



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

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Electronic Application Form Data Exchange Standard 3.0

Supplementary Specification Annex 2 "Variations Form" v.1.26.0.0

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1. Introduction

1.1. How to read this document

In association with this document the **variations.xsd** file contains the XML schema file that provides a 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. This schema file can be found on the esubmissions website under the eAF page at the following link:

<http://esubmission.ema.europa.eu/eaf/index.html>

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 human 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 supplementary specifications document. This again can be found on the esubmissions website under the eAF page.

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 how 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)

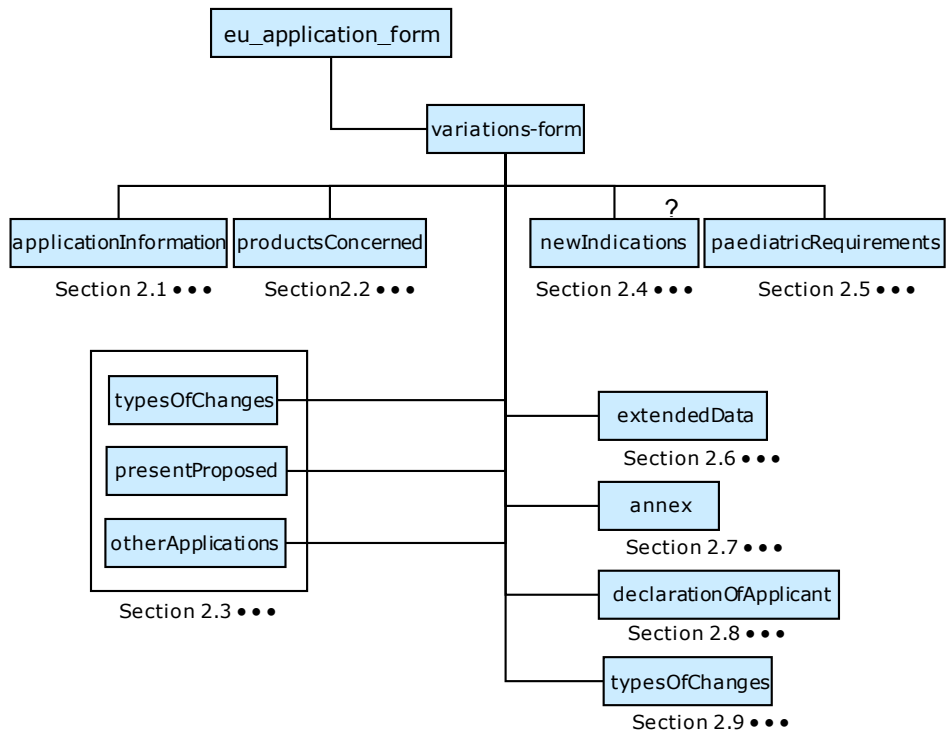
The diagrams of this version reflect the DES 3.0 standard described in the DES 3.0 Supplementary Specifications document. The ETD shows which are the concepts involved in the mapping of all the application form fields in the Element Mapping Table and the hierarchical constraints between them.

2. Variations Form

The “variations-form” 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:variations-form/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E2.1	APPLICATION FOR VARIATION TO A MARKETING AUTHORISATION	maa:applicationInformation/		See Section 2.1
E2.2	PRODUCTS CONCERNED BY THIS APPLICATION	maa:productsConcerned/		See Section 2.2
E2.3	TYPE OF CHANGE(S)	maa:typesOfChanges/ and maa:presentProposed/ and maa:otherApplications/		See Section 2.3
E2.4	Type II Variations – new Indications – orphan medicinal product information	maa:newIndications/		See Section 2.4
E2.5	Type II Variations – Paediatric requirements	maa:paediatricRequirements/		See Section 2.5
E2.6	Type II variations – Extended data/market exclusivity	maa:extendedData/		See Section 2.6
E2.7	The following amended product information proposals are provided in the relevant sections of the EU-CTD format for NTA volume 6B format, where applicable:	maa:annex/		See Section 2.7
E2.8	DECLARATION OF APPLICANT	maa:declarationOfApplicant/		See Section 2.8
E2.9	Signatures	maa:Signatory/		See Section 2.9

Element Tree Diagram

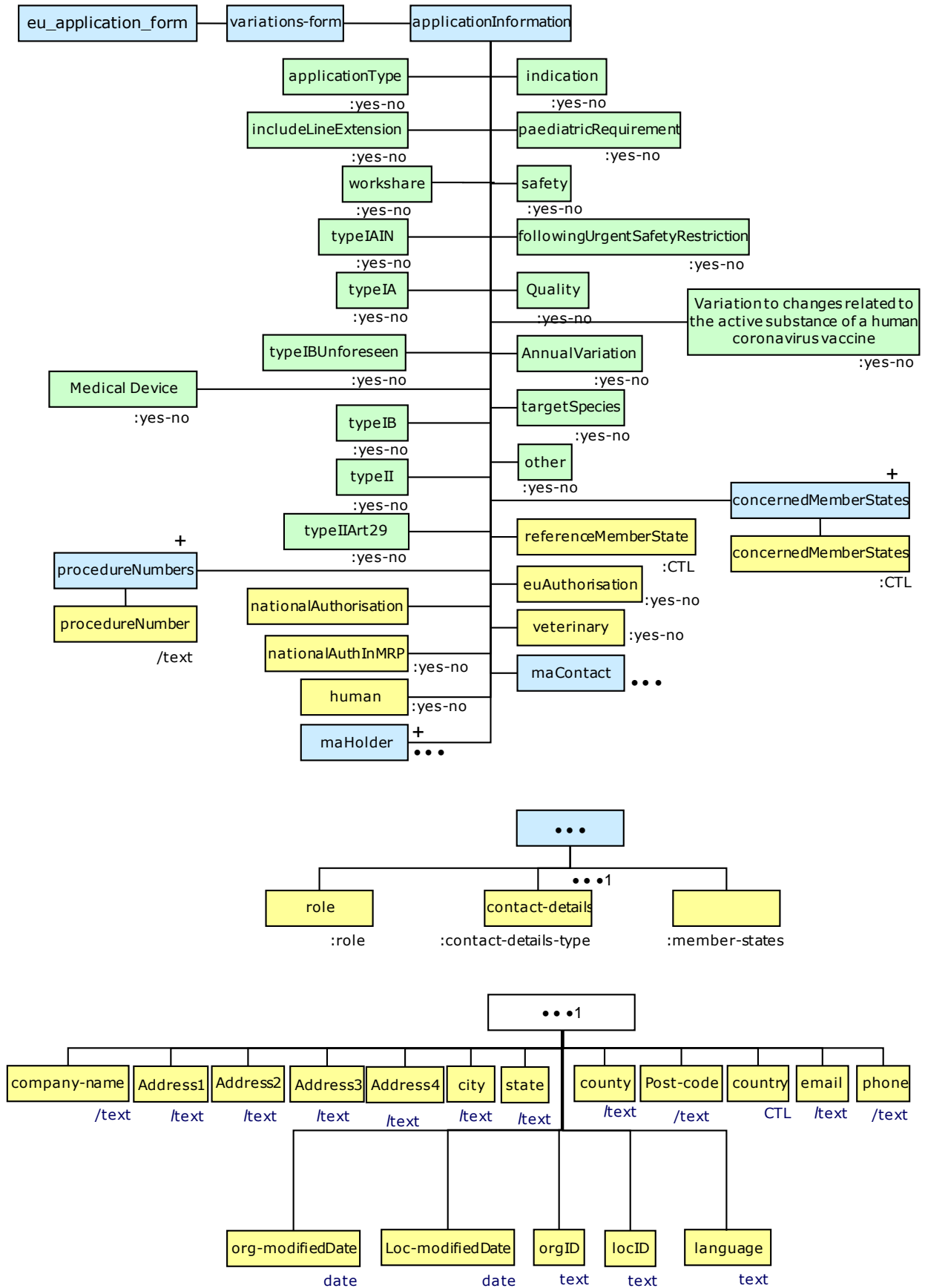


2.1. APPLICATION FOR VARIATION TO A MARKETING AUTHORISATION

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:variations-form/maa:applicationInformation/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E21-1	Human	maa:human	Domain CTL	B21-1
E21-2	Veterinary	maa:veterinary	Domain CTL	B21-1
E21-3	National Authorisation in MRP/DCP	maa:nationalAuthInMRP	Procedure Type CTL	
E21-4	EU Authorisation	maa:euAuthorisation	Procedure Type CTL	
E21-5	National Authorisation	maa:nationalAuthorisation	Procedure Type CTL	
E21-6	Variation procedure number(s) ¹ ;	maa:procedureNumbers/ maa:procedureNumber	MP Procedure > procedure number	
E21-7	Reference Member State / Reference Authority for worksharing	maa:.referenceMemberState	MP Procedure > Role > Country CTL	
E21-8	Concerned Member State(s)	maa:concernedMemberStates/ maa:concernedMemberState	MP Procedure > Role > Country CTL	
E21-9	Type of Application (tick all applicable options)			
E21-10	Single variation	maa:applicationType (value=1)	Var Grouping CTL	B21-2
E21-11	Grouping of variations	maa:applicationType (value=2)	Var Grouping CTL	B21-2, B21-3
E21-12	Include a line extension ³	maa:includeLineExtension	Var Grouping CTL	B21-3
E21-13	Worksharing	maa:workshare	Var Grouping CTL	
E21-14	Type IA _{IN}	maa:typeIAIN	Var Type Summary CTL	B21-4
E21-15	Type IA	maa:typeIA	Var Type Summary CTL	B21-4
E21-16	Type IB unforeseen ²	maa:typeIBUnforeseen	Var Type Summary CTL	B21-4, B21-5,
E21-17	Type IB	maa:typeIBForeseen	Var Type Summary CTL	B21-4, B21-5
E21-18	Type II	maa:typeII	Var Type Summary CTL	B21-4, B21-5, B24-1, B26-2
E21-19	Type II Art. 29 ⁴	maa:typeIIArt29	Var Type Summary CTL	B21-4, B21-5, B24-1, B26-2
E21-20	Change(s) concern(s) (for Type IB and Type II variations only, tick all changes applicable)			
E21-21	Indication	maa:indication	Var Domain Summary CTL	B21-5, B24-1
E21-22	Paediatric Requirements	maa:paediatricIndication	Var Domain Summary CTL	B21-5, B24-1
E21-23	Safety	maa:safety	Var Domain Summary CTL	B21-5, B21-6
E21-24	Following Urgent Safety Restriction	maa:followingUrgentSafetyRestriction	Var Domain Summary CTL	B21-6
E21-25	Quality	maa:Quality	Var Domain Summary CTL	B21-5
E21-26	Annual variation for human influenza vaccines	maa:AnnualVariation	Var Domain Summary CTL	B21-5
E21-27	Non-food producing target species	maa:targetSpecies	Var Domain Summary CTL	B21-5
E21-28	Other	maa:other	Var Domain Summary CTL	B21-5
E21-29a	Variation to changes related to the active substance of a human coronavirus vaccine	maa: covidSubstanceChange ⁶	Var Domain Summary CTL	B21-5
E21-29b	Medical Device	maa:medicalDevice	Var Domain Summary CTL	B21-5
	Name and address of the MA Holder ⁵			
E21-30	Company Name	maa:maHolder/maa:role/rdm:name	Application > Party > Organisation > Name	
E21-31	Address1	maa:maHolder/maa:contact-details/rdm:address1	Application > Role > Party > Contact Details > Address	
		maa:maHolder/maa:contact-details/rdm:address2		
		maa:maHolder/maa:contact-details/rdm:address3		
		maa:maHolder/maa:contact-details/rdm:address4		
E21-32	city	maa:maHolder/maa:contact-details /rdm:city	Application > Role > Party > Contact Details > City	
E21-32a	state	maa:maHolder/maa:contact-details/rdm:state		

E21-32b	county	maa:maHolder/maa:contact-details/rdm:county		
E21-33	PostCode	maa:maHolder/maa:contact-details/rdm:post-code	Application > Role > Party > Contact Details > Post Code	
E21-34	Country	maa:maHolder/maa:contact-details/rdm:country	Application > Role > Party > Contact Details > Address > Country CTL	
E21-34a	orgID	maa:maHolder/maa:contact-details/rdm:orgID		
E21-34b	locID	maa:maHolder/maa:contact-details/rdm:locID		
E21-34c	loc-modifiedDate	maa:maHolder/maa:contact-details/rdm:loc-modifiedDate		
E21-34d	org-modifiedDate	maa:maHolder/maa:contact-details/rdm:org-modifiedDate		
E21-34e	Language	maa:maHolder/maa:contact-details/rdm:language		
E21-35	Name and address of contact person ⁶			
E21-36	Title	maa:maContact/maa:role/rdm:personal-title	Role > Party > Person > Personal Title	
E21-37	First name	maa:maContact/maa:role/rdm:given-name	Role > Party > Person > given name	
E21-38	Surname	maa:maContact/maa:role/rdm:family-name	Role > Party > Person > family name	
E21-39	Company Name	maa:maContact/maa:contact-details/rdm:companyName	Role > Party > Person > Company name	
E21-40	Address1	maa:maContact/maa:contact-details/rdm:address1	Role > Party > Contact Details > Address	
		maa:maContact/maa:contact-details/rdm:address2		
		maa:maContact/maa:contact-details/rdm:address3		
		maa:maContact/maa:contact-details/rdm:address4		
E21-41	city	maa:maContact/maa:contact-details/rdm:city	Role > Party > Contact Details > City	
E21-41a	state	maa:maContact/maa:contact-details/rdm:state		
E21-41b	county	maa:maContact/maa:contact-details/rdm:county		
E21-42	PostCode	maa:maContact/maa:contact-details/rdm:post-code	Role > Party > Contact Details > Post Code	
E21-43	Country	maa:maContact/maa:contact-details/rdm:country	Role > Party > Contact Details > Address > Country CTL	
E21-43a	orgID	maa:maContact/maa:contact-details/rdm:locID		
E21-43b	locID	maa:maContact/maa:contact-details/rdm:locID		
E21-43c	Loc-modifiedDate	maa:maContact/maa:contact-details/rdm:loc-modifiedDate		
E21-43d	Org-modifiedDate	maa:maContact/maa:contact-details/rdm:org-modifiedDate		
E21-43e	Language	maa:maContact/maa:contact-details/rdm:language		
E21-44	Telephone	maa:maContact/maa:contact-details/rdm:phone	Role > Party > Contact Details > Electronic contact > electronic contact	
E21-46	Email	maa:maContact/maa:contact-details/rdm:email	Role > Party > Contact Details > Electronic contact > electronic contact	

Element Tree Diagram

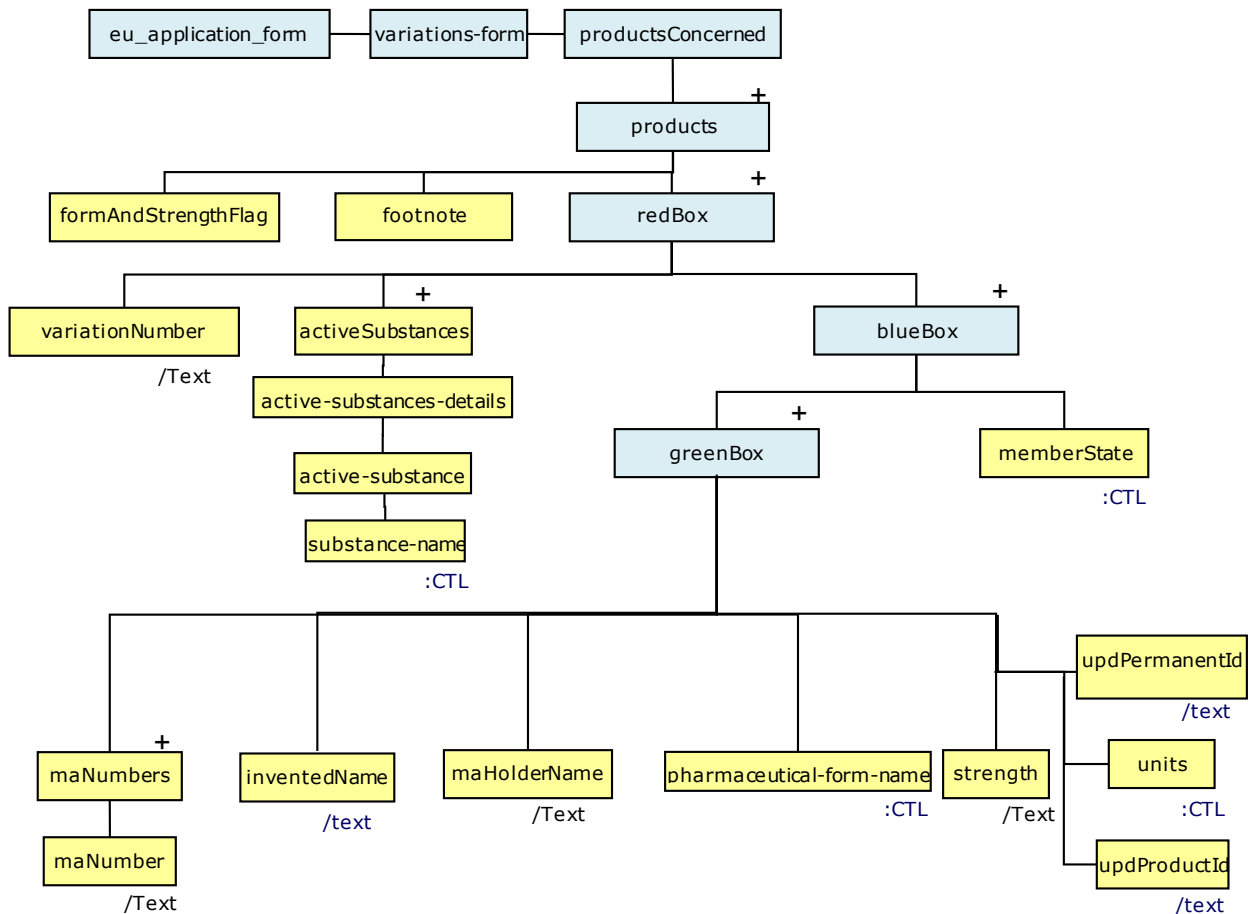


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B21-1	E21-1, E21-2	Mandatory	Mutually Exclusive	
B21-2	E21-10, E21-11	Mandatory	Mutually Exclusive	
B21-3	E21-11, E21-12	E21-11 is mandatory , E21-12 is optional	If E21-11 is selected, then E21-12 is displayed but not required.	
B21-4	E21-14 to E21-19	Mandatory	At least one mandatory, more than one allowed	
B21-5	E21-21 to E21-23, E21-25 to E21-28, E21-16 to E21-19	Mandatory	If any of E21-16 to E21-19 are selected, then at least one of E21-21 to E21-23, E21-25 to E21-28 is required, more than one are allowed	
B21-6	E21-23, E21-24	E21-23 is Mandatory, E21-24 is optional	If E21-23 is selected, then E21-24 is displayed but it is still optional	
B21-6	E39a, E39b	Optional	If E21-1 is selected then E19a and E39b are displayed	

2.2. PRODUCTS CONCERNED BY THIS APPLICATION

Common DES 3.0 Context			Common RDM Entry point	
maa:eu_application_form/maa:variations-form/maa:productsConcerned/maa:products/			Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E22-1	MRP Variation Number ⁸	maa:redBox/maa:variationNumber		
E22-2	Active Substance(s)	maa:redBox/maa:activeSubstances		
E22-3	ActiveSubstance	maa:redBox/maa:activeSubstances/maa:active-substance/maa:substance-name/maa:name		
E22-4	Member States	maa:redBox/maa:bluebox/maa:memberState		
E22-5	MA Number ⁸	maa:redBox/maa:bluebox/maa:greebox/maa:maNumbers/maa:maNumber		
E22-6	Invented Name	maa:redBox/maa:bluebox/maa:greebox/maa:inventedName		
E22-7	MA Holder Name	maa:redBox/maa:bluebox/maa:greebox/maa:maHolderName		
E22-8	Pharmaceutical Form name	maa:redBox/maa:bluebox/maa:greebox/maa:pharmaceutical-form-name		
E22-9	Strength	maa:redBox/maa:bluebox/maa:greebox/maa:strength		
E22-10	Units	maa:redBox/maa:bluebox/maa:greebox/maa:units		
E22-11	UPD Permanent Identifier ⁶ for the concerned national product(s)	maa:redBox/maa:bluebox/maa:greebox/maa:updPermanentId		
E22-12	UPD Product Identifier ⁶ (only relevant for MRP and CP)	maa:redBox/maa:bluebox/maa:greebox/maa:updProductId		

Element Tree Diagram



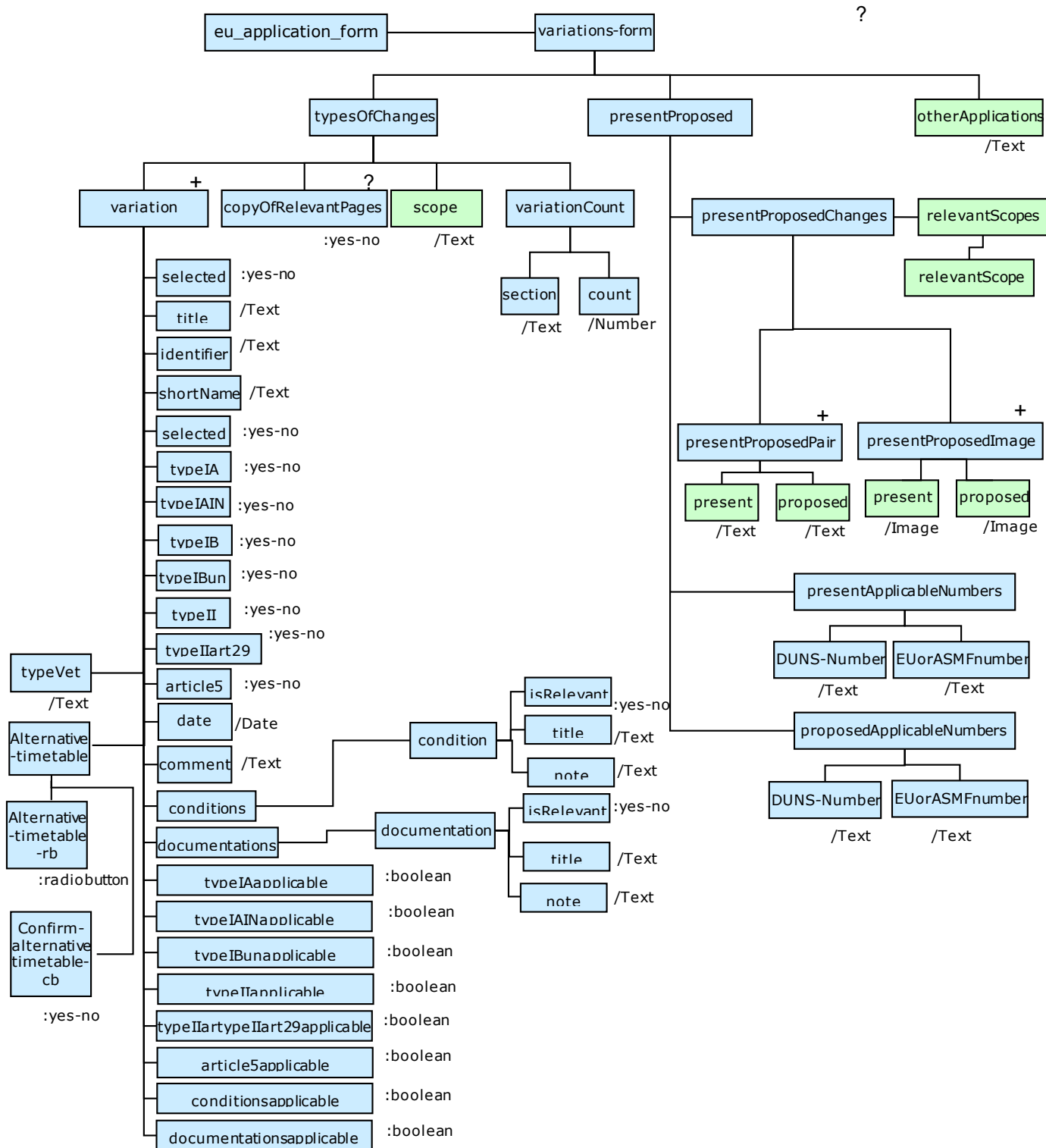
2.3. TYPE OF CHANGE(S)

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:variations-form/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E23-1	Copy of the relevant page(s) from the Guideline for this/these change(s) is attached and the relevant boxes for conditions and documentations (both for Type IA and Type IB) are ticked.	maa:typesofChanges/ maa:copyOfRelevantPages		
E23-2	Variation	maa:typesofChanges/maa:variationCount/ maa:section		
E23-3	Selected	maa:typesofChanges/maa:variationCount/ maa:count		
Variation Items				
E23-4	Selected (no label present)	maa:typesofChanges/maa:variation/rdm:selected		
E23-5	Title	maa:typesofChanges/maa:variation/rdm:title		
E23-6	identifier (No label present)	maa:typesofChanges/maa:variation/rdm:identifier		
E23-7	shortName (No label present)	maa:typesofChanges/maa:variation/rdm:shortName		
E23-8	typeIA	maa:typesofChanges/maa:variation/rdm:typeIA		
E23-9	typeIAIN	maa:typesofChanges/maa:variation/rdm:typeIAIN		
E23-10	typeIB	maa:typesofChanges/maa:variation/rdm:typeIB		
E23-11	typeIBun	maa:typesofChanges/maa:variation/rdm:typeIBun		
E23-12	typeII	maa:typesofChanges/maa:variation/rdm:typeII		
E23-13	typeIIart29	maa:typesofChanges/maa:variation/rdm:typeIIart29		
E23-13a	typeVet	maa:typesofChanges/maa:variation/rdm:typeVet		
E23-13b	Alternative timetable agreed in advance	maa:typesofChanges/maa:variation/alternative-timetable/alternative-timetable-rb		
E23-13c	Please tick to confirm that the correspondence confirming the alternative timetable is included in the submission	maa:typesofChanges/maa:variation/alternative-timetable/confirm-alternative-timetable-cb		
E23-14	article5	maa:typesofChanges/maa:variation/rdm:article5		
E23-15	date	maa:typesofChanges/maa:variation/rdm:date		
E23-16	comment	maa:typesofChanges/maa:variation/rdm:comment		
E23-17a	isRelevant	maa:typesofChanges/maa:variation/rdm:conditions/condition/isRelevant		
E23-17b	title	maa:typesofChanges/maa:variation/rdm:conditions/condition/title		
E23-17c	note	maa:typesofChanges/maa:variation/rdm:conditions/condition/note		
E23-18a	isRelevant	maa:typesofChanges/maa:variation/rdm:documentations/documentation/documentations/isRelevant		
E23-18b	title	maa:typesofChanges/maa:variation/rdm:documentations/documentation/documentations/title		
E23-18c	note	maa:typesofChanges/maa:variation/rdm:documentations/documentation/documentations/note		
E23-19	typeIAapplicable	maa:typesofChanges/maa:variation/rdm:typeIAapplicable		
E23-20	typeIAINapplicable	maa:typesofChanges/maa:variation/rdm:typeIAINapplicable		
E23-21	typeIBapplicable	maa:typesofChanges/maa:variation/rdm:typeIBapplicable		
E23-22	typeIBunapplicable	maa:typesofChanges/maa:variation/rdm:typeIBunapplicable		
E23-23	typeIIapplicable	maa:typesofChanges/maa:variation/rdm:typeIIapplicable		
E23-24	typeIIart29applicable	maa:typesofChanges/maa:variation/rdm:typeIIart29applicable		
E23-25	article5applicable	maa:typesofChanges/maa:variation/rdm:article5applicable		
E23-26	conditionsapplicable	maa:typesofChanges/maa:variation/rdm:conditionsapplicable		

E23-27	documentationsapplicable	maa:typesofChanges/maa:variation/rdm:documentationsapplicable		
E23-28	PRECISE SCOPE AND BACKGROUND FOR CHANGE, AND JUSTIFICATION FOR GROUPING, WORKSHARING AND CLASSIFICATION OF UNFORESEEN CHANGES (if applicable) <i>(include a description and background of all the proposed changes. In case of grouping and worksharing a justification should be provided in a separate paragraph. If a variation concerns an unforeseen change, include a justification for its proposed classification).</i>	maa:typesofChanges/maa:scope	scope, background	
E23-28a	RelevantScope	maa:presentProposed/maa:presentproposedChanges/maa:relevantScopes/maa:relevantScope		
E23-28a	PRESENT	maa:presentProposed/maa:presentproposedChanges/maa:presentProposedPair/maa:present	Application Proposal > present product info	
E23-29	PRESENT- IMAGE	maa:presentProposed/maa:presentproposedChanges/maa:presentProposedImage/maa:present		
E23-30	D-U-N-S number	maa:presentProposed/maa:presentApplicableNumbers/maa: DUNS-number		
E23-31	EU or National ASMF number	maa:presentProposed/maa:presentApplicableNumbers/maa: EUorASMfnumber		
E23-32	PROPOSED	maa:presentProposed/maa:presentProposedPair/maa:proposed	Application Proposal > present product info	
E23-33	PresentAddress	maa:presentProposed/maa:presentProposedAddresses/maa:present		
E23-34	ProposedAddress	maa:presentProposed/maa:presentProposedAddresses/maa:proposed		
E23-35	PROPOSED- IMAGE	maa:presentProposed/maa:presentProposedImage/maa:proposed		
E23-36	D-U-N-S number	maa:presentProposed/maa:proposedApplicableNumbers/maa: DUNS-number		
E23-37	EU or National ASMF number	maa:presentProposed/maa:proposedApplicableNumbers/maa: EUorASMfnumber		
E23-38	OTHER APPLICATIONS	maa:otherApplications	Application > other application	



Element Tree Diagram

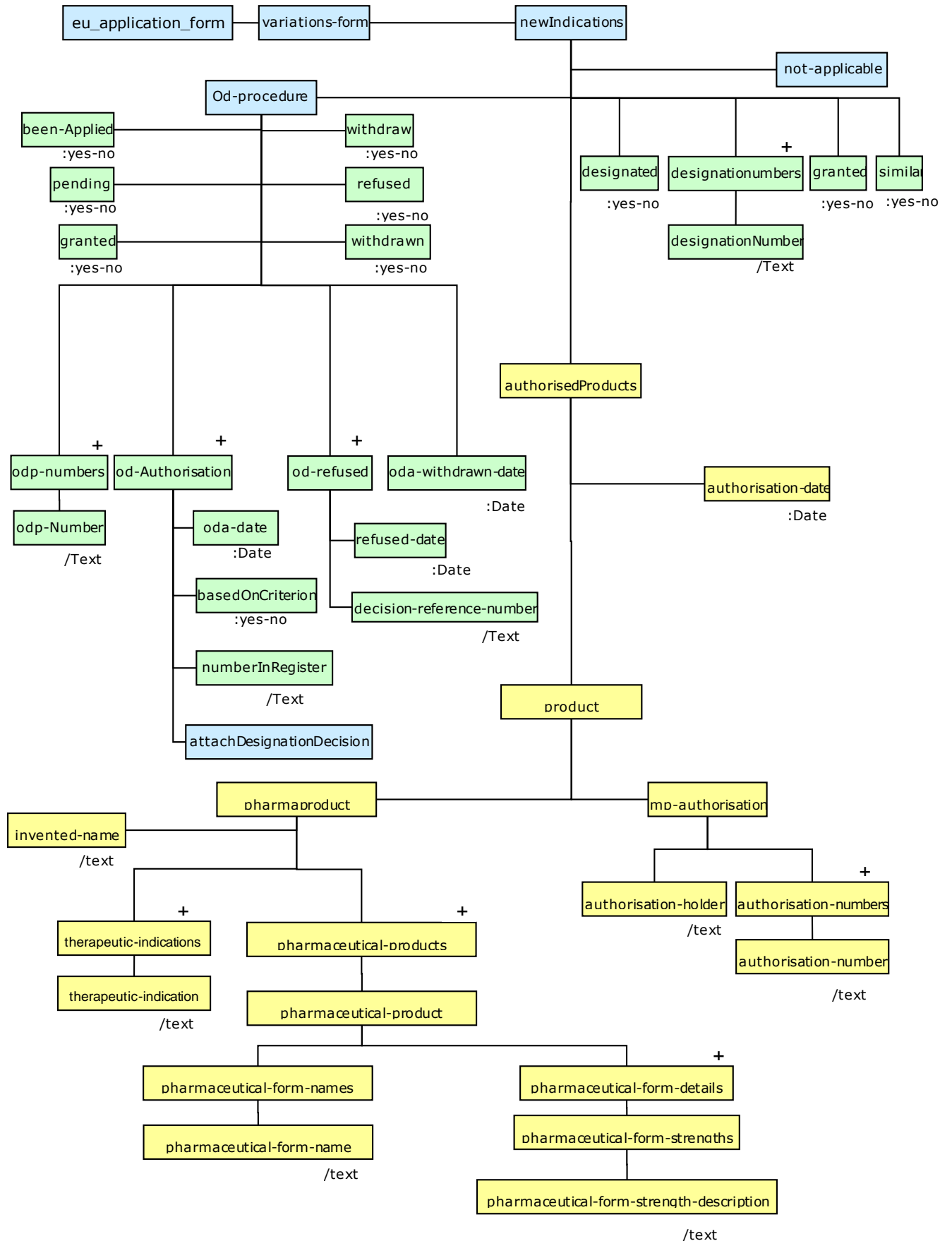


2.4. Type IB & Type II Variations – new Indications – orphan medicinal product information

	Common DES 3.0 Context		Common RDM Entry point	
	maa:eu_application_form/maa:variations-form/maa:newIndications/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E24-1	Select flag if not applicable; section will not be displayed.	maa:not-applicable		B24-7
E24-2	Has orphan designation been applied for, for this new indication?			
E24-3	Yes	maa:been-applied (Value=1)	Orphan Designation > has od applied	B24-2, B24-8
E24-4	No	maa:been-applied (Value=0)	Orphan Designation > has od applied	B24-2
E24-5	Orphan designation procedure number	maa:od-procedure/maa:odp-numbers/maa:odp-number	Orphan Designation > od procedure number	B24-8
E24-6	Pending	maa:od-procedure/maa:pending	Orphan Designation > Authorisation Status CTL	B24-3
E24-7	Orphan designation granted	maa:od-procedure/maa:granted	Orphan Designation > Authorisation Status CTL	B24-3, B24-9
E24-8	Date	maa:od-procedure/maa:od-authorisation/oda-date	Orphan Designation > od status date	B24-9
E24-9	Based on criterion of "significant benefit"			B24-9
E24-10	Yes	maa:od-procedure/maa:od-authorisation/maa:basedOnCriterion (Value=1)	Orphan Designation > is based significant benefit	B24-10, B24-9
E24-11	No	maa:od-procedure/maa:od-authorisation/maa:basedOnCriterion (Value=0)	Orphan Designation > is based significant benefit	B24-10, B24-9
E24-12	Number in the EU Register of Orphan Medicinal Products	maa:od-procedure/maa:od-authorisation /maa:numberInRegister	Orphan Designation > od community reg number	B24-9
E24-13	Attach copy of the Designation Decision	maa:od-procedure/maa:od-authorisation /maa:attachDesignationDecision	od decision ref number	B24-9
E24-14	Orphan Designated Refused	maa:od-procedure /maa:refused	Orphan Designation > Authorisation Status CTL	B24-3, B24-13
E24-15	Date	maa:od-procedure/maa:od-refused/maa:oda-refused-date	Orphan Designation > od status date	B24-13
E24-16	Commission Decision Reference number	maa:od-procedure/maa:od-refused/oda-decision-reference-number	Orphan Designation > od decision ref number	B24-13
E24-17	Orphan Designation Withdrawn	maa:od-procedure/maa:withdrawn	Orphan Designation > Authorisation Status CTL	B24-3, B24-14
E24-18	Date	maa:od-procedure/maa:oda-withdrawn-date	Orphan Designation > od status date	B24-14
E24-19	Information relating to orphan market exclusivity			
E24-20	Has any medicinal product been designated as an Orphan medicinal product for a condition relating to the new indication proposed in this variation application?			
E24-21	Yes	maa:designated (Value=1)	Orphan Designation > has mp designated orphan	B24-4, B24-11
E24-22	No	maa:designated (Value=0)	Orphan Designation > has mp designated orphan	B24-11
E24-23	Please specify the EU Orphan Designation Number(s):	maa:designationNumbers/ maa:designationNumber	OD Number > eu od number	B24-4, B24-11
E24-24	Has any of the designated Orphan medicinal product(s) been granted a marketing authorisation in the EU?			B24-11
E24-25	Yes	maa:granted (Value=1)	Orphan Designation > has mp granted ma	B24-5, B24-11, B24-12
E24-26	No	maa:granted (Value=0)	Orphan Designation > has mp granted ma	B24-5, B24-11, B24-12
E24-27	Please specify for authorised product:			B24-12
E24-28	Product (invented) name	maa:authorisedProducts/maa:product/rdm:pharmaproduct/rdm:invented-name	Medicinal Product > medicinal product name	B24-12
E24-29	Pharmaceutical form(s)	maa:authorisedProducts/maa:product/rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-	Medicinal Product > Pharmaceutical Dose Form CTL, Pharmaceutical	B24-12

		product/rdm:pharmaceutical-form-names/rdm:pharmaceutical-form-name	Product > Pharmaceutical Dose Form CTL	
E24-30	Strength(s)	maa:authorisedProducts/maa:product/rdm:pharmaproduct/rdm:pharmaceutical-products/rdm:pharmaceutical-product/rdm:pharmaceutical-form-details/rdm:pharmaceutical-form-strengths/rdm:pharmaceutical-form-strength-description	Ingredient	B24-12
E24-31	Marketing authorization holder	maa:authorisedProducts/maa:product/rdm:mp-authorisation/rdm:authorisation-holder	Role>Party>Person>given name	B24-12
E24-32	Marketing authorisation number	maa:authorisedProducts/maa:product/rdm:mp-authorisation/rdm:authorisation-numbers/rdm:authorisation-number	MP Authorisation > authorisation number	B24-12
E24-33	Date of authorization	maa:authorisedProducts/maa:authDate	MP Procedure > procedure number	B24-12
E24-34	Therapeutic indication	maa:authorisedProducts/rdm:product/rdm:pharmaproduct/rdm:therapeutic-indications/rdm:therapeutic-indication		B24-12
E24-35	Is the medicinal product, subject of this application, considered as "similar" to any of the authorised Orphan medicinal product(s)? (as defined in Article 3 of Commission Regulation (EC) No 847/2000)			
E24-36	No (module 1.7.1 to be completed)	maa:similar(Value=0)	Orphan Designation > is mp similar to authorised	B24-6
E24-37	Yes (modules 1.7.1 and 1.7.2 to be completed)	maa:similar(Value=1)	Orphan Designation > is mp similar to authorised	B24-6

Element Tree Diagram

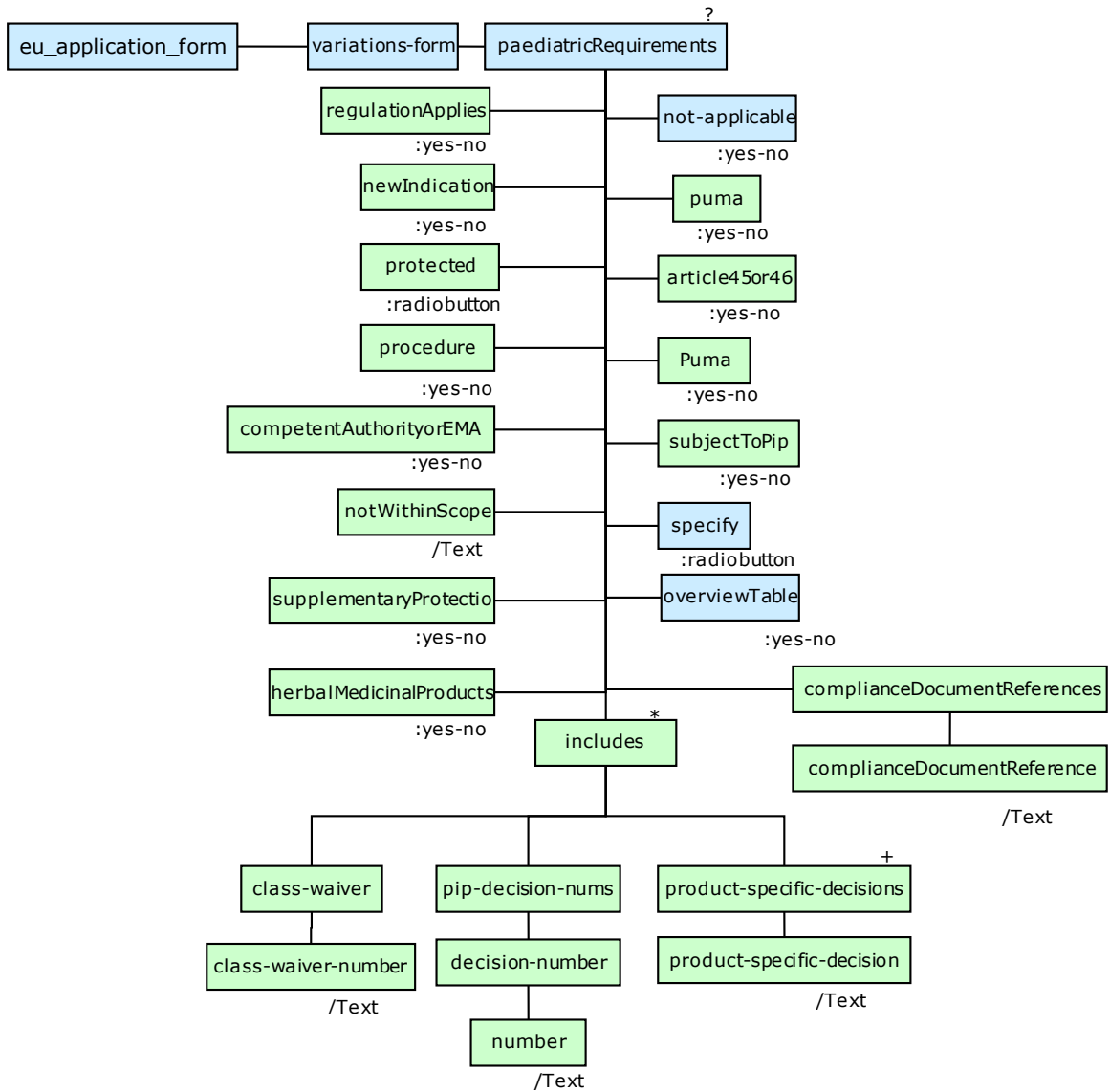


Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B24-1	E21-18, E21-19, E21-21, E21-22	This section is optional	If E21-18 or E21-19 are selected, and E21-21 or E21-22 is selected, then this section(2.4) is available	
B24-2	E24-3, E24-4	Mandatory	Mutually Exclusive	
B24-3	E24-6, E24-7, E24-14, E24-17	Mandatory	Mutually Exclusive	
B24-4	E24-21, E24-22	Mandatory	Mutually Exclusive	
B24-5	E24-20, E24-21	Mandatory	Mutually Exclusive	
B24-6	E24-36, E24-37	Mandatory	Mutually Exclusive	
B24-7	E24-1	Mandatory	If selected, this section(2.4) is not available	
B24-8	E24-3, E24-5	E24-3 is Mandatory, E24-5 is Optional	If E24-3 is selected, then E24-5 is required	
B24-9	E24-7, E24-8 to E24-13	E24-7 is Mandatory, E24-8 is Optional	If E24-7 is selected, then E24-8 to E24-11 are required.	
B24-10	E24-10, E24-11	Mandatory	Mutually Exclusive	
B24-11	E24-21, E24-23 to E24-21	E24-21 is Mandatory, E24-23 to E24-26 are Optional	If E24-21 is selected, then E24-23 to E24-26 are required	
B24-12	E24-25, E24-26, E24-27 to E24-34	E24-25, E24-26 is Mandatory, E24-27 to E24-34 are Optional	If E24-25 is selected, then the rest are required.	
B24-13	E24-14, E24-15 and E24-16	E24-14 is Mandatory, E24-15 and E24-16 is Optional	If E24-14 is selected, then E24-15 and E24-16 are required.	
B24-14	E24-17, E24-18	E24-17 is Mandatory, E24-18 is Optional	If E24-17 is selected, then E24-18 is required.	

2.5. Type IB & Type II variations - Paediatric requirements

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:variations-form/ maa:paediatricRequirements/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E25-1	Select flag if not applicable; section will not be displayed.	maa:not-applicable		B25-3
E25-2	Article 8 of paediatric regulation applies to this variation application since.	maa:regulationApplies	Paed Regulation App CTL	B25-1,B25-4
E25-3	This application relates to a new indication for an authorised medicinal product which:	maa:newIndication	Paed Regulation App CTL	B25-4, B25-5, B25-6
E25-4	is protected by a supplementary protection certificate under Regulation (EC) No 469/2009.	maa:protected (Value=1)	Paed Regulation App CTL	B25-6, B25-7
E25-5	is protected by a patent which qualifies for the granting of the supplementary protection certificate.	maa:protected (Value=2)	Paed Regulation App CTL	B25-6, B25-7
E25-6	This application relates to a previous/ongoing/parallel procedure which triggered Article 8 requirement.	maa:procedure	Paed Regulation App CTL	B25-4, B25-5
E25-7	Competent authority/EMA procedure number	maa:competentAuthorityor EMA	Paed Regulation App CTL	B25-9
E25-8	Article 8 of the paediatric regulation does not apply to this application, since.	maa:notWithinScope	Paed Regulation App CTL	B25-1, B25-8
E25-9	The authorised medicinal product is not protected by a supplementary protection certificate under Regulation (EC) No 469/2009 or by a patent which qualifies for the granting of the supplementary protection	maa:supplementaryProtec tion	Paed Regulation App CTL	B25-8
E25-10	It relates to a well-established use, generic, hybrid, bio-similar marketing authorisations or traditional herbal medicinal products	maa: herbalMedicinalProducts	Paed Regulation App CTL	B25-8
E25-11	This application relates to a new indication for a paediatric use marketing authorisation (PUMA).	maa:puma	Paed Regulation App CTL	
E25-12	This application relates to paediatric studies submitted according to Article 45 or 46 of the paediatric regulation.	maa:article45or46	Paed Regulation App CTL	
E25-13	This application relates to paediatric studies included in a paediatric investigation plan	maa:articlePIP	Paed Regulation App CTL	B25-1
E25-14	This application includes:			
E25-15	PIP Decision Number	maa:includes/ maa:pip-decision-nums/ maa:decision-number/ maa:number	Paediatric Designation > art 7 pip decision number	
E25-16	Product-Specific Waiver Decision Number	maa:includes/maa:product -specific-decisions/ maa:product-specific- decision	Paediatric Designation > art 7 product waiver number	
E25-17	Class Waiver decision Number	maa:includes/ maa:class-waiver/ maa:class-waiver-number	Paediatric Designation > art 7 class waiver number	
E25-18	Has this application been subject to PIP compliance verification?			
E25-19	Yes	maa:subjectToPip (Value=1)	Paediatric Designation > subject to pip compliance	B25-2
E25-20	No	maa:subjectToPip (Value=2)	Paediatric Designation > subject to pip compliance	
E25-21	Please specify			B25-2
E25-22	The compliance document reference	maa:complianceDocument References/ maa:complianceDocument Reference	Paediatric Designation > nca emea doc reference	B25-2
E25-23	Please provide the overview table of PIP results in Module 1.10	maa:overviewTable		

Element Tree Diagram

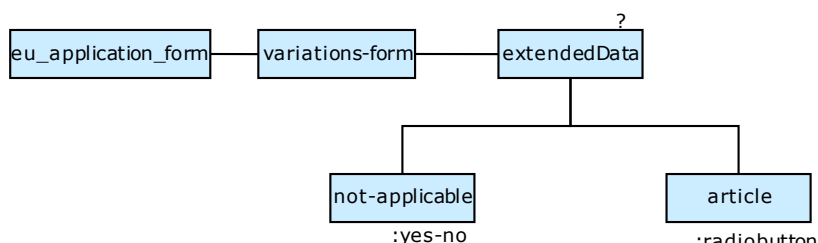


BusinessRules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B25-1	E25-2, E25-8, E25-13	Mandatory	Mutually Exclusive	
B25-2	E25-19, E25-22	Optional	If E25-19 is selected, then E25-22 is required	
B25-3	E25-1	Optional section	If E25-1 is selected, then this section is not displayed	
B25-4	E25-2, E25-3, E25-6	E25-2 is Mandatory, rest are optional	If E25-2 is selected, then the rest are required.	
B25-5	E25-3, E25-6	Optional	Mutually Exclusive	
B25-6	E25-3, E25-4, E25-5	Optional	If E25-3 is selected, then E25-4 and E25-5 are mandatory.	
B25-7	E25-4, E25-5	Optional	Mutually Exclusive	
B25-8	E25-8, E25-9, E25-10	E25-8 is Mandatory, rest are Optional	If E25-8 is selected, then E25-9 and E25-10 are required	
B25-9	E25-6, E25-7	E25-7 is optional	If E25-6 is selected, then E25-7 is required	

2.6. Type II variations – Extended data exclusivity/market protection

Common DES 3.0 Context			Common RDM Entry point	
maa:eu_application_form/maa:variations-form/maa:extendedData/			Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E26-1	Select flag if not applicable; section will not be displayed.	maa:not-applicable		B26-1
E26-2	Consideration of this application is also requested under the following article in directive 2001/83/EC or regulation (EC) No 726/2004:			
E26-3	Article 10(1) of Directive 2001/83/EC / Article 14(11) of Regulation (EC) No 726/2004 (one year of market protection for a new indication)	maa:article (Value=1)		
E26-4	Article 10(5) of Directive 2001/83/EC (one year of data exclusivity for a new indication)	maa:article (Value=2)		
E26-5	Article 74(a) of Directive 2001/83/EC (one year of data exclusivity for a change in classification)	maa:article (Value=3)		

Element Tree Diagram



BusinessRules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B26-1	E26-1	Optional section	If E26-1 is selected, then this section is not displayed	
B26-2	E21-18, E21-19	Optional Section	If any of E21-18 and E21-19 are selected, then this section is displayed	

2.7. Change to the design or indented purpose of the medical device component, or a new device/ device constituent part

Common DES 3.0 Context			Common RDM Entry point	
maa:eu_application_form/maa:variations-form/maa:VariationPurpose/			Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
	Does this variation application refer to:			
E27-1	a change to the design or intended	maa: change-to-		

	purpose of a device component previously listed in the marketing authorisation of the medical product. Please explain the purposed changed in present/purposed section	device/ change-to-device-cb		
	What is the type of device affected by this variation?			
	Note: Please tick the applicable statement and duplicate this section as needed for each device component used with the medicinal product.			
	a) medical device which incorporates, as an integral part, a medicinal product and the action of that medicinal product is principal and not ancillary to that of the device (Art 1(8), second subparagraph of Regulation (EU) 2017/745)			
E27-2	Yes	maa: change-to-device/Med-devices-checkboxes/Med-devices-checkbox-a		
E27-3	No	maa: change-to-device/Med-devices-checkboxes/Med-devices-checkbox-a		
		maa: change-to-device/Med-devices-checkboxes/Med-devices-subSections-a/.. (common table)		
	b) medical device intended to administer a medicinal product where they form a single integral product which is intended exclusively for use in the given combination and which is not reusable (Art 1(9) second subparagraph of Regulation (EU) 2017/745)			
E27-4	Yes	maa: change-to-device/Med-devices-checkboxes/Med-devices-checkbox-b		
E27-5	No	maa: change-to-device/Med-devices-checkboxes/Med-devices-checkbox-b		
		maa: change-to-device/Med-devices-checkboxes/Med-devices-subSections-b/.. (common table)		
	<i>Note: in accordance with Annex I, Section 3.2, point 12 to Directive 2001/83/EC as amended by Article 117 of Regulation (EU) 2017/745, conformity of the device part with the general safety and performance requirements of Annex I to Regulation 2017/745 should be demonstrated by providing a manufacturer's EU declaration of conformity, a EU certificate issued by a Notified Body or a Notified Body opinion where applicable.</i>			
	c) medical device incorporated as integral part of an ATMP (article 2 (d) of Regulation 1394/2007)			
E27-6	Yes	maa: change-to-device/Med-devices-checkboxes/Med-devices-checkbox-c		
E27-7	No	maa: change-to-device/Med-devices-checkboxes/Med-devices-checkbox-c		
		maa: change-to-device/Med-devices-checkboxes/Med-		

		devices-subSections-c/.. (common table)		
	d) medical device is co-packaged with the medicinal product. <i>Note: the device must comply with Regulation (EU) 2017/745 including being CE-marked.</i>			
E27-8	Yes	maa: change-to-device/Med-devices-checkboxes/Med-devices-checkbox-d		
E27-9	No	maa: change-to-device/Med-devices-checkboxes/Med-devices-checkbox-d		
		maa: change-to-device/Med-devices-checkboxes/Med-devices-subSections-d/.. (common table)		
	e) medical device which is supplied separately but referenced in the product information of the medicinal product			
	<i>Note: the device must comply with Regulation (EU) 2017/745, including being CE-marked</i>			
E27-10	Yes	maa: change-to-device/Med-devices-checkboxes/Med-devices-checkbox-e		
E27-11	No	maa: change-to-device/Med-devices-checkboxes/Med-devices-checkbox-e		
		maa: change-to-device/Med-devices-checkboxes/Med-devices-subSections-e/.. (common table)		
E27-12	a new device introduced in the marketing authorisation of the medical product	maa:new-device/new-device-cb		
	What is the type of device affected by this variation?			
	Note: Please tick the applicable statement and duplicate this section as needed for each device component used with the medicinal product.			
	a) medical device which incorporates, as an integral part, a medicinal product and the action of that medicinal product is principal and not ancillary to that of the device (Art 1(8), second subparagraph of Regulation (EU) 2017/745)			
E27-13	Yes	maa:new-device/Med-devices-checkboxes/Med-devices-checkbox-a		
E27-14	No	maa:new-device/Med-devices-checkboxes/Med-devices-checkbox-a		
		maa:new-device/Med-devices-checkboxes/Med-devices-subSections-a/.. (common table)		
	b) medical device intended to administer a medicinal product where they form a single integral product which is intended exclusively for use in the given combination and which is not reusable (Art 1(9) second subparagraph of Regulation (EU) 2017/745)			
E27-15	Yes	maa:new-device/Med-devices-checkboxes/Med-		

E27-16	No	devices-checkbox-b maa:new-device/Med-devices-checkboxes/Med-devices-checkbox-b		
		maa:new-device/Med-devices-checkboxes/Med-devices-subSections-b/.. (common table)		
	<i>Note: in accordance with Annex I, Section 3.2, point 12 to Directive 2001/83/EC as amended by Article 117 of Regulation (EU) 2017/745, conformity of the device part with the general safety and performance requirements of Annex I to Regulation 2017/745 should be demonstrated by providing a manufacturer's EU declaration of conformity, a EU certificate issued by a Notified Body or a Notified Body opinion where applicable.</i>			
	c) medical device incorporated as integral part of an ATMP (article 2 (d) of Regulation 1394/2007)			
E27-17	Yes	maa:new-device/Med-devices-checkboxes/Med-devices-checkbox-c		
E27-18	No	maa:new-device/Med-devices-checkboxes/Med-devices-checkbox-c		
		maa:new-device/Med-devices-checkboxes/Med-devices-subSections-c/.. (common table)		
	d) medical device is co-packaged with the medicinal product. <i>Note: the device must comply with Regulation (EU) 2017/745 including being CE-marked.</i>			
E27-19	Yes	maa:new-device/Med-devices-checkboxes/Med-devices-checkbox-d		
E27-20	No	maa:new-device/Med-devices-checkboxes/Med-devices-checkbox-d		
		maa:new-device/Med-devices-checkboxes/Med-devices-subSections-d/.. (common table)		
	e) medical device which is supplied separately but referenced in the product information of the medicinal product			
	<i>Note: the device must comply with Regulation (EU) 2017/745, including being CE-marked</i>			
E27-21	Yes	maa:new-device/Med-devices-checkboxes/Med-devices-checkbox-e		
E27-22	No	maa:new-device/Med-devices-checkboxes/Med-devices-checkbox-e		
		maa:new-device/Med-devices-checkboxes/Med-devices-subSections-e/.. (common table)		

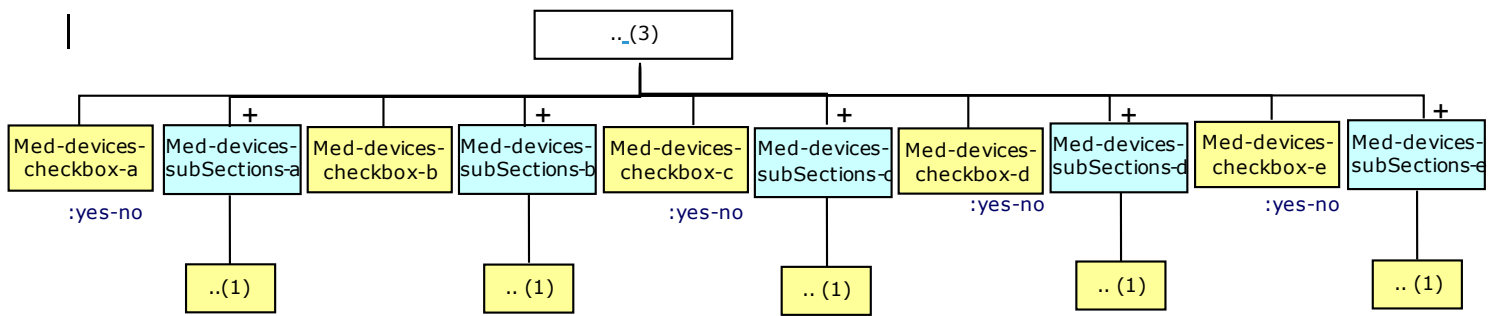
Common table for a,b,c,d,e

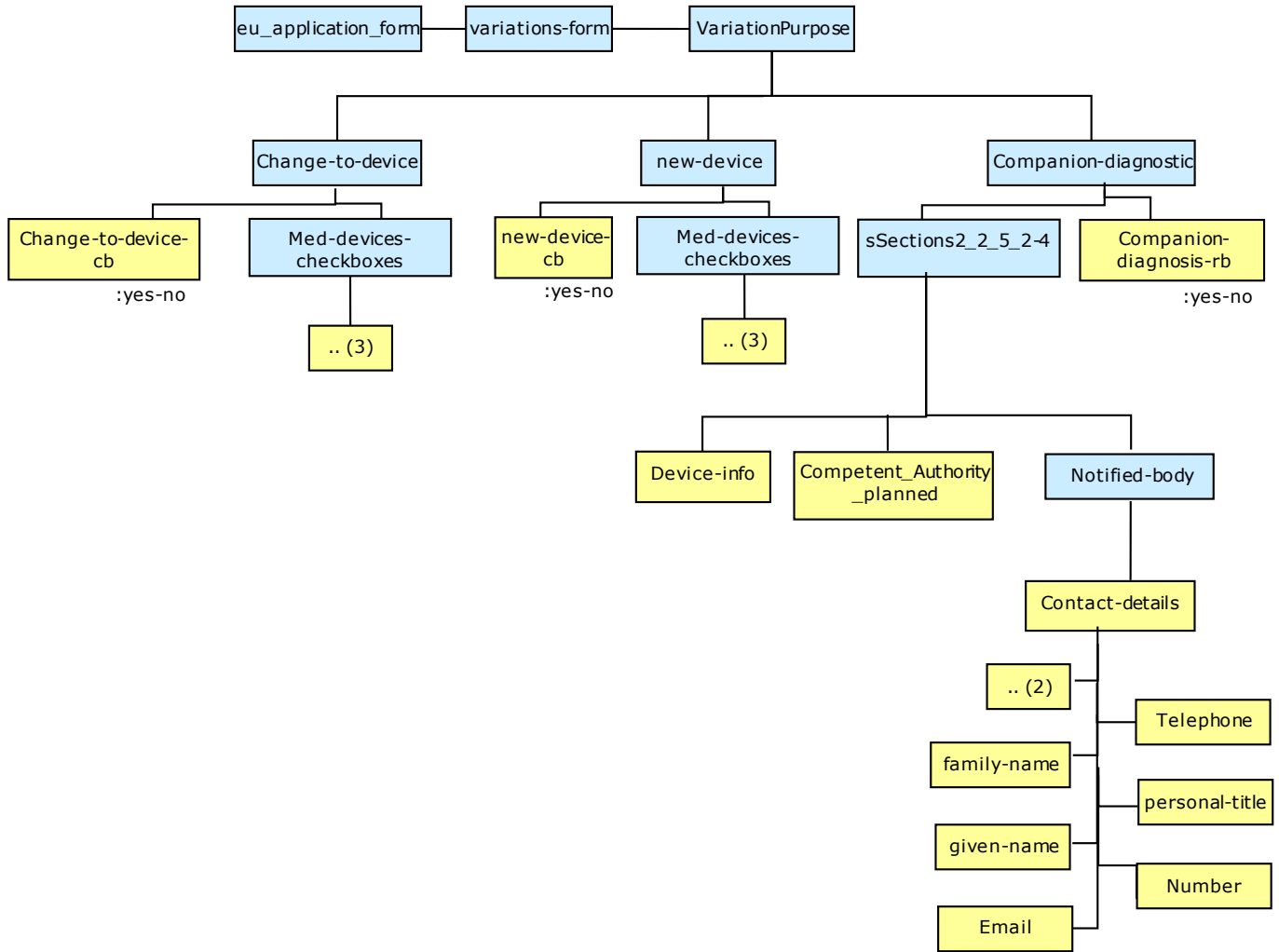
	Device(s) identification and classification			
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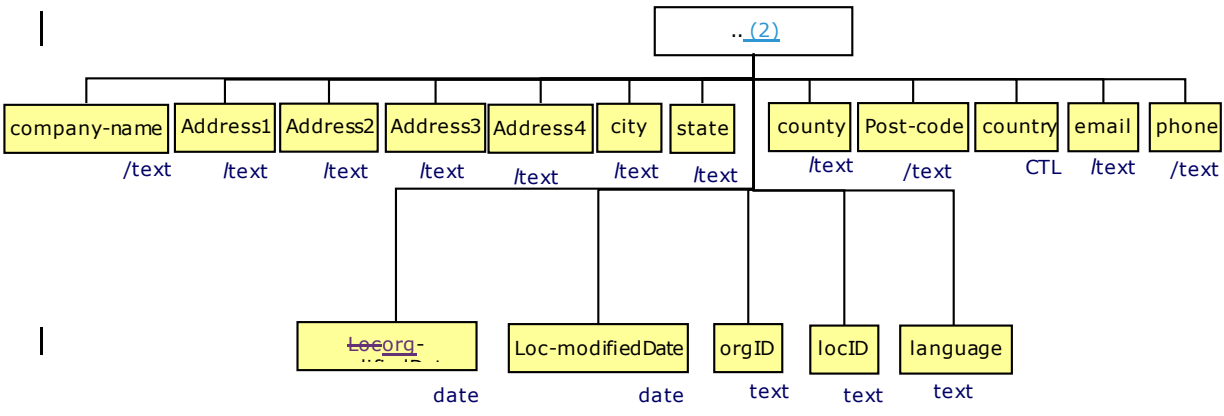
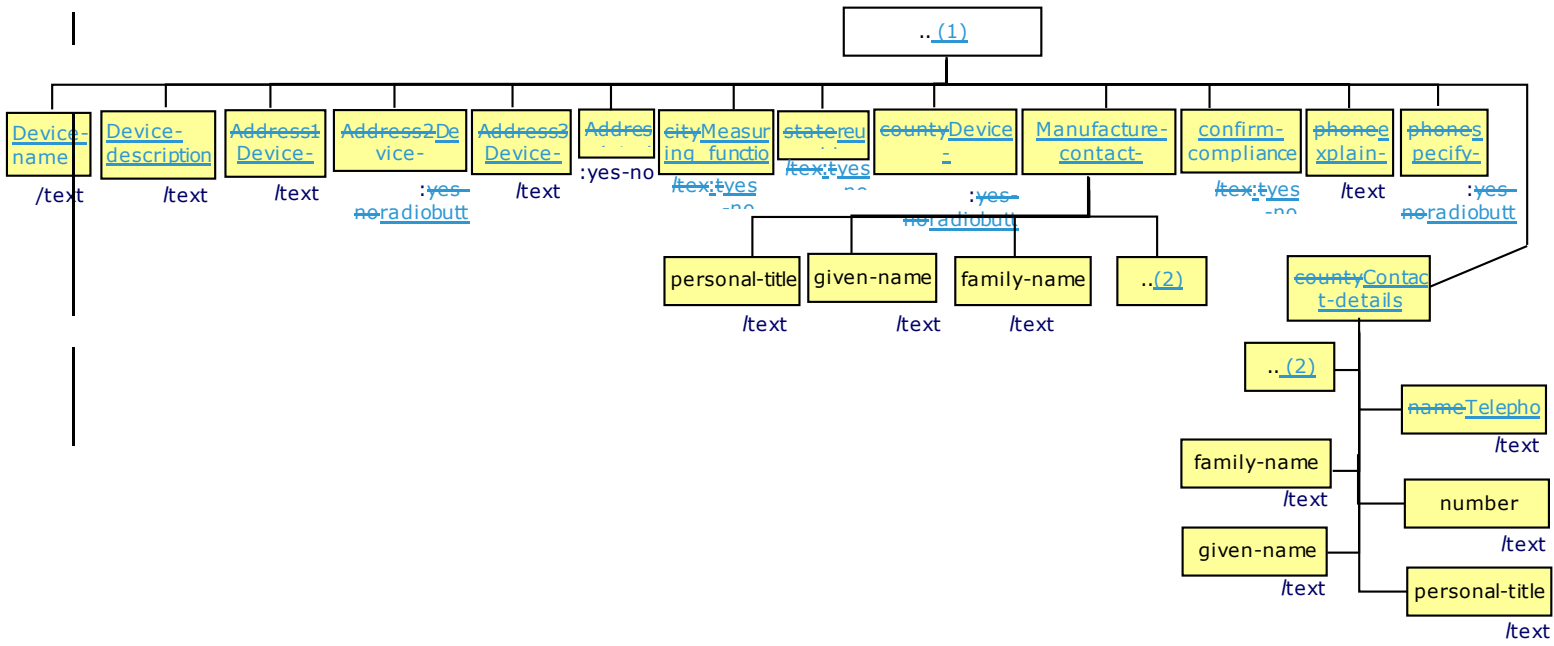
E27-23	Name of the device(s)	../Device-name		
E27-24	Brief description of the device	../Device-description		
E27-25	Intended purpose of the device	../Device-purpose		
	Classification			
E27-26	Class I	../Device-classification(value=1)		
E27-27	Class IIa	../Device-classification(value=2)		
E27-28	Class IIb	../Device-classification(value=3)		
E27-29	Class III	../Device-classification(value=4)		
E27-30	Serial number / unique device identifier (UDI) or other indications necessary to delimit precisely the device incorporated, if applicable	../Device-serial-no		
E27-31	Sterile	../sterile		
E27-32	with measuring function	../measuring_function		
E27-33	reusable surgical instrument	../reusable		
	Manufacturer of the device			
	<i>(Note: for manufacturers outside the EEA, please provide details of the authorised representative instead)</i>			
E27-34	EU Manufacturer	../Device-manufacturer(value = 1)		
E27-35	EU Authorized representative	../Device-manufacturer(value = 2)		
	Name of contact person			
E27-36	Title	../Manufacture-contact-details/ rdm:personal-title	Role>Party>Person> Personal Title	B2321-1
E27-37	First name	../Manufacture-contact-details/ rdm:given-name	Role>Party>Person> given name	B2321-1
E27-38	Surname	../Manufacture-contact-details/ rdm:family-title	Role>Party>Person> family name	B2321-1
E27-39	Company name	../Manufacture-contact-details/ rdm:company-name		
E27-40	Address	../Manufacture-contact-details/ rdm:address1	Role>Party> Contact Details >Address	B2321-1
		../Manufacture-contact-details/ rdm:address2		
		../Manufacture-contact-details/ rdm:address3		
		../Manufacture-contact-details/ rdm:address4		
E27-41	City/Locality/Town/Village	../Manufacture-contact-details/ rdm:city	Role>Party>Contact Details > city	B2321-1
E27-42	State	../Manufacture-contact-details/ rdm:state		
E27-43	County	../Manufacture-contact-details/ rdm:country		
E27-44	Postcode	../Manufacture-contact-details/ rdm:post-code	Role>Party>Contact Details > postcode	B2321-1
E27-45	Country	../Manufacture-contact-details/ rdm:country	Role>Party>Contact Details > Address > Country CTL	B2321-1
E27-46	OrgID	../Manufacture-contact-details/ rdm:orgID		
E27-47	LocID	../Manufacture-contact-details/ rdm:locID		

E27-48	Loc-modifiedDate	../Manufacture-contact-details/ rdm:loc-modifiedDate		
E27-49	Org-modifiedDate	../Manufacture-contact-details/ rdm:org-modifiedDate		
E27-50	language	../Manufacture-contact-details/ rdm:language		
E27-51	Telephone	../Manufacture-contact-details/ rdm:phone	Role>Party>Contact Details > Electronic Contact > electroniccontact	B2321-1
E27-52	E-mail	../Manufacture-contact-details/ rdm:email	Role>Party>Contact Details > Electronic Contact > electroniccontact	B2321-1
	Documentation to confirm compliance to the Medical Device Regulation (EU) 2017/745			
	Does this application include a Manufacturer's EU declaration of conformity, an EU certificate issued by a Notified Body or a Notified Body opinion, if applicable?			
E27-54	No, please explain at which timepoint the relevant document will be provided	../confirm-compliance(Value=0)		
E27-57	Yes	../confirm-compliance(Value=1)		
E27-58		../explain-no		
	If yes, please explain			
E27-59	Manufacturer's EU declaration of conformity (for medical devices not requiring the involvement of a notified body in the conformity assessment)	../specify-yes(Value=1)		
E27-60	EU certificate issued by a Notified Body	../specify-yes(Value=2)		
E27-61	Notified Body opinion	../specify-yes(Value=3)		
	<i>Please note, the above mentioned documents (as applicable) should be provided in module 3.2.R of the EU-CTD.</i>			
	Notified Body			
E27-62	Name of contact person			
E27-63	Title	../Contact-details/ rdm:personal-title	Role>Party>Person> Personal Title	B2321-1
E27-64	First name	../Contact-details/ rdm:given-name	Role>Party>Person> given name	B2321-1
E27-65	Surname	../Contact-details/ rdm:family-name	Role>Party>Person> family name	B2321-1
E27-66	Telephone	../Contact-details/ rdm:Telephone	Role>Party>Contact Details > Electronic Contact > electroniccontact	B2321-1
E27-67	E-mail	../Contact-details/ rdm:Email	Role>Party>Contact Details > Electronic Contact > electroniccontact	B2321-1
E27-68	Notified Body number	../Contact-details/ rdm:Number		
E27-69	Name of the Notified Body	../Contact-details/ rdm:company-name		
E27-70	Address	../Contact-details/ rdm:address1	Role>Party>Contact Details > Address	B2321-1
		../Contact-details/ rdm:address2		
		../Contact-details/ rdm:address3		
		../Contact-details/ rdm:address4		
E27-71	City/Locality/Town/Village	../Contact-details/ rdm:city	Role>Party>Contact Details > city	B2321-1
E27-72	State	../Contact-details/ rdm:state		
E27-73	County	../Contact-details/ rdm:coutry		
E27-74	Postcode	../Contact-details/ rdm:post-code	Role>Party>Contact Details > postcode	B2321-1
E27-75	Country	../Contact-details/ rdm:country	Role>Party>Contact Details > Address > Country CTL	B2321-1
E27-76	OrgID	../Contact-details/ rdm:orgID		

E27-77	LocID	../Contact-details/rdm:locID		
E27-78	Loc-modifiedDate	../Contact-details/rdm:loc-modifiedDate		
E27-79	Org-modifiedDate	../Contact-details/rdm:org-modifiedDate		



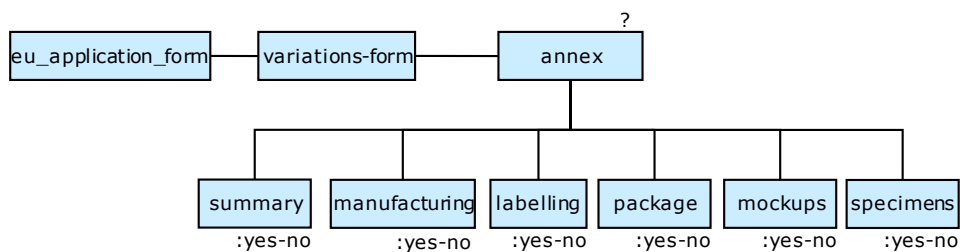




The following amended product information proposals are provided in the relevant sections of the EU-CTD format or NTA volume 6B format, where applicable:

Common DES 3.0 Context		Common RDM Entry point		
	maa:eu_application_form/maa:variations-form/maa:annex/	Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E27-1	Summary of product characteristics	maa:summary		
E27-2	Manufacturing Authorisation Holder responsible for batch release and conditions of the Marketing Authorisation ¹⁵	maa:manufacturing		
E27-3	Labelling	maa:labelling		
E27-4	Package leaflet	maa:package		
E27-5	Mock-ups ¹⁶	maa:mockups		
E27-6	Specimens ¹⁶	maa:specimens		

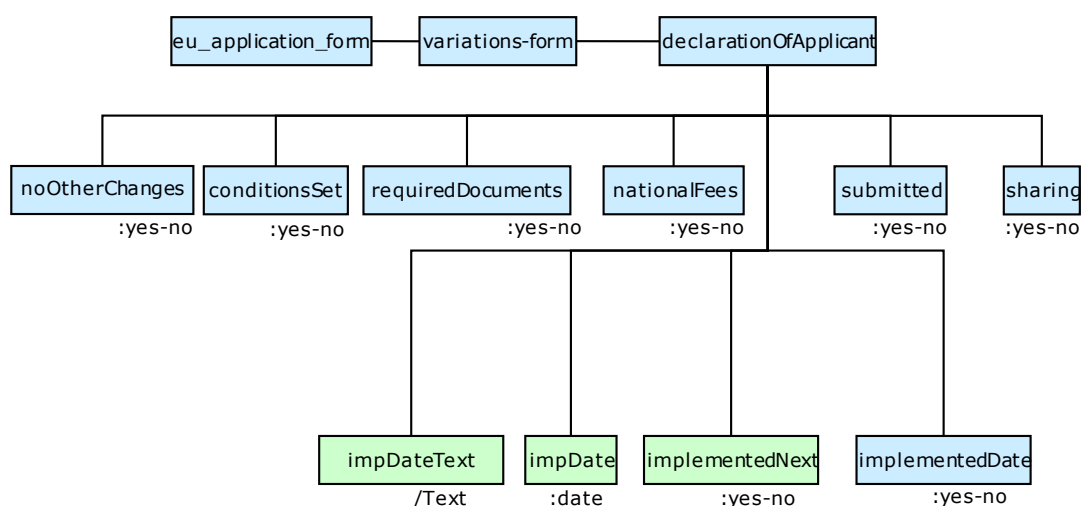
Element Tree Diagram



2.8. DECLARATION OF APPLICANT

Common DES 3.0 Context			Common RDM Entry point	
	maa:eu_application_form/maa:variations-form/maa:declarationOfApplicant/		Application >	
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E28-1	I hereby submit a notification/application for the above Marketing Authorisation(s) to be varied in accordance with the proposals given above. I declare that <i>(Please tick appropriate declarations)</i> :			
E28-2	There are no other changes than those identified in this application (except for those addressed in other variations submitted in parallel);	maa:noOtherChanges		
E28-3	Where applicable, all conditions as set for the variation(s) concerned are fulfilled;	maa:conditionsSet		
E28-4	For type IA notifications: the required documents as specified for the changes concerned have been submitted;	maa:requiredDocuments		
E28-5	Where applicable, national fees have been paid;	maa:nationalFees		
E28-6	This notification/application has been submitted simultaneously in RMS and all CMSs (for products within the Mutual Recognition Procedure and worksharing) or both to EMA and (Co-)Rapporteur (for products within the Centralised Procedure) or, in case of worksharing involving the EMA, to the relevant National Competent Authorities and/or RMS/CMS (as applicable) and the EMA;	maa:submitted		
E28-7	For worksharing or grouped variations affecting more than one MA: the MAs concerned belong to the same MAH.	maa:sharing		
E28-8	Change(s) will be implemented from ¹⁷ :			
E28-9	Next production run/next printing	maa:implementedNext	change from next production	B28-1
E28-10	Date	maa:implementedDate		B28-1,B28-2
E28-11	Date	maa:impDate	change from date	B28-2
E28-12	Free Text field	maa:impDateText		B28-2

Element Tree Diagram



Business Rules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B28-1	E28-9, E28-10	Optional	To be completed only for Type IB and Type II applications	
B28-2	E28-10, E28-12	E28-10 is mandatory, E28-11 and 12 is optional	If E28-10 is clicked, then E28-11 is required	

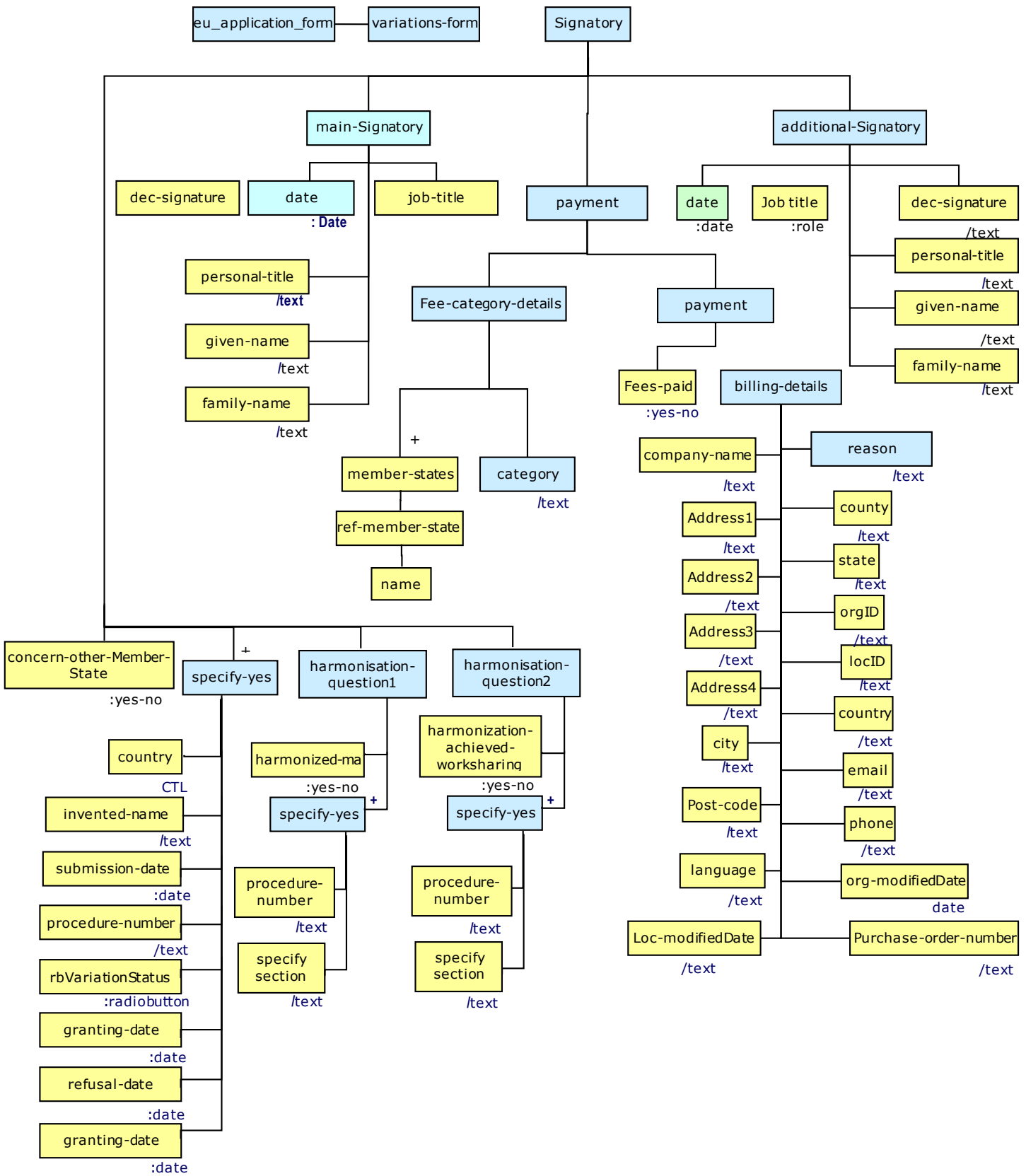
2.9. Signatures

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form/maa:variations-form/maa:Signatory/		Application >		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E29-1	Proof of payment (when relevant)			B29-1
E29-2	Have all relevant fees been prepaid to competent authorities?			B29-1,
E29-3	Yes	maa:payment/maa:payment/maa:fees-paid (1)		B29-1, B29-2, B29-3
E29-4	No	maa:payment/maa:payment/maa:fees-paid (0)		B29-1, B29-2, B29-3
E29-5	For Member States	maa:payment/maa:fee-category-details/rdm:member-states/rdm:ref-member-state/rdm:name		B29-1, B29-2, B29-3
E29-6	Please specify fee category under National rules	Maa:payment/maa:fee-category-details/maa:category		B29-1, B29-2, B29-3
E29-7	Please specify the reasons according to National requirements (exemption or later payment).	maa:payment/maa:payment/maa:billingdetails/maa:reason		B29-1, B29-2, B29-3
E29-8	Billing address (when relevant)			B29-1, B29-2, B29-3
E29-9	For Member States	maa:payment/maa:payment/maa:billing-details/rdm:member-states/rdm:ref-member-state/rdm:name		B29-1, B29-2, B29-3
E29-10	Company Name	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/maa:company-name		B29-1, B29-2, B29-3
E29-11	VAT Number	maa:payment/maa:payment/maa:billing-details/maa:VAT-number		B29-1, B29-2, B29-3
E29-12	Address1	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:address1	Role>Party>Contact Details > Address	B29-1, B29-2, B29-3
		maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:address2		
		maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:address3		
		maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:address4		
E29-13	city	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:city	Role>Party>Contact Details > city	B29-1, B29-2, B29-3
E29-13a	state	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:state		
E29-13b	county	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:county		
E29-14	Postcode	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:postcode	Role>Party>Contact Details > postcode	B29-1, B29-2, B29-3
E29-15	Country	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:country	Role>Party>Contact Details > Address > Country CTL	B29-1, B29-2, B29-3
E29-15a	orgID	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:orgID		
E29-15b	locID	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:locID		

E29-15c	Loc-modifiedDate	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:loc-modifiedDate		
E29-15d	Org-modifiedDate	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:org-modifiedDate		
E29-15e	Language	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:language		
E29-16	Telephone	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:phone	Role>Party>Contact Details > Electronic Contact > electronic contact	B29-1, B29-2, B29-3
E29-18	E-mail	maa:payment/maa:payment/maa:billing-details/rdm:contact-details-type/rdm:email	Role>Party>Contact Details> Electronic Contact > electronic contact	B29-1, B29-2, B29-3
E29-19	Purchase Order (PO) Number	maa:payment/maa:payment/maa:billing-details/Purchase-order-number		B29-1, B29-2, B29-3
	Declaration of the applicant about submission(s) of the same type II variation application for the same product in other Member States – for MRP/DCP/purely nationally authorised products			
	Does the same variation application to the same product concern any other Member State(s)?			
E29-20	Yes	maa:payment/concem-other-Member-State(Value=1)		
E29-21	No	maa:payment/concem-other-Member-State(Value=0)		
E29-22	Country	maa:payment/specify-yes/country		
E29-23	Invented name	maa:payment/specify-yes/invented-name		
E29-24	Date of submission	maa:payment/specify-yes/submission-date		
E29-25	Procedure number (if applicable)	maa:payment/specify-yes/procedure-number		
E29-26	Variation pending	maa:payment/specify-yes/rbVariationStatus (Value=1)		
E29-27	Variation accepted	maa:payment/specify-yes/rbVariationStatus (Value=2)		
E29-28	Variation refused	maa:payment/specify-yes/rbVariationStatus (Value=3)		
E29-29	Variation withdrawn	maa:payment/specify-yes/rbVariationStatus (Value=4)		
E29-30	Date of granting	maa:payment/specify-yes/granting-date		
E29-31	Date of refusal	maa:payment/specify-yes/refusal-date		
E29-32	Date of withdrawal	maa:payment/specify-yes/withdrawal-date		
	Information on harmonisation of product information for MRP/DCP/purely nationally authorised products:			
	1) Has/have the concerned MA(s) been harmonised or partially harmonised, by an Article 30 or 31(1) referral?			
E29-33	Yes	maa:payment/harmonization-question1/harmonized-ma(Value=1)		
E29-34	No	maa:payment/harmonization-question1/harmonized-ma(Value=0)		
E29-35	Procedure(s) reference number	maa:payment/harmonization-question1/specify-		

E29-36	Specify the section(s) of SmPC/PIL/labeling	yes/procedure-number maa:payment/harmonization-question1/specify-yes/specify-section		
	2) Has harmonisation of a section/some sections of the SmPC/PIL/labelling been achieved through a variation worksharing?			
E29-37	Yes	maa:payment/harmonization-question2/harmonized-achieved-worksharing (Value=1)		
E29-38	No	maa:payment/harmonization-question2/harmonized-achieved-worksharing (Value=0)		
E29-39	Procedure(s) reference number	maa:payment/harmonization-question2/specify-yes/procedure-number		
E29-40	Specify the section(s) of SmPC/PIL/labeling	maa:payment/harmonization-question2/specify-yes/specify-section		

Element Tree Diagram

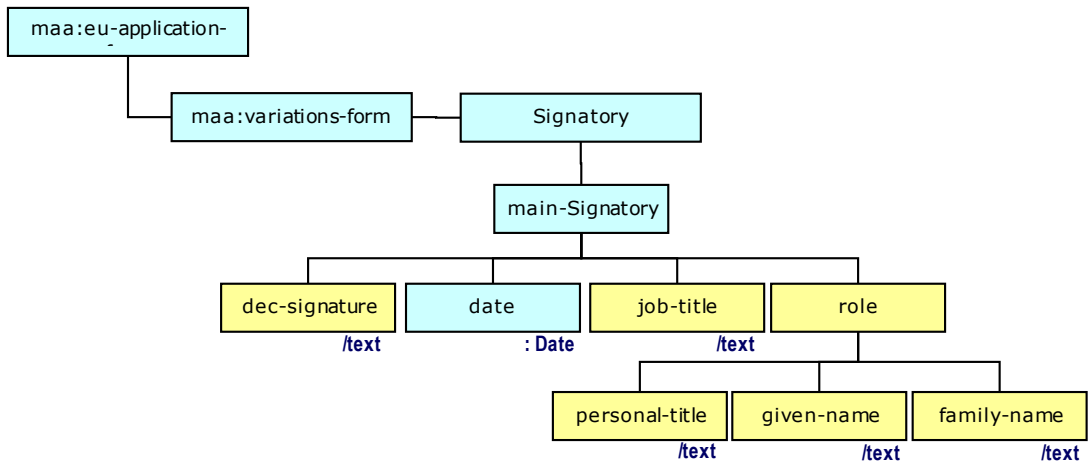


BusinessRules				
Rule ID	Element id(s)	Default BR	Rule	Effect(s)
B29-1	E29.1, E29.2 - E29.19		If E29.1 is selected (value = 1), then E29.2 - E29.19 are visible.	
B29-2	E29.3, E261.4, E29.19		If E29.3 is selected (value = 1), then E29.5 - E29.6 is visible.	If E29.3 is not selected (value = 0), then E29.7 - E29.19 is invisible, rest are visible
B29-3	E29.3, E29.4, E29.19		If E29.4 is selected (value = 0), then E29.7 - E29.19 is visible	If E29.4 is not selected (value = 1), then E29.5 - E29.6 is invisible, rest are visible
B29-4	E29.20 to E29.32		If E21-3 or E21-5 (or both) are selected and at least E21-18 is selected then E29.20 to E29.32 is visible	
B29-5	E29-33 to E29.40		If E21-3 or E21-35 (or both) are selected then E29-33 to E29.40 is visible	

Main Signatory Section

		Common DES 3.0 Context	Common RDM Entry point	
		maa:eu_application_form: variations-form /maa:Signatory/rdm:main-Signatory		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E262.1	Main Signatory	rdm: dec-signature	Application>Role>Party>Person>given name	Is this the same as First name?
E262.2	Title	rdm:role/rdm:personal-title	Application>Role>Party>Person>personal title	
E262.3	First name	rdm:role/rdm:given-name	Application>Role>Party>Person>given name	Is this the same as Main Signatory?
E262.4	Surname	rdm:role/rdm:family-name	Application>Role>Party>Person>family name	
E262.5	Status (Job Title)	rdm:job-title	Application>Role>Party>Person>job title	
E262.6	For worksharing/grouping for more than one MA: the main signatory confirms authorisation to sign on behalf of the designated contacts as specified in section 2.4.3 in Part IA/Module 1 Application Form for each of the MAs concerned.	rdm:signOnBehalf		
E262.7	Date	rdm:date	Application > signature date	

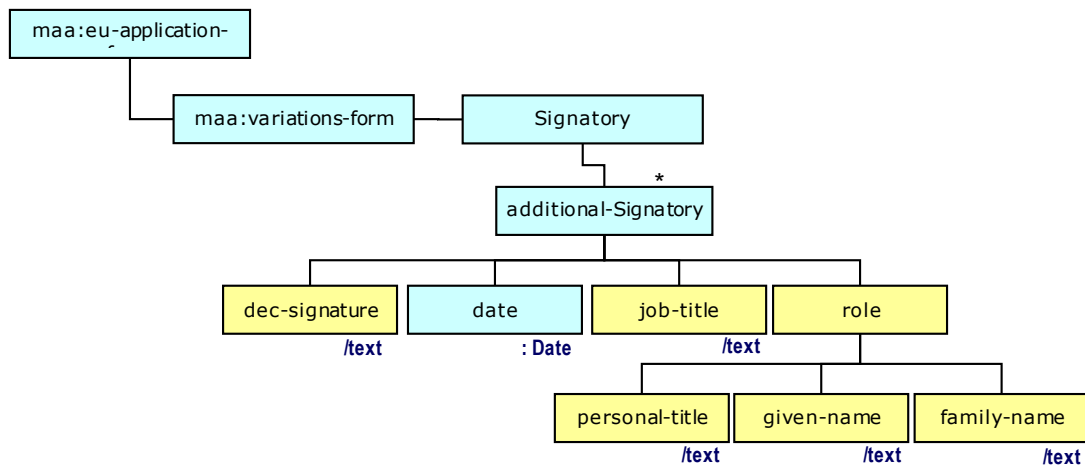
Element Diagram



Additional Signatories Section

Common DES 3.0 Context		Common RDM Entry point		
maa:eu_application_form		maa: variations - form/maa:Signatory/rdm:additional-Signatory		
Elem Id	Label	DES 3.0 Mapping	RDM 3.0 Mapping	Remarks
E263.1	Signatory	rdm: dec-signature	Application>Role>Party>Person>given name	Is this the same as First name?
E263.2	Title	rdm:role/rdm:personal-title	Application>Role>Party>Person>personal title	
E263.3	First name	rdm:role/rdm:given-name	Application>Role>Party>Person>given name	Is this the same as Signatory?
E263.4	Surname	rdm:role/rdm:family-name	Application>Role>Party>Person>family name	
E263.5	Status (Job Title)	rdm:job-title	Application>Role>Party>Person>job title	
E262.6	Date	rdm:date	Application > signature date	

Element Diagram



ABOUT THIS DOCUMENT

Definitions, Acronyms, and Abbreviations

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	eXtensible Markup Language
XSL	XML Stylesheet Language