



GOVERNMENT OF THE PEOPLE'S REPUBLIC OF BANGLADESH

MINISTRY OF HEALTH & FAMILY WELFARE
DIRECTORATE GENERAL OF DRUG ADMINISTRATION
OUSHAD BHABAN, MOHAKHALI
DHAKA-1212, BANGLADESH
www.dgda.gov.bd



CERTIFICATE OF GOOD MANUFACTURING PRACTICE (GMP) FOR MEDICAL DEVICE(S) & PHARMACEUTICAL PRODUCT(S)

This certificate conforms to the format recommended by the World Health Organization (WHO)

Certificate Number: DGDA/6-247/2020/5549 Date: 22-03-2022

It is hereby certified that M/S. Get Well Ltd., a Pharmaceutical Products, Medical Devices & Surgical Products Manufacturing & marketing organization, has been given license to manufacture & sell its products freely in the People's Republic of Bangladesh as lawfully required & granted in pursuance of The Drugs Act, 1940 (XXIII of 1940) & The Drugs (Control) Ordinance, 1982 and The Drugs Control (Amendment) Act. 2006.

On the basis of the inspection carried out on 07.12.2020 & 10.03.2021, we certify that the site indicated in this certificate complies with Good Manufacturing Practices for the capacity forms, categories and activities listed in table - 1.

- 1. Name & address of site : Get Well Ltd. Olipur, Shajibazar, Shaistaganj, Habiganj
2. Manufacturer's license number : Biological- 305, Non-Biological- 518
Date of Issue: Biological: 30.12.2015 Non-Biological:19.04.2018

3. Table: 1

Table with 3 columns: Dosage Form(s) of Product (s), Category (ies), and Activity (ies). It lists details for Coated Tablets and Powder For Suspension (PFS).

<b>Granules For Suspension (GFS)</b>	Quinolones Antibiotics	<ul style="list-style-type: none"> <li>• Quality Assurance of finished product.</li> <li>• Storage of finished product.</li> </ul>
<b>Capsule</b>	Cephalosporin (Third generation) Antibiotics	
<b>Pediatric Drops</b>	Cephalosporin (Third generation) Antibiotics	
<b>Hand Sanitizer</b>	Antiseptics	Quality Control of Incoming Materials, In-process Control, Manufacturing, Assembling, Ribbon Packing/ Blistering, Packing, Sterilizing, Finished Product Analysis, Finished Product Release, Storage and Finished Product Delivery.
<b>Disposable Syringe</b>	Disposable Syringe 1 ml Disposable Syringe 2 ml Disposable Syringe 3 ml Disposable Syringe 5 ml Disposable Syringe 10 ml Disposable Syringe 20 ml Disposable Syringe 30 ml Disposable Syringe 50 ml	Quality Control of Incoming Materials, In-process Control, Manufacturing, Assembling, Ribbon Packing/ Blistering, Packing, Sterilizing, Finished Product Analysis, Finished Product Release, Storage and Finished Product Delivery.
<b>Insulin Syringe</b>	Insulin Syringe 1ml (U-40) Insulin Syringe 1ml (U-100)	Quality Control of Incoming Materials, In-process Control, Manufacturing, Assembling, Ribbon Packing/ Blistering, Packing, Sterilizing, Finished Product Analysis, Finished Product Release, Storage and Finished Product Delivery.

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I.V. Infusion Set	Intravenous Infusion Set	Quality Control of Incoming Materials, In-process Control, Manufacturing, Assembling, Ribbon Packing/ Blistering, Packing, Sterilizing, Finished Product Analysis, Finished Product Release, Storage and Finished Product Delivery.
Scalp Vein Set	Scalp Vein Set	Quality Control of Incoming Materials, In-process Control, Manufacturing, Assembling, Ribbon Packing/ Blistering, Packing, Sterilizing, Finished Product Analysis, Finished Product Release, Storage and Finished Product Delivery.
Disposable Needle	Disposable Needle	Quality Control of Incoming Materials, In-process Control, Manufacturing, Assembling, Ribbon Packing/ Blistering, Packing, Sterilizing, Finished Product Analysis, Finished Product Release, Storage and Finished Product Delivery.
First Aid Bandage	First Aid Bandage	Quality Control of Incoming Materials, In-process Control, Manufacturing, Assembling, Ribbon Packing/ Blistering, Packing, Sterilizing, Finished Product Analysis, Finished Product Release, Storage and Finished Product Delivery.

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The responsibility for the quality of the individual batches of the medical devices & pharmaceuticals products manufactured through this process lies with the manufacturer.

The manufacturing plant in which the medical devices & pharmaceutical products are produced is subject to inspection at suitable intervals.


The manufacturer conforms to the requirements for good practices in the manufacturer & quality control (GMP) of medical devices & drugs, as required under law in this country, as well as recommended by the World Health Organization (WHO) in respect of medical devices & pharmaceutical products to be manufactured, sold or distributed within the country of origin or to be exported.

**This certificate remains valid for a period of 2 (two) years from the date of issue.** 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.

Address of certifying authority : Directorate General of Drug Administration  
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Telephone : +88-022222-80803  
E-mail : [dgda.gov@gmail.com](mailto:dgda.gov@gmail.com)  
Web-site : [www.dgda.gov.bd](http://www.dgda.gov.bd)

Name of Authorized Person : Major General Md Mahbubur Rahman

Stamp & Date:

 21 MAR 2021

**Major General Md Mahbubur Rahman**  
**Director General**  
**Directorate General of Drug Administration**  
&  
**Licensing Authority (Drugs)**  
**Government of the People's Republic of Bangladesh**