatory.	
B2355-6	E2355-23, E2355-24	Mandatory.	Mutually Exclusive.	
B2355-7	E2355-23, E2355-25 to E2355-26	Optional.	If E2355-23 is selected, then the rest are required.	
B2355-8	E2355-25, E2355-26	Optional.	Mutually Exclusive.	
B2355-9	E2355-24, E2355-25 to E2355-26	Invisible	If E2355-24 is selected, then the rest of the fields are invisible	
B2355-10	E2355-28, E2355-30	Optional	If E2355-28 is selected, then E2355-30 is mandatory	
B2355-11	E2355-28, E2355-30	Invisible.	If E2355-29 is selected, then E2355-30 is visible.	
B2355-12	E2355-1 to E2355-12	mandatory and visible	E2355-1 is yes then E2355-1a to E2355-12 are visible else E2355-1a to E2355-7 are hidden	

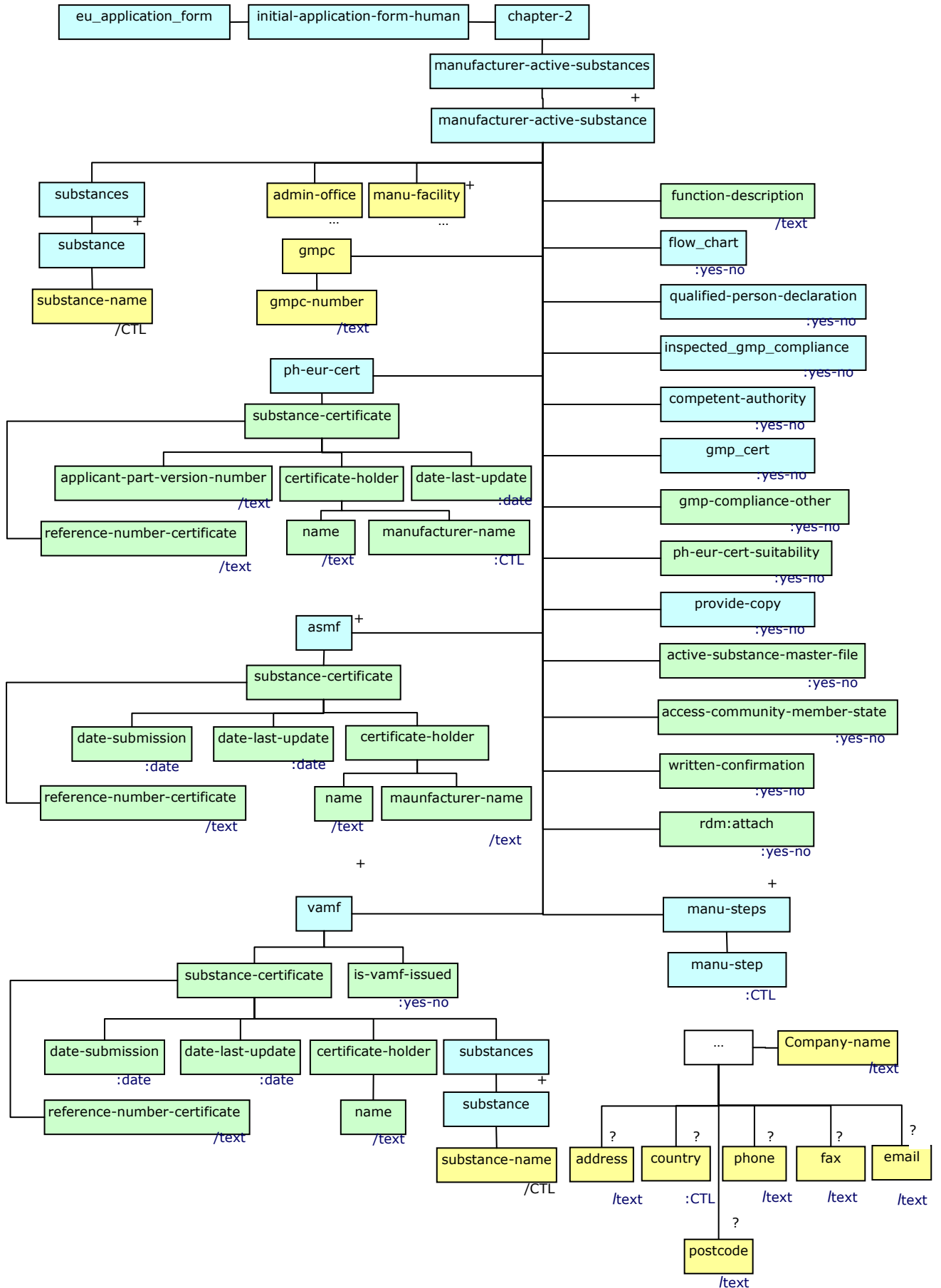
2.3.5.4. Manufacturer(s) of the active substance(s) and site(s) of manufacture *Note: All manufacturing sites involved the manufacturing process of each source of active substance, including quality control/ in-process testing sites, should be listed. Broker or supplier details alone are not acceptable. For biotech products include all sites of storage of master and working cell bank and preparation of working cell banks.*

	Common DES 3.0 Context		Common RDM Entry point	
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:manufacturer-active-substances/maa:manufacturer-active-substance-vet/		Application > Manufacturer Substance	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2356-1	Active Substance	rdm:active-substances/ rdm:active-substance/ rdm:substance-name	Medicinal Product > Pharmaceutical Product > Ingredient > Substance CTL > term id	
E2356-2	Do you have admin address and manufacturer address	maa:role/maa:contact- details/rdm:admin-manu- address		B2356-12
E2356-2a	Company Name	rdm:/contact-details/ rdm:company-name	Role > Party > Organisation > Name	B2356-12
E2356-3	Admin Office Address 1	rdm:/contact-details/ rdm:admin-office/ rdm:admin-office/rdm:address	Role>Party>Contact Details > Address	B2356-12
E2356-3a	Admin Office Address 2	rdm:/contact-details/ rdm:admin-office/rdm:city	Role>Party>Contact Details > city	B2356-12
E2356-3b	Postcode	maa:role/rdm:contact- details/rdm: admin- office/rdm:postcode	Role> Party > Contact Details > electronic contact	B2356-12
E2356-4	Admin Office Country	rdm:/contact-details/ rdm:admin-office/rdm:country	Role>Party>Contact Details > Address > Country CTL	B2356-12
E2356-5	Admin Office Telephone	rdm:/contact-details/ rdm:admin-office/rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	B2356-12
E2356-6	Admin Office Telefax	rdm:/contact-details/ rdm:admin-office/rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	B2356-12
E2356-7	Admin Office E-mail	rdm:/contact-details/ rdm:admin-office/rdm:email	Role>Party>Contact Details > Electronic Contact > electronic contact	B2356-12
E2356-7a	Company Name	rdm:/contact-details/ rdm:manu-facility/ rdm:company-name	Role > Party > Organisation > Name	B2356-12
E2356-8	Manufacturing Facility Address 1	rdm:/contact-details/ rdm:manu-facility/ rdm:address	Role>Party>Contact Details > Address	B2356-12
E2356-8a	Manufacturing Facility Address 2	rdm:/contact-details/ rdm:manu-facility/ rdm:city	Role>Party>Contact Details > City	B2356-12
E2356-8b	Postcode	rdm:/contact-details/ rdm:manu-facility/ rdm:postcode	Role>Party>Contact Details > Postcode	B2356-12
E2356-9	Manufacturing Facility Country	rdm:/contact-details/ rdm:manu-facility/ rdm:country	Role>Party>Contact Details > Address > Country CTL	B2356-12
E2356-10	Manufacturing Facility Telephone	rdm:/contact-details/ rdm:manu-facility/rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	B2356-12
E2356-11	Manufacturing Facility Telefax	rdm:/contact-details/ rdm:manu-facility/rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	B2356-12
E2356-12	Manufacturing Facility E-mail	rdm:/contact-details/ rdm:manu-facility/rdm:email	Role>Party>Contact Details > Electronic Contact > electronic contact	B2356-12
E2356-13	Brief description of manufacturing steps performed by manufacturing site: (note: please see the 'Compilation of Union Procedures on Inspections and Exchange of Information' document, (see pages - Interpretation of the Union Format for Manufacturer/Importer Authorisation): http://www.ema.europa.eu/docs/en_GB/document_library/Regulatory_and_procedural_guideline/2009/10/WC500004706.pdf	rdm:function-description	Manufacturing steps	

	Site(s)			
E2356-13a	Manufacturer steps	rdm:manu-steps/rdm: manu-step		
E2356-14	Attach flow-chart indicating the sequence and activities of the different sites involved in the manufacturing process, including batch control sites (Annex 5.8)	rdm:flow_chart		
E2356-15	For each active substance, attach a Qualified Person declaration that the active substance is manufactured in compliance with the principles and Guidelines on good manufacturing practice for starting materials (Annex 5.22)	rdm:qualified-person-declaration		
E2356-16	Has the site been inspected for GMP compliance by an EEA authority or by an authority of countries where MRA or other European Union arrangements apply within the terms of agreement?			
E2356-17	Yes	rdm:inspected_gmp_compliance (Value=1)	inspected by eea authority	B2356-1, B2356-2
E2356-18	No	rdm:inspected_gmp_compliance (Value=0)	inspected by eea authority	B2356-1
E2356-19				
E2356-20				
E2356-21	Attach latest GMP certificate in Annex 5.9	rdm:gmp_cert		B2356-2, B2356-3
E2356-22	EudraGMP certificate reference number	rdm:gmpc/rdm:gmpc-number	Eudragmp certificate ref no	B2356-2
E2356-23	Has the site been inspected for GMP compliance by any other authority (including those of countries where MRA or other Community arrangements apply but not within their respective territory)?			
E2356-24	Yes	rdm:gmp-compliance-other (Value=1)	inspected by other authority	B2356-4, B2356-5
E2356-25	No	rdm:gmp-compliance-other (Value=0)	inspected by other authority	B2356-4
E2356-26	If yes, please provide summary information in Annex 5.9 (and, if available a GMP certificate or a statement from the competent authority which carried out the inspection)	rdm:summary-information		B2356-5
E2356-27	Has a Ph.Eur. Certificate of suitability been issued for the active substance(s):			
E2356-28	Yes	rdm:ph-eur-cert-suitability (Value=1)	Ph Eur Certificate > has certificate issued	B2356-6, B2356-7
E2356-30	No	rdm:ph-eur-cert-suitability (Value=0)	Ph Eur Certificate > has certificate issued	B2356-6
E2356-31	Provide copy in Annex 5.10	rdm:provide-copy		
E2356-32	Name of the CEP holder	rdm:ph-eur-cert/rdm:substance-certificate/rdm:certificate-holder/rdm:name	Role > Party > Organisation> Name	B2356-7
E2356-33	Name of the manufacturer	rdm:ph-eur-cert/rdm:substance-certificate/rdm:certificate-holder/rdm:manufacturer-name	Role > Party > Organisation> Name	B2356-7
E2356-34	CEP number	rdm:ph-eur-cert/rdm:substance-certificate/rdm:reference-number-certificate	PH Eur Certificate > reference number	B2356-7
E2356-35	Date of last update	rdm:ph-eur-cert/rdm:substance-certificate/rdm:date-last-update	PH Eur Certificate > last update date	B2356-7
E2356-36	Is an Active Substance Master File (European Drug Master File) to be used for the active substance(s) reference/original?			
E2356-38	Yes	rdm:active-substance-master-file (Value=1)	European Drug Master File > used for active substance	B2356-8, B2356-9
E2356-39	No	rdm:active-substance-master-file (Value=0)	European Drug Master File > used for active substance	B2356-8
E2356-40	Name of the ASMF holder	rdm:asmf/rdm:substance-certificate/rdm:certificate-holder/rdm:name	European Drug Master File > Substance CTL	B2356-9
E2356-41	Name of the manufacturer	rdm:asmf/rdm:substance-certificate/rdm:certificate-holder/rdm:manufacturer-name	Role > Party > Organisation> Name	B2356-9
E2356-42	Reference number for EMEA/competent authority	rdm:asmf/rdm:substance-certificate/rdm:reference-number-certificate	European Drug Master File > Reference number	B2356-9
E2356-43	Applicant part version number	rdm:asmf/rdm:substance-certificate/rdm:Applicant-part-version-number	European Drug Master File > Version ?	B2356-9
E2356-44	Date of submission	rdm:asmf/rdm:substance-certificate/rdm:date-submission	European Drug Master File > Submission date	B2356-9
E2356-45	Date of last update	rdm:asmf/rdm:substance-	European Drug Master File >	B2356-9

E2356-46	Attach letter of access for Community/Member State authorities where the application is made (see "European ASMF procedure for active ingredients") (Annex 5.10)	certificate/rdm:date-last-update rdm:access-community-member-state	Last update date	B2356-9
E2356-47	Attach copy of written confirmation from the manufacturer of the active substance to inform the applicant in case of modification of the manufacturing process or specifications according to Annex 1 of Directive 2001/82/EC (Annex 5.11)	rdm:written-confirmation		B2356-9
E2356-48	Is an EMEA certificate for a Vaccine Antigen Master File (VAMF) issued or submitted in accordance with Directive 2001/83/EC Annex I, Part III, being used for this MAA?			
E2356-49	Yes	rdm:vamf/rdm:is-vamf-issued (Value=1)	Vaccine Antigen Master File>is certificate issued	B2356-10
E2356-50	No	rdm:vamf/rdm:is-vamf-issued (Value=0)	Vaccine Antigen Master File>is certificate issued	B2356-10
E2356-51	Active Substance	rdm:vamf/rdm:substance-certificate/rdm:active-substances/ rdm:active-substance/ rdm:substance-name	Vaccine Antigen Master File > Medicinal Product > Pharmaceutical Product > Ingredient > Substance CTL > term id	
E2356-52	Name of the VAMF Certificate Holder/VAMF Applicant	rdm:vamf/rdm:substance-certificate/rdm:certificate-holder/rdm:name	Role > Party > Organisation > Name	
E2356-53	Reference number of Application/Certificate	rdm:vamf/rdm:substance-certificate/rdm:reference-number-certificate	Vaccine Antigen Master File>reference Number	
E2356-54	Date of submission (if pending)	rdm:vamf/rdm:substance-certificate/rdm:date-submission	Vaccine Antigen Master File > submission date	
E2356-55	Date of approval or last update (if approved)	rdm:vamf/rdm:substance-certificate/rdm:date-last-update	Vaccine Antigen Master File > last update date	
E2356-56	Provide copy in (Annex 5.20)	rdm:attach		

Element Tree Diagram

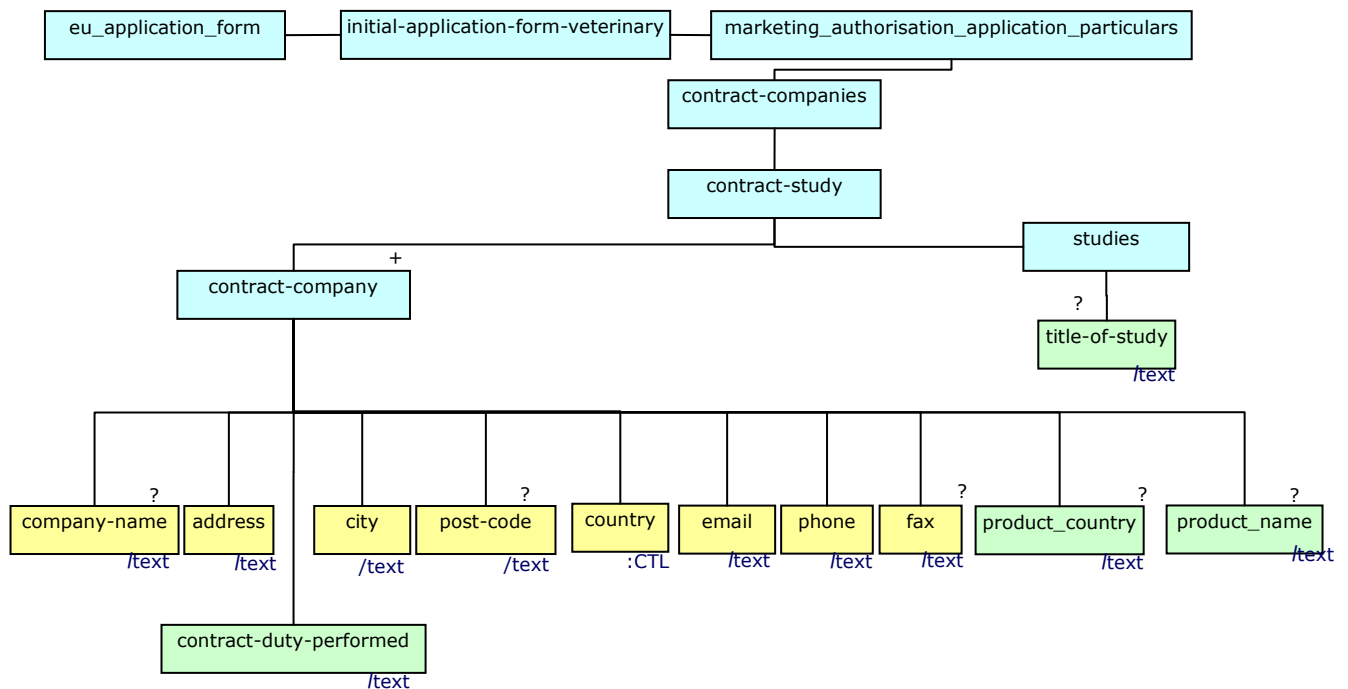


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2356-1	E2356-17, E2356-18	Mandatory.	Mutually Exclusive.	
B2356-2	E2356-17, E2356-20 to E2356-22	E2356-20 to E2356-22 are optional, E2356-17 is mandatory.	If E2356-17 is selected, then E2356-20 to E2356-22 are required.	
B2356-3	E2356-20, E2356-21	Optional.	Mutually Exclusive.	
B2356-4	E2356-24, E2356-25	Mandatory.	Mutually Exclusive.	
B2356-5	E2356-24, E2356-26	Mandatory.	If E2356-24 is selected, then E2356-26 is required.	
B2356-6	E2356-28, E2356-29	Mandatory.	Mutually Exclusive.	
B2356-7	E2356-28, E2356-32 to E2356-35	E2356-28 is Mandatory, Rest are optional.	If E2356-28 is selected, the rest are required.	
B2356-8	E2356-38, E2356-39	Mandatory.	Mutually Exclusive.	
B2356-9	E2356-38, E2356-40 to E2356-46	E2356-38 is Mandatory, Rest are optional.	If E2356-38 is selected, the rest are required.	
B2356-10	E2356-48, E2356-49	Mandatory.	Mutually Exclusive.	
B2356-11	E2356-48, E2356-50 to E2356-56	E2356-48 is Mandatory, Rest are optional.	If E2356-48 is selected, the rest are required.	
B2356-12	E2356-2 to E2356-12	mandatory and visible	E2356-2 is yes then E2356-2a to E2356-12 are visible else E2356-2a to E2356-7 are hidden	

2.3.5.5. Contract companies used for clinical trial(s), bioavailability or bioequivalence trials. For each contract company, state where analytical tests are performed and where efficacy data are collected and give

Common DES 3.0 Context			Common RDM Entry point	
maa:eu_application_form/maa:initial-application-form-veterinary/ maa:marketing_authorisation_application_particulars/maa:contract-companies/maa:contract-study/			Application > Contract Company CT >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2357-1	Company Name	rdm:contract-company/ rdm:company-name	Role > Party > Organisation > Name	
E2357-2	Address	rdm:contract-company/ rdm:address	Role>Party>Contact Details > Address	
E2357-3	Address2	rdm:contract-company/ rdm:city	Role>Party>Contact Details > Address > City	
E2357-4	PostCode	rdm:contract-company/ rdm:post-code	Role>Party>Contact Details > Address > post code	
E2357-5	Country	rdm:contract-company/ rdm:country	Role>Party>Contact Details > Address > Country CTL	
E2357-6	Telephone	rdm:contract-company/ rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	
E2357-7	Telefax	rdm:contract-company/ rdm:fax	Role>Party>Contact Details > Electronic Contact > electronic contact	
E2357-8	E-mail	rdm:contract-company/ rdm:email	Role>Party>Contact Details > Electronic Contact > electronic contact	
E2357-9	Duty performed according to contract	rdm:contract-company/ rdm:contract-duty-performed	Duty performed	
E2357-10	Name of product	rdm:contract-company/ rdm:product_name	Protocol code	
E2357-11	Country of product	rdm:contract-company/ rdm:product_country	Eudract number	
E2357-11	For each contract company, state where analytical tests are performed and where clinical data are collected and give:	rdm:studies/rdm:title-of-study	Study title	

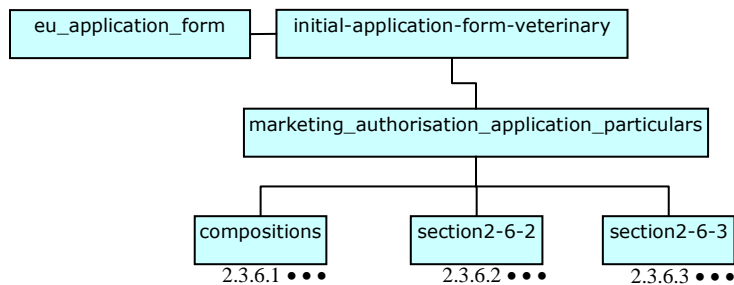
Element Tree Diagram



2.3.6. QUALITATIVE AND QUANTITATIVE COMPOSITION

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/ maa:marketing_authorisation_application_particulars/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E236-1	Qualitative and Quantitative composition in terms of the active substance(s) and the excipient(s)	maa:compositions/		See Section 2.3.6.1
E236-2	List of materials of animal origin contained or used in the manufacturing process of the veterinary medicinal product?	maa:section2-6-2/		See Section 2.3.6.2
E236-3	Does the veterinary medicinal product contain or consist of Genetically Modified Organisms(GMOs) within the meaning of Directive 2001/18/EC?	maa:section2-6-3		See Section 2.3.6.3

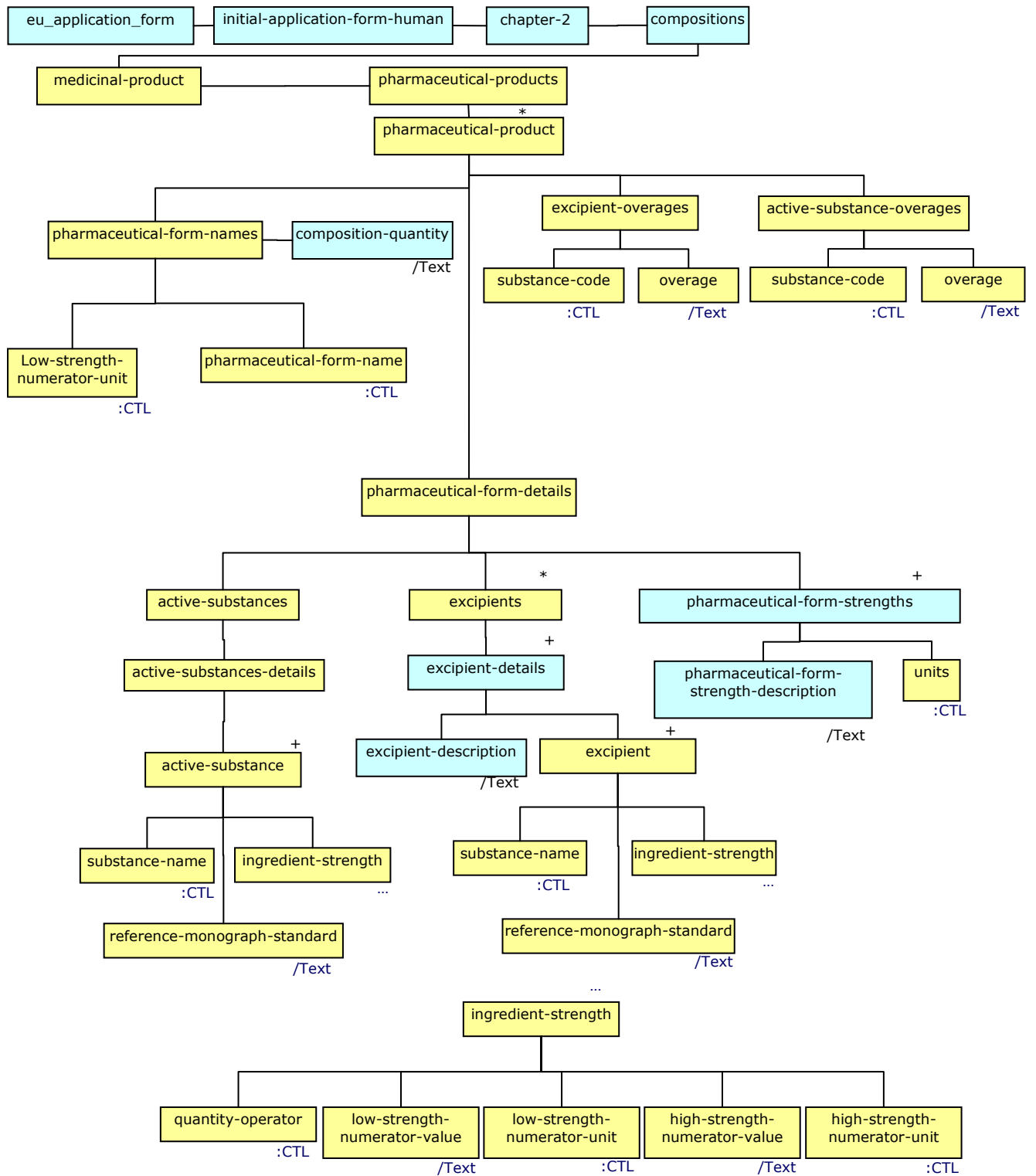
Element Tree Diagram



2.3.6.1. Qualitative and Quantitative composition in terms of the active substance(s) and the excipient(s)

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-human/maa:chapter-2/maa:compositions/maa:composition/rdm:medicinal-product/rdm:pharmaceutical-products/rdm:pharmaceutical-product/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2361-1	A note should be given as to which quantity the composition refers (e.g. 1 capsule)	rdm:pharmaceutical-form-names/rdm:composition-quantity		
E2361-2	composition-unit	rdm:pharmaceutical-form-names/rdm:Low-strength-numerator-unit		
E2361-3	Pharmaceutical Form	rdm:pharmaceutical-form-names/rdm:pharmaceutical-form-name/	Pharmaceutical Dose Form CTL > term id	
E2361-4	Strength	rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strength-description		
E2361-4a	Units	rdm:pharmaceutical-form-details /rdm:units		
E2361-5	Name of active substance	rdm:pharmaceutical-form-details/rdm:active-substances/ rdm:active-substances-details/rdm:active-substances-details/rdm:active-substance/rdm:substance-name	Ingredient > Substance CTL	
E2361-6	Quantity / Unit	rdm:pharmaceutical-form-details/rdm:active-substances/ rdm:active-substances-details/rdm:active-substance/rdm:ingredient-strength	Ingredient > Unit CTL	
E2361-7	Reference / Monograph Standard	rdm:pharmaceutical-form-details/rdm:active-substances/rdm:active-substances-details/rdm:active-substance/rdm:reference-monograph-standard	??	
E2361-8	Name of Excipient	rdm:pharmaceutical-form-details/rdm:excipients /rdm:excipients-details/rdm:excipient /rdm:substance-name	Ingredient > Substance CTL	
E2361-9	Quantity / Unit	rdm:pharmaceutical-form-details/rdm:excipients /rdm:excipients-details/rdm:excipient /rdm:ingredient-strength	Ingredient > Unit CTL	
E2361-10	Reference / Monograph Standard	rdm:pharmaceutical-form-details/rdm:excipients /rdm:excipients-details/rdm:excipient /rdm:reference-monograph-standard	??	
E2361-11	Active Substance	rdm:pharmaceutical-form-details/rdm:active-substance-overages/rdm:substance-code	Ingredient > Substance CTL	
E2361-12	Overage	rdm:active-substance-overages/rdm:overage	Ingredient > Overage	
E2361-13	Excipient	rdm:excipient-overages/rdm:excipient-code	Ingredient > Substance CTL	
E2361-14	Overage	rdm:excipient-overages /rdm:overage	Ingredient > Overage	

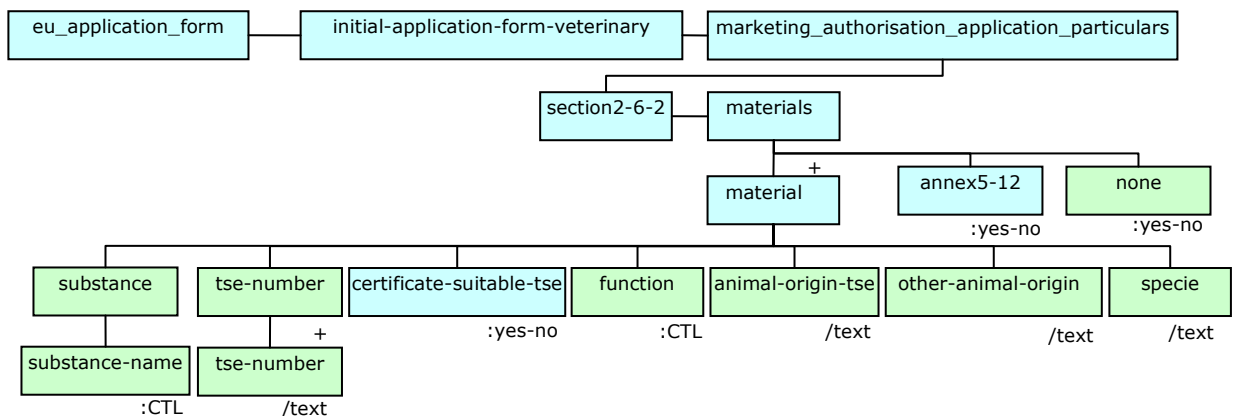
Element Tree Diagram



2.3.6.2. List of materials of animal origin contained or used in the manufacturing process of the veterinary medicinal product?

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:section2-6-2/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2362-1	NONE	maa:materials/maa:none	Manufacturing Material > is material used	B2362-1
E2362-2	Active Substance	maa:materials/maa:material/maa:substance/rdm:substance-name	Manufacturing Material > material name	B2362-1
E2362-3				
E2362-4	Active Substance	maa:materials/maa:material/function (Value=1)	Material Function CTL	B2362-1, B2362-2
E2362-5	Excipient (incl. starting materials used in manufacture of the active substance/excipient),	maa:materials/maa:material/function (Value=2)	Material Function CTL	B2362-1, B2362-2
E2362-6	Reagent / Culture Medium (incl. those used in the preparation of master and working cell banks)	maa:materials/maa:material/function (Value=3)	Material Function CTL	B2362-1, B2362-2
E2362-7	Animal Origin Susceptible to TSE as defined in section 2 (scope) of the Note for Guidance on minimising the risk of transmitting animal spongiform encephalopathy agents via human and veterinary medicinal products.	maa:materials/maa:material/animal-origin-tse	Material Origin CTL	B2362-1, B2362-3
E2362-8	Other Animal Origin	maa:materials/maa:material/other-animal-origin	Material Origin CTL	B2362-1, B2362-3
E2362-9	Species	maa:materials/maa:material/maa:specie		B2362-1, B2362-3
E2362-10	Certificate of suitability for TSE	maa:materials/maa:material/maa:certificate-suitable-tse		B2362-1, B2362-4
E2362-11	TSE number	maa:materials/maa:material/maa:tse-number	Manufacturing Material > certificate number	B2362-1, B2362-4
E2362-12	If a Ph. Eur. Certificate of suitability for TSE is available according to the Resolution AP/CSP(99)4 of the Council of Europe attach it in Annex 5.12	maa:materials/maa:annex5-12		

Element Tree Diagram

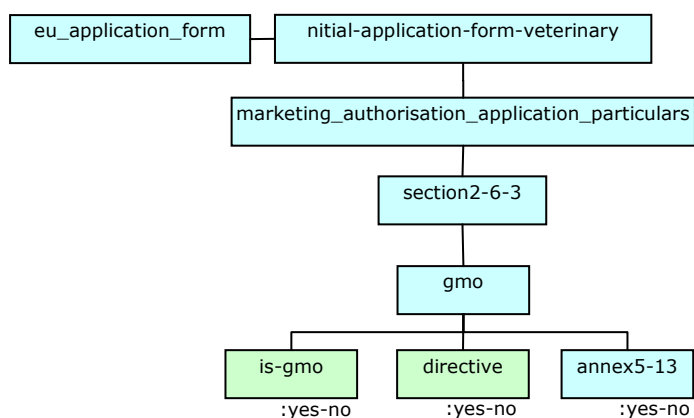


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B2362-1	E2362-1, E2362-2 to E2362-11	Mandatory.	If E2362-1 is selected then the rest are optional.	
B2362-2	E2362-4 to E2362-6	Optional.	Mutually Exclusive.	
B2362-3	E2362-7 to E2362-9	Optional.	Mutually Exclusive.	
B2362-4	E2362-10, E2362-11	Optional.	If E2362-10 is selected, E2362-11 is required.	

2.3.6.3. Does the veterinary medicinal product contain or consist of Genetically Modified Organisms (GMOs) within the meaning of Directive 2001/18/EC?

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:initial-application-form-veterinary/maa:marketing_authorisation_application_particulars/maa:section2-6-3/maa:gmo/	Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2364-1	Yes	maa:is-gmo (Value=1)	App GMO > mp consist of gmo	B2364-1
E2364-2	No	maa:is-gmo (Value=0)	App GMO > mp consist of gmo	B2364-1
E2364-3	If yes, does the product comply with Directive 2001/18/EC?			
E2364-4	Yes	maa:directive	App GMO > comply with directive	
E2364-5	No	maa:directive	App GMO > comply with directive	
E2364-6	Attach a copy of any written consent(s) of the competent authorities to the deliberate release into the environment of the GMOs for research and development purposes where provided for by Part B of the above-mentioned Directive (Annex 5.13)	maa:annex5-13		

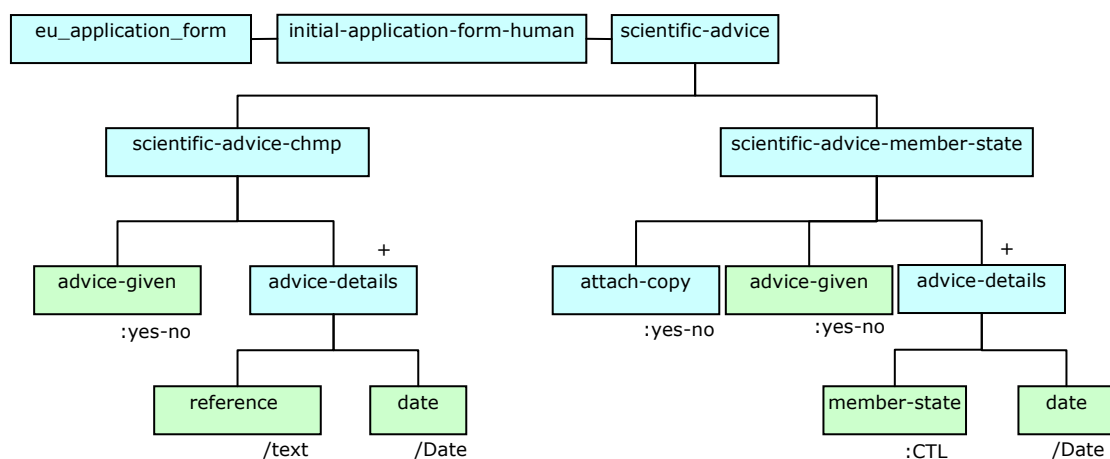
Element Tree Diagram



2.4. SCIENTIFIC ADVICE

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:initial-application-form-veterinary/maa:scientific-advice/		Application > App Scientific Advice >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E24-1	Was there formal scientific advice given by the CVMP for this veterinary medicinal product?			
E24-2	Yes	rdm:scientific-advice-chmp/rdm:advice-given	was scientific advice given	B24-1, B24-2
E24-3	No	rdm:scientific-advice-chmp/rdm:advice-given	was scientific advice given	B24-1
E24-4	Date	rdm:scientific-advice-chmp/rdm:advice-details/rdm:date	scientific advice date	B24-2
E24-5	Reference of the scientific advice letter	rdm:scientific-advice-chmp/rdm:advice-details/rdm:reference	scientific advice ref	B24-2
E24-6	Attach copy of scientific advice(s) (Annex 5.14)	rdm:scientific-advice-member-state/rdm:attach-copy		
E24-7	Was there scientific recommendation(s) given by Member State(s) for this veterinary medicinal product?			
E24-8	Yes	rdm:scientific-advice-member-state/rdm:advice-given	scientific Advice Source CTL	B24-3,B24-4
E24-9	No	rdm:scientific-advice-member-state/rdm:advice-given	scientific Advice Source CTL	B24-3
E24-10	Date	rdm:scientific-advice-member-state/rdm:advice-details/rdm:date	scientific advice date	B24-4
E24-11	Member State	rdm:scientific-advice-member-state/rdm:advice-details/rdm:member-state	Country CTL	B24-4

Element Tree Diagram

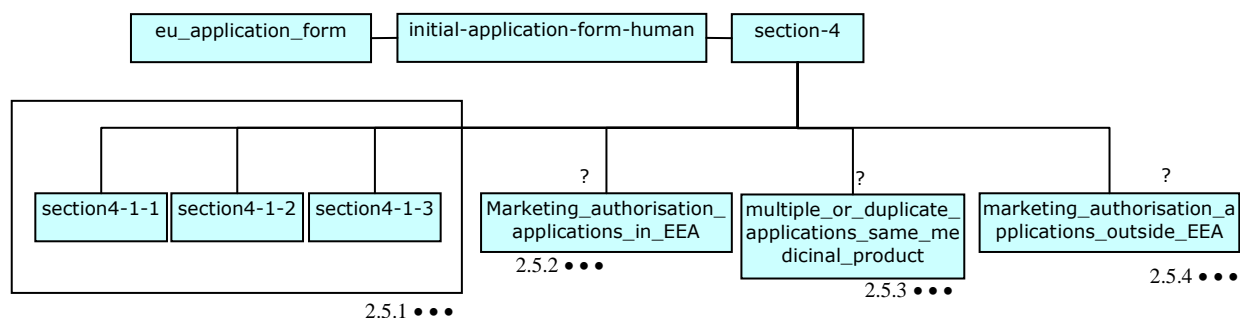


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B24-1	E24-2, E24-3	Mandatory	Mutually Exclusive	
B24-2	E24-2, E24-4, E24-5	Mandatory	If E24-2 is selected, then the others are required	
B24-3	E24-8, E24-9	Mandatory	Mutually Exclusive	
B24-4	E24-8, E24-10 to E24-11	E24-8 is Mandatory, rest are optional	If E24-8 is selected, then the other fields are required.	

2.5. OTHER MARKETING AUTHORISATION APPLICATIONS

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/ maa:initial-application-form-veterinary/ maa:other_marketing_authorisation_applications/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E25-1	FOR NATIONAL/MRP/DCP APPLICATIONS, PLEASE COMPLETE THE FOLLOWING IN ACCORDANCE WITH ARTICLE 8(j)-(l) OF DIRECTIVE 2001/82/EC	maa:section4-1-1 and maa:section4-1-2 and maa:section4-1-3		See Section 2.5.1 B25-1
E25-2	Marketing authorisation applications for the same product in the EEA (same qualitative and quantitative composition in active substance(s) and having the same pharmaceutical form from applicants belonging to the same mother company or group of companies OR which are "licensees").	maa:Marketing_authorisation_applications_in_EEA		See Section 2.5.2
E25-3	For multiple/duplicate applications of the same veterinary medicinal product:	maa:multiple_or_duplicate_applications_same_medicinal_product		See Section 2.5.3
E25-4	Marketing authorisation applications for the same product outside the EEA (i.e from applicants belonging to the same mother company or group of companies OR which are "licensees". Same qualitative and quantitative composition in active substance(s) and having the same pharmaceutical form).	maa:marketing_authorisation_applications_outside_EEA		See Section 2.5.4

Element Tree Diagram

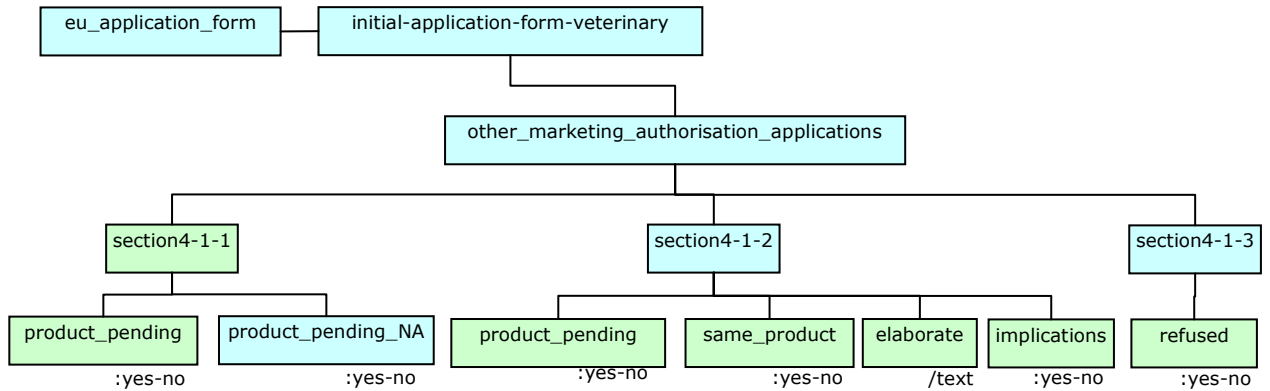


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B25-1	E25-1, E221-4	Optional	If E221-4 is selected, then E25-1 is required.	

2.5.1. FOR NATIONAL APPLICATIONS ONLY, PLEASE COMPLETE THE FOLLOWING IN ACCORDANCE WITH ARTICLE 12(1) OF DIRECTIVE 2001/82/EC

	Common DES 3.0 Context		Common RDM Entry point	
	maa:eu_application_form/ maa:initial-application-form-veterinary/ maa:other_marketing_authorisation_applications/		Application > Other MA Application	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E251-1	Is there another Member State(s) where an application for the same ⁴ product is pending?			
E251-2	Yes	maa:section4-1-1/maa:product_pending (Value=1)	app pending in other ms	B251-1, B252-1
E251-3	No	maa:section4-1-1/maa:product_pending (Value=0)	app pending in other ms	B251-1
E251-3a	Not applicable	maa:section4-1-1/maa:product_pending_NA		B251-1
E251-4	Is there another Member state(s) where an authorisation is granted for the same ⁴ product?			
E251-5	Yes	maa:section4-1-2/maa:same_product (Value=1)	is auth granted in other ms	B251-2, B251-3, B252-1
E251-6	No	maa:section4-1-2/maa:same_product (Value=0)	is auth granted in other ms	B251-2
E251-7	Are there any differences which have therapeutic implications between this application and the applications/authorisations for the same product in other Member States (for national applications, Article 21 or 22 of Directive 2001/82/EC shall apply)?			B251-3
E251-8	Yes	maa:section4-1-2/maa:implications (Value=1)	are there differences	B251-3, B251-4, B251-5
E251-9	No	maa:section4-1-2/maa:implications (Value=0)	are there differences	B251-3, B251-4
E251-10	If yes, please elaborate:	maa:section4-1-2/maa:elaborate	differences description	B251-3, B251-5
E251-11	Is there another Member State(s) where an authorisation was refused/suspended/revoked by competent authorities for the same ⁴ product?			
E251-12	Yes	maa:section4-1-3/maa:refused (Value=1)	was auth refused	B251-6, B252-1
E251-13	No	maa:section4-1-3/maa:refused (Value=0)	was auth refused	B251-6
E251-14	<p><i>Note: * "same product" means same qualitative and quantitative composition in active substance(s) and having the same pharmaceutical form from applicants belonging to the same mother company or group of companies OR which are "licensees".</i></p> <p><i>** This is covering applications submitted at an earlier time or in parallel to this application if not already listed under 1.1.2 or 1.1.3</i></p>			

Element Tree Diagram



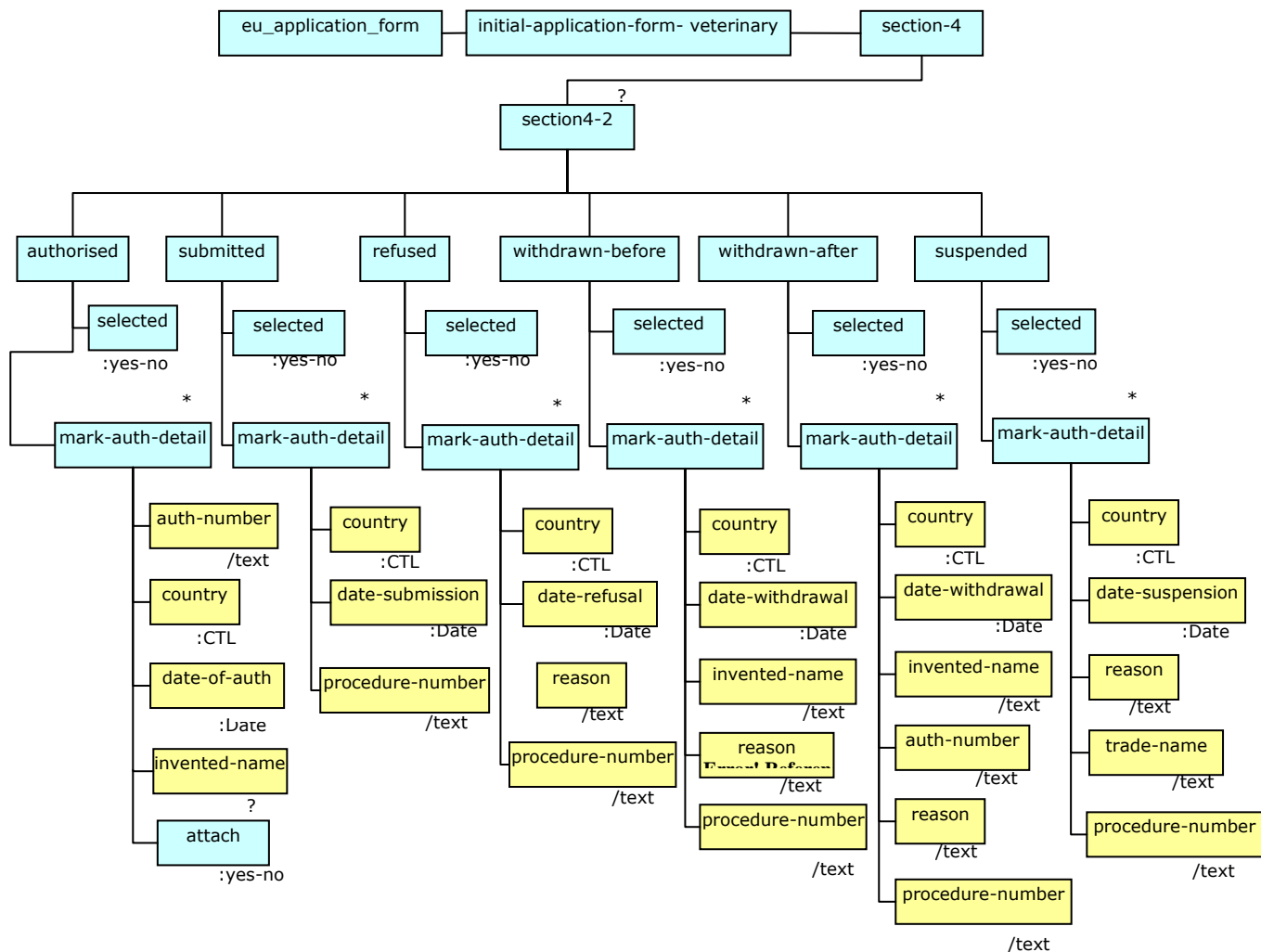
Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B251-1	E251-2, E251-3a	Mandatory.	Mutually Exclusive.	
B251-2	E251-5, E251-6	Mandatory.	Mutually Exclusive.	
B251-3	E251-5, E251-7 to E251-10	E251-5 is mandatory, rest are optional.	If E251-5 is selected, then the other fields are required.	
B251-4	E251-8, E251-9	Mandatory.	Mutually Exclusive.	
B251-5	E251-8, E251-10	Optional.	If E251-8 is selected, then E251-10 is required.	
B251-6	E251-12, E251-13	Mandatory.	Mutually Exclusive.	

2.5.2. Marketing authorisation applications for the same product in the eea (same qualitative and quantitative composition in active substance(s) and having the same pharmaceutical form from applicants belonging to the same mother company or group of companies or which are "licensees").

	Common DES 3.0 Context		Common RDM Entry point	
	maa:eu_application_form/ maa:initial-application-form-veterinary/ maa:other_marketing_authorisation_applications/ maa:Marketing_authorisation_applications_in_EEA/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E252-1	Authorised	maa:authorised/rdm:selected	Authorisation Status CTL	B252-2
E252-2	Country	maa:authorised/rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B252-2
E252-3	Date of authorisation	maa:authorised/rdm:mark-auth-detail/rdm:date-of-auth	MP Authorisation > authorisation date	B252-2
E252-4	Invented name	maa:authorised/rdm:mark-auth-detail/rdm:invented-name	Medicinal Product Group > invented name	B252-2
E252-5	Authorisation number	maa:authorised/rdm:mark-auth-detail/rdm:auth-number	MP Authorisation > authorisation number	B252-2
E252-6	Attach marketing authorisation (Annex 5.15)	maa:authorised/rdm:attach		B252-2
E252-7	Submitted	maa:pending/rdm:selected	Authorisation Status CTL	B252-3
E252-8	Country	maa:pending/rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B252-3
E252-9	Date of submission	maa:pending/rdm:mark-auth-detail/rdm:date-submission	Application > submission date	B252-3
E252-10	Procedure number of MRP/DCP	maa:pending/rdm:mark-auth-detail/procedure-number	MP Authorisation > MP Procedure > procedure number	B252-3
E252-11	Refused	maa:refused/selected	Authorisation Status CTL	B252-4
E252-12	Country	maa:refused/rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B252-4
E252-13	Date of refusal	maa:refused/rdm:mark-auth-detail/rdm:date-refusal	MP Authorisation > authorisation date	B252-4
E252-14	Reason for refusal	maa:refused/rdm:mark-auth-detail/rdm:reason	MP Authorisation > authorisation description	B252-4
E252-15	Procedure number of MRP/DCP	maa:pending/rdm:mark-auth-detail/procedure-number	MP Authorisation > MP Procedure > procedure number	B252-4
E252-16	Withdrawn (by applicant before authorisation)	maa:withdrawn-before/rdm:selected	Authorisation Status CTL	B252-5
E252-17	Country	maa:withdrawn-before/rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B252-5
E252-18	Date of withdrawal	maa:withdrawn-before/rdm:mark-auth-detail/rdm:date-withdrawal	MP Authorisation > authorisation date	B252-5
E252-19	Invented name	maa:withdrawn-before/rdm:mark-auth-detail/rdm:invented-name	Medicinal Product Group > invented name	B252-5
E252-20	Reason for withdrawal	maa:withdrawn-before/rdm:mark-auth-detail/rdm:reason	MP Authorisation > authorisation description	B252-5
E252-21	Procedure number of MRP/DCP	maa:pending/rdm:mark-auth-detail/procedure-number	MP Authorisation > MP Procedure > procedure number	B252-5
E252-22	Withdrawn (by applicant after authorisation)	maa:withdrawn-after/rdm:selected	Authorisation Status CTL	B252-6
E252-23	Country	maa:withdrawn-after/rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B252-6
E252-24	Date of withdrawal	maa:withdrawn-after/rdm:mark-auth-detail/rdm:date-withdrawal	MP Authorisation > authorisation date	B252-6
E252-25	Authorisation number	maa:withdrawn-after/rdm:mark-auth-detail/rdm:auth-number	MP Authorisation > authorisation number	B252-6
E252-26	Invented name	maa:withdrawn-after/rdm:mark-auth-detail/rdm:invented-name	Medicinal Product Group > invented name	B252-6
E252-27	Reason for withdrawal	maa:withdrawn-after/rdm:mark-auth-detail/rdm:reason	MP Authorisation > authorisation description	B252-6
E252-28	Procedure number of MRP/DCP	maa:pending/rdm:mark-auth-detail/procedure-number	MP Authorisation > MP Procedure > procedure number	B252-6
E252-29	Suspended/revoked (by competent authority)	maa:suspended/rdm:selected	Authorisation Status CTL	B252-7
E252-30	Country	maa:suspended/rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B252-7
E252-31	Date of suspension/revocation	maa:suspended/rdm:mark-auth-detail/rdm:date-suspension	MP Authorisation > authorisation date	B252-7
E252-32	Reason for suspension/revocation	maa:suspended/rdm:mark-auth-detail/rdm:reason	MP Authorisation > authorisation description	B252-7

E252-33	Invented name	maa:suspended/rdm:mark-auth-detail/rdm:invented-name	Medicinal Product Group > invented name	B252-7
E252-34	Procedure number of MRP/DCP (if applicable)	maa:pending/rdm:mark-auth-detail/procedure-number	MP Authorisation > MP Procedure > procedure number	B252-7

Element Tree Diagram

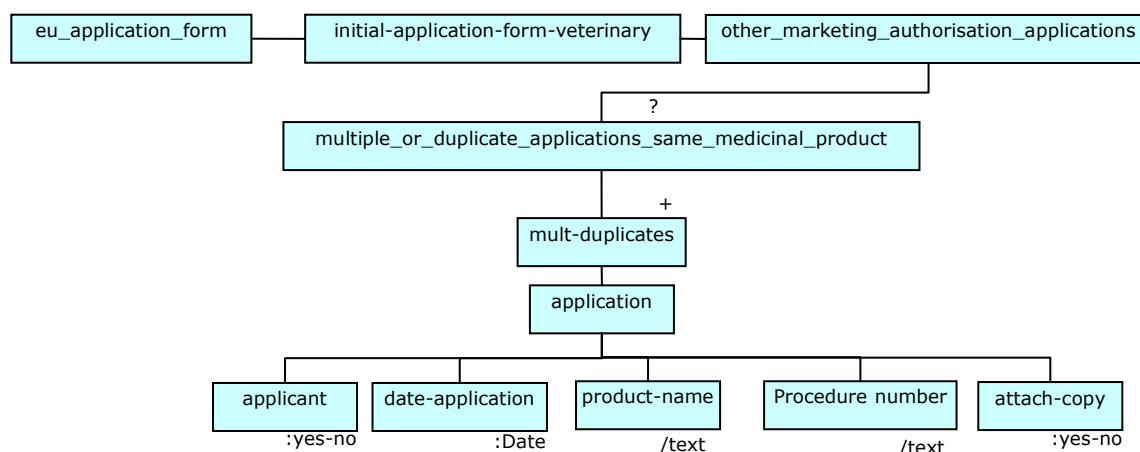


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B252-1	E251-1, E251-5, E251-12	Mandatory.	If one of the fields are selected, section 2.5.2 is mandatory.	
B252-2	E252-1 to E252-6	E252-1 mandatory, rest are optional.	If E252-1 is selected, then the rest are mandatory.	
B252-3	E252-7 to E252-10	E252-7 mandatory, rest are optional.	If E252-7 is selected, then the rest are mandatory.	
B252-4	E252-11 to E252-15	E252-13 mandatory, rest are optional.	If E252-11 is selected, then the rest are mandatory.	
B252-5	E252-16 to E252-21	E252-16 mandatory, rest are optional.	If E252-16 is selected, then the rest are mandatory.	
B252-6	E252-22 to E252-28	E252-22 mandatory, rest are optional.	If E252-22 is selected, then the rest are mandatory.	
B252-7	E252-29 to E252-34	E252-29 mandatory, rest are optional.	If E252-29 is selected, then the rest are mandatory.	

2.5.3. FOR MULTIPLE APPLICATIONS OF THE SAME VETERINARY MEDICINAL PRODUCT

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/ maa:initial-application-form-veterinary/ maa:other_marketing_authorisation_applications/ maa:multiple_or_duplicate_applications_same_medicinal_product/	Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E253-1	Multiple/duplicate applications for:			
E253-2	Name of other product	maa:mult-duplicates/ maa:application/maa:product-name	Medicinal Product > medicinal product name	
E253-3	Date of application(s)	maa:mult-duplicates/ maa:application/maa:date-application	Application > Submission date	
E253-4	Applicant	maa:mult-duplicates/ maa:application/maa:applicant	Role > Party > Organisation> Name	
E253-5	Procedure number for MRP/DCP (if applicable)	maa:mult-duplicates/ maa:application/maa:procedure- number		
E253-5	Attach copy of letter from the Commission services, for centralised procedures only (Annex 5.16)	maa:mult-duplicates/ maa:application/maa:date-application /maa:attach-copy		

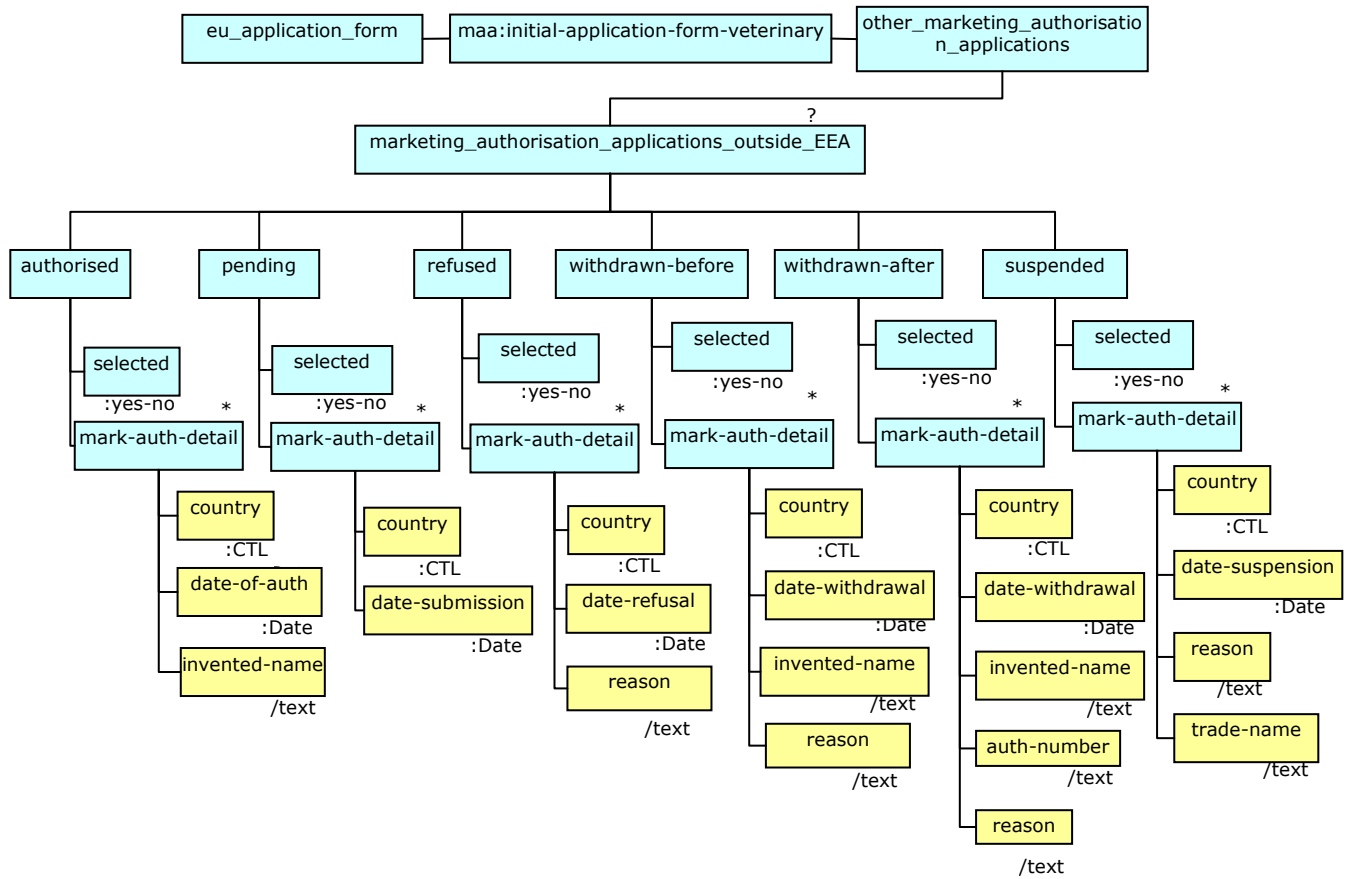
Element Tree Diagram



2.5.4. Marketing authorisation applications for the same product outside the eea (i.e. From applicants belonging to the same mother company or group of companies or which are "licensees". Same qualitative and quantitative composition in active substance(s) and having the same pharmaceutical form).

	Common DES 3.0 Context		Common RDM Entry point	
	maa:eu_application_form/ maa:initial-application-form-veterinary/ maa:other_marketing_authorisation_applications/ maa:marketing_authorisation_applications_outside_EEA/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E254-1	Authorized	maa:authorised/rdm:selected	Authorisation Status CTL	B254-1
E254-2	Country	maa:authorised/rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B254-1
E254-3	Date of authorization	maa:authorised/rdm:mark-auth-detail/rdm:date-of-auth	MP Authorisation > authorisation date	B254-1
E254-4	Invented name	maa:authorised/rdm:mark-auth-detail/rdm:invented-name	Medicinal Product Group > invented name	B254-1
E254-6	Pending	maa:pending/rdm:selected	Authorisation Status CTL	B254-2
E254-7	Country	maa:pending/rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B254-2
E254-8	Date of submission	maa:pending/rdm:mark-auth-detail/rdm:date-submission	Application > submission date	B254-2
E254-9	Refused	maa:refused/selected	Authorisation Status CTL	B254-3
E254-10	Country	maa:refused/rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B254-3
E254-11	Date of refusal	maa:refused/rdm:mark-auth-detail/rdm:date-refusal	MP Authorisation > authorisation date	B254-3
E252-12	Reason for refusal	maa:refused/rdm:mark-auth-detail/rdm:reason	MP Authorisation > authorisation description	B252-3
E254-13	Withdrawn (by applicant before authorization)	maa:withdrawn-before/rdm:selected	Authorisation Status CTL	B254-4
E254-14	Country	maa:withdrawn-before/rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B254-4
E254-15	Date of withdrawal	maa:withdrawn-before/rdm:mark-auth-detail/rdm:date-withdrawal	MP Authorisation > authorisation date	B254-4
E254-16	Invented name	maa:withdrawn-before/rdm:mark-auth-detail/rdm:invented-name	Medicinal Product Group > invented name	B254-4
E254-17	Reason for withdrawal	maa:withdrawn-before/rdm:mark-auth-detail/rdm:reason	MP Authorisation > authorisation description	B254-4
E254-18	Withdrawn (by applicant after authorization)	maa:withdrawn-after/rdm:selected	Authorisation Status CTL	B254-5
E254-19	Country	maa:withdrawn-after/ rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B254-5
E254-20	Date of withdrawal	maa:withdrawn-after/ rdm:mark-auth-detail/rdm:date-withdrawal	MP Authorisation > authorisation date	B254-5
E254-21	Authorization number	maa:withdrawn-after/ rdm:mark-auth-detail/rdm:auth-number	MP Authorisation > authorisation number	B254-5
E254-22	Invented name	maa:withdrawn-after/ rdm:mark-auth-detail/rdm:invented-name	Medicinal Product Group > invented name	B254-5
E254-23	Reason for withdrawal	maa:withdrawn-after/ rdm:mark-auth-detail/rdm:reason	MP Authorisation > authorisation description	B254-5
E254-24	Suspended/revoked (by competent authority)	maa:suspended/rdm:selected	Authorisation Status CTL	B254-6
E254-25	Country	maa:suspended/rdm:mark-auth-detail/rdm:country	MP Authorisation > Country CTL	B254-6
E254-26	Date of suspension/revocation	maa:suspended/rdm:mark-auth-detail/rdm:date-suspension	MP Authorisation > authorisation date	B254-6
E254-27	Reason for suspension/revocation	maa:suspended/rdm:mark-auth-detail/rdm:reason	MP Authorisation > authorisation description	B254-6
E254-28	Trade name	maa:suspended/rdm:mark-auth-detail/rdm:invented-name	Medicinal Product Group > invented name	B254-6

Element Tree Diagram

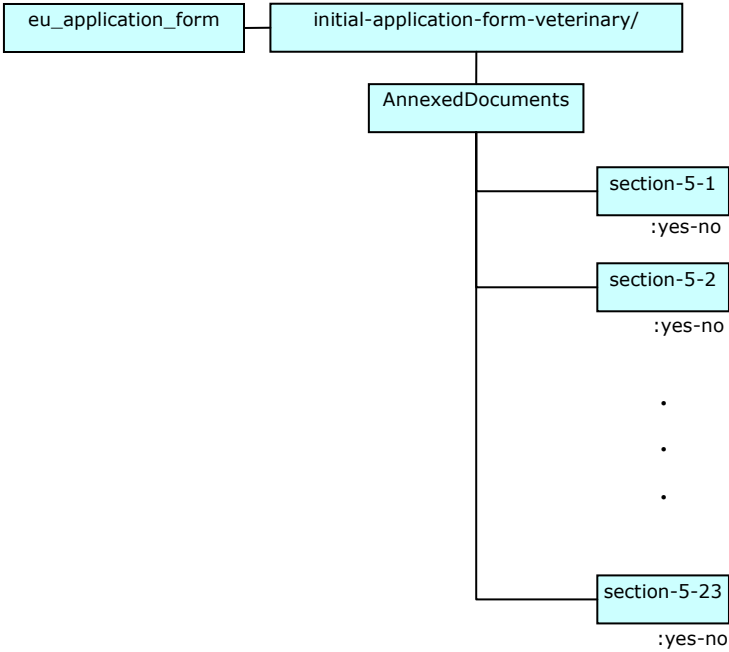


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B254-1	E254-1 to E254-4	E254-1 mandatory, rest are Optional.	If E254-1 is selected, then the rest are mandatory.	
B254-2	E254-6 to E254-8	E254-6 mandatory, rest are Optional.	If E254-7 is selected, then the rest are mandatory.	
B254-3	E254-9 to E254-12	E254-9 mandatory, rest are Optional.	If E254-9 is selected, then the rest are mandatory.	
B254-4	E254-13 to E254-17	E254-12 mandatory, rest are Optional.	If E254-12 is selected, then the rest are mandatory.	
B254-5	E254-18 to E254-23	E254-18 mandatory, rest are Optional.	If E254-18 is selected, then the rest are mandatory.	
B254-6	E254-24 to E254-28	E254-24 mandatory, rest are Optional.	If E254-24 is selected, then the rest are mandatory.	

2.6. ANNEXED DOCUMENTS (WHERE APPROPRIATE)

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/ maa:initial-application-form-veterinary/ maa:AnnexedDocuments		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E26-1	Proof of payment	maa:section5-1		
E26-2	Informed consent letter of marketing authorisation holder of authorised veterinary medicinal product.	maa:section5-2		
E26-3	Proof of establishment of the applicant in the EEA.	maa:section5-3		
E26-4	Letter of authorisation for communication on behalf of the applicant/MAH.	maa:section5-4		
E26-5	(empty)			
E26-6	Manufacturing Authorisation required under Article 44 of Directive 2001/82/EC (or equivalent, outside of the EEA where MRA or other Community arrangements apply). A reference to EudraGMP will suffice when available.	maa:section5-6		
E26-7	(empty)			
E26-8	Flow-chart indicating all sites involved in the manufacturing process of the veterinary medicinal product or active substance (including sites involved in sampling and testing for batch release of products manufactured in third countries). Note: ALL manufacturing and control sites mentioned throughout the dossier MUST be consistent regarding their names, detailed addresses and activities	maa:section5-8		
E26-9	Statement (or GMP Certificate issued by an EEA inspectorate, when available) from the competent authority which carried out the inspection of the manufacturing site(s) (not older than 3 years). References to Eudra GMP will suffice when available. Where applicable a summary of the other GMP inspections performed in the last 2 years.	maa:section5-9		
E26-10	Letter(s) of access to Active Substance Master File(s) (Drug Master File(s)) or copy of Ph. Eur. Certificate(s) of suitability.	maa:section5-10		
E26-11	Copy of written confirmation from the manufacturer of the active substance to inform the applicant in case of Modification of the manufacturing process or specifications according to Annex I of Directive 2001/82/EC	maa:section5-11		
E26-12	Ph. Eur. Certificate(s) of suitability for TSE.	maa:section5-12		
E26-13	Written consent(s) of the competent authorities regarding GMO release in the environment.	maa:section5-13		
E26-14	Scientific Advice given by CVMP or Member State.	maa:section5-14		
E26-15	Copy of Marketing Authorization(s) required under Article 44 of Directive 2001/82/EC in the EEA and the equivalent in third countries on request (a photocopy of the pages which give the marketing authorization number, the date of authorisation and the page which has been signed by the authorizing competent authority will suffice).	maa:section5-15		
E26-16	Letter from Commission services regarding multiple applications.	maa:section5-16		
E26-17	List of Mock-ups or Samples/specimens sent with the application, as appropriate (see EMA/CMDv website).	maa:section5-17		
E26-18	List of proposed (invented) names and marketing authorisation holders in the concerned member states.	maa:section5-18		
E26-19	For each active substance, attach a declaration(s) from the Qualified Person of the manufacturing authorisation holder in Section 2.5.1 and from the Qualified Person of each of the manufacturing authorisation holders (i.e. located in EEA) listed in Section 2.5.2 where the active substance is used as a starting material that the active substance is manufactured in compliance with the detailed guidelines on good manufacturing practice for starting materials. Alternatively, such declaration may be signed by one Qualified Person on behalf of all QPs involved (provided this is clearly indicated).	maa:section5-19		
E26-20	Detailed description of the Pharmacovigilance system and, where appropriate, the risk management system that the Applicant will put in place.	maa:section5-20		
E26-21	Copy of 'Qualification of SME Status'.	maa:section5-21		
E26-22	Evidence and justification to support the claim of new active substance status in the Union for applications based on Article 12(3) of Directive 2001/82/EC	maa:section5-22		
E26-23	Copy of EMA certificate for a Vaccine Antigen Master File	maa:section5-23		

Element Tree Diagram



3. About this Document

3.1. Definitions, Acronyms, and Abbreviations

3.1.1. Acronyms

Name	Definition
CMS	Concerned Member State
DCP	DeCentralised Procedure
DTD	Data Type Definition
ETD	Element Tree Diagram
EU	European Community
MA	Marketing Authorisation
MRP	Mutual Recognition Procedure
NP	National Procedure
RDM	Reference Data Model
RMS	Reference Member State
TSE	Transmissible Spongiform Encephalopathy
XML	eXtended Markup Language
XSL	XML Stylesheet Language