The Arab Republic of Egypt
Egyptian Drug Authority
Central administration of Operations
General Directorate of factories Inspection
Inspection Administration of Human, Herbal and
Veterinary and Disinfectants Pharmaceuticals
Factories.



جمهورية مصر العربية هيئة الدواء المصرية الإدارة المركزية للعمليات الإدارة العامة للتفتيش على المصانع الإدارة العامة للتفتيش على المصانع إدارة التفتيش على مصانع المستحضرات الصيدلية والعشبية والبيطرية والمطهرات

Certificate of Good Manufacturing Practices

This certificate conforms to the format recommended by the World Health Organization

- 1. <u>Purpose of this certificate:</u> To be introduced to Ministry of Health of **El Salvador** for the purpose of Export.
- 2. Certificate No.: 1322/2021
- 3. Name and address of site: EVA Pharma for pharmaceutical and medical appliances (176 Elsadat st., Kafr El-Gabl, El. Giza).
 - On the basis of the inspection, we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the dosage forms, categories and activities listed in the Table 1 below
- 4. Manufacturer's license number: 112018062600008 (License issued from Industrial Development Authority)

Dosage form(s)	Last	Category(ies)	Activities carried out by the
	Inspection		company
Sterile: Vials & ampoules (Liquid - suspension) - Solid dosage forms (Tablets			
- Hard gelatin capsule - effervescent and non-effervescent powder packed in			
sachet - effervescent and non-effervescent powder packed in bottles) - capsule	2019		
for inhalation – bilayer tablet) –			
		General	
Eye drops (multidose & unidose), Ophthalmic gel – water for injection - Single	2020	Category(ies)	Production and packaging
dose inhalation -		Categorytics	
Non-sterile: Semi-Solid dosage forms (Cream - Ointment - Gel) -	2021		
Transdermal patches - oral dispersible film (ODF)-vet powder filled in sachets			
& cans - Vet. Suspensions & solution - liquid dosage form (syrup-oral drop-			
suspension) for internal use & (lotion - shampoo) for external use			

• The responsibility for the quality of the individual batches of the pharmaceutical products manufactured through this process lies with the manufacturer.

• This certificate remains valid until $\frac{21/12/2022}{11/12/2022}$ It becomes invalid if the activities and/or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.

Dr. Amira Mahgoub

Manager of the Inspection Administration of All Human, Herbal and Veterinary and Disinfectants

Pharmaceuticals Factories.

Authenticated

Dr. Yasin Ragaey

Head of Central Administration of Operations/

Note: Not valid without stamp

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