



Republic of the Philippines  
Department of Health  
**FOOD AND DRUG ADMINISTRATION**



04 JUN 2021

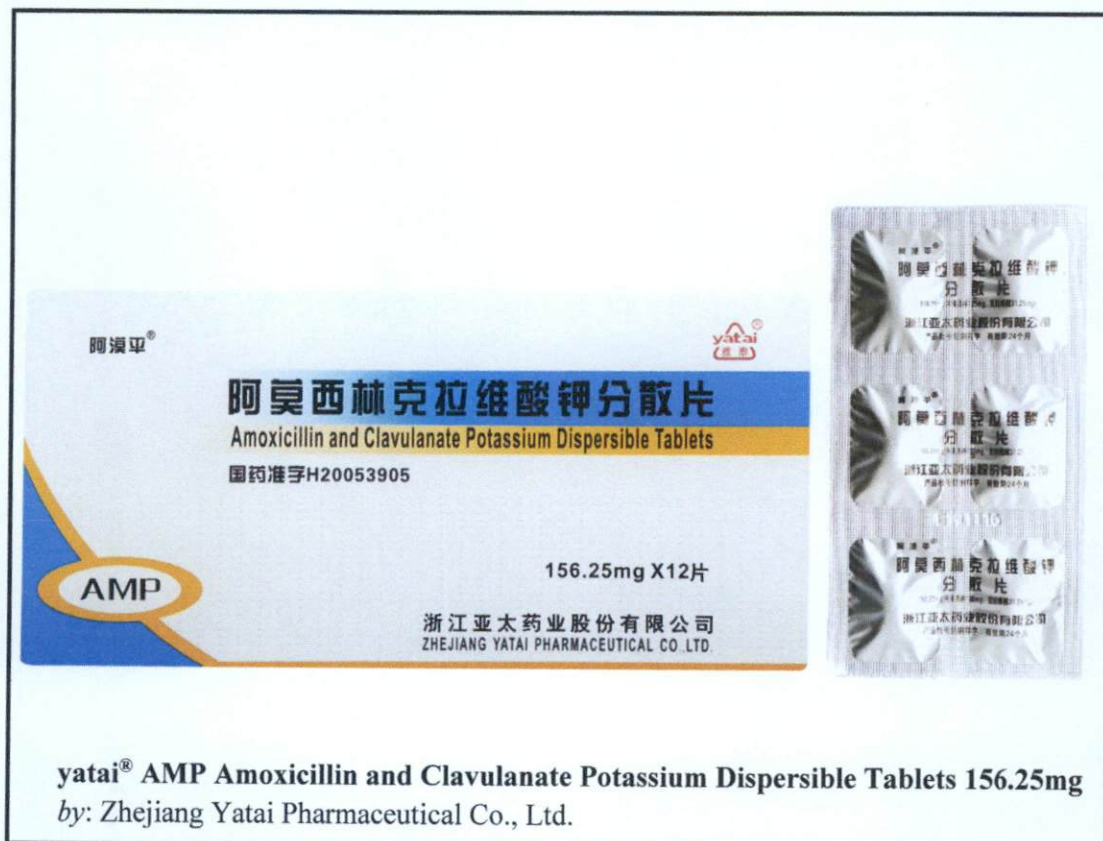
FDA ADVISORY  
No. **2021-1220**

**TO : ALL HEALTHCARE PROFESSIONALS AND THE GENERAL PUBLIC**

**SUBJECT : Public Health Warning Against the Purchase and Use of the Following Unregistered Drug Products:**

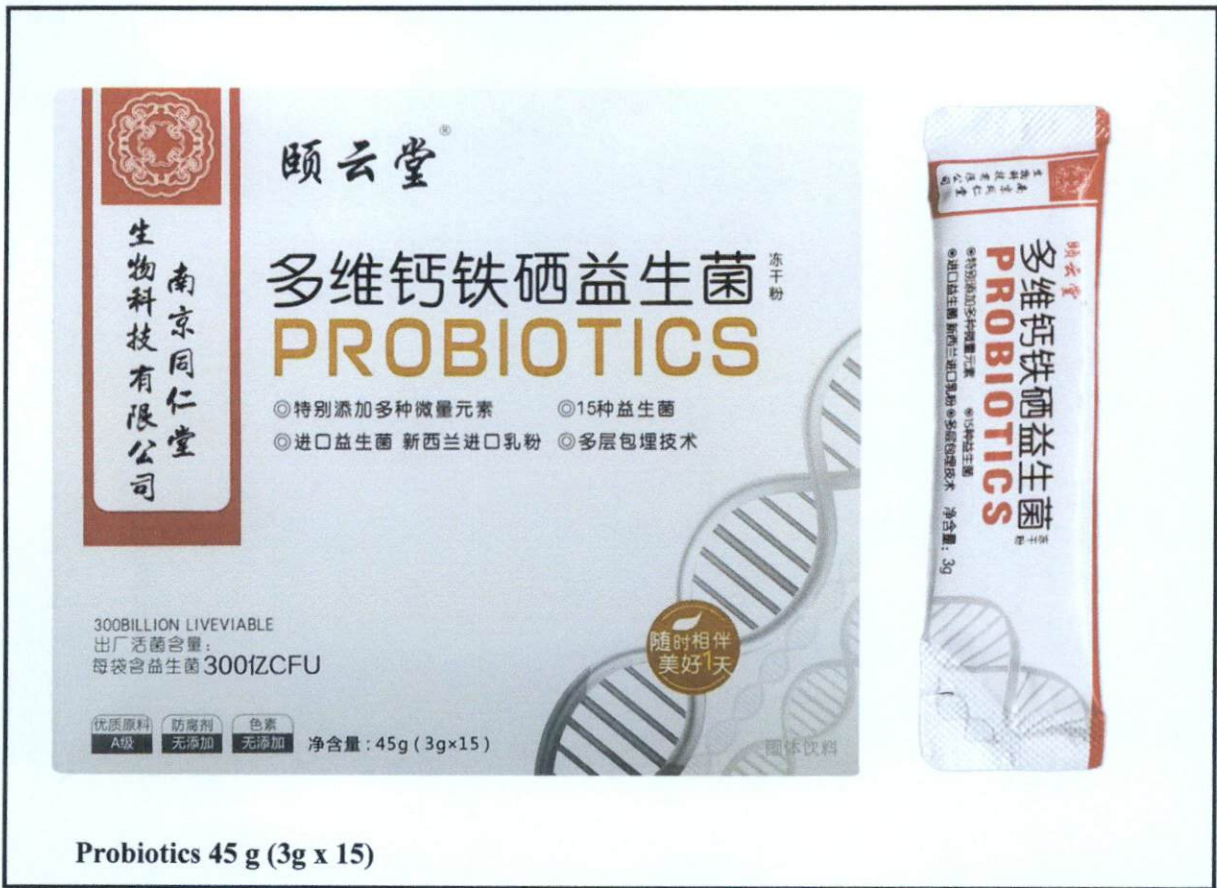
1. yatai® AMP Amoxicillin and Clavulanate Potassium Dispersible Tablets 156.25mg
2. Probiotics 45g (3g x 15)
3. Complex Protein Peptide 60g (4g x 15)
4. OTC Baofukang Shuan [as reflected in the package insert]
5. Sanyoujianan Medicine™ Nuan Gong Tie

The Food and Drug Administration (FDA) advises the public against the purchase and use of the following unregistered drug products:



yatai® AMP Amoxicillin and Clavulanate Potassium Dispersible Tablets 156.25mg  
by: Zhejiang Yatai Pharmaceutical Co., Ltd.

Figure 1. Unregistered drug product



Probiotics 45 g (3g x 15)

Figure 2. Unregistered drug product



Complex Protein Peptide 60g (4g x 15)

Figure 3. Unregistered drug product



OTC Baofukang Shuan [as reflected in the package insert]  
by: Hainan Bikai Pharmaceutical Co., Ltd.

Figure 4. Unregistered drug product



Sanyoujianan Medicine™ Nuan Gong Tie  
by: Anhui naford pharmaceutical technology Co., Ltd

Figure 5. Unregistered drug product

FDA Post-Marketing Surveillance (PMS) activities have verified that the abovementioned drug products have not gone through the registration process of the Agency and have not been issued with proper authorization in the form of Certificate of Product Registration. Thus, the Agency cannot guarantee its quality, safety and efficacy. Therefore, consumption of such violative products may pose potential danger or injury to health.


Pursuant to Republic Act No. 9711, otherwise known as the "Food and Drug Administration Act of 2009", the manufacture, importation, exportation, sale, offering for sale, distribution, transfer, promotion, advertising or sponsorship of health products without proper authorization from FDA is prohibited.

All concerned establishments and/or entities are warned not to distribute the above-identified violative drug products until they have already been covered by the appropriate authorization, otherwise, regulatory actions and sanctions shall be strictly pursued.

All Local Government Units (LGUs) and Law Enforcement Agencies (LEAs) are requested to ensure that these products are not sold or made available in their localities or areas of jurisdiction.

For more information and inquiries, please e-mail us at [info@fda.gov.ph](mailto:info@fda.gov.ph). To report continuous sale or distribution of unregistered health products, kindly e-mail us via [ereport@fda.gov.ph](mailto:ereport@fda.gov.ph). You may also call the Center for Drug Regulation and Research at telephone number (02) 8809-5596. For any suspected adverse drug reaction (ADR), report immediately to FDA through this link: <https://primaryreporting.who-umc.org/Reporting/Reporter?OrganizationID=PH> and fill out all the required fields.

Dissemination of the information to all concerned is requested.

  
ROLANDO ENRIQUE D. DOMINGO, MD  
Director General

DTN:



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