



Fimvo – Counterforce against falsified medicines



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Upcoming invoicing for 2026

The Fimvo Board has decided the annual fee for 2026 to be **4 300 € per MAH**.

If you need to have your PO number on the invoice, please let us know the PO by the end of November by filling the [Appendix I and II form](#). Please fill it completely and send it to us by email (info@fimvo.fi). You'll receive the invoice in the first week of January and the due date will be at the beginning of February. Fimvo does not grant reduced fees. Thank you for making the invoicing go smoothly!

Updating Product Master Data (PMD) for All Target Markets

EMVO has identified instances where Product Master Data (PMD) is not being correctly updated to reflect all target markets.

All intended markets for your product(s) must be included in the Product Master Data (PMD) uploaded to the EU Hub. This is essential to ensure that the associated batch data can be correctly transmitted to the relevant national systems.

If you are adding a new market to an existing multi-market pack, please ensure that you create **a new version of the PMD** and include all relevant target markets.

For detailed guidance on completing the Product Master Data and Product Pack Data, please consult the [EMVS Master Data Guide](#).

Should you have any questions, please do not hesitate to contact EMVO's Helpdesk (helpdesk@emvo-medicines.eu) or consult EMVO's Self-Service Portal available on their website [here](#).



Nordic recommendations on the use of status 'Locked' in the EMVS

The functionality to mark a pack as locked can provide added value and security in preventing medicines in quarantine or under investigation entering in or going back into the supply chain.

The functionality is not a legal requirement and the use of it is voluntary.

The Nordic medicines verification organisations have published a Best Practice Guide for using this feature in the system.

[Find the Guide here](#)

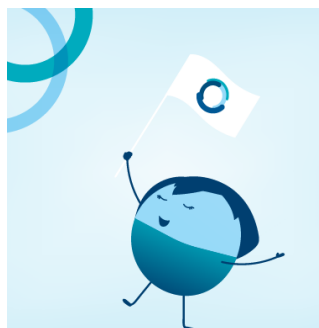
Contact us!

Regarding contracts and invoices:

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