BADAN POM

Daftar MoU Bilateral Tahun 2025

No	Negara/Mitra	Judul MoU	Tanggal Penandatanganan dan Masa Berlaku	Tanggal Berakhir	Ruang Lingkup Kerja Sama dalam MoU
1	India/Ministry of Ayush of The Republic of Indonesia	Memorandum of Understanding between Indonesian Food and Drug Authority of the Republic of Indonesia and Pharmacopoeia Commission for Indian Medicine & Homoeopathy, Ministry of Ayush of The Republic of India on Cooperation in The Field of Traditional Medicine Quality Assurance	21 Januari 2025/Masa Berlaku 5 Tahun (Diperpanjang otomatis untuk jangka waktu 5 Tahun berikutnya)	21 Januari 2030 (Dapat diperpanjang otomatis sampai 21 Januari 2035)	1) Exchange of information, experience and knowledge regarding the regulatory 2) Exchange of experts for capacity building activites including seminars, 3) Technical visit related to familiarisation of regulatory processes of both 4) Participation in international events organised by either Party 5) Joint training programmes in mutually agreed areas, either for the Parties or 6) Any other forms of co-operation as agreed by the Parties
2	Sudan / The National Medicines and Poisons Board	Memorandum of Understanding between The Food And Drug Authority of The Republic of Indonesia and the National Medicines and Poisons Board of The Republic of the Sudan on Cooperation in The Fields of Pharmaceutical Products, Natural Medicinal Products, Health Supplements and Cosmetics Control	3 Juli 2025/Masa Berlaku 5 Tahun (Akan diperpanjang otomatis untuk jangka waktu 5 Tahun)	3 Juli 2030 (Dapat diperpanjang sampai 3 Juli 2035)	1. Exchange and sharing of information, best practices, knowledge and skills regarding standards, guidelines and regulations; 2. Facilitate information sharing to support Sudan NMPB in the implementation of regulatory reliance on the Indonesian FDA, in accordance with the terms mutually agreed upon by the Parties; 3. Undertake capacity building programs (technical assistance), including workshop and training; 4. Exchange of information on the status of Good Manufacturing Practice (GMP) inspection results as may be requested by one of the Parties; 5. Provide safety alerts for new ingredients in cosmetics or natural medicinal products, particularly those that could potentially affect the public health of the other Party; 6. Exchange of visits of high-level delegations and experts;

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					7. Participation in international events organized by either Party
3	Amerika Serikat/The United States Pharmacopeial Convention	Memorandum of Understanding between The Indonesian Food And Drug Authority of The Republic of Indonesia and The United States Pharmacopeial Convention	7 Mei 2025/Masa Berlaku 5 Tahun (Akan diperpanjang jangka waktu 5 Tahun melalui nota diplomatik)	7 Mei 2030 (Dapat diperpanjang sampai 7 Mei 2035)	3.1 Organize joint activities including but not limited to research collaborations on the development of analytical methods and/or participate in each other's scientific fora, roundtables, expert panels or other suitable platforms such as USP Convention Regional Chapter meetings in selected areas of scientific interest identified by the Parties.
					3.2 exchange of information and best practices on pharmaceutical products and processed food, as applicable for the development and strengthening of policy and regulation.
					3.3 develop and publish scientific publications or reports resulting from scientific workshops, fora, roundtables, expert panels, joint projects or other platforms.
					3.4 capacity building for Regulators and Industry in Indonesia, including but not limited to, technical assistance, exchange of information and knowledge, short courses, training, workshops, seminars, internships/fellowships, and research collaborations including potential facilitation of preferential access to selected USP educational resources.
					3.5 facilitation of preferential access to the newest edition of the compendia of both Pharmacopeias (USP-NF, USP-NF in English) upon separate written agreement by both Parties.
					3.6 develop advocacy and awareness programs and tools to support the use of public standards for processed foods and pharmaceutical products.
					3.7 assist the development of the laboratory systems of the Indonesian FDA, including but not limited to systems, equipment, infrastructure and laboratory digitization, the work of which is entirely contingent upon availability of requisite 3rd party donor funding and other forms of cooperation to be mutually agreed in writing by the Parties.

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					3.8 USP may grant the Indonesian FDA restricted rights to translate, copy, adapt, and incorporate the intellectual property of monographs, general chapters, methods, tests, and assays from the United States Pharmacopeia and the National Formulary (“USP-NF”) into publication and regulation published by Indonesian FDA, by way signing a separate written Agreement.
					3.9 exploring and executing other forms beneficial projects in the areas of mutual interest to be agreed in writing by the Parties.