

Union Product Database (UPD) release notes Referring to version 1.7.2424

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Acronym key and glossary terms

Application Programming Interface	-		
	PET	Veterinary medicinal products intended for animals which are exclusively kept as pets: aquarium or pond animals, ornamental fish, cage birds, homing pigeons, terrarium animals, small rodents, ferrets and rabbits	
API Manager	PMS	Product Management Service	
Availability Status	PSMF	Pharmacovigilance System Master File	
Competent Authority	QPPV	Qualified Person Responsible For Pharmacovigilance	
Centrally Authorised Products	RMS	Reference Member State	
Coordination group for mutual recognition and decentralised procedures for veterinary medicinal products	RN	Release Notes	
Concerned Member State	SIAMED	EMA product information and application tracking system	
Comma-separated values	SIT	System Integration Testing	
Decentralised Procedure	SMS	Substances Management Services	
EMA Account Management	SPOR	Substances, Products, Organisations and Referentials	
European Commission	SRP	Subsequent Recognition Procedure	
European Economic Area	UAT	User Acceptance Testing	
European Medicines Agency	UC	User Case	
End Point	UI	User Interface	
European Union Implementation Guide	UPD	Union Product Database	
Fast Healthcare Interoperability Resources	NCA	National Competent Authority	
Hot Fix	NP	National Procedure	
Health Level Seven	OMS	Organisation Management Service	
JavaScript Object Notation	URN	Uniform Resource Names	
Location identifier	UUID	Universally Unique Identifier	
Marketing Authorisation Holder	VNeeS	Veterinary Non eCTD Electronic Submission	
Master Data Management	VNRA	Variations not requiring assessment	
Mutual Recognition Procedure	VoS XML	Volume of Sales eXtensible Markup Language	
Member State			
Nationally Authorised Products			
	Availability StatusCompetent AuthorityCentrally Authorised ProductsCoordination group for mutual recognition and decentralised procedures for veterinary medicinal productsConcerned Member StateComma-separated valuesDecentralised ProcedureEMA Account ManagementEuropean CommissionEuropean Economic AreaEuropean Medicines AgencyEnd PointEuropean Union Implementation GuideFast Healthcare Interoperability ResourcesHot FixHealth Level SevenJavaScript Object NotationLocation identifierMarketing Authorisation HolderMutual Recognition ProcedureMember State	Availability StatusPSMFCompetent AuthorityQPPVCentrally Authorised ProductsRMSCoordination group for mutual recognition and decentralised procedures for veterinary medicinal productsRNConcerned Member StateSIAMEDComma-separated valuesSITDecentralised ProcedureSMSEMA Account ManagementSPOREuropean CommissionSRPEuropean Medicines AgencyUCEnd PointUIEuropean Union Implementation GuideUPDFast Healthcare Interoperability ResourcesNCAHot FixNPHealth Level SevenOMSJavaScript Object NotationURNLocation identifierUUIDMarketing Authorisation HolderVNeeSMaster Data ManagementVOS XMLMutual Recognition ProcedureVoS XML	

The structure of these release notes has been refined and simplified for enhanced accessibility to all users. The document contains now 3 sections and 4 annexes. It should be noted that specific segments have been excised, owing to their availability within other documents (such as the EU IG).

Overview of key changes:

With every new release, the UPD release notes are updated to highlight to the user the changes compared to previous versions by detailing new/updated functionalities and/or issues that have been resolved, are known, and/or are newly reported.

Resolved issues since the previous release (UPD version 1.7.2417, released on 24 April 2024)	36
Known Issues	30
Next release's expected date	1 July 2024 (for further details please refer to Annex 3)

Overview of new functionality(ies):

- VNRA Submissions (MAHs)
 - MAHs can save an incomplete VNRA submission via a user interface and resume it within 30 days. For further details, watch the "How to save and resume draft VNRA submissions" video tutorial available on <u>UPD video tutorials for MAHs</u>.

Notes:

- A new form called "*Email Configuration*" might be visible to the MAHs & CAs Super users, but these users **should not use it** yet, because the development of "Receiving UPD notifications by email" is underway and will be available to use soon.
- In case of receiving an error file after the Availability Status (AvS) submission, MAHs are advised to follow these steps:
 - If the errors in the file are due to business validations (see section 4.3.2 of <u>Vet EU</u> <u>IG Chapter 7</u>), fix the errors and resubmit the file.
 - **If the file contains ER.36** (see section 4.3.1 of <u>Vet EU IG Chapter 7</u>), then you may receive two types of error files:
 - In the first case, no updates have been processed successfully. This can be evidenced by the fact that the last column in the error report only contains ER.36 and values of type 'N/A'. In this case, capture the ER.36 errors in an Excel or CSV file and submit it as a ticket to EMA Service Now: https://support.ema.europa.eu/, and then resubmit the part of the file containing values 'N/A'.
 - In the second case, some updates have been processed successfully. This can be evidenced by the fact that the last column in the error report contains ER.36 values and values of the type 'Database updated -Submission 0000 - Product 00000'. In this case, just capture the ER.36 errors in an Excel or CSV file and submit it as a ticket to EMA Service Now: https://support.ema.europa.eu/. No need to resubmit the part of the file containing values type 'Database updated - Submission 0000 - Product 00000'.
 - Once the errors of type ER.36 have been addressed, incorporate the AvS of those products into the next submission, and if you again receive any error repeat all the above steps.

Over time, as ER.36 issues are cleaned up, the size of the carry forward from month to month should diminish in size and eventually disappear.

For information:

- Between August and September 2024, marketing authorisation holders will be required to manually add in UPD the relevant email addresses of the QPPV for their products, without submitting a VNRA. Currently, UPD only contains the name and location of the QPPV. For the purpose of Regulation (EU) 2024/568 on fees and charges payable to the European Medicines Agency which will become applicable from 1 January 2025, advice notes, chargeable units line listing(s) and communications will be sent to the QPPV email address available in UPD. Please note that the feature will be available for a limited time and is only for the addition of the QPPV email address. Any other change to QPPV details (name and/or location) must follow the C.1 VNRA process. After this period, any necessary changes to QPPV details including the addition or update of email address, must follow the C.1 VNRA process. Detailed information will be communicated in advance via email to all industry users registered in the UPD in the upcoming weeks. A separate information will be sent to all NCA users too.
- New naming convention for releases: after version 1.6.42, the new format will include the year, quarter and sprint. For example, in release 1.7.2413, 24 stands for the year; 1 for the quarter, and 3 for the sprint.
- VNRA supergrouping: due to the complexity of the functionality, MAHs are strongly advised to read the revised <u>CMDv Best Practice Guide for Variations Not Requiring</u> <u>Assessment</u> and watch the "How to submit VNRA Supergrouping" video tutorial available on <u>EMA's UPD webpage</u>.
- **Updates of legacy data**: for some of the products approved under DCP/MRP, it could be the case that only one RMS and no CMS(s) are involved in the process. Given that the current implementation of the UPD does not support this scenario, the workaround for recording and updating these products will be as follows:
 - **Step 1)** the RMS creates the DCP adding as CMS a country belonging to EEA (this country should preferably have very few CMSs and no RMS products).
 - **Step 2)** to prevent the product from being available to the general public and the MAH, the CMS will not update the national part of the product
 - **Step 3)** finally, the CMS product will be nullified by the CMS once UPD allows having these products with only one RMS.

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1. Summary of issues

1.1. Resolved issues

Use Case	Affects API and/or UI	Issue reference (ADO)	Resolved issues
All UC	NCA UI & MAH UI	83277	Clarification: in the previous release notes for version 1.7.2417 published on 24 April 2024 it is written that this ticket has been closed because the issue is considered obsolete. This is not the case and therefore the following clarification is required: Some CAPs with status of Withdrawn or Surrendered have been loaded into UPD from EMA's source system (SIAMED) with status of Valid. This ticket has been replaced by the User Story 162162 (see Annex 2 below) which will cover all cases where for CAP products the MA status has been loaded to UPD incorrectly.
UC01 Create product	API	82249	Validation in all resources of URN UUID for full URL attribute: letters allowed are only a to f to form the hexadecimal set from 0 to f pattern of 8-4-4-4-12 The post may not be rejected or may not give an error message that clearly identifies this as being the issue. This ticket has been closed as it has never been reported by an API user since UPD went live.
UC01 Create product	API	168950	Create Homeopathic Product via API: if payload contains attributes with CMS information this is accepted and the information stored. These attributes should either give validation error or be ignored as not applicable for this procedure type.
UC01 Create product	NCA UI	171860	When creating a product with multiple ingredients, the unit of presentation dropdown does not present any options for the subsequent ingredients.
UC03 Search product	API	123745	API user was not able to search and view products and receives 403 invalid query and 403 Product is NOT in user affiliations response.

UC03 Search product	API	83332	API user only: A search of products using parameters of _lastUpdated works as expected. User can define an date interval on the search.
UC01 Create product UC08 Update product	API & NCA UI	82570	UC01 Create & UC08 Update Product - POST should be valid where Reference Strength is populated but there is no Substance Strength; or if specify Substance Strength a Reference Substance and no Reference Substance Strength. Instead, there is a validation error and Substance Strength must always be specified. This error has not been reported by any user and so was considered obsolete.
UC01 Create product UC08 Update product	NCA UI	83327	If product contains two or more Pharmaceutical products, the labels are not properly formatted on the View product screen. The case where two or more Pharmaceutical products should link to the same Ingredient to be considered and review documentation. This was considered not a bug and Data Quality Framework will be dealing with products currently facing this situation.
UC01 Create product UC08 Update product	API & NCA UI	147296	CA should only be able to add Availability status entry for the same country as the Authorisation country of their product.
UC01 Create product UC08 Update product	NCA UI	82452	All procedure types - leading and trailing spaces in free-text fields should be removed by the system before validation.
UC01 Create product UC08 Update product	API & NCA UI	82761	The Manufactured Item Quantity will be truncated to 2 decimal places. It should be possible to enter greater precision if required of up to 8 decimal places.
UC03 Search product	NCA UI & MAH UI	173475	MA Number is not displayed in the search results view, if defined at package level.
UC03 Search product	NCA UI & MAH UI	162057	The value for the attribute strength will appear as 'N/A' in the search results table for the cases where the strength of an ingredient was provided in terms of concentration.
UC05 View product	NCA UI & MAH UI	173232	In Package section - (5.6.2.) Manufactured item quantity is missing the Units of measurement in the view
UC06 Submit VNRA	MAH UI	161519	Foreseen Decision Maker dropdown list was not displaying all countries (options).
UC06 Submit VNRA	MAH UI	83112	This ticket has been closed as it is considered obsolete since the expected behaviour of the system today is that a MAH will be blocked by the system when trying to submit VNRAs for automated codes if any of the products within the submission

			belonging to a DC/MR/SR procedure are missing mandatory national data.
UC09 Approve/Reject VNRA	NCA UI	84163	CMS NCA was able to select Approve/Reject checkbox when viewing a VNRA, although the Submit button correctly remained disabled.
UC08 Update product	NCA API	172176	It is possible for a user to remove common data in a national update via API
UC08 Update product	NCA UI	175359	An error was raised while updating a PET product, stating field resource.basis was not available
UC08 Update product	NCA UI	180599	Common data updates were failing for products with no Authorization Status defined.
UC08 Update product	NCA UI	180887	On an National Data update, a valid payload was altered by the system (duplicating Manufactured Item Definitions) rendering it invalid
UC08 Update product	NCA UI	180943	National data update was failing with error message after updating common data by deleting Pharmaceutical Product of DCP product
UC08 Update product	API & NCA UI	109885	Products that had previously been affected by Bug 89511 (replacing Pharmaceutical product removed the Ingredients) cannot be further updated. Datafix for affected products was executed.
UC09 Approve/Reject VNRA	NCA UI	168118	Decision table is not updated if information is added at VNRA level.
UC19 Nullify product	API	82811	API Manager Nullification endpoint: when Try It option is selected the Content-Type request header defaults to application/json and it should be application/fhir+json. Using the default value will give an error.
UC21 Manage Notifications	NCA UI & MAH UI	83254	Date format inconsistent between different actions. Ticket deemed obsolete after implementation of cognitive search.
UC21 Manage Notifications	NCA UI & MAH UI	175866	Notifications for failed VNRA submissions are not generated
UC21 Manage Notifications	NCA UI & MAH UI	171293	Search by product owner is returning wrong results
UC25 Update Availability status	MAH UI	153934	When Availability Status is submitted by MAHs, if an ER.36 is thrown by the system (see Vet EUIG-Chapter 7), the description of the error provided in the csv file does not comply with the format expected, even though is correct

UC25 Update Availability statusMAH UI161155A submission of Availability Status for a DCP/MRP/SR where a CMS was removed and then re-added will faUC25 - Update availability status and placing on the market via UIMAH UI160241Surrendered packages were included in the download file.	il as invalid
	ded AVS csv
UC28 View VNRANCA UI & 175982VNRA PDF information missing for the new codes: C.MAH UIC.10.e)	10.d) and
UC33 Manage Third Country Product MAH UI 159277 Submission of Third country data of the DCP/SRP Pro Names if one of the CMS products is nullified or has a CMS w marketing status set to suspended.	
UC07 - Download Packages and MAH UI 158743 Surrendered packages are included in the downloade file. Submission of Volume Sales via Form Form File.	d VoS csv
Data Fix NCA UI & 83291 Data Fix Parallel Trade products: where Source Memily product had two or more Ingredients, the first Ingred that product was duplicated in the new Parallel trade	dient from
UC33 Manage Third Country ProductMAH UI180823'View Submission of third country product names' retNamesfor new and previous submissions	urns results

1.2. New issues since last release

This table is ordered by Use Case number. This section lists known issues in this release that have not previously been included in the Release Notes. Some issues had existed in a previous release, and some are new issues in this new release.

Use Case	Affects API and/or UI	Issue reference (ADO)	New Issue description	Workaround
UC01	NCA UI	183496	When resuming a product draft, the validation of ATC vet code flag is incorrect	User needs to interact with the element to reset the validation. For instance, remove and re-add the ATC Vet Code check in order to unblock the create button
UC01 Create Product	NCA UI	183127	When creating an Homeopathic product via API, chapter 4 validation rules are incorrect and allow the user to create a product without a package	

UC08 Update Product	NCA UI	183038	Incorrect validation error is raised when adding a package to a homeopathic product	
UC08 Update Product	NCA UI	184011	Timeout waiting for connection from pool	
UC08 Update Product	NCA UI & API	184794	A common update to an MRP product will fail if a user had defined a Marketing Authorization Number at product level, given that the original reference NAP product had Marketing Authorization Number at package level	
UC08 Update Product	NCA API	180967	When updatating a product via API, if MedicinalProductDefinition is not the first resource entry in the payload the version date displays {{date}} in the UI view	With MedicinalProductDefinition set as the first resource in the payload there will be no issues.
UC08 Update Product	NCA UI	182508	On a common data update, a payload with reference to deleted Ingredient in Administrable Product Definition is accepted and invalid data is incorrectly saved	
UC21 Manage Notifications	NCA UI	180324	Notifications are not being generated for the migrated CAP products	
UC21 Manage Notifications	NCA UI	180398	Nullify of a CAP product triggers 2 notifications, the second one incorrectly with an UPDATE action	
UC25 Availability Status	MAH UI	183756	The download starts automatically if the user leaves the VNRA Submission page and opens the AvS download page	
UC28 View VNRA	NCA UI	185240	Incorrect number of submissions shown on VNRA View table when navigating to further pages of the results	When navigating to page 2, system loses the filter by Submission Status. As a workaround the user can explicity select this value to be filterer.

2. User support

API and UI users may seek support by contacting the User Support via <u>EMA Service Now:</u> <u>https://support.ema.europa.eu/</u>.

For the technical team to address your query in a timely manner, please include the following information as appropriate:

- UI: Print screen of the information entered to create a veterinary product (go to your browser settings, select Print (or press Control + P) and "Save as PDF" on your computer
- API: Operational outcome of the unsuccessful task; the request URL and request headers; and for a Create or Update the request body

2.1. Available training materials and guidance

- Webinars
- <u>Video tutorials</u>
- Guidance for National Competent Authorities
- Guidance for Marketing Authorisation Holders
- <u>EU Implementation Guide</u>
- <u>Release notes</u>

3. References

- 1. Registration Process for UPD (See under section 'How to register') (PDF document)
- 2. SPOR API Specification V2 R5 (europa.eu) API specifications for SMS and PMS, based on FHIR
- 3. <u>HL7 FHIR Release 5 Preview 2: the authoritative source for the FHIR specifications used by EMA</u> to implement SMS and PMS API
- 4. Referentials Management System
- 5. <u>Additional information</u> on the Referentials Management System
- 6. Organisations Management System
- 7. Additional information on the Organisations Management System

Annex 1: Overview of functionality and business value

Functionalities provided in this release

	RMS can create DCP products (data and documents)
	RMS can create MRP products (data and documents)
	RMS can create SRP products (data and documents)
	RMS and CMS can complement DCP/MRP/SRP products with national DCP/MRP/SRP data and documents
	RMS can update Common data for DCP/MRP/SRP products (data and documents)
	NCA can create and update NAP products (data and documents)
API	NCA can create & update Registered Homeopathic products (data and documents)
E	NCA can create & update Parallel Trade products (data and documents)
	NCA can create & update Pet products (data and documents)
	NCA can Nullify product
	NCA can Search/view product (data and documents)
	NCA can Search, View and Approve/Reject VNRA submissions
	NCA can Approve/Reject VNRA submissions on behalf of other NCAs when a Supergouping VNRA submission applies
	NCA can View Volume of Sales data

	RMS can create DCP products (data and documents)
	RMS can create MRP products (data and documents)
	RMS can create SRP RMS can create SRP products (data and documents)
	RMS and CMS can complement DCP/MRP/SRP products with national DCP/MRP data (including documents)
	RMS can update Common data for DCP/MRP/SRP products (data and documents)
	NCA can create and update NAP products (data and documents)
	NCA can create & update Registered Homeopathic products (data and documents)
	NCA can create & update Parallel Trade products (data and documents)
NCA UI	NCA can create & update Pet products (data and documents)
	NCA can save and retrieve drafts for product submissions
	NCA can Nullify product
	NCA can Bulk Upload Documents
	NCA can Transfer Marketing Authorisation
	Search/view/export products (data and documents)
	Notifications for Create and Update of products and Other Post-Authorisation Data actions
	View Volume of Sales information
	Search, View and Approve/Reject VNRA submissions
	NCA can Approve/Reject VNRA submissions on behalf of other NCAs when a Supergouping VNRA submission applies
	EMA and EC staff can update CAP products

	Search/view/export products (data and documents)
	Notifications for Create and Update of products and Other Post-Authorisation Data actions
	Download, Submit, and View Volume of Sales information
	Submit VNRA and View VNRA submissions
	Submit Supergouping VNRAs with the selection of the Foreseen Decision Maker that will approve/reject the whole submission on behalf of the others NCAs involved
	Submit updates for Marketing authorisation status
	Download and Submit updates for Availability status
	Submit Products Grouping
	Submit 3 rd country product names

MAH Validation UI

Validate Volume of Sales submission file

Authorisation	Integration with EMA Account Management (EAM) system for CA and Industry (MAH) roles
for NCA & MAH UI	CA users may search and view all Vet products
	MAH users may search and view only products under the responsibility of the organisations the user represents



EMA can maintain messages to appear in banner of UPD UI

Functionality not included in this release

The following functionality is not included in this release.

NCA UI:

• A new form called "Email Configuration" is visible to the Super users, but these users should not use it, because the development of "Receiving UPD notifications by email" is underway and will be available soon

MAH UI:

• A new form called "Email Configuration" is visible to the Super users, but these users should not use it, because the development of "Receiving UPD notifications by email" is underway and will be available soon

Annex 2: Known issues

Issue reference is an internal number used by the UPD Project team when managing issues. It has been included as User Support may refer to this reference number when responding to your queries. In addition, you can include this reference number when contacting user support on this topic and seeking clarification.

Filter the columns to find those tickets relevant to your role and for NCAs whether you are an API or NCA User or both.

This table is ordered by Use Case number.

Use Case	Affects user	Issue reference (ADO)	Known Issue Description	Workaround
All UC	API & NCA UI & MAH UI	143996	CAP: there are now two products for Exzolt and expected there just to be one.	
API Manager	API	82994	API Manager has duplicate Products listed for "UPD API" (v1 and v3 versions of EP); and exposes many EP not intended to be used by API NCA Users. There should only be the one product at this time with v1 Endpoints.	
UC01 Create product	API	83042	Create parallel trade product via API: the GET OperationOutcome response is populating in the DCP format and it was expected would use same pattern as NAP.	
UC01 Create product UC08 Update product	NCA UI & API	154083	(Marketing authorisation application) Legal basis: Vet EU IG Chapter 2 section 1.7.1 : some terms from the RMS list are missing.	
UC03 Search product	MAH UI	107914	After organization merge, MAH cannot find their products in General search, VNRA submission screen or VoS csv (UPD, VNRA or OPAD databases). Note: even though this issue has been in UPD and known for a while, but it has not been well documented in a ticket. The issue is expected to be resolved in the near future.	

UC03 Search product	NCA UI & MAH UI	83234	Search limitations due to FHIR limitation or MS FHIR limitation.	
UC05 View product	NCA UI & MAH UI	83259	When View product QPPV displays as N/A even although the product does have a LOC-ID populated. This affects only some products and may be due to some Data Quality issue in the affected products.	
UC08 Update product	API & NCA UI	79977	All procedure types: if product does not contain any existing value for Responsible Authority or Product Owner, when an update is submitted the new LOC-ID is not saved.	
UC08 Update product	API & NCA UI	152242	CAP product only - after updating product in UPD there is a duplicated Pack size attribute. This duplicate attribute is only seen view Retrieve product via API. Subsequent updates via UPD for affected products are successful.	
UC08 Update product	API	82437	Change to procedure number not saved if existing inline attribute id is not included in the request body.	
UC08 Update product	NCA UI & MAH UI	83142	For CAP products: there are examples where two products have been created and expected just one. This may occur when a new package has been added or package information has been updated. The cause of the issue will be resolved and affected products corrected.	
UC08 Update product	API	82569	UC08 Update SC2 NAP - should reject update with validation error message if MedicinalProductDefinition.id is not populated.	
UC08 Update product	API & NCA UI	144350	Update Common Data DCP/MRP/SRP to remove the last remaining country from the list of Concerned MS fails with Validation error when submitting via NCA UI. There is no validation error if submitted via API. Acceptance criteria and validation required to be reviewed so that API and NCA UI are aligned.	
UC08 Update product	API & NCA UI	83203	Update of product fails with error in GET OperationOutcome of ERR-1003. An error from previous failed update with an incorrect payload prevents a subsequent update.	

UC08 Update product	ETL for CAP products	162162	Retrieve from UPD correct information on 2.5. Authorisation status (MA status) and 2.6. Date of authorisation status change for CAP products	
UC19 Nullify product	API	132758	Nullify Product via API - OperationOutcome ID now has suffix of "-Patch" which is not expected and is potentially a breaking change for API users. When submitting GET OperationOutcome/ID the response code is 499 Client Closed Request. Therefore it is not possible to nullify a product via the API in this release and NCA UI will need to be used.	
UC25 Update Availability status	MAH UI	177951	Submissions of Availability Status fails for some CAP products with Error 36	
UC28 View VNRA	NCA UI	83344	For a VNRA submitted for a product where the Responsible Authority is not correctly populated (for example may have incorrectly been populated with MAH LOC-ID): an NCA User for that Authorisation country is not able to view the VNRA Submission even after the Responsible Authority has been corrected in the product(s) included in the submission.	
UC31 Manage VNRA Submissions via API	API	142804	All of the endpoints for VNRA API https://spor.azure- api.net/upd/api/vnra/v3/vnra-submission fail with 404 Resource Not Found.	

Annex 3: Release Schedule

Environment	Closed from	Closed to	Expected to be open	Description
PROD	23 April 2024	23 April 2024	24 April 2024	Upgrade of UPD to 1.7.2417
UAT	21 May 2024	21 May 2024	21 May 2024	Upgrade of UPD to 1.7.2417-6 (this update contains only hot fixes and does not include other bug fixes nor new features).
PROD	30 May 2024	30 May 2024	31 May 2024	Upgrade of UPD to 1.7.2417-6 (this update contains only hot fixes and does not include other bug fixes nor new features).
UAT	4 June 2024	4 June 2024	5 June 2024	Upgrade of UPD to 1.7.2424
PROD	12 June 2024	12 June 2024	13 June 2024	Upgrade of UPD to 1.7.2424
UAT	20 June 2024	20 June 2024	20 June 2024	Upgrade of UPD to 1.7.2424-5 (additional security in the API Manager layer)
PROD	1 July 2024	1 July 2024	2 July 2024	Upgrade of UPD to 1.7.2424-5 (additional security in the API Manager layer)
UAT/PROD				Q3 2024 releases will be considered at the next Program Increment planning to be held on 3-4 July 2024 and communicated in due course.

Annex 4: Guidance for API users

4.1 UPD API to Maintain Products and Product Documents

4.1.1. Scope of this release for API

- Create DCP based on Chapter 4 Legacy or Chapter 2 rules
- Create MRP based on Chapter 4 Legacy or Chapter 2 rules
- Create SRP based on Chapter 4 Legacy or Chapter 2 rules
- RMS can update Common Data for products under DCP/MRP/SRP (data and documents)
- RMS and CMS can complement DCP/MRP/SRP product with national data
- Create NP & Registered Homeopathic based on Chapter 4 Legacy or Chapter 2 rules
- Update NP & Registered Homeopathic product based on Chapter 4 Legacy or Chapter 2 rules
 - Edit existing, add new, or delete an existing non-mandatory attribute
 - Add new resources. For example: add an Ingredient or add another Package
 - Delete an existing non-mandatory resource. For example: remove an Ingredient
- Create & Update Parallel trade based on Chapter 4 Legacy or Chapter 2 rules
- Create & Update Pet products based on Chapter 4 Legacy or Chapter 2 rules
- Search and retrieve products
- Nullify product
- Upload, search, retrieve, and update Documents (for product under any procedure type)

4.1.2. UPD API supported Product Service endpoints

EP302 Search Product Part and EP305 Get Product Part endpoints are no longer available.

SPOR API Specification v2	API Manager
EP301 Search Product	GET MedicinalProductDefinition - Search for a MedicinalProductDefinition resource or resources
EP303 Get Product	GET MedicinalProductDefinition - Get a MedicinalProductDefinition ID
EP304 Get Product Full	GET Everything Current - Get \$everything for a MedicinalProductDefinition ID
EP306 Get Product Version	GET MedicinalProductDefinition Version - Get version of MedicinalProductDefinition ID
EP306a Get Product Version Full	GET Everything Versioned - Get \$everything for a version of MedicinalProductDefinition ID
EP307 Get Product Versions	GET MedicinalProductDefinition - Get history of MedicinalProductDefinition ID

SPOR API Specification v2	API Manager
EP309 Create Product	 NAP: POST Bundle - Create/Update resources in the bundle DCP: POST dcp-bundle - Submit a Create DCP payload MRP: POST mrp_bundle - Submit a Create MRP payload SRP: POST srp_bundle - Submit a Create SRP payload Registered homeopathic: POST Bundle - Create/Update resources in the bundle Parallel trade: POST ptp-bundle - Create/Update resources in the bundle Pet: POST pet-bundle - Create/Update resources in the bundle Refer to 4.1.5.2. Create and Update endpoints
EP309 Create Product EP311 Update Product for use with any Create or Update	GET OperationOutcome - Get a resource by ID Note: use this to query the outcome of Create or Update when response to Post is "202 Accepted"
EP311 Update Product	NAP: POST Bundle - Create/Update resources in the bundle Update National Data: POST /upd/api/v1/national-data-bundle/ - Submit an Update National Data payload for DCP/MRP/SRP products Update Common Data: POST /upd/api/v1/common-data-bundle/ - Submit an Update Common Data payload for DCP/MRP/SRP products

SPOR API Specification v2	API Manager
EP318 Validate Product	POST Validate Bundle – To validate a bundle and the resources in the bundle Used for all procedure types; for both chapter 2 or legacy validation rules; and for both Create & Update
EP UC19 Nullify Product	POST /upd/api/v1/vmp-nullification/
EP401 Search document	GET DocumentReference - Search for DocumentReference No
EP402 Get/Retrieve document by Id	GET DocumentReference - Get a DocumentReference by Id Note
EP403 Create document	POST DocumentReference - Create a DocumentReference
EP404 Update document by Id	PUT DocumentReference - Update a DocumentReference Please note: API Manager method shows as PUT however please use POST with request header is_update=true.

4.1.3. API Manager product subscription

Any new API users should register a user and subscribe to the product Authorised - UPD API - Milestone 3 (UPD 1.03) in API Manager.

The credentials for this new product can be used for all supported endpoints as listed in section 4.1.2. UPD API supported Product Service endpoints

Refer to the document UPD 01.03 Registration Process for UPD API in Production/UAT listed in the References section.

4.1.4. Apply Chapter 4 Legacy or Chapter 2 Validation rules

When submitting a POST for EP309 Create Product or EP311 Update Product, there is a Request header that is used to specify which validation rules are to be applied.

Please note that each type of update may use a different value for the Key.

Value	Validation rules applied
Request header not included	Vet EUIG Chapter 2
false	Vet EUIG Chapter 2
true	Vet EUIG Chapter 4 Legacy

4.1.5. API EP309 Create, EP311 Update & Nullify product endpoints

4.1.5.1. Request headers applicable for all Create, Update & Nullify POST

When submitting a POST for EP309 Create Product or EP311 Update or Nullify Product, the same Request headers are used for all endpoints that specify the format for the request and response.

Request Header: Key	Values	Purpose
Content-type	application/fhir+xml application/fhir+json	Specifies the format of the request body that is being submitted

Request Header: Key	Values	Purpose
Accept	application/fhir+xml application/fhir+json	Specifies the format for the response body of the POST if there are any validation or other errors

4.1.5.2. Create and Update endpoints

- As specified in SPOR API v2 Specification section 6.4.12
- Refer to API Manager developer portal
- The Request body is a Bundle (type=transaction) of MedicinalProductDefinition and other resources
- For all the Update endpoints, the Bundle should be based on all data in the existing product. This includes Update Common Data DCP/MRP/SRP where all existing National data should also be included in the bundle even although it is only Common data that will be updated
- Create MRP is an update to an existing NP product. The Bundle should be based on all national data in that product, with the additional Common data added, and the procedure type updated to MRP
- Create SRP is an update to an existing DCP/MRP/SRP product. The Bundle should be based on all national data in that product, with the additional Common data added
- Please refer to the example bundles and recommended approach sections

Type and Procedure	POST Endpoint	Request header Key for validation rules	Additional Request header
Create NP	/pms/api/v2	chapter4	
Update NP	/pms/api/v2	chapter4	is update = true
Create DCP	/upd/api/v1/dcp-bundle/	chapter4	

Type and Procedure	POST Endpoint	Request header Key for validation rules	Additional Request header
Update Common Data DCP/MRP/SRP	/upd/api/v1/common-data- bundle/	chapter4	is update = true
Update National Data DCP/MRP/SRP	/upd/api/v1/national-data- bundle/	chapter4	is update = true
Create MRP	/upd/api/v1/mrp-bundle/	chapter4	
Create SRP	/upd/api/v1/srp-bundle/	chapter4	
Create Registered Homeopathic	/pms/api/v2	homeopathicschapter2 = true OR homeopathicschapter4 = true	
Update Registered Homeopathic	/pms/api/v2	<pre>homeopathicschapter2 = true OR homeopathicschapter4 = true</pre>	is update = true
Create Parallel Trade	/upd/api/v1/ptp-bundle/	parallelchapter2 = true OR	

Type and Procedure	POST Endpoint	Request header Key for validation rules	Additional Request header
		parallelchapter4 - true	
Update Parallel Trade	/upd/api/v1/ptp-bundle/	parallelchapter2 = true OR parallelchapter4 - true	is update = true
Create Pet	/upd/api/v1/pet-bundle/	chapter4	
Update Pet	/upd/api/v1/pet-bundle/	chapter4	is update = true
To Validate any Create or Update bundle	/pms/api/v2/\$Validate	Use appropriate request header to apply validation rules based on the procedure type	 Use is update = true when validating the following bundles: Update NP Update Registered Homeopathic Update Parallel Trade Update Pet Update Common Data DCP/MRP/SRP Update National Data DCP/MRP/SRP Create MRP Create SRP

4.1.5.3. Nullify endpoint

Type and Procedure	POST Endpoint	Request header Key for validation rules	Additional Request header
Nullify product	/upd/api/v1/vmp-nullification/	not required	

Content- Type	Request body	
JSON	{	
	"permanentId": "Permanent Identifier"	
	}	
	For example:	
	{	
	"permanentId": "600011984989"	
	}	
XML	<root><permanentid> Permanent Identifier </permanentid></root>	
	For example:	
	<root><permanentid>600011353107</permanentid></root>	

Response to POST:

- Response code 202 Accepted indicates the nullification has been successfully submitted
- Response code 400 Bad request indicates there is a validation error and the Response body will contain error message. For example: "Resource type 'Bundle' with id '600011984989' couldn't be found."

4.1.5.4. Response to POST for Create, Update or Nullify and use of Get OperationOutcome

When POST for Create, Update or Nullify is successful and it cannot be honoured timely it is automatically queued. The Response header **Content-Location** contains an id that can be used to obtain the status of the operation.

Content-Location has two parts: **post-operation/operation-outcome-id**

The status of the operation can be consulted, it is one of:

- QUEUED
- IN_PROGRESS
- MSG_CREATED
- ERROR

Upon successful creation, update or nullification of the medicinal product, the operation outcome will show a status of MSG_CREATED along with the unique Permanent identifier(s) of the product(s).

The endpoint GET OperationOutcome/**operation-outcome-id** is used to query the status of the operation and this should be repeated until it is successful with MSG_CREATED or has ERROR.

The format of the Content-Location is showing in the following table, and the response value can be used for Get OperationOutcome.

POST	Content Location example showing format of the operation-outcome-id	
Create NP	OperationOutcome/baab996e-8e58-4825-89d1-90a8f30458db	
Update NP	OperationOutcome/c2e2275c-141c-4631-a42e-045726d95adb	

POST	Content Location example showing format of the operation-outcome-id
Create DCP	Release 1.6.16 and prior: dcp-operation-outcome/ddb9f96b-10f5-4428-9503-170feb5c58db-DCP
	Release 1.6.20 is now: OperationOutcome/ddb9f96b-10f5-4428-9503-170feb5c58db-DCP
Update Common Data	Release 1.6.16 and prior: common-data-operation-outcome/f4d76850-358a-48f1-a9bb-3fb4b1615bdb-CD
DCP/MRP/SRP	Release 1.6.20 is now: OperationOutcome/ f4d76850-358a-48f1-a9bb-3fb4b1615bdb-CD
Update National Data	Release 1.6.16 and prior: national-data-operation-outcome/b371f2db-dd29-4c60-b6ab-63b0abf95bdb-ND
DCP/MRP/SRP	Release 1.6.20 is now: OperationOutcome/ b371f2db-dd29-4c60-b6ab-63b0abf95bdb-ND
Create MRP	Release 1.6.16 and prior: mrp-operation-outcome/2f89089c-3ad7-4427-9311-7ea491395ddb-MRP
	Release 1.6.20 is now: OperationOutcome/2f89089c-3ad7-4427-9311-7ea491395ddb-MRP
Create SRP	Release 1.6.16 and prior: srp-operation-outcome/cf7af9a9-b34d-4db9-a551-89d40c077306-SRP
	Release 1.6.20 is now: OperationOutcome/cf7af9a9-b34d-4db9-a551-89d40c077306-SRP
Create & Update Registered Homeopathic	OperationOutcome/a588416b-7a0b-40b1-8d03-a88ea4668f8f
Create & Update	OperationOutcome/04b5bc00-16f4-4ea0-b33e-1a95029d8f8f-PTP
Parallel Trade	
Create & Update Pet	OperationOutcome/2664fdf2-6aef-4540-8254-b6df6451b8af-PET

4.1.5.5. Creating products for DCP or Update Common Data if national data is provided

When the RMS submits a request bundle to create DCP products, they should only provide Common Data. Refer to Annex 1 of Vet EU IG Chapter 2.

If any National data attributes are populated in the create request bundle this does not result in a validation error. The products for the RMS and each CMS will be created, and any national data entered will be silently ignored.

The same applies for Update Common Data. The RMS should populate the complete Update bundle for their RMS product containing all existing Common and National Data. Only Common Data will be updated to the RMS product and the CMS products under the Product identifier.

4.1.5.6. Key changes in valid request bundle for create and update

Attribute	Change
None	

4.1.6. API EP309 Create product example request bundles

Examples for EP309 Create Product for NP and DCP. Please note that the purpose of these examples is as illustration of the FHIR attributes to be populated.

The value for MedicinalProductDefinition as a cross referenced product is a valid permanent identifier from UAT.

Please note: example files still to be updated and re-released taking into account that pack size is now mandatory.

Procedure type	Validation rules	Example file
DCP	Chapter 2	UPD_1.6.5-6_DCP_Chpt2_C2_Mandatory_VetIG.JSON
		UPD_1.6.5-6_DCP_Chpt2_C2_Mandatory_VetIG.XML
		UPD_1.6.5-6_DCP_Chpt2_C110_VetEUIG_AllData.JSON
		UPD_1.6.5-6_DCP_Chpt2_C110_VetEUIG_AllData.XML
DCP	Chapter 4 Legacy	UPD_1.6.5-6_DCP_Legacy_C2_Mandatory_VetIG.JSON
		UPD_1.6.5-6_DCP_Legacy_C2_Mandatory_VetIG.XML

Procedure type	Validation rules	Example file
		UPD_1.6.5-6_DCP_Legacy_C110_VetEUIG_AllData.JSON
		UPD_1.6.5-6_DCP_Legacy_C110_VetEUIG_AllData.XML
NAP	Chapter 2	2.2 Authorisation/registration/entitlement number is specified at Product level
		UPD_1.6.1-4_NAP_Chpt2_C2_Mandatory_VetIG_MANumber_AtMedicinalProductLevel.JSON
		UPD_1.6.1-4_NAP_Chpt2_C2_Mandatory_VetIG_MANumber_AtMedicinalProductLevel.XML
		UPD_1.5.1-0_NAP_Chpt2_C110_VetEUIG_AllData_MANumber_AtMedicinalProductLevel.JSON
		UPD_1.5.1-0_NAP_Chpt2_C110_VetEUIG_AllData_MANumber_AtMedicinalProductLevel.XML
		5.5 Marketing authorisation (package level)
		UPD_1.5.1-0_NAP_Chpt2_C111_VetEUIG_AllData_MANumber_AtPackageLevel.JSON
		This example contains 2 packages.
		There are 3 RegulatedAuthorization resources:
		• One with subject reference = MedicinalProductDefinition resource; populated with attributes from Section 2 (Vet EUIG Chapter 2), excluding the marketing authorisation number
		 One with subject reference = 1st PackagedProductDefinition resource; populated with the Marketing authorisation number for Package 1
		 One with subject reference = 2nd PackagedProductDefinition resource; populate with the Marketing authorisation number for Package 2
NAP	Chapter 4 Legacy	UPD_1.6.1-4_NAP_Legacy_C2_Mandatory_VetIG_MANumber_AtMedicinalProductLevel.JSON

Procedure type	Validation rules	Example file
		UPD_1.6.1-4_NAP_Legacy_C2_Mandatory_VetIG_MANumber_AtMedicinalProductLevel.XML
		$eq:upd_1.5.1-0_NAP_Legacy_C110_VetEUIG_AllData_MANumber_AtMedicinalProductLevel.JSON$
		UPD_1.5.1-0_NAP_Legacy_C110_VetEUIG_AllData_MANumber_AtMedicinalProductLevel.XML
NAP	Chapter 4 Legacy	 UPD_1.5.1-0_NAP_Legacy_Cx_ManyAttributesAndResources_MANumberAtMedicinalProductLevel.XML This example contains: 2 or more values for those attributes that are repeatable. For example, Product name, ATC Vet Code, Manufacturing Business Operation
		 2 Packages (PackagedProductDefinition)
		2 Manufactured Items (ManufacturedItemDefinition)
		3 Ingredients (Ingredient)
NAP	Chapter 2	$eq:upd_1.5.1-0_NAP_Chpt2_ExampleForStrengthAsPresentationOrConcentration.XML$
		This example contains Ingredient resources that illustrate how to specify Substance and Reference Strength as either Presentation or Concentration.
NAP	Chapter 2	NAP_Chpt2_Create_BR-178_StrengthFreeTextExample_1.6.22-6.XML
		F178: This example contains Ingredient resources that illustrate how to specify free-text substance or reference substance strength
Registered	Chapter 2	UPD_1.6.1-4_HOM_Chpt2_C2_Mandatory_VetIG_MANumber_AtMedicinalProductLevel.JSON
Homeopathic		UPD_1.6.1-4_HOM_Chpt2_C110_VetEUIG_AllData_MANumber_AtMedicinalProductLevel.JSON
Parallel Trade	Chapter 2	UPD_1.6.8-4_PAT_Chpt2_C2_Mandatory_VetIGI.JSON

Procedure type	Validation rules	Example file
		UPD_1.6.8-4_PAT_Chpt2_C110_VetEUIG_AllData.JSON
Pet	Chapter 2	PET_Chpt2_C2_Mandatory_VetIG_MANumber_AtMedicinalProductLevel_1.6.34-5.json
		PET_Chpt2_C110_AllData_VetIG_MANumber_AtMedicinalProductLevel_1.6.34-5.json

4.1.6.1. Recommended approach to prepare update request bundle

The recommended approach for preparing a request bundle to update a product (any procedure type) is:

- Use the response from EP304 GET MedicinalProductDefinition/{permanent identifier}/\$everything as a starting point
- Add Bundle.entry.request for each resource and update Bundle.type

Attribute	Change
Bundle.type	Must be "transaction"
For every	Bundle.entry.request must also be populated.
Bundle.entry	Bundle.entry.request.method should be:
	PUT to update an existing resource
	POST to add a new resource
	Bundle.entry.request.url should be:
	Same value as Bundle.entry.fullUrl

For example:

	ersion="1.0" encoding="utf-8" <mark>?></mark> xmlns="http://hl7.org/fhir">
	value="600000022531" />
<me< th=""><th>ta></th></me<>	ta>
	<pre><versionid value="1"></versionid></pre>
	<lastupdated value="2021-07-07T08:52:51.607+00:00"></lastupdated>
<th>eta></th>	eta>
<ty< th=""><th>pe value="transaction" /></th></ty<>	pe value="transaction" />
<en< th=""><th>try></th></en<>	try>
	<fullurl value="MedicinalProductDefinition/600000022531"></fullurl>
	<resource></resource>
	<medicinalproductdefinition></medicinalproductdefinition>
	<request></request>
	<method value="PUT"></method>
	<pre><url value="MedicinalProductDefinition/600000022531"></url></pre>
<td>ntry></td>	ntry>
<en< td=""><td>try></td></en<>	try>
	<fullurl value="PackagedProductDefinition/170427"></fullurl>
	<resource></resource>
	<packagedproductdefinition></packagedproductdefinition>
	<request></request>
	<method value="PUT"></method>
	<url value="PackagedProductDefinition/170427"></url>

• DO NOT edit or remove the IDs for each resource and in-line within each resource in the EP304 Get \$everything response

4.1.6.2. How to use Update NP product endpoint and example bundle

Create product via API	POST Bundle	Sample XML bundle used:
---------------------------	-------------	-------------------------

		UPD_1.5.1- 0_NAP_Legacy_C110_VetEUIG_AllData_MANumber_AtMedicinalProductLevel.XML
Check operation outcome	MSG_CREATED message expected containing Permanent identifier	
EP304 Get Product Full	Prepare update bundle based on the response by updating Bundle.type to transaction and adding Bundle.entry.request.method for each resource. Edit the payload e.g. - modify product name - add another ATC Vet code - add another ManufacturedItemDefinition including this into the existing PackagedProductDefinition	Sample XML of Get Everything response used as a starting point: UPD_1.5.1-0_EP311_UpdateProduct_GetEverything_version1.XML Update bundle prepared: UPD_1.5.1-0_EP311_UpdateProduct_RequestBundle.XML
Update product via API	 POST Bundle with request headers to /pms/api/v2 "is_update=true" "chapter4" = true or false for the validation rules to apply 	
Check operation outcome	MSG_CREATED message expected containing Permanent identifier	
EP304 Get Product Full	Check the response for modifications	Sample XML of GET everything after update: UPD_1.5.1-0_EP311_UpdateProduct_GetEverything_version2.XML

4.1.6.3. How to use Update National Data DCP/MRP/SRP product endpoint and example bundle

EP304 Get Product Full	Prepare update bundle based on the response by updating Bundle.type to transaction and adding Bundle.entry.request.method for each resource.	Create DCP using this example file: UPD_1.6.16-5_CreateDCPForUpdateNationalData.XML Product Identifier: d0f4414c-cd65-478b-921e-f107c66f7a85 CMS for Italy Permanent identifier: 600000251886
	Edit the payload and add national data e.g. - Product name - Legal status of supply (product level) - Package description - Marketing authorisation number (product level) - Marketing authorisation status & dates - Responsible authority	Sample XML of Get Everything response used as a starting point: UPD_1.6.16_DCP_UpdateNationalData_600000251886_GetEverything_v1.XML Update bundle prepared: UPD_1.6.16_DCP_UpdateNationalData_600000251886_BasedOn_v1.XML
Update product via API	 POST Bundle with request headers to /upd/api/v1/national-data-bundle/ "is_update=true" "chapter4" = true or false for the validation rules to apply 	
Check operation outcome	MSG_CREATED message expected containing Permanent identifier	
EP304 Get Product Full	Check the response for modifications	Sample XML of GET everything after update: UPD_1.6.16_DCP_UpdateNationalData_600000251886_GetEverything_v2.XML

4.1.6.4. How to use Update Common Data DCP/MRP/SRP product endpoint and example bundle

EP304 Get	Prepare update bundle based on the	Sample XML of Get Everything response used as a starting point:
Product Full	response by updating Bundle.type to	
	transaction and adding	UPD_1.5.3-
	Bundle.entry.request.method for each resource.	4_DCP_UpdateCommonData_Product_600000149642_GetEverything_Version1.XML
	Edit the payload	Update bundle prepared:
	e.g.	
	 modify common product name 	UPD_1.5.3-
	- add another ATC Vet code	4_DCP_UpdateCommonData_Product_600000149642_UpdateBundleBasedOnVersion1 XML
	Important: any national data that has	
	been populated should be also included	
	in the update bundle.	
Update product	POST Bundle with request headers to	
via API	/upd/api/v1/common-data-bundle/	
	"io undoto truo"	
	 "is_update=true" "chapter4" = true or false for 	
	the validation rules to apply	
Check	MSG_CREATED message expected	
operation	containing Permanent identifiers	
outcome		
EP304 Get	Only the Common data in the RMS and	Please refer to Known issues section for any outstanding issues where national data
Product Full	CMS products under that Product	submitted when updating common data is not being ignored.
	Identifier will be updated	

4.1.6.5. How to use Create MRP product endpoint and example bundle

EP304 Get Product Full	Prepare update bundle based on the response by updating Bundle.type to transaction and adding Bundle.entry.request.method for each resource.	Sample XML of Get Everything response used as a starting point: UPD_1.5.3-4_CreateMRP_NP_600000184179_GetEverything_version1.XML
Prepare Create MRP Bundle	 Change procedure type from NP to MRP Add Common Name with Country = EU and Language = English Add Reference member state and Concerned member state Add Common package description in English (if doesn't exist) 	Create MRP bundle prepared: UPD_1.5.3-4_CreateMRP_BasedOn_NP_version1.XML
Create MRP via API	 POST Bundle with request headers to /upd/api/v1/mrp-bundle/ "chapter4" = true or false for the validation rules to apply 	
Check operation outcome	MSG_CREATED message expected containing Permanent identifiers for RMS NP product and products created for each CMS	
EP304 Get Product Full	RMS: • Contains the Common data that was added CMS:	

Each new product is only populated with Common data, with status of Provisional

4.1.6.6. How to use Create SRP product endpoint and example bundle

EP304 Get Product Full	Prepare update bundle based on the response by updating Bundle.type to transaction and adding Bundle.entry.request.method for each resource.	Sample XML of Get Everything response used as a starting point: UPD_1.6.1-4_CreateSRP_RMSProduct_GetEverything_version1.XML
Prepare Create SRP Bundle	 Add new Concerned member state(s) Update common data as required 	Create SRP bundle prepared: UPD_1.6.1-4_CreateSRP_BasedOnRMSProduct_version1.XML
Create SRP via API	 POST Bundle with request headers to /upd/api/v1/srp-bundle/ "chapter4" = true or false for the validation rules to apply 	
Check operation outcome	MSG_CREATED message expected containing Permanent identifiers for existing RMS & CMS products and products created for each new CMS	
EP304 Get Product Full	 RMS & existing CMS: Contains the new CMS Procedure type remains unchanged Contains the Common data that was updated 	

4.1.7. API Manage document

4.1.7.1. EP403 Create document

Resource Information

Endpoint	POST /pms/api/v2/DocumentReference	
Request		
Accept	application/fhir+xml application/fhir+json	
Body	<documentreference </documentreference 	
Content-type	application/fhir+xml application/fhir+json	

Response	
Body	Document with version 1 and document ID returned
	Nata, ID amostad formation and a
	Note: ID expected format example:
	3c46270e-3c3d-4869-a73c-ad4d7c3f2893

Query Parameters

None

Example Request

For UAT environment: POST <u>https://spor-uat.azure-api.net/pms/api/v2/DocumentReference</u>

Example file for request body: UPD_1.6.1-4_Doc_EP403_CreateDocument.XML

PDF document that was converted to base64: EP403_UploadDocument.PDF

- Document status value is case-sensitive (e.g.: current will work; CURRENT will fail)
- Document language value is case-sensitive (e.g.: en will work; EN will fail)

4.1.7.2. EP401 Search document

Resource Information

Endpoint	GET /pms/api/v2/DocumentReference?{ param}={value}[&{param}={value}]	
Request		
Accept	application/fhir+xml application/fhir+json	
Body	n/a	
Content-Type	n/a	
Response		
Body	Bundle of <documentreference>(s)</documentreference>	
	e.g. Bundle Total value=N [entry {DocumentReference Resource Type}] *	

Path Parameters

Name	Description
Version	Service version number
	Example value:
	2

Query Parameters

Name	Description
related	Permanent identifier of the product the document is related to
type	Type of document
_summary	Boolean set to true or false.
	If set to true, the contents of the document is not populated in the response in DocumentReference.content.atttachement,data.
	There is a url provided but it is not intended that you can use this to retrieve the document.

Example request

GET /pms/api/v2/DocumentReference?related=MedicinalProductDefinition/600000216133

- GET /pms/api/v2/DocumentReference?type=100000155538
- GET /pms/api/v2/DocumentReference?related=MedicinalProductDefinition/600000216133&_summary=true

4.1.7.3. EP402 Get/retrieve document

Resource Information

Endpoint GET /pms/api/v2/DocumentReference/{document-id}
--

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Request		
Accept	application/fhir+xml application/fhir+json	
Body	n/a	
Content-Type	n/a	
Response		
Body	Resource of type MedicinalProductDefinition	

Path Parameters

Name	Description
Document id	A unique document identifier UUID
	Example value:
	7a88176d-10f9-4db3-8fa0-4e4ae4594df7
version	Service version number
	Example value:
	2

Query Parameters

None

Example Request

GET /v2/DocumentReference/3c46270e-3c3d-4869-a73c-ad4d7c3f2893

4.1.7.4. EP404 Update document

Resource Information

Endpoint	POST /pms/api/v2/DocumentReference		
Request			
Accept	application/fhir+xml application/fhir+json		
Body	<documentreference> <id value="fcd2c31c-0ef9-455c-99a0-75149b888a27"></id> </documentreference>		
Content-type	application/fhir+xml application/fhir+json		
is_update	true		
Response	Response		
Body	Document with version number incremented by 1		

Query Parameters

None

Example Request

For UAT environment: POST https://spor-uat.azure-api.net/pms/api/v2/DocumentReference

Example file for request body:

- GET of document before update: UPD_1.6.1-4_Doc_EP402_GetDocument_version1.XML
- Update posted: UPD_1.6.1-4_Doc_EP404_UpdateDocument_BasedOnVersion1.XML
- Response to POST: UPD_1.6.1-4_Doc_EP404_ResponseAfterUpdate.XML
- GET of document after update: UPD_1.6.1-4_Doc_EP402_GetDocument_AfterEP404Update_version2.XML

4.1.7.5. Changes for Create and Update document payload

• There are no changes to payload

4.2. UPD API for VNRA

4.2.1. Scope of this release for VNRA API

UPD-UC31 Manage VNRA Submissions via API

- Search and Retrieve VNRA
- Approve/Reject VNRA

4.2.2. UPD API supported VNRA endpoints

4.2.2.1. Query / Retrieve VNRA Submission

Query / Retrieve VNRA	GET	Returns the complete collection of submissions which the caller is entitled to view /vnra-submission?permanentIdentifier <u>={</u> permanentId}
Submission		upd/api/vnra/v3/vnra-submission?permanentIdentifier =600013438271
APIM entry point	UAT	https://spor-uat.azure-api.net/upd/api/vnra/v3/vnra-submission?permanentIdentifier=600013438271
APIM entry point	PROD	https://spor.azure-api.net/upd/api/vnra/v3/vnra-submission?permanentIdentifier=600013438271
Query Parameters		Query Parameters (All Are Optional) Note: Calls to base url, (without parameters) /vnra-submission will return the complete collection of submissions which the caller is entitled to view 1. productName : Product name – free text field and case insensitive 2. productIdentifier : Product identifier – free text field 3. permanentIdentifier : Permanent identifier – free text field

4. mah : OMS LOC_ID of Product owner – LOC-100005358
5. responsibleAuthority : OMS LOC_ID of Responsible authority (organisation) – LOC-100001603
6. maNumber : Authorisation/registration/entitlement number – free text field
7. procedureType : Procedure type – RMS Code
8. procedureNumber : Procedure number – free text field with "Starts with" and "Contains" and
case insensitive
9. submissionIdentifier : Submission identifier – free text field
10. submissionStatus : Submission status – PENDING APPROVED PARTIALLY_APPROVED
REJECTED
11. dateFrom : Date From-To – calendar field to add interval "from"
12. dateTo : Date From-To – calendar field to add interval "to"
13. vnraStatus : VNRA Status – single selection field with list of VNRA status -PENDING
APPROVED REJECTED
14. vnraClassificationIdentifier : vnraClassificationIdentifierClassification – field with list of VNRA
classifications - RMS Code

Headers

Headers

The following Headers will be provided / injected by APIM -

- 1. APIM-Correlation-ID Generated and Included with all POST / PUT / DELETE by default. For future
- BAM requirements include with GET
- 2. APIM-User-ID ==> From User's bearer token.
- 3. APIM-Org-ID ==> org affiliations are included.

Security Headers (Mandatory)

v3 of the API require a mandatory Bearer Token which is passed via the Authorization header

Oauth Bearer Token

curl -X GET \

-H "Authorization: Bearer \$(oauth-access-token)" \ https://spor.azure-api.net/upd/api/vnra/v3/vnra-submission

Pagination

Pagination

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Pagination is implemented using Spring Boot Pagination which returns the following standard **Pagination Payload.** submission data are returned with in "content": [...],

PageSize is set using the _size parameter.

Iterating through the pages is managed via _page=x totalPages: y evaluation, If totalPages=y and the consumer searches for the last page, then _number should be set to y-1. <u>https://spor.azure-api.net/upd/api/vnra/v3/vnra-submission?</u>_size=5 https://spor.azure-api.net/upd/api/vnra/v3/vnra-submission? size=5& page=2

Pagination Payload

```
"content": [...],
   "pageable": {
 "sort": {
   "empty": false,
   "sorted": true,
  "unsorted": false
 },
 "offset":0,
 "pageNumber": 0,
 "pageSize":1,
 "paged": true,
 "unpaged": false
},
"totalPages": 485,
"totalElements": 485,
"last": false,
"sort":{
 "empty": false,
 "sorted": true,
 "unsorted": false
},
"size":1,
"number": 0,
"first": true,
"numberOfElements":1,
```

"empty": false
}

Sample		
Payload	"content": [
	"submissionId": 1588,	
	"submissionDate":1694433983143,	
	"submissionComment": "NoComments",	
	"submissionStatus": "PENDING",	
	"products": [
	{	
	"permanentId": "600001120431",	
	"procedureType": "100000155062",	
	"productRelationships": [
	{	
	"organisationId": "ORG-100004089",	
	"relationship": "Holder"	
	},	
	$\{$	
	"organisationId": "ORG-100003944",	
	"relationship": "Regulator"	
	}	
	1 / ·	
	"permanentId": "600001120431",	
	"procedureType": "100000155062",	
	"productRelationships": [
	"organisationId": "ORG-100004089",	
	"relationship": "Holder"	
	},	
	{	
	"organisationId": "ORG-100003944",	
	"relationship": "Regulator"	

```
]
],
"pageable":{
 "sort": {
  "empty": false,
  "sorted": true,
  "unsorted": false
 },
 "offset":0,
 "pageNumber":0,
 "pageSize":1,
 "paged": true,
 "unpaged": false
},
"totalPages":485,
"totalElements": 485,
"last": false,
"sort":{
 "empty": false,
 "sorted": true,
 "unsorted": false
},
"size":1,
"number":0,
"first": true,
"numberOfElements":1,
"empty": false
```

1

Union Product Database (UPD) Release Notes

4.2.2.2. Retrieve a VNRA Submission

Retrieve a VNR	Α	Retrieve a specific VNRA submission identified by its submissionId
Submission	GET	<u>/vnra-submission/<submissionid></submissionid></u> ?summary <u>={</u> true false}
		upd/api/vnra/v3/vnra-submission/456?summary=true
APIM entry point	UAT	https://spor-uat.azure-api.net/upd/api/vnra/v3/vnra-submission/456?summary=false
APIM		https://spor.azure-api.net/upd/api/vnra/v3/vnra-submission/456?summary=false
entry point	PROD	
Path Parameter		/vnra-submission/ <submissionid> <submissionid> is the ID of the submission to retrieve</submissionid></submissionid>
Query Parameters		Query Parameter (All Are Optional) summary (Optional) : _(true false) Returns a summary view of the submission else a full view_
Headers		Headers The following Headers will be provided / injected by APIM - 1. APIM-Correlation-ID Generated and Included with all POST / PUT / DELETE by default. For future BAM requirements include with GET 2. APIM-User-ID ==> From User's bearer token. 3. APIM-Org-ID ==> org affiliations are included. Security Headers (Mandatory) v3 of the API require a mandatory Bearer Token which is passed via the Authorization header Oauth Bearer Token curl -X GET \ -H "Authorization: Bearer \$(oauth-access-token)" \ https://spor.azure-api.net/upd/api/vnra/v3/vnra-submission/456?summary=false
Sample Payload		{ "submissionId":1596, "submissionDate":1694450625907,
Summary=false	2	"submissionComment": "Submit VNRA For NAP 11/09/2023", "submissionStatus": "APPROVED",

```
"variations": [
  "variationId": 16517,
  "vnraGroup": "a458cce6-5553-4efb-b974-7147069d13fc",
  "productName": "Automation Test Create NAP CH2 2023-09-11 GYxEGh",
  "productIdentifier": "926d544f-3fd6-44a3-9150-48bbb277fed6",
  "permanentIdentifier": "600001120724",
  "procedureNumber": "EMEA/V/C/777777",
  "responsibleAuthority": "LOC-100000065",
  "authorisationCountry": "10000000535",
  "marketingAuthorisationNumber": "EMEA/V/C/777777",
  "vnraCode": "200000018624",
  "implementationDate": 1694390400000,
  "decisionDate": 1694390400000,
  "decisionAuthor": "Beyond Automation",
  "decisionMaker": "ORG-100003944",
  "decisionComment": "Comment Beyond Automation",
  "status": "APPROVED",
  "marketingAuthorisationHolder": "LOC-100002851",
  "fieldChanges": []
],
"vnessFileName": "Test.zip"
```

Sample
Payload
{
 "submissionId":1596,
 "submissionDate":1694450625907,
 "submissionComment":"Submit VNRA For NAP11/09/2023",
 "submissionStatus": "APPROVED",
 "vnessFileName": "Test.zip"
}

4.2.2.3. Download a VNeeS

Download a		Download a VNeeS linked to a VNRA Submission		
VNeeS	GET	<u>/vnra-submission/<submissionid></submissionid></u> /vness upd/api/vnra/v3/vnra-submission/456/vness		
APIM entry point	UAT	https://spor-uat.azure-api.net/upd/api/vnra/v3/vnra-submission/vness_		
APIM entry point		https://spor.azure-api.net/upd/api/vnra/v3/vnra-submission/456/vness		
Path Parameter		/vnra-submission/ <submissionid> <submissionid> is the ID of the submission to retrieve</submissionid></submissionid>		
Query Parameters		None		
Headers		Headers The following Headers will be provided / injected by APIM - 1. APIM-Correlation-ID Generated and Included with all POST / PUT / DELETE by default. For future BAM requirements include with GET 2. APIM-User-ID ==> From User's bearer token. 3. APIM-Org-ID ==> org affiliations are included.		
		Security Headers (Mandatory) v3 of the API require a mandatory Bearer Token which is passed via the Authorization header Oauth Bearer Token		

curl -X GET \ -H "Authorization: Bearer \$(oauth-access-token)" \ <u>https://spor.azure-api.net/upd/api/vnra/v3/vnra-submission/456?summary=false</u>

4.2.2.4. Submit a decision for the VNRA

Submit a	VNRA submit decision - Approve/Reject VNRA /vnra-submission/ <submissionid>/decision</submissionid>
decision for _{PUT} the VNRA	<u>1</u> upd/api/vnra/v3/vnra-submission/456/decision
APIM entry point	https://spor-uat.azure-api.net/upd/api/vnra/v3/vnra-submission/456/decision_
APIM entry point PROD	https://spor.azure-api.net/upd/api/vnra/v3/vnra-submission/456/decision
Path Parameter	/vnra-submission/ <submissionid> <submissionid> is the ID of the submission containing the variation to approve</submissionid></submissionid>
Query Parameters	None
Headers	Headers The following Headers will be provided / injected by APIM - 1. APIM-Correlation-ID Generated and Included with all POST / PUT / DELETE by default. For future BAM requirements include with GET 2. APIM-User-ID ==> From User's bearer token. 3. APIM-Org-ID ==> org affiliations are included. Security Headers (Mandatory) v3 of the API require a mandatory Bearer Token which is passed via the Authorization header Oauth Bearer Token curl -X GET \ -H "Authorization: Bearer \$(oauth-access-token)" \ https://spor.azure-api.net/upd/api/vnra/v3/vnra-submission/456/decision.
Sample Payload	<pre>{ "vnraDecisionItems": [{ "variationId": 3711, "vnraDecision": "APPROVED", "decisionComment": "Submission-decision-approve-all test case", "decisionAuthor": "Beyond Automation", "decisionDate": "2022-05-03T12:00:00Z",</pre>

```
"decisionMaker": "ORG-100003944",
    "utils": {}
    }
]
```

4.2.2.5. User registration for VNRA API

Access to the VNRA API is requested by the Super user of an NCA (i.e. user with the role "UPD - CA Super User"); who will request a new role of "UPD CA API".

On receipt of the email confirming API role has been approved, the API credentials can be used to obtain the OAuth bearer token required to use with the VNRA API endpoints.

Refer to the document Registration guide: Union product database for veterinary medicinal products listed in the References section.

4.3. UPD API for Volume of Sales Data

4.3.1. Scope of this release for Volume of Sales API

• Retrieve Volume of Sales Data

4.3.2. Endpoint, Authorisation header, Query Parameters, Pagination

Endpoint

- UAT GET https://spor-uat.azure-api.net/upd/api/vos/v3/vos-sales-json?
- PROD GET <u>https://spor.azure-api.net/upd/api/vos/v3/vos-sales-json</u>?

Request Security Header (Mandatory)

This endpoint requires a mandatory OAuth Bearer Token which is passed via the Authorization header

Query Parameters

Note: Calls to the base url without any parameters will return the complete collection of sales data for all products.

/upd/api/vos/v3/vos-sales-json?permanentId= {permanentID}&yearFrom={yearFrom}&yearTo={yearTo}&modifiedDate={modifiedDate}

permanentId (optional) :- Permanent identifier of Medicinal Product. Will return sales for the provided Permanent identifier e.g. permanentId=600000225806

yearFrom (optional) :- yearFrom={year-month} Start date for range of sales data to be returned

yearTo (optional) :- yearTo={year-month} End date for range of sales data to be returned

e.g. yearFrom=2020-01&yearTo=2021-07

modifiedDate (optional) :- Modified Date of Sales data of Medicinal Product. Will return sales modified since a date The following prefixes apply to date comparisons against a stored (modified date) value. If no prefixes are specified, the default is eq.

- eq: equals, the exact stored value is inside the range defined by the precision of the parameter value
- gt: the exact stored value is greater than the exact parameter value

e.g. modifiedDate=2023-03-01 or with prefix modifiedDate=gt2023-03-01

Examples:

GET https://spor-uat.azure-api.net/upd/api/vos/v3/vos-sales-json?permanentId=600000225806&yearFrom=2020-01&yearTo=2021-07&modifiedDate=gt2023-01-01

GET https://spor-uat.azure-api.net/upd/api/vos/v3/vos-sales-json?yearFrom=2020-01&yearTo=2021-07

GET https://spor-uat.azure-api.net/upd/api/vos/v3/vos-sales-json?permanentId=600000225806

Pagination

Pagination is implemented using Spring Boot Pagination which returns the following standard Pagination Payload.

- sales data is returned within "content": [...],
- pageSize is set using the _size parameter
- iterating through the pages is managed using the _page=x parameter
- totalPages: y evaluation: If totalPages=y and the consumer searches for the last page, then page number parameter should be set to y 1.

Examples:

{

GET https://spor-uat.azure-api.net/upd/api/vos/v3/vos-sales-json?_size=5

GET https://spor-uat.azure-api.net/upd/api/vos/v3/vos-sales-json?_size=5&_page=2

Pagination Payload

```
"content": [...],
"pageable": {
    "sort": {
        "empty": false,
        "sorted": true,
        "unsorted": false
    },
    "offset": 0,
    "pageNumber": 1,
    "pageSize": 100,
    "paged": true,
    "unpaged": false
```

```
},
"totalPages": 6,
"totalElements": 596,
"last": false,
"sort": {
    "empty": false,
    "sorted": true,
    "unsorted": false
},
"size": 100,
"number": 0,
"numberOfElements": 100,
"first": true,
"empty": false
```

Sample Response Payload

}

```
{
 "content":
  {
   "productIdentifier": "c74a510c-1689-4f46-bdce-f3a5dd84b1da",
   "productName": "TEST-PRODUCT-NAME2-95363f02-c9b9-442b-8bdf-21a54bf15b2e-VOS",
   "permanentIdentifier": "600013438271",
   "authorisationProcedureNumber": "VOS/TEST/HOLDER-NAME2/TEST/EMEA/H/C/000175",
   "packageIdentifier": "be7bfd42-df3f-45e2-8af9-3d96a870f5f7",
   "packageDescription": "PACKAGE3-TEST-PRODUCT-NAME2-95363f02-c9b9-442b-8bdf-21a54bf15b2e-VOS",
   "packSizeNumericValue": "94",
   "packSizeUnitOfPresentation": "Capsule",
   "packSizeUnitOfPresentationIdentifier": "20000002113",
   "country": "European Union",
   "countryIdentifier": "10000000390",
   "marketingAuthorisationNumber": "VOS/TEST/HOLDER-NAME2-1591819011837",
   "creationDateOfProduct": "2021-11-12",
```

```
"yearMonth": "2021-03",
   "volumeOfSales": "111",
  "speciesIdentifier": "100000108926",
  "speciesPercent": "100.00",
  "doseFactor": "1.00",
  "comment": "Mandatory",
  "modifiedDate": "2023-06-14 09:06:28.047"
 }
],
"pageable": {
 "sort": {
  "empty": true,
  "unsorted": true,
  "sorted": false
 },
 "offset": 0.
 "pageNumber": 0,
 "pageSize": 1,
 "paged": true,
 "unpaged": false
},
"totalElements": 5,
"totalPages": 5,
"last": false,
"sort": {
 "empty": true,
 "unsorted": true,
 "sorted": false
},
"size": 1,
"number": 0,
"first": true,
"numberOfElements": 1,
```

```
"empty": false
}
```

4.3.3. User registration for Volume of Sales Data API

Access to the Volume of Sales API is requested by the Super user of an NCA (i.e. user with the role "UPD - CA Super User"); who will request a new role of "UPD CA API".

On receipt of the email confirming API role has been approved, the API credentials can be used to obtain the OAuth bearer token required to use with the VoS API endpoint.