HP VITARAN — English Fact Sheet (Yaksu Medi)

GLOBAL English only · 2 September 2026 · Not an IFU · Wholesale research synthesis

1. Identity

Brand: HP VITARAN / HP Cell VITARAN — PN intradermal skin booster (medical device)
Manufacturer: BR PHARM Co., Ltd., Wonju, Gangwon, Republic of Korea — https://www.brpharm.com/
Active: Polynucleotide sodium (PN). Not HA filler. Not a PDRN drug injectable.
Concentration (sourced): 20 mg/mL (2%) PN
Korea MFDS skin family: Jeheo 23-784 (Jul 2023) — temporary improvement of adult crow's feet
Related joint SKU: HP VITARAN J — MFDS Jeheo 22-99 (separate licence)

2. Official skin SKUs

HP VITARAN — typically 1 mL x 2 PFS / box (BR Pharm TradeKorea listing)
HP VITARAN I (i / Eye) — same MFDS 23-784 family
HP VITARAN S — same MFDS 23-784 family
Storage 1–30 C, light-protected · Shelf life up to 3 years from manufacture
Reseller “VITARAN II 2x2 mL” is not a core MFDS SKU name without manufacturer IFU

3. CE / EU status (sourced)

CE status for the HP VITARAN family: YES (manufacturer-stated EU CE MDR).
Primary source — BR Pharm certification gallery caption:
“HP VITARAN product family, EU CE MDR certification”
(also: “HP cell VITARAN J product family, EU CE MDR certification”)
https://www.brpharm.com/rnd/certification
Authorised EU distributor ODA Medical (English page): Class III CE certified medical device
https://odamedical.pl/en/hp-vitaran-en/
Notified-body number 2265 appears on some EU professional Class III PN listings
(example: Biomedico card “Numer jednostki notyfikowanej 2265”).
Print as NB 2265 only if needed — do NOT invent a full certificate serial / Basic UDI-DI.
EUDAMED DoC PDF not independently downloaded this pass.
FDA: injectable NOT approved. MoCRA cosmetic listing ≠ injectable clearance.

4. Clinical snapshot

Jeon et al., Aesth Plast Surg 2026 — HP Vitaran I ionization pain study
PMID 41361144 · n=15 · VAS 3.39 vs 5.04 ($p<0.005$) · BR Pharm-funded
Crow's-feet confirmatory narrative (MFDS/symposium): n=250 · ~93.2% responders
Full English CSR / NCT not retrieved — licence/symposium narrative only
Duration months: not independently verified

5. Do-not-print

Any non-English catalog (Polish, Turkish, etc.) on the GLOBAL website
Invented CE certificate serial · FDA seal · MoCRA as syringe approval
G-prime / BDDE ppm · “PDRN drug” label · sister OEM carton names as this SKU

Replaces non-English distributor katalogs for Yaksu Medi GLOBAL English use.

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