

SYNTHETIC DEMONSTRATION

Partner intake pack.

A structured dossier before regulatory review.

This fictitious example shows how NaruDesk organises documents, versions, gaps and open questions before a Korean cosmetic dossier reaches an EU Responsible Person or qualified safety assessor.

DEMO PRODUCT Barrier Cloud Serum 30 mL Demo Brand Korea	Preparation status Preparation in progress
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8	5	4
documents received	items missing	open questions

NaruDesk coordinates preparation. This sample is not a compliance determination, safety assessment, CPNP submission or legal opinion.

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01. Document register

One version-controlled view of what was received, what is current and what still needs to be requested.

Document	Version / date	Status	Owner / next step
Product brief	v03 / 2026-07-22	RECEIVED	Brand
Final INCI list	v05 / 2026-07-24	QUESTION	Manufacturer: confirm final
Qualitative CAS/EC index	v02 / 2026-07-24	RECEIVED	Manufacturer
Quantitative formula	Not held by NaruDesk	DIRECT ROUTE	Send to assessor
GMP statement	2026-03-11	RECEIVED	Manufacturer
Stability summary	Draft / 2026-06-30	UPDATE	Lab: final signed copy
Challenge test	Not supplied	MISSING	Brand to request
Packaging specification	v01 / 2026-02-19	QUESTION	Confirm pump supplier
EU artwork	Draft 02	UPDATE	Brand + RP review
Claims evidence set	3 files supplied	GAP	Add moisturising support

Illustrative status only. SHA-256 hashes and source receipt dates would be recorded in the operational register.

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02. Cross-document gaps

Open points are separated by type so the regulatory partner can review evidence without reconstructing the dossier.

Type	Observed item	Why it matters	Resolution owner
FACT	Product brief names "Barrier Cloud Serum". Artwork says "Barrier Serum".	Two names are in circulation.	Brand
GAP	Pack specification says 32 mL nominal fill. Artwork states 30 mL.	The final pack record is not aligned.	Manufacturer
QUESTION	INCI v05 is dated after the latest stability summary.	Confirm which formula version the test covered.	Manufacturer + lab
GAP	"48-hour hydration" appears in the draft claims sheet.	No matching evidence file was identified in the supplied set.	Brand
HUMAN	Draft EU artwork contains a placeholder RP address.	Final wording and address require partner review.	EU RP

Regulatory pointers for partner review

The pack routes product information, label and claims questions to the relevant official sources. It does not convert those pointers into a final verdict.

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03. Action list by owner

Each request has one owner, one deliverable and one target date. This is the part that removes repeated clarification loops.

Priority	Owner	Requested item	Target	State
P0	Brand	Confirm one commercial product name	2026-07-30	OPEN
P0	Manufacturer	Confirm final INCI and linked formula version	2026-07-30	OPEN
P0	Lab	Issue signed final stability summary	2026-08-03	OPEN
P1	Brand	Provide challenge-test report or explain availability	2026-08-03	OPEN
P1	Brand	Provide evidence supporting “48-hour hydration”	2026-08-05	OPEN
P1	EU RP	Review draft artwork questions and RP address	After intake	HUMAN
P1	Assessor	Receive confidential quantitative formula directly	Agreed route	HUMAN

Confidential-data route

The quantitative formula and underlying confidential data are not uploaded to NaruDesk or Hermes. They move directly from the manufacturer to the EU RP or qualified safety assessor through the route that partner approves.

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04. Handoff summary

The handoff gives the European partner a clean intake view, while leaving every regulatory and safety decision with the qualified party.

Ready for handoff	Still open
Document register with versions and receipt dates	Final formula/version confirmation
Qualitative ingredient index supplied by manufacturer	Final signed stability summary
Artwork and claims questions grouped by owner	Challenge-test report
Direct route proposed for confidential formula	Claims support and final RP address

Proposed next step

The RP or safety assessor reviews the open questions, confirms the confidential-data route and identifies any additional evidence required. NaruDesk then coordinates requests and version updates on the Korean side.

Official reference points

- Regulation (EC) No 1223/2009 on cosmetic products, consolidated text of 2026-05-01, accessed 2026-07-28:
eur-lex.europa.eu/eli/reg/2009/1223/2026-05-01/eng
- Commission Regulation (EU) No 655/2013 on common criteria for cosmetic claims, accessed 2026-07-28:
eur-lex.europa.eu/legal-content/EN/TXT/?uri=CELEX:32013R0655
- European Commission CPNP information page, accessed 2026-07-28:
single-market-economy.ec.europa.eu/sectors/cosmetics/cosmetic-product-notification-portal_en



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