

Form 11

(r.  
6)

**THE PHARMACY AND POISONS ACT**  
**THE PHARMACY AND POISONS (REGISTRATION OF DRUGS)**  
**RULES**

**REGISTRATION OF DRUGS**  
**CERTIFICATE**

H2020/CTD5511/1363ER

It is hereby certified that the drug as described hereunder, has been registered subject to the conditions indicated hereunder:

1.	<b>Trade name under which marketed</b>	ESOPRA 40
2.	<b>Approved name</b>	ESOPRA 40
3.	<b>Form of preparation</b>	Tablet
4.	<b>Active Ingredients and quantities per unit</b>	INN Esomeprazole magnesium as Dihydrate  API Strength per Dosage: Esomeprazole magnesium as Dihydrate Eqi. To Esomeprazole ... 40 mg
5.	<b>Condition(s) under which is registered</b>	To be retained annually
6.	<b>Name and business address of manufacturer</b>	Associated Biotech.
7.	<b>Registered in the name of</b>	Rena exports
	<b>Local Technical Representative</b>	
8.	<b>Date of Registration</b>	Thu May 28 12:33:41 EAT 2020
8.	<b>Date of Issue</b>	2020-05-26 00:00:00
9.	<b>Expiry date of Registration</b>	Annual Retention
	<b>Date of Viewing</b> Thu Nov 5 12:36:52 EAT 2020	
		C.E.O

**CONSIDER THIS DATE**