Publications

Filter: Doc Categories Reset Show All entries

Search:

Title	Date
2024.03.01 Guideline for Registration of Medicine	June 14, 2024
2023.09.06 Guideline-for-WHO-Pre-qualified-Medicines-through-Collaborative-Registration-Procedure-	June 14, 2024
2023.12.20 GUIDELINE FOR NON-ROUTINE REGISTRATION	June 14, 2024
2023.11.02 Guideline on Variation Application to Registered Medicines	June 14, 2024
2023.12.22 Guidelines on Reliance for Regulatory Decision Making	June 14, 2024
2023.12.21 Guideline on Medical Products Special import permit	June 14, 2024
2024.01.11 Guideline for control of advertising and promotion	June 14, 2024
2024.01.19 Guidance for Registration of Radiopharmaceuticals	June 14, 2024
2024.01.19 Guideline of Medicinal Gases Registration	June 14, 2024
2024.01.20 Guideline for Medicinal product information	June 14, 2024
2024.01.20 Guideline for preparation and publication of public assessment reports	June 12, 2024
2024.01.20 Guideline for Renewal of MA	June 12, 2024
2024.01.29 Guideline on Variation Applications to Registered Vaccines	June 12, 2024

https://t.me/pharmacyh

Title	Date
2024.01.31 Guidelines for Conditional Approval	June 12, 2024
2024.01.31.Guideline-for-Registration-of-Antiseptics-and-Disinfectants	June 12, 2024
2024.01.31.Guideline-for-Registration-of-oral-care-skin-care-products-with-therapeutic-claim	June 12, 2024
2024.02.27 Guidance-on-waiver-of-in-vivo-bioequivalence-requirements	June 12, 2024
2024.04.16 Good RevPractice Guideline	June 12, 2024
2024.01.13 Guideline for describing the role and responsibilities of MA	June 12, 2024
2024.01.30 Guidelines for Registration of Biotherapeutic Proteint Products	June 12, 2024
2024.01.29 Guideline-for-Registration-of-Vaccine	June 12, 2024
2024.01.30 Guidelines on evaluation of similar biotherapeutics	June 12, 2024
1003- ማብራሪያ – የትንባሆ ምርት አወጋገድ መመሪያ ቁጥር 1003-2016 Description TOBACCO PRODUCTS DISPOSAL DIRECTIVE 1003-2016	May 31, 2024
TOBACCO PRODUCTS DISPOSAL DIRECTIVE 1003-2016	May 31, 2024
Pharmacovigilance newsletter Q 3 Issue 1_Final__2024	May 31, 2024
Ethiopian National Tobacco Control Strategic Plan From 2023 – 2031	May 27, 2024
Sinopharm Covid 19 vaccine active surveillance report	April 16, 2024
EFDA Voice Newsletter April 2024	April 11, 2024
1000 – የመድኃኒት ማምረቻ ተቋማት የብቃት ማረጋገጫ አሰጣጥ እና ቁጥጥር መመሪያ ቁጥር 1000-2016	April 3, 2024
999 – የመድኃኒት የመልካም አመራረት ሥርዓት ቁጥጥር መመሪያ ቁጥር 999-2016	April 3, 2024
Rational Drug Use of insulin	April 3, 2024
Rational Drug Use 2	April 3, 2024
Product Registration	April 3, 2024
Jimma Branch	April 3, 2024
Counterfeit Drugs	April 3, 2024

Title	Date
Pharmacovigilance newsletter Q 2 Issue 1	March 29, 2024
Guideline on Variation Applications to Registered Vaccines	March 26, 2024
Guideline of Medicinal Gases Registration	March 26, 2024
Guideline for Renewal of MA	March 26, 2024
Guideline for preparation and publication of public assessment reports	March 26, 2024
Guideline for Non-Routine registration	March 26, 2024
Guideline for Medicinal product information	March 26, 2024
Guideline for describing the role and responsibilities of MA	March 26, 2024
Guidance on waiver of in vivo bioequivalence requirements	March 26, 2024
Guidance for Registration of Radiopharmaceuticals	March 26, 2024
Good RevPractice Guideline	March 26, 2024
Guidelines for Conditional Approval	March 26, 2024
Guideline for Registration of oral care skin care products with therapeutic claim	March 26, 2024
Guideline-for-Registration-of-Antiseptics-and-Disinfectants	March 26, 2024
Guideline Variation Application to Registered Medicines	March 26, 2024
EFDA Voice Newsletter December 2023	January 16, 2024
Traditional medicine	January 15, 2024
Tobacco	January 15, 2024
Tobacco control Addis Ababa 3	January 15, 2024
Tobacco control Addis Ababa 2	January 15, 2024
Tobacco control Addis Ababa 1	January 15, 2024
Self medication	January 15, 2024
Seeding trees	January 15, 2024
Press conference	January 15, 2024

Title	Date
Prescription and non- prescription drug	January 15, 2024
Pharmacovigilance	January 15, 2024
Milk safety	January 15, 2024
Iodized salt	January 15, 2024
Good Dispensing practice	January 15, 2024
Food Adulteration	January 15, 2024
ERIS	January 15, 2024
Anti-microbial resistance	January 15, 2024
Clinical Trial Application Form	November 30, 2023
Routine Clinical Trial Application screening form	November 30, 2023
Non routine clinical trial application screening form	November 30, 2023
Clinical Trial Authorization Guideline (4th edition)	October 31, 2023
EFDA Voice Newsletter September 2023	October 3, 2023
Overview of barcode use for pharmaceutical products in Ethiopia	July 25, 2023
Identification and Labeling of Pharmaceutical Products with barcode	July 25, 2023
Tenofovir alafenamide fumarate 25 mg _Vemlidy_Gilead Sciences, Inc	July 12, 2023
Sofosbuvir 400 mg and Velpatasvir 100 mg_Epclusa_Gilead Sciences, Inc	July 12, 2023
Emtricitabine and Tenofovir alafenamide fumarate_Descovy 200 mg-25 mg _Gilead Sciences, Inc	July 12, 2023
Emtricitabine and Tenofovir alafenamide fumarate_Descovy 200 mg-10 mg _Gilead Sciences, Inc	July 12, 2023
Artemether injection 80 mg_KPC Pharmaceuticals, Inc_SmPC	July 12, 2023
Artemether injection 40 mg_KPC Pharmaceuticals, Inc_SmPC	July 12, 2023

Title	Date
Artemether 20 mg and Lumefantrine 120 mg tablet_Comether_KPC Pharmaceuticals, Inc	July 12, 2023
Amphotericin B 50mg _AmBisome_Gilead Sciences, Inc.	July 12, 2023
List of Medicines for Community Pharmacy	June 9, 2023
List of Medicines for Drug Shop	June 9, 2023
CEM-ART drugs-2016-2021	June 8, 2023
A blueprint for strengthening pharmacovigilance systems in resource limited countries	June 8, 2023
Prevalence, Intensity, and Correlates of Schistosomiasis and Soil-Transmitted Helminth Infections after Five Rounds of Preventive Chemotherapy among School Children in Southern Ethiopia	June 8, 2023
Clinical Trial Authorization Guideline	June 8, 2023
Guideline on Medical Products Special import permit	June 8, 2023
Health Policy	June 7, 2023
Guidelines for Registration of Similar Biotherapeutic Products (SBPs)	June 7, 2023
Guidelines for Registration of Biotherapeutic Protein Products	June 7, 2023
Clinical Trial application processing flow diagram	June 7, 2023
List of recalled medicines and medical devices	June 7, 2023
Medicines Waste Management Disposal Directive	June 7, 2023
Report of Pfizer vaccine AEFI in July 2022	June 6, 2023
Janssen AEFI Active surveillance	June 6, 2023
Efficacy and safety of praziquantel preventive chemotherapy in Schistosoma mansoni infected school children in Southern Ethiopia A prospective cohort study	June 6, 2023
COVISHIELD vaccine report	June 6, 2023
Comparative Assessment of the National Pharmacovigilance Systems	June 6, 2023
Baseline assessment of pharmacovigilance	June 6, 2023

Title	Date
Adverse Drug Events Reported on DTG	June 6, 2023
ADE-aDSM-Preliminary EFDA	June 6, 2023
MQCD ISO/IEC 17025:2017	June 6, 2023
Definition of Powers and Duties of the Executive Organs Proclamation No. 1263-2021	June 5, 2023
Medicine and Medical Device Import, Export and Wholesale Control Directive 872-2022	June 5, 2023
SUMMARY OF PRODUCT CHARACTERISTICS TEMPLATE FOR Applicants	June 5, 2023
PATIENT INFORMATION LEAFLET (PIL)TEMPLATE FOR Applicants	June 5, 2023
AEFI SURVEILLANCE COURSE FOR HEALTH PROFESSIONALS	June 5, 2023
PV Trainer Guide May 2022	June 5, 2023
AEFI Training Manual for HCP-EFDA	June 5, 2023
PV Training for Healthcare Professionals Participant's Manual 2022	June 5, 2023
list of GCP Inspectors	June 4, 2023
List of GMP inspectors	June 4, 2023
List of pharmaceutical Importer and Wholesaler inspector	June 4, 2023
Guidelines for Consumer Reporting of Side effects of medicines	June 4, 2023
National Pharmacovigilance Guideline	June 4, 2023
TB PV guideline	June 2, 2023
HIV PV guideline	June 2, 2023
Antimalarial PV guideline	June 2, 2023
NTD PV guideline	June 2, 2023
NCD PV guideline	June 2, 2023
EFDA Vaccine Safety Risk and Crisis Communication Guideline	June 2, 2023
EFDA GMP Guideline for pharmaceutical products second edition	June 2, 2023

Title	Date
EFDA Voice Newsletter May 2023	May 24, 2023
EFDA Voice Newsletter December 2022	May 24, 2023
EFDA Voice Newsletter August 2022	May 24, 2023
EFDA Voice Newsletter April 2022	May 24, 2023
GUIDELINE FOR PREPARATION AND PUBLICATION OF PUBLIC ASSESSMENT REPORTS FOR APPROVED MEDICINES	May 23, 2023
Brochure Results on ADR Reports on DTG containing ARV regimens	May 23, 2023
Magnesium sulfate-Kalceks 500 mg per ml solution for injection_ Public assessment summary report_Package label	May 17, 2023
OCTANATE 500 and OCTANATE 1000 _SUMMARY OF PRODUCT CHARACTERISTICS	May 17, 2023
OCTANATE 500 and OCTANATE 1000 _ Public assessment summary report	May 17, 2023
Utrogestan Vaginal 100 mg and 200 mg Soft capsules _ Public assessment summary report_Package label	May 17, 2023
Utrogestan Vaginal 100 mg and 200 mg Soft capsules_SUMMARY OF PRODUCT CHARACTERISTICS	May 17, 2023
Minox 5, 50 mg ml, cutaneous solution_SUMMARY OF PRODUCT CHARACTERISTICS	May 16, 2023
Minox 5, 50 mg ml, cutaneous solution_ Public assessment summary report_Package label	May 16, 2023
Delstrigo 100 mg 300 mg 245 mg film-coated tablets_ Public assessment summary report_Package label-1	May 16, 2023
CELMANTIN 10 mg and 20 mg film-coated tablets_public assessment summary report_Package label	May 16, 2023
Delstrigo100 mg 300 mg 245 mg film-coated tablets_SUMMARY OF PRODUCT CHARACTERISTICS	May 16, 2023
CELMANTIN 10 mg and 20 mg film-coated tablets_SUMMARY OF PRODUCT CHARACTERISTICS	May 16, 2023

Title	Date
Verquvo 2.5 mg, 5 mg 10 mg film-coated tablets Summary of product Characteristics	May 15, 2023
Verquvo 2.5 mg, 5 mg 10 mg film-coated tablets_Public assessment summary report and packaging material	May 15, 2023
Guideline for Renewal of Medicines Marketing Authorization	May 15, 2023
J&J Vaccine Brochure	May 9, 2023
Medicine Marketing Authorization Directive No. 963-2023 የመድኃኒት የገበያ ፈቃድ አሰጣጥ መመሪያ ቁጥር 963-2015	May 4, 2023
CLINICAL TRIAL DIRECTIVE No. 964 2023 የህክምና ሙከራ መመሪያ ቁጥር 964-2015	May 4, 2023
Brochure-MDR medicines aDSM-TICs analysis report	April 27, 2023
Brochure-COVID-19 (Covishield-astrazeneca) vaccine active safety monitoring report	April 27, 2023
Brochure-Medsafety mobile app for ADR reporting user guide	April 27, 2023
Pfizer Vaccine Brochure	April 27, 2023
Guidelines on Reliance for Regulatory Decision Making	April 25, 2023
Pharmacovigilance newsletter 2015 Q3 March	April 24, 2023
Pharmacovigilance newsletter 2015 Q2 December	April 24, 2023
Pharmacovigilance newsletter 2015 1st quarter	April 24, 2023
Guideline for Emergency use Authorization of Medicines for Public Emergency Situations	April 3, 2023
Guideline for control of medicine advertising and promotion	April 3, 2023
List of Authorized Clinical Trial Applications V1, 2015 E.C	March 10, 2023
Definition of Organization, Powers and Duties of the Ethiopian Food and Drug Authority Council of Ministers Regulation No. 531/2023	March 8, 2023
957 Traditional medicinal products Manufacturing Certificate of Competence and Market Authorization Directive	February 1, 2023
Medicine Donation Control Directive No. 958/2023	February 1, 2023

Title	Date
Summary of product Characteristics of Sildenafil Citrate 50	January 3, 2023
Public assessment summary report of sildenafil	January 3, 2023
Pharmacovigilance newsletter Quarter 1 ,Issue 1 Sep, 2022	December 21, 2022
Pharmacovigilance newsletter Volume 1, Issue 1, 2022	December 21, 2022
Droxiderm SmPC	November 24, 2022
Droxiderm PAsRv1	November 24, 2022
Diclofenac Sodium Injection 75 mg/2 ml Public assessment summary report	November 2, 2022
SUMMARY OF PRODUCT CHARACTERISTICS DICLOFENAC SODIUM INJECTION 75MG/2ML	November 2, 2022
AEFI standard reporting form Ethiopia	October 21, 2022
Addendum to AEFI Surveillance Guideline 2022	October 21, 2022
ADE-ADR-ME-reporting form-EFDA	October 21, 2022
AEFI-Surviellance & response of AEFI guideline-3rd edition	October 21, 2022
AEFI-Surviellance & response of AEFI guideline-3rd edition	October 19, 2022
ADE-ADR-ME-reporting form-EFDA	October 19, 2022
Addendum to AEFI Surveillance Guideline 2022	October 19, 2022
AEFI standard reporting form Ethiopia	October 19, 2022
Pharmacovigilance Directive No. 932/2022 932 የመድኃኒት ጎጂ ባህሪያት ክትትል ሥርዓት መመሪያ ቁጥር 932-2015	October 17, 2022
Summary of product characteristics for Ritonavir Tablets USP 100 mg	October 17, 2022
Summary of product characteristics for Pyrazinamide 150mg Tablets	October 14, 2022
Summary of product characteristics for DOVPRELA (Pretomanid Tablets 200 mg)	October 14, 2022

Title	Date
Public assessment summary report for DOVPRELA (Pretomanid Tablets 200 mg)	October 14, 2022
Summary of product characteristics for Isoniazid 100mg and 300mg tablets	October 10, 2022
Public assessment summary report for Isoniazid 100 and 300 mg tablets	October 10, 2022
PRIVACY POLICY	October 8, 2022
Guidelines for Software as Medical Devices	September 28, 2022
Guidelines for Medical Device Refurbishment	September 28, 2022
Guidelines for Medical Device Donations	September 16, 2022
Emergency medicines list	September 2, 2022
OTC medicines list	September 2, 2022
Guidelines for Medical devices Clinical Investigation Authorization	July 26, 2022
Medical devices Decommissioning and Disposal Guideline	July 18, 2022
Medical Devices Recall Guideline	July 18, 2022
Medical devices Post Marketing Surveillance Guideline	July 18, 2022
Guidelines for Medical Devices GMP Inspection	May 18, 2022
Ethiopia's Medicine and Medicine Facility Inspection Process Achieved ISO/IEC 17020:2012 Accreditation	May 10, 2022
Pharmaceutical Products Barcoding Guideline	May 6, 2022
Guidelines for Medical device Labeling	April 28, 2022
Hemlibra_Emicizumab_ Summary of product characteristics v1	April 6, 2022
Hemlibra_Emicizumab_ Public assessment summary reportv1	April 6, 2022

Title	Date
Lumerax DT 20120_ Artemether 20 mg & Lumefantrine 120 mg Dispersible_public assessment summary report	March 21, 2022
Lumerax DT 20120_ Artemether 20 mg & Lumefantrine 120 mg Dispersible_Summary of product characteristics	March 21, 2022
የኮቪድ 19 ወረርሽኝ ለመከላከልና ለመቆጣጠር ስለሚወሰዱ ክልከላዎች እና ስለሚጣሉ ግዴታዎች ለመወሰን የወጣ መመሪያ ቁጥር 803	March 10, 2022
ፕሪከርሰር ኬሚካሎችን ለመቆጣጠር እና በአግባቡ ጥቅም ላይ ለማዋል የወጣ መመሪያ ቁጥር 393_2012	March 10, 2022
ጉድለት የተገኘበት መድኃኒት እና የህክምና መሳሪያ አሰባሰብ መመሪያ ቁጥር_392_2013	March 10, 2022
በባለስልጣኑ ስር ለሚገኙ የቴክኒክ ኮሚቴዎች የአበል ክፍያ አፈጻጸም መመሪያ 377/2013	March 10, 2022
የምግብ መድሀኒትና የጤና ክብካቤ አስተዳደርና ቁጥጥር ባለስልጣን የአንድ ጊዜ የመድኃኒት ግዢ መመሪያ 374	March 10, 2022
የአልኮል ማስታወቂያና ሽያጭ መመሪያ_372_2013	March 10, 2022
የናርኮቲክ መድኃኒቶችንና የሣይኮትሮፒክ ንጥረ ነገር ማዘዣ ወረቀት ለመቆጣጠር የወጣ መመሪያ ቁጥር 369	March 10, 2022
የምግብ ጨው ቁጥጥር መመሪያ 361-2013	March 10, 2022
የምግብ ላኪ፣ አስመጪና አከፋፋይ ቁጥጥር መመሪያ ቁጥር 357_2013	March 10, 2022
የመድኃኒትና የህክምና መሣሪያ ማስተዋወቅ መመሪያ ቁጥር 353_2013	March 10, 2022
የመድኃኒት ችርቻሮ ድርጅቶች ቁጥጥር መመሪያ ቁጥር 349_2013	March 10, 2022
ከጨርቅ የተሰራ የፊት መሸፈኛ ጭንብል አምራች የብቃት ማረጋገጫ የምስክር ወረቀት እና የገበያ ፍቃድ 347	March 10, 2022
አግባባዊ የመድኃኒት አጠቃቀም ቁጥጥር መመሪያ ቁጥር 346 2013	March 10, 2022
Administrative Measure Taking and Complaint Handling Directive No_345_2013	March 10, 2022
Pharmaceutical Manufacturer GMP Inspection Directive No 338	March 10, 2022
Medical Equipment Donation Directive No. 336-2020	March 10, 2022
Infant Formula and Follow up Formula Directive No 335 2020	March 10, 2022
Graphic_Health_Warning_directive_No 334_2013	March 10, 2022

Title	Date
Food Supplement Directive No. 333-2020	March 10, 2022
Guidelines for Medical Devices Good Clinical Practices	March 8, 2022
Master data Guideline _2nd Edition 2022	February 7, 2022
Ethiopian AMR PC_Strategic Plan 2021_2025	February 2, 2022
Risk Management Plan Guideline for COVID-19 Vaccines	January 27, 2022
General Guidelines for Medical devices Marketing Authorization	January 25, 2022
Guideline on requirements of Medical Devices Clearance at ports of entry	January 24, 2022
Guidelines for Medical Device GDP and GSP	January 20, 2022
Global Trade Item Number (GTIN) Allocation Guideline	January 17, 2022
Global Location Number (GLN) Allocation Guideline	January 17, 2022
GMP Application form	December 30, 2021
INIGAST 40 Public assessment summary report v1	December 15, 2021
INIGAST 40 Summary of product characteristics v1	December 15, 2021
ALLTERA 50 Summary of product characteristics v1	December 15, 2021
ALLTERA 50 Public assessment summary report v1	December 15, 2021
ALLTERA 50 Public assessment summary report v1	December 14, 2021
Baby food control directive 840-2021	November 25, 2021
Guideline for Medical Devices Good Manufacturing Practice	November 1, 2021
Regulatory Preparedness and Mitigation Strategy for Emergency Health Threats	October 11, 2021

Title	Date
Guidelines for Borderline Medical devices_EFDA	October 8, 2021
Guideline for Registration of Medicines 2020	September 13, 2021
የመድኃኒት የመልካም አመራረት ስርዓት ቁጥጥር አሰራር መመሪያ 830 2013	September 13, 2021
Medicine Good Manufacturing Practice (GMP) Inspection Procedure Directive 830 2021	September 13, 2021
Guideline for Registration of Insecticide Treated Net 2021	August 4, 2021
Guideline for Classification of In Vitro Diagnostic Medical Devices_EFDA	July 21, 2021
Guideline for Classification of Medical devices other than IVD Medical devices_EFDA	July 21, 2021
Guideline for Application of Accessories and Spare parts_EFDA	July 12, 2021
Guideline for Medical Devices Bundling for Marketing authorization application	May 6, 2021
Tobacco Control Directive Number 771/2021	April 20, 2021
Tobacco Control Directive Number 771/2021	April 20, 2021
National-pharmacovigilance-Roadmap-2020	April 13, 2021
Guideline for Medical device Post-approval Change Notification EFDA	April 5, 2021
Guideline for Marketing Authorization of Low risk Medical devices_EFDA	April 5, 2021
Covid-19 vaccine AEFI Reporting Form	March 12, 2021
Covid-19 vaccine AEFI Reporting Form	March 12, 2021
EFDA_Guidelines for IVD Registration Requirements	February 19, 2021
EFDA_Guidelines for Non-IVD Registration Requirements	February 19, 2021
Guideline for COVID-19 Vaccine donation	February 15, 2021
Guidance for Emergency Use Authorization of COVID-19 Vaccine	January 29, 2021

Title	Date
Traditional Medicine Clinical Trial guidance 2020	November 9, 2020
Supplementary guideline for Clinical trial Authorization 2020	November 9, 2020
Guideline for WHO Pre-qualified Medicines through Collaborative Registration Procedure	October 1, 2020
Guideline for Registration of Low-risk Medicines	October 1, 2020
Risk-based Guideline for Post-Marketing Quality Surveillance	September 25, 2020
Food Consignment Technical guideline	September 22, 2020
PV Newsletter V7 ISSUE 2 Feb 2020	July 8, 2020
PV Newsletter V7 Issue 1 Oct 2019	July 8, 2020
Annexes to cosmetic product manufacturing Directive No. 49/2020	June 17, 2020
Food and Medicine Administration Proclamation No.1112/2019	June 10, 2020
Internal quality Food import export	June 5, 2020
Traditional Medicine Guidance	May 26, 2020
የኮሮና ቫይረስ ግብአት አምራች፣ አስመጪና አከፋፋይ ድርጅት የብቃት ማረጋገጫ የምስክር ወረቀት አሰጣጥ ጊዜያዊ መመርያ	April 13, 2020
Temporary COVID-19 Medical Product Approval and Import Permit Authorization Directive	April 10, 2020
Annexes to Cosmetics Import, Export and Wholesale Control Directive No. 48/2020	April 1, 2020
Cosmetics Import, Export and Wholesale Control Directive No 331-2020	April 1, 2020
የኮሮና ቫይረስ ወረርሺኝን ለመከላከል ኅብረተሰቡ የሚጠቀምባቸው የመከላከያ ምርቶች አምራች፣ አስመጪ እና አከፋፋይ የብቃት ማረጋገጫ የምስክር አሰጣጥ ጊዜያዊ የአሠራር መመሪያ። 2012	March 25, 2020
Strategy for the Prevention and Containment of AMR in Ethiopia Oct 2015	March 7, 2019
Strategic plan For Traceability Implementation	March 7, 2019

Title	Date
STG -Primary Hospital	March 7, 2019
STG -Health Center	March 7, 2019
STG-General Hospital.	March 7, 2019
Standard operating procedures for pharmaceuticals good distribution and storage practices	March 7, 2019
RDV Formulary	March 6, 2019
PV NL 8	March 6, 2019
PV Newsletter VI ISSUE I Nov 2010	March 6, 2019
PV_Newsletter_V5_ISSUE 3_Dec_2015	March 6, 2019
PV Newsletter V2 ISSUE 1 Sept 2012	March 6, 2019
PV Newsletter V2 issue 2	March 6, 2019
PV Newsletter V1 ISSUE 3 Nov 2011	March 6, 2019
PV Newsletter V1 ISSUE 2 June 2011	March 6, 2019
PV Newsletter V5 ISSUE 2 Sep 2015	March 6, 2019
PV newsletter V3 Issue 4	March 6, 2019
PV newsletter V3 Issue 1	March 6, 2019
Proclamation 661	March 6, 2019
Pharmacovigilance newsletter No 15	March 6, 2019
Pharmacovigilance newsletter No 10	March 6, 2019
pharmacovigilance newsletter No 11	March 6, 2019
pharmacovigilance newsletter no 7	March 6, 2019
Pharmacovigilance newsletter 16	March 6, 2019
Pharmacovigilance newsletter 14	March 6, 2019
Pharmaceutical Assessment 2016	March 6, 2019
OVER THE COUNTER MEDICINES LIST FOR ETHIOPIA 2nd edition	March 6, 2019

Title	Date
NPS List and report format	March 6, 2019
NPS FORMULARY	March 6, 2019
National Medical Instruments List with minimum specification 2013	March 6, 2019
Narcotic Manual final	March 6, 2019
Medicines Market Authorization Strategy 2017	March 6, 2019
Medicines Good Dispensing Manual second edition 2012	March 6, 2019
Medicine Retail outlet model 2013	March 6, 2019
Medicine For Ethiopia NDL	March 6, 2019
Medicines Good Prescribing Manual second edition 2012	March 6, 2019
LIST OF ORPHAN MEDICINES FOR ETHIOPIA 2014	March 6, 2019
LIST OF MEDICINES FOR RURAL DRUG VENDOR 2011	March 6, 2019
List of Medicine for Health Centers	March 6, 2019
LIST OF ESSENTIAL MEDICINES FOR ETHIOPIA	March 6, 2019
LDHC	March 6, 2019
Inspection Manual for Inspectors	March 6, 2019
Health Institution Promotion	March 6, 2019
Handbook on Substance Abuse For Trainers	March 6, 2019
Guidelines on evaluation of similar bio-therapeutics	March 6, 2019
Guidelines For Registration Of Medical Devices or Supplies	March 6, 2019
Guidelines for Registration of Bio-therapeutic Protein Products	March 6, 2019
Guideline For Drug Information Center	March 6, 2019
Good prescribing manual for Ethiopia new 2	March 6, 2019
Good Dispensing manual new 2	March 4, 2019
GMP Guideliens	March 4, 2019

Title	Date
GCP Guideline	March 4, 2019
Finalized_Promotion_Control_guideline_2nd_Edition	March 4, 2019
EXECUTIVE SUMMARY GATS Ethiopia Oct 20 2017	March 4, 2019
Ethiopia Tobacco Control Strategic Plan	March 4, 2019
Ethiopian Pharma co-vigilance Guideline 2014	March 4, 2019
Ethiopia_National Drug Control Master Plan 2017	March 4, 2019
Ethiopia GATS FactSheet Aug 2017	March 4, 2019
DIB_V5_issue2_new	March 4, 2019
Citizen Charter	March 4, 2019
Bulletin_2007_Issue_1	March 4, 2019
Bulletin_2006_Issue_3	March 4, 2019
Bulletin_2006_Issue_2	March 4, 2019
ART Study	March 4, 2019
AMR Baseline Survey	March 4, 2019
Allergy_card_scan	March 4, 2019
Adverse Drug Event Reporting Form	March 4, 2019
ADR_Study	March 4, 2019
4th_LMDS_List_for_Drug_Shop	March 4, 2019
Medicines Market Authorization Strategy 2017	February 26, 2019
Rate of service fees regulation no 370 2015	February 26, 2019
Food Medicine and Healthcare Administration and Control Councils of Ministers Regulation No 299 2013	February 26, 2019
FCTC Ratification proclamation	February 26, 2019

Title	Date
Guideline for Registration of Vaccine	February 26, 2019
Guideline for Registration of Medical devices 2014	February 26, 2019
Ethiopian Medicines Formulary 2013	February 26, 2019
Drug Administration And Control Proclamation No. 176/99	January 8, 2019
National Drug Policy	January 8, 2019

Showing 1 to 353 of 353 entries

OUR VISION

To be a center of excellence in food and health products regulation in Africa.

OUR MISSION

To protect and promote public health by ensuring the safety, effectiveness, quality and proper use of regulated products through licensing, inspection, registration, laboratory testing, post-marketing surveillance, community participation, and provision of up-to-date regulatory information.

Contact Information

Address:
Africa Avenue, near Wolosefer, Kirkos sub city, 02/03 kebelle, 02

Business hours:
Mon - Fri: 8:30 AM - 5:30 PM

Phone number:
Tel: +25111-5-524122, Fax: +251115521392 & +25111552411 E-Mail: contactefda@efda.gov.et

Recent News

More than 2.7 million birr worth of illegal drugs, food, and tobacco products were collected and disposed of, along with food items adulterated with foreign substances.
November 7, 2023

The Hawassa branch office of the Authority has announced that more than 1.6 thousand tons of food and more than 978 thousand Birr worth of drugs and medical equipment have been imported into the country in the past three months at the Moyale entry and exit port.
November 7, 2023

The Ethiopian Food and Drug Authority has stated that it will collaborate with interested parties to provide effective drug safety monitoring.
May 24, 2023

Like Us On Facebook

Ethiopian Fooc
18,978 followers

Follow Page